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(54) Title: MUTATED LENTIVIRAL ENV PROTEINS AND THEIR USE AS DRUGS

(57) Abstract: The present invention relates to a pharmaceutical composition comprising, as active substance a mutated lentiviral ENV protein, substantially devoid of immunosuppressive properties or a variant of said mutated lentiviral ENV protein or a fragment of the above proteins, in association with a pharmaceutically acceptable carrier.

MUTATED LENTIVIRAL ENV PROTEINS AND THEIR USE AS DRUGS

The present invention relates to mutated lentiviral ENV proteins and their use as drugs.

Any discussion of the prior art throughout the specification should in no way be considered as an admission that such prior art is widely known or forms part of common general knowledge in the field.

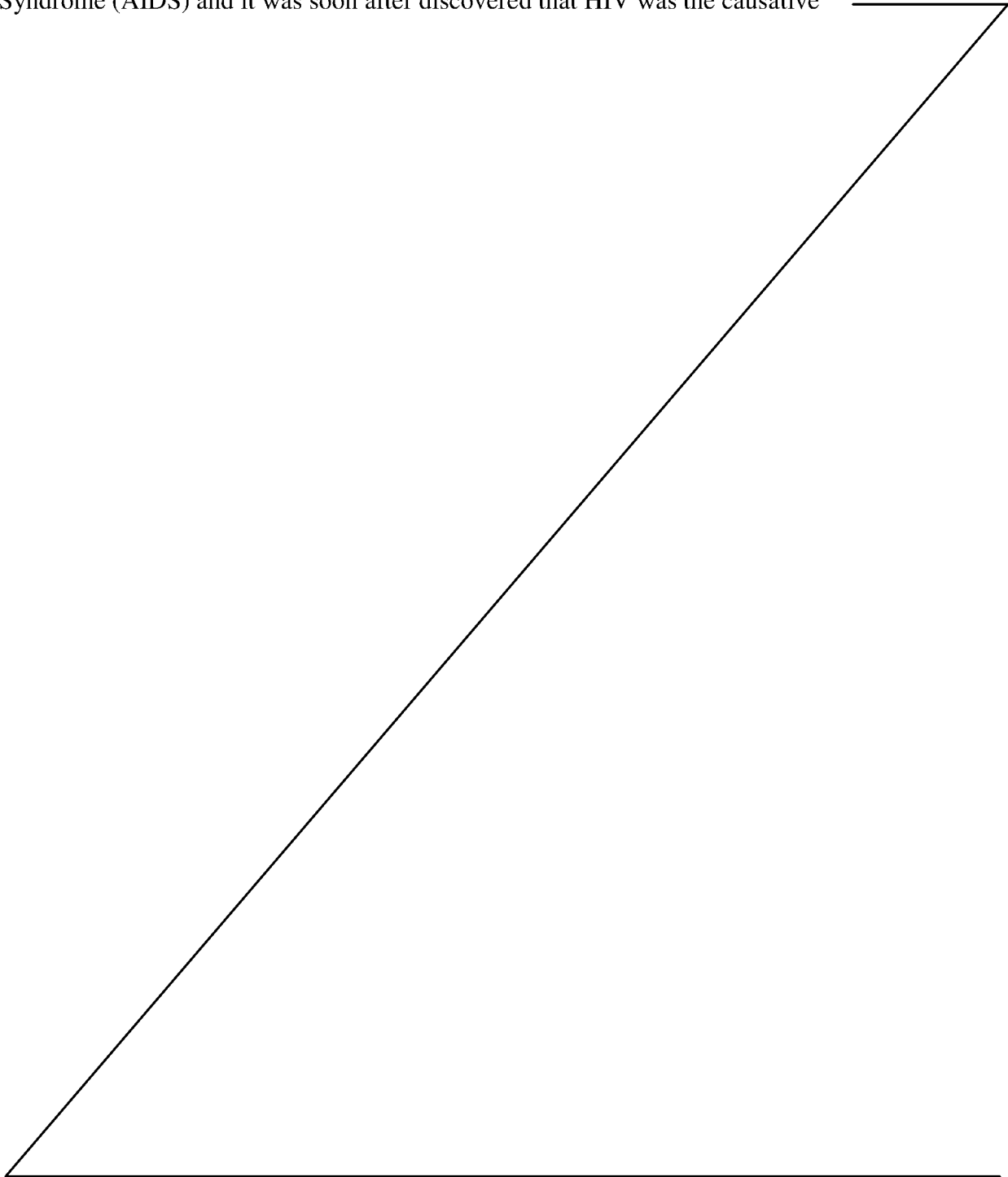
Lentivirus is a genus of slow viruses of the Retroviridae family, characterized by a long incubation period. Lentiviruses can deliver a significant amount of genetic information into the DNA of the host cell and have the unique ability among retroviruses of being able to replicate in non-dividing cells, so they are one of the most efficient methods of a gene delivery vector. HIV, SIV, and FIV are all examples of lentiviruses.

Human immunodeficiency virus (HIV) is according to WHO one of the most serious health crisis the world faces today. AIDS has killed more than 25 million people since 1981. In the most severely affected countries, the average life expectancy is now declining to 49 years of age - 13 years less than in the absence of AIDS. According to UNAIDS an estimated number of 39.5 million people were living with HIV virus in 2006 and 4.3 million were infected in 2006. In many regions new infections are heavily concentrated in the younger generations (15-24 years of age). Access to treatment and care has greatly increased in recent years. Determining real time trends to HIV incidence and in particular the impact of prevention programmes ideally requires long studies of a large number of people. Given the practical difficulties of conducting such studies focus has been placed on young women and their infants. Children living with HIV typically acquire infection through a mother-to-child-transmission (MTCT), which occur during pregnancy, delivery or during breastfeeding. Renewed efforts are urgently required to increase access to comprehensive and integrated programs to prevent HIV infection in infants and young children, which will indicate a route to HIV-free generations.

There are two known types of HIV; HIV-1 and HIV-2 that infect humans. They belong to a group of retroviruses called Lentiviruses and a virus similar to HIV has been found in African monkeys. Retroviruses transfer their genes from a producer cell to a target cell as a genomic

RNA transcript, which is reverse-transcribed after infection and integrated into the DNA genome of the target cell.

The first person with a documented HIV-infection died in 1959. In the early 1980s doctors in the US become aware, that more and more patients suffered from abnormal infections and showed signs of immune failure. The syndrome was named Acquired Immune Deficiency Syndrome (AIDS) and it was soon after discovered that HIV was the causative



agent for the observed destruction of the immune system. Initially patients were offered a treatment based solely on pain relief and almost all inevitably died. In mid 1990s there were two important breakthroughs in treatment. Firstly, a new group of antiretroviral agents were discovered and secondly it became possible to measure the amount of HIV virus in blood. These two advances made it possible to treat patients with a combination of different agents and doctors were able to check, whether the treatment actually worked. The result was that the immune system of infected patients gradually became normal and patients lived longer. Today infected people in Western countries are having the same level of quality of life as those not infected and they are able to have children, although the economical and psychological consequences of having HIV are huge. The situation is, however, even more severe in developing countries, where more than 95% of those people infected with HIV/AIDS are living. Worldwide more than 25 million people have died from AIDS in the last 25 years.

Approximately 95% of the people who get infected today live in the developing countries, where expensive antiviral drugs are not available. Therefore, there is an urgent need for an effective vaccine - the only effective solution to the uncontrolled HIV pandemic. During the last few years research has brought up new knowledge on the fundamental biology of HIV-virus which is leading to new antiviral drugs and strategies for vaccine design. In spite of these substantial advances, an effective vaccine does not yet exist. Only attenuated (that is live but weakened) HIV-strains has been able to provide immunity in primate studies even though they will never reach a required safety profile suitable for mass vaccination.

The replication process for HIV-1 has an error rate of about one per 10,000 base pairs. Since the entire viral genome is just under 10,000 base pairs, it is estimated that on average one error is introduced into the HIV-1 genome at each viral replication cycle.

This high mutation rate contributes to extensive variability of the viruses inside any one person and an even wider variability across populations.

This variability has resulted in three HIV-1 variants being described, and the subspecies of virus called "clades." The distinctions are based on the structure of the envelope proteins, which are especially variable. The M (for major) variant is by far the most prevalent world wide. Within the M variant are clades A, B, C, D, E, F, G, H, I, J and K, with clades A through E representing the vast majority of infections globally. Clades A, C and D are dominant in Africa, while clade B is the most prevalent in Europe, North and South

America and Southeast Asia. Clades E and C are dominant in Asia. These clades differ by as much as 35%. Another variant is clade O, which is observed in Cameroun isolates of HIV-1. The greatest variation in structure is seen in the envelope proteins gp120 and gp41.

5 There are two important results from the very high mutation rate of HIV- 1 that have profound consequences for the epidemic. First, the high mutation rate is one of the mechanisms that allow the virus to escape from control by drug therapies. These new viruses represent resistant strains. The high mutation rate also allows the virus to escape the patient's immune system by altering the structures that are recognized by immune components. An added consequence of this extensive variability is that the virus can also
10 escape from control by vaccines, and therefore makes it difficult to find vaccines based on envelope proteins which are effective.

Moreover, the virus produces proteins having immunosuppressive properties, allowing to escape the patient's immune system survey. Thus, cell expressing such proteins become “invisible” to the immune system.

15 Consequently for a vaccine, there is a need to provide proteins as antigens having lost, or substantially lost, their immunosuppressive functions, in order to generate an efficient response. This will enable the individuals once infected by the virus to allow the immune system to destroy the infected cells and prevent/cure the infection..

Prior art has already intended to provide such proteins.

20 For instance, the international application WO 2005/095,442 (Inventors : Renard, Mangeney & Heidmann) discloses mutations in the immunosuppressive domains of endogenous retroviruses (ERV) or onco retroviruses, such as HTLV or FeLV, ENV proteins. This document demonstrates that mutations at a specific position abolish the immunosuppressive properties of ENV proteins of ERV or onco retroviruses. However, the
25 international application WO 2005/095, 442 never mentions or suggests that the mutations made in the immunosuppressive domain of ERV- or onco retroviruses ENV proteins can be transposable to lentiviral ENV proteins.

The international application WO 2010/022,740 discloses an extremely wide consensus
30 sequence of a region of HIV ENV protein, described as follows:

X(1-22)-C(23)-X(24-28)-C(29)-(X30-50)

wherein the amino acid residues of the consensus sequence are selected from the groups of residues consisting of:

- X(1): L, S, R, P, F, A, V, M, and I; and
X(2): Q, R, K, H, L, M, and P; and
X(3): A, T, V, H, S, R, Q, G, M, and E; and
X(4): R, K, G, E, T, S, C, M, and H; and
5 X(5): V, I, L, D, A, S, F, M, and G; and
X(6): L, Q, V, M, P, W, T, and I; and
X(7): A, S, T, V, L, G, F, D, M, and E; and
X(8): V, L, I, M, A, W, K, G, and E; and
X(9): E, K, G, D, A, V, M, and F; and
10 X(10): X; and
X(11): Y, L, F, H, C, I, T, M, and N; and
X(12): L, I, V, M, Q, P, T, Y, and A; and
X(13): K, R, Q, G, S, E, H, W, T, V, M, N, Z, Y, A, P, and C; and
X(14): D, N, G, E, Y, V, S, H, A, M, and I; and
15 X(15): Q, R, H, K, P, L, M, and N; and
X(16): Q, K, R, T, H, E, S, P, M, and L; and
X(17): L, F, I, R, V, P, S, M, and H, and
X(18): L, M, P, I, H, and S; and
X(19): X; and
20 X(20): I, L, M, V, S, F, T, D, A, R, P, and J; and
X(21): W, R, G, F, L, M, and T; and
X(22): G, D, A, R, M, and C; and
X(24): X; and
X(25): G, R, E, N, A, M, and D; and
25 X(26): K, R, N, E, Q, T, S, I, M, and G; and
X(27): L, H, I, T, V, F, R, Q, S, P, A, J, M, and Y; and
X(28): I, V, T, L, R, F and M; and
X(30): T, P, Y, A, N, S, I, V, R, L, M, and H; and
X(31): T, S, P, N, M and I; and
30 X(32): A, N, T, S, D, R, FQ, P, I, E, V, M, L, K, H, C, and B; and
X(33): V, A, L, M, G, R, and C; and
X(34): X; and
X(35): W, R, G, L, M, and P; and

X(36): N, S, D, B, K, E, R, Q, M, and G; and

X(37): S, T, A, N, D, V, I, E, Y, K, L, R, G, P, M, F, W, H, Q, B, and C; and

X(38): S, T, N, I, G, R, L, C, A, W, M and E; and

X(39): W, G, A, R, E, C, Y, V, S, M, and H; and

5 X(40): X; and

X(41): N, G, K, S, D, E, T, R, H, P, A, B, V, Q, Y, M, and I; and

X(42): K, R, N, D, S, T, G, E, I, V, Y, Q, P, H, A, W, M, and C, and

X(43): S, T, N, K, I, R, D, E, P, L, A, W, G, M, H, Y, F, V, and C,

X(44): L, Y, Q, F, E, H, S, V, K, M, T, I, W, N, D, R, P, A, and G; and

10 X(45): D, E, N, S, T, K, G, L, A, Q, H, I, Y, B, R, V, P, M, F, W, Z, and C; and

X(46): E, D, Q, Y, K, N, T, S, A, W, H, M, R, I, G, L, V, Z, F, B, and P ; and

X(47): I, D, E, M, G, T, Q, S, W, L, N, Y, K, V, R, F, A, P, and H, and

X(48): W, I, T, N, D, E, L, G, S, Y, R, V, K, H, A, Q, M, and F; and

X((49): D, N, E, G, W, Q, K, H, L, B, S, I, Y, T, A, R, M, Z, and V; and

15 X(50): N, D, T, K, S, H, L, G, E, W, I, Q, M, R, B, Y, P, and A;

This consensus sequence contains 50 amino acids, in which the specific amino acids in position 10, 19, 24, 34 and 40 are defined as affecting the immunogenic properties of a HIV-1 envelope polypeptide, and the 45 remaining positions are randomly defined including the most common amino acids of wild-type HIV ENV proteins.

20

In fact, the teaching of WO 2010/022,740 is a transposition from endogenous retroviruses (ERV) or onco retroviruses to lentivirus on the basis of the teaching of WO 2005/095,442 (Inventors : Renard, Mangeney & Heidmann), but said transposition is inappropriate in the case of lentivirus, as shown by the Inventor of the present invention.

25

Briefly speaking, Dr Heidmann is an Inventor in WO 2005/095,442 and in the present invention. As a matter of fact, the effects of the mutations described in WO 2005/095,442 were also tested in lentivirus by the Inventor of the present invention, but no effect was observed when the mutations identified in endogenous retroviruses or onco retroviruses were transposed into ENV protein of lentivirus.

30

Moreover, since any amino acid can be assigned to the positions 10, 19, 24, 34 or 40 in the consensus sequence, WO 2010/022,740 teaches that such mutations can be effective using

any amino acid residue. This teaching is in contradiction with the present invention, showing that the immunosuppressive properties of HIV-1 ENV protein are only affected by specific mutations that are defined not only by their position, but also by the nature of the substituted amino acid residues.

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Furthermore, WO 2010/022,740 discloses experimental results for only one specific mutation within the immunosuppressive domain of HIV ENV protein, as defined by the 50 amino acids consensus sequence. The mutation, a substitution by R as the only one exemplified in the international application WO 2010/022,740, occurs at the amino acid in position 19, which again is equivalent to the position disclosed in the international application WO 2005/095,442 if one simply aligns ENV sequences (see Fig 3 in WO 2010/022,740, which is a copy of Figure 3 in *Benit et al. 2001, Journal of Virology, Vol. 75, No. 23, p.11709-11719*) (Of note not only the position of the amino acid, but also the nature of the substitution (by arginine) is similar to the one described in WO 2005/095,442).

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More specifically, when aligning the ENV sequences respectively of an endogenous retrovirus or onco retrovirus and a lentivirus, according to Figure 3 of Benit et al. :

Endogenous or onco retrovirus :

L Q N R R G L D L L F L K (E) G G L C A A L (WO 2005/095,442)

Lentivirus:

L A V E R Y L K D Q Q L L (G) I W G C S G K (WO 2010/022,740)

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it appears that the position 19 in lentivirus corresponds to the position in WO 2005/095,442 where a specific substitution into arginine, E→R for endogenous or onco retrovirus, results in loss of immunosuppressive activity.

25

The international application WO 2010/022,740 discloses that said mutated HIV ENV protein inhibits proliferation of PBMC *ex vivo*, but such *ex vivo* result has no *in vivo* predictive value.

However, WO 2010/022,740 never discloses or suggests that mutants are efficient in vivo, i.e. that cell expressing mutants are detected by the immune system.

Moreover, as disclosed hereafter, such mutations are not efficient in vivo. Indeed, results disclosed hereafter in the example section relative to the present invention demonstrate that said substitution G19R does not inhibit in vivo the immunosuppressive properties of the ENV protein.

As a consequence, WO 2010/022,740 raises the same technical problem as the present invention, but does not offer an appropriate technical solution. This prior art reveals the difficulties to overcome the identification of the effective mutations affecting the immunosuppressive properties of the lentiviral ENV proteins.

Thus, the provision of in vivo effective non immunosuppressive lentiviral ENV proteins remains.

One aim of the invention is to provide new mutated ENV proteins devoid of immunosuppressive properties.

Another aim of the invention is to provide a new pharmaceutical composition efficient for treating lentiviral infection.

Another aim of the invention is to provide an efficient vaccine.

According to a first aspect, the invention provides a pharmaceutical composition comprising as active substance :

- a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein having at least 90% identity, to one sequence chosen from the group consisting of SEQ ID NO : 216, SEQ ID NO : 420 and SEQ ID NO : 421,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

in association with a pharmaceutically acceptable carrier,

said substantial absence of immunosuppressive activity of the above mentioned mutated human or simian lentiviral ENV protein or of the above defined fragment being liable to be assessed by the fact that in an in vivo assay involving engrafted tumor cells rejection,

said tumor cells being transduced either so as to express said mutated ENV protein or said fragment (mutated ENV tumor cells),

or said tumor cells being transduced so as to express said wild type ENV protein or a fragment thereof (wild type ENV tumor cells),

or said tumor cells being not transduced (normal tumor cells),

the following ratio :

immunosuppression index of said mutated ENV protein or of said fragment ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV protein ($i_{\text{wild type env}}$) is less than 0.5,

$i_{\text{mutated env}}$ being defined by : (maximum area reached by mutated ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells), and

$i_{\text{wild type env}}$ being defined by : (maximum area reached by wild type ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells).

According to a second aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention, consisting of modifying the immunosuppressive property of:

a wild-type human or simian lentiviral ENV protein,

or a fragment of said wild-type human or simian lentiviral ENV protein, said fragment comprising at least 40 amino acids,

said ENV protein or fragment thereof presenting a transmembrane subunit (TM) comprising an immunosuppressive domain (ISU) containing the following amino acid sequence :

$$\text{A-[I/V]-E-[K/R]-X}'_a\text{-X}'_b\text{-X-D-Q (SEQ ID NO: 427),}$$

wherein

X represents any amino acid,

$X}'_a$ is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and

$X}'_b$ is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

said method comprising a step of introduction of at least one mutation of $X}'_a$ and/or $X}'_b$,

to obtain:

an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein having at least 90% identity, to one sequence chosen from the group consisting of SEQ ID NO : 216, SEQ ID NO : 420 and SEQ ID NO : 421,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

said substantial absence of immunosuppressive activity of the above mentioned mutated human or simian lentiviral ENV protein or of the above defined fragment being liable to be assessed by the fact that in an in vivo assay involving engrafted tumor cells rejection, said tumor cells being transduced either so as to express said mutated ENV protein or said fragment (mutated ENV tumor cells),

or said tumor cells being transduced so as to express said wild type ENV protein or a fragment thereof (wild type ENV tumor cells),

or said tumor cells being not transduced (normal tumor cells),

the following ratio :

immunosuppression index of said mutated ENV protein or of said fragment ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV protein ($i_{\text{wild type env}}$) is less than 0.5,

$i_{\text{mutated env}}$ being defined by : (maximum area reached by mutated ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells), and

$i_{\text{wild type env}}$ being defined by : (maximum area reached by wild type ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells).

According to a third aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention, consisting of modifying the immunosuppressive property of :

a wild-type human or simian lentiviral ENV protein,

or a fragment of said wild-type human or simian lentiviral ENV protein, said fragment comprising at least 40 amino acids,

said ENV protein or fragment thereof presenting a transmembrane subunit (TM) comprising an immunosuppressive domain (ISU) containing the following amino acid sequence :

A-[I/V]-E-[K/R]-Y-L-X-D-Q (SEQ ID NO : 1),

wherein

X represents any amino acid,

said method comprising a step of introduction of at least one mutation of Y in position 5 and/or L in position 6,

to obtain:

an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R.

According to a fourth aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R.

According to a fifth aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R, or

X_a is F or L, and X_b is F, G or R, or

X_a is A, G or R, and X_b is A.

According to a sixth aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is F, G or R, or

X_a is F or L, and X_b is F, G or R, or

X_a is A, G or R, and X_b is A, or

X_a is A, G or R, and X_b is F, G or R.

According to a seventh aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R.

According to an eighth aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is F, G or R.

According to a ninth aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
 or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
 said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is R, or

X_a is R, and X_b is R.

According to a tenth aspect, the invention provides a method to obtain the active substance of a pharmaceutical composition, as defined in the invention,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising an additional mutation in one at least of the amino acids at positions, 29, 36 and 37 of SEQ ID NO: X :

A[I/V]E[K/R]X_aX_bX₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPX_eX₁₀Z₁Z₂Z₃Z₄
 Z₅X_dX_e[S/T] (SEQ ID NO: 426)

wherein

X_a and X_b are as defined in any one of claims 1 to 14,

X₁ to X₁₀ represent any amino acid,

Z₁ to Z₅ represent no amino acid or any amino acid, independently from each other such that

- X_c is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V or Y, preferably A, D, or N,
- X_d is A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, Y or W, preferably A, G, S or Y,
- X_e is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, T, V, Y or W, preferably A, D or N.

According to an eleventh aspect, the invention provides the active substance when prepared according to method of the invention.

Unless the context clearly requires otherwise, throughout the description and the claims, the words “comprise”, “comprising”, and the like are to be construed in an inclusive sense as opposed to an exclusive or exhaustive sense; that is to say, in the sense of “including, but not limited to”.

The invention relates to a pharmaceutical composition comprising as active substance:

- a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,
said mutated human or simian lentiviral ENV protein having at least 70% identity, preferably at least 80% identity, to one sequence chosen from the group consisting of SEQ ID NO : 216, SEQ ID NO : 420 and SEQ ID NO : 421,
said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein, _____



X represents any amino acid,

and either

X_a is A, F, G, L, R or deleted, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

5 X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G, R or deleted, or

X_a is A, F, G, L, R or deleted, and X_b is A, F, G, R or deleted,

or

10 b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids, said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

15 A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

20 X_a is A, F, G, L, R or deleted, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G, R or deleted, or

X_a is A, F, G, L, R or deleted, and X_b is A, F, G, R or deleted,

25 in association with a pharmaceutically acceptable carrier,

said substantial absence of immunosuppressive activity of the above mentioned mutated human or simian lentiviral ENV protein or of the above defined fragment being liable to be assessed by the fact that in an *in vivo* assay involving engrafted tumor cells rejection,

30 said tumor cells being transduced either so as to express said mutated ENV protein or said fragment (“mutated ENV tumor cells”),

or said tumor cells being transduced so as to express said wild type ENV protein or a fragment thereof (“wild type ENV tumor cells”),

or said tumor cells being not transduced (“normal tumor cells”),

the following ratio :

5 immunosuppression index of said mutated ENV protein or of said fragment ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV protein ($i_{\text{wild type env}}$) is less than 0.5,

10 $i_{\text{mutated env}}$ being defined by : (maximum area reached by mutated ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells), and

$i_{\text{wild type env}}$ being defined by : (maximum area reached by wild type ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells).

15 The present application is based on the unexpected observation made by the Inventors that some specific amino acids of the immunosuppressive domain of a lentiviral ENV protein can be mutated conferring to said lentiviral ENV protein essentially no immunosuppressive properties, or no immunosuppressive properties, while retaining its antigenicity, the three-dimensional structure of the immunosuppressive domain, and its expression at the plasma
20 membrane. Moreover, the mutated lentiviral ENV protein according to the invention does not alter the infectivity of a virus expressing it.

In the invention, the “mutated simian or human lentiviral ENV proteins” means that the ENV proteins derive from the expression of an *env* gene of a lentivirus of human or simian.

25

Lentiviruses according to the invention encompassed by the invention are HIV-1 and 2 and Simian immunodeficiency virus (SIV).

Because of the high mutation rate of HIV-1, HIV-2 and SIV viruses, the “mutated ENV protein”, as defined in the invention, encompasses two meanings.

30 According to the first meaning, the said “mutated ENV protein” is the unnatural result of the intervention of human beings.

According to the second meaning, the mutated ENV protein also encompasses naturally occurring variants for which up to now the non immunosuppressive properties remain unknown.

This second meaning takes into consideration the natural variability of HIV and SIV variants inside a same infected individual, wherein the said “mutated ENV protein” might be non immunosuppressive but its property is undetectable because an HIV infected patient always carries many HIV variants, the majority of which is immunosuppressive.

The three following proteins correspond to wild type sequences of the ENV protein of HIV-1, HIV-2 and SIV respectively. In the invention, they are considered as reference sequences of wild type ENV proteins.

<p>SEQ ID NO: 417 wild type HIV-1</p>	<p>MRVKEKYQHLWRWGWKWGTMLLGILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDPNP QEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNNTGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCS GKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYTS LIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYIK IFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPLSFQTHLPTPRGPD RPEGIEEGGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRLR DLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVSL LNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
<p>SEQ ID NO: 418 Wild type HIV-2</p>	<p>MCGRNQLFVASLLASACLIYCVQYVTVFYGVPVWRNASIPLFC ATKNRDTWGTIQCLPDNDYQEIALNVTEAFDAWNNTVTEQA VEDVWVSLFETSIKPCVKLTPLCVAMRCNSTTAKNTTSTPTTTT ANTTIGENSSCIRTDNCTGLGEEEMVDCQFNMTGLERDKKKLY NETWYSKDVVCESENKTKKEKTCYMNHCNTSVITESCDKHYW DTMRFRYCAPPGFALLRCNDTNYSGFEPNCSKVVAATCTRMM ETQTSTWFGFNGTRAENRTYIYWHGRDNRTIISLNFYNTLVH CKRPGNKTVVPITLMSGLVFHSQPINRRPRQAWCWFKGEWKE AMKEVKLTLAKHPRYKGTNDTEKIRFIAPGERSDPEVAYMWT NCRGEFLYCNMTWFLNWFENRTNQTQHNYVPCHIKQIINTWH KVGKNVYLPREGQLTCNSTVTSIHANIDGGENQTNITFSAEVA</p>

	<p>ELYRLELGDYKLIEVTPIGFAPTPVKRYSSAPVRNKRGVFVLGF LGFLTTAGAAMGAASLTLSAQSRLLAGIVQQQQQLLDVVKR QQEMLRLTVWGTKNLQARVTAIEKYLKDQAQLNSWGCAFRQ VCHTTVPWVNDLTPDWNMTWQEWQIRNLEANISESLEQ AQIQQEKMYELQKLNSWDVFGNWFDLTSWIKYIQYGVYIVV GIIVLRIVIYVVQMLSRRLKGYRPVFSPPAYFQQIHIHKDREQP AREETEEDVGNVGDNWWPWPIRYIHFLIRQLIRLLNRLYNICR DLLSRSFQTLQLISQSLRRALTAVRDWLRFNTAYLQYGGEWIQ EAFRAFARATGETLTNAWRGFWGTLGQIGRGILAVPRRIRQGA EIALL</p>
<p>SEQ ID NO: 419 wild type SIV mac239</p>	<p>MGCLGNQLLIAILLVSVYGIYCTLYVTVFYGVPAWRNATIPLFC ATKNRDTWGTTQCLPDNGDYSEVALNVTESFDAWNNTVTEQ AIEDVWQLFETSIPKCVKLSPLCITMRCNKSETDRWGLTKSITT TASTTSTTASAKVDMVNETSSCIAQDNCTGLEQEQMISCKFNM TGLKRDKKKEYNETWYSADLVCEQGNNTGNESRCYMNH CNT SVIQESCDKHYWDAIRFRYCAPPGYALLRCNDTNYSGFMPKCS KVVVSSCTRMMETQTSTWFGFNGTRAENRTYIYWHGRDNRTI ISLNKYYNLTMKCRRPGNKTVLPVTIMSGLVFHSQPINDRPKQ AWCWFGGKWKDAIKEVKQTIVKHPRYTGTNNTDKINLTAPGG GDPEVTFMWTNCRGEFLYCKMNWFLNWVEDRNTANQKPKE QHKRNYVPCHIRQIINTWHKVGKNVYLPPREGDLTCNSTVTSLI ANIDWIDGNQTNITMSAEVAELYRLELGDYKLVEITPIGLPTD VKRYTTGGTSRNKRGVFVLGFLGFLATAGSAMGAASLTTLAQ SRTLLAGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTA IEKYLKDQAQLNAWGCAFRQVCHTTVPWPNASLTPKWNNET WQEWERKVDLFLEENITALLEEAQIQQEKMYELQKLNSWDVF GNWFDLASWIKYIQYGVYIVVGVILLRIVYIVQMLAKLRQGY RPVFSPPSYFQQTHIQDPALPTREGKERDGGEGGGNSSWPW QIEYIHFLIRQLIRLLTWLFSNCR TLLSRVYQILQPILQRLSATLQ RIREVLRTELTYLQYGWSYFHEAVQAVWRSATETLAGAWGDL WETLRRGGRWILAIARRIRQGLELTL</p>

Variant in the invention encompasses SIV, HIV-1 and HIV-2 ENV proteins.

5 Variants of the HIV-1 mutated ENV proteins according to the invention have at least 70%, preferably at least 80%, more preferably at least 90% of identity with the wild type amino acid sequence of the HIV-1 ENV protein, and comprises the mutations as described above, and harbour no immunosuppressive activity.

10 Variants of the HIV-2 mutated ENV proteins according to the invention have at least 70%, preferably at least 80%, more preferably at least 90% of identity with the wild type amino acid sequence of the HIV-2 ENV protein, and comprises the mutations as described above, and harbour no immunosuppressive activity.

Variants of the SIV mutated ENV proteins according to the invention have at least 70%, preferably at least 80%, more preferably at least 90% of identity with the wild type amino acid sequence of the SIV ENV protein, and comprises the mutations as described above, and harbour no immunosuppressive activity.

5

The three following proteins (SEQ ID NO : 216, 420 and 421) correspond to three mutated sequences of the ENV protein of HIV-1, HIV-2 and SIV respectively. In the invention, they are considered as reference sequences of the mutated ENV proteins.

More specifically, SEQ ID NO : 216 corresponds to the SEQ ID NO : 417 in which the amino acid residue Y in position 5 (X_a) of the sequence A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q has been substituted by R.

SEQ ID NO : 420 corresponds to the SEQ ID NO : 418 in which the amino acid residue Y in position 5 (X_a) of the sequence A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q has been substituted by R.

15 SEQ ID NO : 421 corresponds to the SEQ ID NO : 419 in which the amino acid residue Y in position 5 (X_a) of the sequence A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q has been substituted by R.

SEQ ID NO: 216 mutated HIV-1	MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDNP QEVVLNVNVTENFNMWKNMVEQMHEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLLNGSLAEEVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNNTSGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLLGIWGCS GKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYTS LIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYIK IFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSFQTHLPTPRGPD RPEGIEEGGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRLR DLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVSL LNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL
SEQ ID NO: 420 mutated HIV-2	MCGRNQLFVASLLASACLIYCVQYVTVFYGVPVWRNASIPLFC ATKNRDTWGTIQCLPDNDYQEIALNVTEAFDAWNNTVTEQA

	<p>VEDVWSLFETSIPKCVKLTPLCVAMRCNSTTAKNTTSTPTTTTT ANTTIGENSSCIRTDNCTGLGEEEMVDCQFNMTGLERDKKKLY NETWYSKDVVCESENDRKKEKTCYMNHCONTSVITESCDKHYW DTMRFRYCAPPGFALLRCNDTNYSGFEPNCSKVVAATCTRMM ETQTSTWFGFNGTRAENRTYIYWHGRDNRTIISLNKFYNLTVH CKRPGNKTVVPITLMSGLVFHSSQPINRRPRQAWCWFKGEWKE AMKEVKLTLAKHPRYKGTNDTEKIRFIAPGERSDPEVAYMWT NCRGEFLYCNMTWFLNWVENRTNQTQHNYVPCHIKQIINTWH KVGKNVYLPPREGQLTCNSTVTSIIANIDGGENQTNITFSAEVA ELYRLELGDYKLIEVTPIGFAPTPVKRYSSAPVRNKRGVFVLGF LGFLTTAGAAMGAASLTLSAQSRLLAGIVQQQQQLLDVVKR QQEMLRLTVWGTKNLQARVTAIEKRLKDQAQLNSWGCAFRQ VCHTTVPWVNDTLTPDWNNMTWQEWQIRNLEANISESLEQ AQIQQEKNMYELQKLNSWDVFGNWFDLTSWIKYIQYGVYIVV GIIVLRIVYVQMLSRRLKGYRPFVSSPPAYFQQIHIHKDREQP AREETEEDVGNSVGDNWWPWPIRYIHFLIRQLIRLLNRLYNICR DLLSRSFQTLQLISQSLRRALTAVRDWLRFNTAYLQYGGEWIQ EAFRAFARATGETLTNAWRGFWGTLGQIGRGILAVPRRIRQGA EIALL</p>
<p>SEQ ID NO: 421 mutated SIV mac239</p>	<p>MGCLGNQLLIAILLSSVYGIYCTLYVTVFYGVPAWRNATIPLFC ATKNRDTWGTTQCLPDNGDYSEVALNVTESFDAWNNTVTEQ AIEDVWQLFETSIPKCVKLSPLCITMRCNKSETDRWGLTKSITT TASTTSTTASAKVDMVNETSSCIAQDNCTGLEQEQMISCKFNM TGLKRDKKKEYNETWYSADLVCEQGNNTGNESRCYMNHCONT SVIQESCDKHYWDAIRFRYCAPPGYALLRCNDTNYSGFMPKCS KVVVSSCTRMMETQTSTWFGFNGTRAENRTYIYWHGRDNRTI ISLNKYYNLTMKCRPGNKTVLPVTIMSGLVFHSQPINDRPKQ AWCWFGGKWKDAIKEVKQTIVKHPRYTGTNNTDKINLTAPGG GDPEVTFMWTNCRGEFLYCKMNWFLNWVEDRNTANQKPKE QHKRNYVPCHIRQIINTWHKVGKNVYLPPREGDLTCNSTVTSLI ANIDWIDGNQTNITMSAEVAELRLELGDYKLVEITPIGLPTD VKRYTTGGTSRNKRGVFVLGFLGFLATAGSAMGAASLTTLTAQ SRTLLAGIVQQQQQLLDVVKRQQELLRLTVWGTKNLQTRVTA IEKRLKDQAQLNAWGCAFRQVCHTTVPWPNASLTPKWNNT WQEWERKVDLFLEENITALLEEAQIQQEKNMYELQKLNSWDVF GNWFDLASWIKYIQYGVYIVVGVILLRIVYIVQMLAKLRQGY RPVSSPPSYFQQTHIQQDPALPTREGKERDGGEGGGNSSWPW QIEYIHFLIRQLIRLLTWLFSNCRLLSRVYQILQPILQRLSATLQ RIREVLRTYLYLQYGWSYFHEAVQAVWRSATETLAGAWGDL WETLRRGGRWILAIIPRRIRQGLELTL</p>

The invention also encompasses the variants of the “mutated simian or human lentiviral ENV protein”, harbouring the above mentioned mutations, and conferring a lack of immunosuppressive properties to said variant.

- 5 Variants of the HIV-1 mutated ENV proteins according to the invention have at least 70%, preferably at least 80%, more preferably at least 90% of identity with the reference

mutated sequence of HIV-1 ENV protein (SEQ ID NO : 216), and comprises the mutations as described above, and harbour no immunosuppressive activity.

Variants of the HIV-2 mutated ENV proteins according to the invention have at least 70%, preferably at least 80%, more preferably at least 90% of identity with the reference mutated sequence of the HIV-2 ENV protein (SEQ ID NO : 420), and comprises the mutations as described above, and harbour no immunosuppressive activity.

Variants of the SIV mutated ENV proteins according to the invention have at least 70%, preferably at least 80%, more preferably at least 90% of identity with the reference mutated sequence of the SIV ENV protein (SEQ ID NO : 421), and comprises the mutations as described above, and harbour no immunosuppressive activity.

The immunosuppressive domain (ISU) of the lentivirus according to the invention can be delimited by the sequence SEQ ID NO: 6,

xGIVQQQxxLLxxxxxxQxxLxLxxWGxKxLQxRxxA[I/V]E[K/R]YLxDQxxLx (SEQ ID NO: 6)

in which x represents any amino acid.

SEQ ID NO: 6 corresponds to the non mutated ISU domain.

In this SEQ ID NO : 6, “x” (in small letters) is to be considered independently from “X” (in capital letters) used for the first time in SEQ ID NO : 416 of the present invention.

SEQ ID NO: 6 comprises SEQ ID NO: 1.

The most advantageous immunosuppressive domains of the wild type ENV proteins according to the invention comprise the following sequences:

HIV-1 ENV protein comprises the amino acid sequence AVERYLKDQ (SEQ ID NO: 7),

HIV-2 ENV protein comprises the amino acid sequence AIEKYLKDQ (SEQ ID NO: 8), and

SIV ENV protein comprises the amino acid sequence SEQ ID NO: 7 or 8.

When applying the mutations as defined above, the ISU domain loses its immunosuppressive properties.

In said SEQ ID NO: 6, the underlined amino acids are the essential amino acids.

In the invention, the ISU domain, as defined by SEQ ID NO: 6, is the minimal essential domain of the ENV protein according to the invention which harbours an immunosuppressive activity.

Examples of wild type ISU domains:

5

HIV-1

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLLG (SEQ ID NO : 366)

SGIVQQQNNLLRAIEAQQHLLKLTWGIKQLQARILAVERYLKDQQLLG (SEQ ID NO : 367)

10

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARVLAVERYLKDQQLLG (SEQ ID NO : 368)

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLLG (SEQ ID NO : 369)

15

SGIVQQQNNLLRAIEAQQQMLQLTVWGIKQLRARVLAVERYLRDQQLLG (SEQ ID NO : 370)

SGIVQQQSNNLRAIQARQHMLQLTVWGIKQLQARVLAVERYLRDQQLLG (SEQ ID NO : 371)

SIV

20

SGIVQQQNNLLKAIEAQQHLLQLSIWGVKQLQARLLAVERYLQDQQILG (SEQ ID NO : 372)

SGIVQQQNNLLRAIEAQQHLLQLSVWGIKQLQARVLAIERYLQDQQILG (SEQ ID NO : 373)

AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVSAIEKYLKDQAQLN (SEQ ID NO : 374)

25

AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKYLKDQAQLN (SEQ ID NO : 375)

HIV2

AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKYLKDQAQLN (SEQ ID NO : 376)

30

AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKYLKDQALLN (SEQ ID NO : 377)

The localization of an ISU domain can be determined in all ENV proteins of viruses as described in *Benit et al. 2001, Journal of Virology, Vol. 75, No. 23, p.11709-11719*. In a broad meaning, the ISU domain is defined by its structure and its localization, irrespective of the fact that it possesses or not an immunosuppressive activity.

- 5 In the invention, the ISU domain refers to a specific domain in which a mutation can affect the immunosuppressive property of the ENV protein.

The immunosuppressive property of the ENV protein is preferably measured using *in vivo* procedures, which are representative of the physiological environment.

- 10 The immunosuppressive properties of the mutated ENV proteins according to the invention are measured according to an *in vivo* procedure to assay the immunosuppressive activity of a ENV protein disclosed previously [*Mangeny and Heidmann Proc Natl Acad Sci USA 1998;95: 14920-14925; Mangeny et al. Proc Natl Acad Sci USA, 2007, 104(51):20534-9*].

- 15 As a physiological test, this *in vivo* procedure is performed using ENV proteins, or fragment thereof, which are not associated to another component or carrier proteins, such as BSA.

- Briefly, a wild-type (wild type lentiviral ENV protein) or modified nucleic acid expressing the protein to be tested (mutated lentiviral ENV protein) is transfected in tumour cell lines
20 such as MCA 205 or C18.1 cell lines by known transfection methods. The tumour cells expressing the protein to be tested are then injected especially subcutaneous (s.c.) injection to a host, generally mice. Following said injection, the establishment of tumour or, to the contrary, its rejection, is determined and the tumour area is measured. Tumor establishment was determined by palpation and tumor area (mm^2) was determined by
25 measuring perpendicular tumor diameters. Immunosuppression index is defined as $i = (S_{\text{env}} - S_{\text{none}}) / S_{\text{none}}$, wherein S_{env} is the maximum area reached by a tumour expressing an envelope protein and S_{none} is the maximum area reached by a tumour not expressing ENV protein (negative control).

- According to an embodiment of the invention, the above defined ratio relative to the
30 immunosuppressive index can be less than 0.2, and can even have a negative value (see Figures 2 and 3).

In vitro assay could be carried out, using high doses of synthetic peptides but they are indirect and less convincing, since the expression “immunosuppressive” is relevant when applied to animals possessing a complete immune system and not to cell lines.

An additional difficulty for the functional characterization of an ISU domain relies on the fact that the ISU carried by the retroviral ENV proteins is a highly structured proteic domain, with trimer formation within the complete ENV proteins (Caffrey M., *Biochimica et Biophysica Acta*, 1536:116-122, 2001; Caffrey et al., *The EMBO Journal*, Vol. 17, No. 16, p.4572-4584, 1998). Such structures are not naturally formed with ISU peptides of limited length, and this is most probably why most studies carried out with peptides provide irrelevant results and/or are dependent on specific coupling of the peptides to carrier proteins (such as BSA, e.g. Denner et al., *Current Science, AIDS 1994*, 8:1063-1072).

As mentioned above, the ENV proteins according to the invention are mutated. This mutation is made *in vitro*. Thus, the mutated ENV proteins according to the invention are isolated, and does not correspond to naturally occurring counterpart.

As mentioned above, the lentiviral mutated ENV proteins are substantially devoid of immunosuppressive properties. This means that the mutated ENV proteins according to the invention have no, or have reduced immunosuppressive properties with respect to the natural non mutated ENV protein from a virus of the same species. For instance, a mutated HIV ENV protein according to the invention has reduced immunosuppressive properties with respect to the wild type HIV ENV protein.

In the invention, the terms “substantially devoid of immunosuppressive properties” means that the mutated ENV proteins according to the invention have an immunosuppressive index less than about 0.2 [*Mangeney and Heidmann Proc Natl Acad Sci USA 1998*;95:14920-14925; *Mangeney et al. Proc Natl Acad Sci USA, 2007*, 104(51):20534-9].

In the invention, structures responsible for the antigenicity of the mutated lentiviral ENV protein are essentially preserved.

As intended herein, the expression “structures responsible for antigenicity” relates to structures of the protein which are liable to interact with components of the immune system such as antibodies or membrane receptors of immune cells, in particular T cells.

The mutation(s) within the immunosuppressive domain of the lentiviral ENV proteins is (are) sufficient to decrease the immunosuppressive activity of the mutated lentiviral ENV

protein with respect to the corresponding wild type ENV. However, it might be advantageous that another amino acid be also mutated because it ensures that the structure of the mutated ENV protein is essentially conserved with respect to the corresponding wild type ENV protein.

- 5 The mutated lentiviral ENV protein has substantially retained the structure, especially the antigenic structure, e.g., immunogenic determinants, of the original determined lentiviral ENV protein, i.e. the wild type non mutated lentiviral ENV protein.

These properties can be evaluated by measuring the fusion and/or infectious properties of said mutated lentiviral ENV with respect to the same properties in the wild type non mutated lentiviral ENV protein (see example).

10

Generally speaking, the mutated ENV protein involved in the present invention has an average length of about 700 to about 950 amino acids.

- 15 The invention encompasses fragments of the mutated ENV protein as defined above, provided that said fragment:

- comprises at least the sequence SEQ ID NO: 2, as defined above,
- comprises at least 40 amino acids, preferably comprises at least 50 amino acids,
- is substantially devoid of immunosuppressive properties, as defined above,
- 20 - preferably, comprises the extracellular parts of the ENV protein,
- retains the structure of the ENV protein from which it derives,
- harbours the same epitopes as the corresponding fragment in the wild type ENV protein.

- 25 According to a particular embodiment, the fragment of the mutated ENV protein of the invention can comprise about 40, 50, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650 or 700 amino acids. These values are given only in an illustrative way, as the man skilled in the art will understand that the fragment can comprise any number of amino acids comprised between 40 and 700.

30

Advantageously, the fragments according to the invention are such that, while retaining the antigenic structure of the full length mutated ENV protein, and thus of the wild type ENV

protein, they have lost major antigenic regions that are responsible for antigenicity in another region than the region corresponding to the immunosuppressive domain.

The invention also relates to a pharmaceutical composition wherein the active substance is an isolated non naturally occurring mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, as above defined, or fragments thereof.

In a particular embodiment, the invention relates to a pharmaceutical composition as defined above comprising as active substance :

10 a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

15 said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

20 and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

25 or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing

30 the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

5

in association with a pharmaceutically acceptable carrier.

The invention relates to a pharmaceutical composition as defined above comprising as
10 active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially
no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the
transmembrane (TM) subunit of a wild type human or simian lentiviral ENV
15 protein,

said mutated human or simian lentiviral ENV protein comprising a mutated
immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

20

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

25

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV
protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing
30 the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

5

in association with a pharmaceutically acceptable carrier.

According to a particular embodiment, the invention relates to a pharmaceutical composition as defined above comprising as active substance :

10 a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

15 said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

20

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

or

X_a is F or L, and X_b is F, G or R,

25

or

X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G or R,

30

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

5 X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

10 or

X_a is F or L, and X_b is F, G or R,

or

X_a is A, G or R, and X_b is A,

or

15 X_a is A, G or R, and X_b is F, G, R,

in association with a pharmaceutically acceptable carrier.

According to a particular embodiment, the invention relates to a pharmaceutical
20 composition as defined above comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV
25 protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

30 X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

or

X_a is F or L, and X_b is F, G or R,

or

5 X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G or R,

or

10 b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids, said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

15 A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

20 or

X_a is Y, I, H, C or T, and X_b is F, G or R,

or

X_a is F or L, and X_b is F, G or R,

or

25 X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G or R,

in association with a pharmaceutically acceptable carrier.

30

According to another particular embodiment, the invention also relates to a pharmaceutical composition as defined above comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

in association with a pharmaceutically acceptable carrier.

According to another particular embodiment, the invention relates to a pharmaceutical composition as defined above, comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

in association with a pharmaceutically acceptable carrier.

According to another particular embodiment, the invention relates to a pharmaceutical composition as defined above,

wherein

X_a is R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is R, or

X_a is R, and X_b is R.

5 According to another particular embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the following amino acid sequences :

A-I-E-K- X_a - X_b -X-DQ (SEQ ID NO: 422),

10 A-I-E-R- X_a - X_b -X-DQ (SEQ ID NO: 423),

A-V-E-K- X_a - X_b -X-DQ (SEQ ID NO: 424),

A-V-E-R- X_a - X_b -X-DQ (SEQ ID NO: 425).

15 In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein the resulting immunosuppressive domain, contained in the mutated lentiviral protein, comprises the amino acid sequence of the list consisting of: SEQ ID NO :9 to SEQ ID NO :37.

In the invention “SEQ ID NO : 9 to SEQ ID NO :37” encompass SEQ ID NO : 9, SEQ ID
 20 NO : 10, SEQ ID NO : 11, SEQ ID NO : 12, SEQ ID NO : 13, SEQ ID NO : 14, SEQ ID NO : 15, SEQ ID NO : 16, SEQ ID NO : 17, SEQ ID NO : 18, SEQ ID NO : 19, SEQ ID NO : 20, SEQ ID NO : 21, SEQ ID NO : 22, SEQ ID NO : 23, SEQ ID NO : 24, SEQ ID NO : 25, SEQ ID NO : 26, SEQ ID NO : 27, SEQ ID NO : 28, SEQ ID NO : 29, SEQ ID NO : 30, SEQ ID NO : 31, SEQ ID NO : 32, SEQ ID NO : 33, SEQ ID NO : 34,
 25 SEQ ID NO : 35, SEQ ID NO : 36 and SEQ ID NO : 37.

The correspondence is the following one:

AVERALKD	SEQ ID NO : 9
AVERFLKD	SEQ ID NO : 10
AVERGLKD	SEQ ID NO : 11
AVERLLKD	SEQ ID NO : 12
AVERRLKD	SEQ ID NO : 13

AVERYAKD	SEQ ID NO : 14
AVERYFKD	SEQ ID NO : 15
AVERYGKD	SEQ ID NO : 16
AVERYRKD	SEQ ID NO : 17
AVERAAKD	SEQ ID NO : 18

AVERAFKD	SEQ ID NO : 19
AVERAGKD	SEQ ID NO : 20
AVERARKD	SEQ ID NO : 21
AVERFAKD	SEQ ID NO : 22
AVERFFKD	SEQ ID NO : 23
AVERFGKD	SEQ ID NO : 24
AVERFRKD	SEQ ID NO : 25
AVERGAKD	SEQ ID NO : 26
AVERGFKD	SEQ ID NO : 27
AVERGGKD	SEQ ID NO : 28

AVERGRKD	SEQ ID NO : 29
AVERLAKD	SEQ ID NO : 30
AVERLFKD	SEQ ID NO : 31
AVERLGKD	SEQ ID NO : 32
AVERLRKD	SEQ ID NO : 33
AVERRAKD	SEQ ID NO : 34
AVERRFKD	SEQ ID NO : 35
AVERRGKD	SEQ ID NO : 36
AVERRRKD	SEQ ID NO : 37

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein the resulting immunosuppressive domain, contained in the mutated lentiviral protein, comprises the amino acid sequence of the list consisting of: SEQ ID NO :38 to SEQ ID NO :66.

In the invention “SEQ ID NO : 38 to SEQ ID NO :66” encompass SEQ ID NO : 38, SEQ ID NO : 39, SEQ ID NO : 40, SEQ ID NO : 41, SEQ ID NO : 42, SEQ ID NO : 43, SEQ ID NO : 44, SEQ ID NO : 45, SEQ ID NO : 46, SEQ ID NO : 47, SEQ ID NO : 48, SEQ ID NO : 49, SEQ ID NO : 50, SEQ ID NO : 51, SEQ ID NO : 52, SEQ ID NO : 53, SEQ ID NO : 54, SEQ ID NO : 55, SEQ ID NO : 56, SEQ ID NO : 57, SEQ ID NO : 58, SEQ ID NO : 59, SEQ ID NO : 60, SEQ ID NO : 61, SEQ ID NO : 62, SEQ ID NO : 63, SEQ ID NO : 64, SEQ ID NO : 65 and SEQ ID NO : 66.

The correspondence is the following one:

AIEKALKD	SEQ ID NO : 38	AIEKYAKD	SEQ ID NO : 43
AIEKFLKD	SEQ ID NO : 39	AIEKYFKD	SEQ ID NO : 44
AIEKGLKD	SEQ ID NO : 40	AIEKYGKD	SEQ ID NO : 45
AIEKLLKD	SEQ ID NO : 41	AIEKYRKD	SEQ ID NO : 46
AIEKRLKD	SEQ ID NO : 42	AIEKAAKD	SEQ ID NO : 47
		AIEKAFKD	SEQ ID NO : 48

AIEKAGKD	SEQ ID NO : 49
AIEKARKD	SEQ ID NO : 50
AIEKFAKD	SEQ ID NO : 51
AIEKFFKD	SEQ ID NO : 52
AIEKFGKD	SEQ ID NO : 53
AIEKFRKD	SEQ ID NO : 54
AIEKGAKD	SEQ ID NO : 55
AIEKGFKD	SEQ ID NO : 56
AIEKGGKD	SEQ ID NO : 57

AIEKGRKD	SEQ ID NO : 58
AIEKLAKD	SEQ ID NO : 59
AIEKLFKD	SEQ ID NO : 60
AIEKLGKD	SEQ ID NO : 61
AIEKLRKD	SEQ ID NO : 62
AIEKRAKD	SEQ ID NO : 63
AIEKRFKD	SEQ ID NO : 64
AIEKRGKD	SEQ ID NO : 65
AIEKRRKD	SEQ ID NO : 66

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein the resulting immunosuppressive domain, contained in the mutated lentiviral protein, comprises the amino acid sequence of the list consisting of: SEQ ID NO : 67 to SEQ ID NO : 95.

In the invention "SEQ ID NO : 67 to SEQ ID NO : 95" encompass SEQ ID NO : 67, SEQ ID NO : 68, SEQ ID NO : 69, SEQ ID NO : 70, SEQ ID NO : 71, SEQ ID NO : 72, SEQ ID NO : 73, SEQ ID NO : 74, SEQ ID NO : 75, SEQ ID NO : 76, SEQ ID NO : 77, SEQ ID NO : 78, SEQ ID NO : 79, SEQ ID NO : 80, SEQ ID NO : 81, SEQ ID NO : 82, SEQ ID NO : 83, SEQ ID NO : 84, SEQ ID NO : 85, SEQ ID NO : 86, SEQ ID NO : 87, SEQ ID NO : 88, SEQ ID NO : 89, SEQ ID NO : 90, SEQ ID NO : 91, SEQ ID NO : 92, SEQ ID NO : 93, SEQ ID NO : 94 and SEQ ID NO : 95.

The correspondence is the following one:

AIERALKD	SEQ ID NO : 67
AIERFLKD	SEQ ID NO : 68
AIERGLKD	SEQ ID NO : 69
AIERLLKD	SEQ ID NO : 70
AIERRLKD	SEQ ID NO : 71
AIERYAKD	SEQ ID NO : 72
AIERYFKD	SEQ ID NO : 73
AIERYGKD	SEQ ID NO : 74
AIERYRKD	SEQ ID NO : 75

AIERAAKD	SEQ ID NO : 76
AIERAFKD	SEQ ID NO : 77
AIERAGKD	SEQ ID NO : 78
AIERARKD	SEQ ID NO : 79
AIERFAKD	SEQ ID NO : 80
AIERFFKD	SEQ ID NO : 81
AIERFGKD	SEQ ID NO : 82

AIERFRKD	SEQ ID NO : 83
AIERGAKD	SEQ ID NO : 84
AIERGFKD	SEQ ID NO : 85
AIERGGKD	SEQ ID NO : 86
AIERGRKD	SEQ ID NO : 87
AIERLAKD	SEQ ID NO : 88
AIERLFKD	SEQ ID NO : 89

AIERLGKD	SEQ ID NO : 90
AIERLRKD	SEQ ID NO : 91
AIERRAKD	SEQ ID NO : 92
AIERRFKD	SEQ ID NO : 93
AIERRGKD	SEQ ID NO : 94
AIERRRKD	SEQ ID NO : 95

As mentioned above, the previous ENV proteins having their ISU comprising the above sequence are devoid of immunosuppressive properties.

Thus, in other words, any ENV protein of human or simian lentivirus having in their ISU an amino acid sequence comprising of the sequences SEQ ID NO: 9 to 95, is devoid of immunosuppressive properties.

In other words, a simian or human lentiviral ENV protein comprising, within its ISU domain, an amino acid sequence selected from SEQ ID NO: 9 to 95 is devoid of immunosuppressive properties.

In particular, the invention relates to a pharmaceutical composition as defined above, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences :

SEQ ID NO : 13, SEQ ID NO : 42, SEQ ID NO : 71,
SEQ ID NO : 9 to 12,
SEQ ID NO : 14 to 41,
SEQ ID NO : 43 to 70, and
SEQ ID NO : 72 to 95.

In particular, the invention relates to a pharmaceutical composition as defined above, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences :

SEQ ID NO : 13, SEQ ID NO : 42, SEQ ID NO : 71,
SEQ ID NO : 9, 11, 15 to 21, 23 to 29, 31 to 38, 40, 44 to 50, 52 to 58, 60 to 67, 69,
73 to 79, 81 to 87, 89 to 95.

In the invention, the fragments of the mutated ENV proteins according to the invention comprise or consist of the following sequences: SEQ ID NO: 67 to 211.

These fragments are also devoid of immunosuppressive properties.

However, these fragments retain the structure and the antigenicity of the corresponding immunosuppressive domain that is not mutated, i.e. the wild type immunosuppressive domain.

In the invention, the fragments of the mutated ENV proteins according to the invention comprise or consist of the following sequences: SEQ ID NO: 96 to 211.

The correspondences are as follows

SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERAL KDQQLG	SEQ ID NO : 96
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERFL KDQQLG	SEQ ID NO : 97
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERGL KDQQLG	SEQ ID NO : 98
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERLL KDQQLG	SEQ ID NO : 99
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERRL KDQQLG	SEQ ID NO : 100
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERYA KDQQLG	SEQ ID NO : 101
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERYF KDQQLG	SEQ ID NO : 102
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERYG KDQQLG	SEQ ID NO : 103
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERYR KDQQLG	SEQ ID NO : 104
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERAA KDQQLG	SEQ ID NO : 105
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERAF KDQQLG	SEQ ID NO : 106
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERAG KDQQLG	SEQ ID NO : 107
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERAR KDQQLG	SEQ ID NO : 108
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERFA KDQQLG	SEQ ID NO : 109
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERFF KDQQLG	SEQ ID NO : 110
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERFG KDQQLG	SEQ ID NO : 111
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERFR KDQQLG	SEQ ID NO : 112
SGIVQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERGA KDQQLG	SEQ ID NO : 113

SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERGF KDQQLLG	SEQ ID NO : 114
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERGG KDQQLLG	SEQ ID NO : 115
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERGR KDQQLLG	SEQ ID NO : 116
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERLA KDQQLLG	SEQ ID NO : 117
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERLF KDQQLLG	SEQ ID NO : 118
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERLG KDQQLLG	SEQ ID NO : 119
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERLR KDQQLLG	SEQ ID NO : 120
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERRA KDQQLLG	SEQ ID NO : 121
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERRF KDQQLLG	SEQ ID NO : 122
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERRG KDQQLLG	SEQ ID NO : 123
SGIVQQQSNLLRAIQARQHMLQLTVWGIKQLQARVLAVERRR KDQQLLG	SEQ ID NO : 124

AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKA LKDQAQLNSWGCAFRQ	SEQ ID NO : 125
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKF LKDQAQLNSWGCAFRQ	SEQ ID NO : 126
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKG LKDQAQLNSWGCAFRQ	SEQ ID NO : 127
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKL LKDQAQLNSWGCAFRQ	SEQ ID NO : 128
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKR LKDQAQLNSWGCAFRQ	SEQ ID NO : 129
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKY AKDQAQLNSWGCAFRQ	SEQ ID NO : 130
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKY FKDQAQLNSWGCAFRQ	SEQ ID NO : 131
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKY GKDQAQLNSWGCAFRQ	SEQ ID NO : 132
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKY RKDQAQLNSWGCAFRQ	SEQ ID NO : 133
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKA AKDQAQLNSWGCAFRQ	SEQ ID NO : 134
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKA FKDQAQLNSWGCAFRQ	SEQ ID NO : 135
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKA GKDQAQLNSWGCAFRQ	SEQ ID NO : 136

AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKA RKDQAQLNSWGCAFRQ	SEQ ID NO : 137
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKF AKDQAQLNSWGCAFRQ	SEQ ID NO : 138
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKF FKDQAQLNSWGCAFRQ	SEQ ID NO : 139
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKF GKDQAQLNSWGCAFRQ	SEQ ID NO : 140
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKF RKDQAQLNSWGCAFRQ	SEQ ID NO : 141
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKG AKDQAQLNSWGCAFRQ	SEQ ID NO : 142
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKG FKDQAQLNSWGCAFRQ	SEQ ID NO : 143
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKG GKDQAQLNSWGCAFRQ	SEQ ID NO : 144
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKG RKDQAQLNSWGCAFRQ	SEQ ID NO : 145
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKL AKDQAQLNSWGCAFRQ	SEQ ID NO : 146
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKL FKDQAQLNSWGCAFRQ	SEQ ID NO : 147
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKL GKDQAQLNSWGCAFRQ	SEQ ID NO : 148
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKL RKDQAQLNSWGCAFRQ	SEQ ID NO : 149
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKR AKDQAQLNSWGCAFRQ	SEQ ID NO : 150
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKR FKDQAQLNSWGCAFRQ	SEQ ID NO : 151
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKR GKDQAQLNSWGCAFRQ	SEQ ID NO : 152
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKR RKDQAQLNSWGCAFRQ	SEQ ID NO : 153

AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKA LQDQARLNSWGCAFRQ	SEQ ID NO : 154
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKF LQDQARLNSWGCAFRQ	SEQ ID NO : 155
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKG LQDQARLNSWGCAFRQ	SEQ ID NO : 156
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKL LQDQARLNSWGCAFRQ	SEQ ID NO : 157
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKR LQDQARLNSWGCAFRQ	SEQ ID NO : 158
AGIVQQQQQLLDVVKRQQEMLRLTVWGTKNLQARVTAIEKY AQDQARLNSWGCAFRQ	SEQ ID NO : 159

AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKY FQDQARLNSWGCAFRQ	SEQ ID NO : 160
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKY GQDQARLNSWGCAFRQ	SEQ ID NO : 161
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKY RQDQARLNSWGCAFRQ	SEQ ID NO : 162
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKA AQDQARLNSWGCAFRQ	SEQ ID NO : 163
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKA FQDQARLNSWGCAFRQ	SEQ ID NO : 164
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKA GQDQARLNSWGCAFRQ	SEQ ID NO : 165
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKA RQDQARLNSWGCAFRQ	SEQ ID NO : 166
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKF AQDQARLNSWGCAFRQ	SEQ ID NO : 167
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKF FQDQARLNSWGCAFRQ	SEQ ID NO : 168
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKF GQDQARLNSWGCAFRQ	SEQ ID NO : 169
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKF RQDQARLNSWGCAFRQ	SEQ ID NO : 170
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKG AQDQARLNSWGCAFRQ	SEQ ID NO : 171
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKG FQDQARLNSWGCAFRQ	SEQ ID NO : 172
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKG GQDQARLNSWGCAFRQ	SEQ ID NO : 173
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKG RQDQARLNSWGCAFRQ	SEQ ID NO : 174
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKL AQDQARLNSWGCAFRQ	SEQ ID NO : 175
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKL FQDQARLNSWGCAFRQ	SEQ ID NO : 176
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKL GQDQARLNSWGCAFRQ	SEQ ID NO : 177
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKL RQDQARLNSWGCAFRQ	SEQ ID NO : 178
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKR AQDQARLNSWGCAFRQ	SEQ ID NO : 179
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKR FQDQARLNSWGCAFRQ	SEQ ID NO : 180
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKR GQDQARLNSWGCAFRQ	SEQ ID NO : 181
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQARVTAIEKR RQDQARLNSWGCAFRQ	SEQ ID NO : 182

AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKA LKDQAQLNAWGCAFRQ	SEQ ID NO : 183
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKF LKDQAQLNAWGCAFRQ	SEQ ID NO : 184
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKG LKDQAQLNAWGCAFRQ	SEQ ID NO : 185
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKL LKDQAQLNAWGCAFRQ	SEQ ID NO : 186
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKR LKDQAQLNAWGCAFRQ	SEQ ID NO : 187
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKY AKDQAQLNAWGCAFRQ	SEQ ID NO : 188
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKY FKDQAQLNAWGCAFRQ	SEQ ID NO : 189
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKY GKDQAQLNAWGCAFRQ	SEQ ID NO : 190
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKY RKDQAQLNAWGCAFRQ	SEQ ID NO : 191
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKA AKDQAQLNAWGCAFRQ	SEQ ID NO : 192
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKA FKDQAQLNAWGCAFRQ	SEQ ID NO : 193
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKA GKDQAQLNAWGCAFRQ	SEQ ID NO : 194
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKA RKDQAQLNAWGCAFRQ	SEQ ID NO : 195
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKF AKDQAQLNAWGCAFRQ	SEQ ID NO : 196
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKF FKDQAQLNAWGCAFRQ	SEQ ID NO : 197
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKF GKDQAQLNAWGCAFRQ	SEQ ID NO : 198
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKF RKDQAQLNAWGCAFRQ	SEQ ID NO : 199
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKG AKDQAQLNAWGCAFRQ	SEQ ID NO : 200
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKG FKDQAQLNAWGCAFRQ	SEQ ID NO : 201
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKG GKDQAQLNAWGCAFRQ	SEQ ID NO : 202
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKG RKDQAQLNAWGCAFRQ	SEQ ID NO : 203
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKL AKDQAQLNAWGCAFRQ	SEQ ID NO : 204
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKL FKDQAQLNAWGCAFRQ	SEQ ID NO : 205
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKL	SEQ ID NO : 206

GKDQAQLNAWGCAFRQ	
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKL RKDQAQLNAWGCAFRQ	SEQ ID NO : 207
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKR AKDQAQLNAWGCAFRQ	SEQ ID NO : 208
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKR FKDQAQLNAWGCAFRQ	SEQ ID NO : 209
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKR GKDQAQLNAWGCAFRQ	SEQ ID NO : 210
AGIVQQQQQLLDVVKRQQEELLRLTVWGTKNLQTRVTAIEKR RKDQAQLNAWGCAFRQ	SEQ ID NO : 211

In particular, the invention relates to a pharmaceutical composition as defined above, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences : SEQ ID NO : 96, 98, 100, 102 to 108, 110 to 116, 118 to 125, 127, 129, 131 to 137, 139 to 145, 147 to 154, 156, 158, 160 to 166, 168 to 174, 176 to 183, 185, 187, 189 to 195, 197 to 203, 205 to 211.

Advantageously, the invention relates to the pharmaceutical composition as defined above, wherein said mutated protein consists of one of the following sequences SEQ ID NO : 212 to 269

HIV-1 LAI

MRVKEYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEEEVVIRSANFTDNAKTIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNGSEIFRPGGDMRDNRSELYKYVVKIEPLGVAPT KAKRRVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNLLRAIEAQQHLLQLTVWGKQLQARIL AVERALKDQQL GIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDREPEGIEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGW	SEQ ID NO : 212
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<p>EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERFLKDQQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDREGEIEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 213</p>
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERGLKDQQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDREGEIEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 214</p>
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK</p>	<p>SEQ ID NO : 215</p>

<p>NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERLLKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERLLKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 216</p>
<p>MRVKEYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT</p>	<p>SEQ ID NO : 217</p>

<p>LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERYAKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGKWTMLLILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERYFKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 218</p>
<p>MRVKEKYQHLWRWGKWTMLLILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERYGKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL</p>	<p>SEQ ID NO : 219</p>

<p>DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGWKWTMLLGILMICSATEKLWVTVY YGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRIKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERYRKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 220</p>
<p>MRVKEKYQHLWRWGWKWTMLLGILMICSATEKLWVTVY YGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRIKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERAAKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 221</p>
<p>MRVKEKYQHLWRWGWKWTMLLGILMICSATEKLWVTVY</p>	<p>SEQ ID NO : 222</p>

<p>YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERAFKDDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHS LIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPD RPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRS LCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGKWKGTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERAGKDDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHS LIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPD RPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRS LCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 223</p>
<p>MRVKEYQHLWRWGKWKGTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV</p>	<p>SEQ ID NO : 224</p>

<p>QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERARKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDREGEIEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWKWTMLLGLMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERFAKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDREGEIEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 225</p>
<p>MRVKEYQHLWRWGWKWTMLLGLMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV</p>	<p>SEQ ID NO : 226</p>

<p>QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERFFKDQQQLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLE DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDRPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERFGKDQQQLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLE DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDRPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 227</p>
<p>MRVKEYQHLWRWGWKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERFRKDQQQLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLE DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDRPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLIVTRIVELLGRRGW</p>	<p>SEQ ID NO : 228</p>

<p>EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERGAKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHS LIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPD RPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRS LCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 229</p>
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERGFKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHS LIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPD RPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRS LCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 230</p>
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK</p>	<p>SEQ ID NO : 231</p>

<p>NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERGGKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRLSLCLFSYHRLRDLLLVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGKWTMLL GILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGKQLQARIL AVERGRKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRLSLCLFSYHRLRDLLLVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 232</p>
<p>MRVKEKYQHLWRWGKWTMLL GILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT</p>	<p>SEQ ID NO : 233</p>

<p>LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQHLLQLTVWGIKQLQARIL AVERLAKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGKWTMLLILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQHLLQLTVWGIKQLQARIL AVERLFKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 234</p>
<p>MRVKEKYQHLWRWGKWTMLLILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQHLLQLTVWGIKQLQARIL AVERLGKDQQLLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL</p>	<p>SEQ ID NO : 235</p>

<p>DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRIKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERLRKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 236</p>
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY YGVVWKEATTTLCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNVTENFNMWKNDMVEQMHEIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLNGSLAEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRIKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERRAKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 237</p>
<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVY</p>	<p>SEQ ID NO : 238</p>

<p>YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERRFKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGKWKGTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERRGKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDDLRSCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 239</p>
<p>MRVKEYQHLWRWGKWKGTMLLGILMICSATEKLWVTVY YGVVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDP NPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKPC VKLTPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIK NCSFNISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTS VITQACPKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVS TVQCTHGIRPVVSTQLLLNGSLAEEEEVVIRSANFTDNAKTIIV</p>	<p>SEQ ID NO : 240</p>

<p>QLNQSVEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQA HCNISRAKWNATLKQIASKLREQFGNNKTIIFKQSSGGDPEIV THSFNCGGEFFYCNSTQLFNSTWFNSTWSTEGSNNTEGSDTIT LPCRKQFINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDG GNNNNGSEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPT KAKRRVVQREKRAVGIGALFLGFLGAAGSTMGARSMTLTV QARQLLSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARIL AVERRRKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQI WNNMTWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLEL DKWASLWNWFNITNWLWYIKIFIMIVGGLVGLRIVFAVLSIV NRVRQGYSPFSQTHLPTPRGPDPEGIEEEGGERDRDRSIRL VNGSLALIWDLLRSLCLFSYHRLRDLLLIVTRIVELLGRRGW EALKYWWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIE VVQGACRAIRHIPRRIRQGLERILL</p>	
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HIV-BH10

<p>MRVKEKYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCR KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERA LKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLIFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPFSQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDLLRSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 241</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCR</p>	<p>SEQ ID NO : 242</p>

<p>KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERF LKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPD RPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDL LLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERG LKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPD RPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDL LLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 243</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNV TENFNMWKNDMVEQM HEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERL LKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS</p>	<p>SEQ ID NO : 244</p>

<p>LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WVNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNTNRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERR LKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WVNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 245</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNTNRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERY AKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WVNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 246</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV</p>	<p>SEQ ID NO : 247</p>

<p>YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERY FKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMT WMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWASL WNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVRQ GYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSLA LIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGWEALKYW WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQGAY RAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERY GKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGWEALKY WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 248</p>
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS</p>	<p>SEQ ID NO : 249</p>

<p>VEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILVERY RKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDREPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDL LLIVTRIVELLGRRGW EALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWRWGTMLLGM LMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQM HEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILVERA AKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDREPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDL LLIVTRIVELLGRRGW EALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 250</p>
<p>MRVKEYQHLWRWGWRWGTMLLGM LMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQM HEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL</p>	<p>SEQ ID NO : 251</p>

<p>LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERA FKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNMT WMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWASL WNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVRQ GYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSLA LIWDDLRLSLCLFSYHRLRDLLIVTRIVELLGRRGWEALKYW WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQGAY RAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERA GKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLIVTRIVELLGRRGWEALKY WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 252</p>
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERA RKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLIVTRIVELLGRRGWEALKY</p>	<p>SEQ ID NO : 253</p>

<p>WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVVERF AKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 254</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVVERF FKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMT WMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWASL WNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVRQ GYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSLA LIWDDLRLSCLFSYHRLRDLLLIVTRIVELLGRRGWEALKYW WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQGAY RAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 255</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI</p>	<p>SEQ ID NO : 256</p>

<p>STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVVERF GKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPD RPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQM HEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVVERF RKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPD RPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 257</p>
<p>MRVKEYQHLWRWGWRWGTMLLGMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQM HEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI</p>	<p>SEQ ID NO : 258</p>

<p>KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERG AKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERG FKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNMT WMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWASL WNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVRQ GYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSLA LIWDDLRLSLCLFSYHRLRDLLLLIVTRIVELLGRRGWEALKYW WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQGAY RAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 259</p>
<p>MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERG GKDQQLGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS</p>	<p>SEQ ID NO : 260</p>

LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL	
MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAV ERG RKDQQL LGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHS�IEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL	SEQ ID NO : 261
MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV YYGVPVWKEATTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAV ERL AKDQQL LGIWGCSGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHS�IEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL	SEQ ID NO : 262
MRVKEKYQHLWRWGWRWGTMLLGMMLMICSATEKLWVTV	SEQ ID NO : 263

<p>YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERL FKDQQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMT WMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWASL WNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVRQ GYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSLA LIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGWEALKYW WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQGAY RAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERL GKDQQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRSCLFSYHRLRDLILLIVTRIVELLGRRGWEALKY WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 264</p>
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQS</p>	<p>SEQ ID NO : 265</p>

<p>VEINCTRPNNNTRKSIRIQRGPGRFVVTIGKIGNMRQAHCNIS RAKWNNTLQKIDSKLREQFGNNKTIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERL RKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDREPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRFVVTIGKIGNMRQAHCNIS RAKWNNTLQKIDSKLREQFGNNKTIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERR AKDQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDK WAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDREPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRLSLCLFSYHRLRDLLLIVTRIVELLGRRGWEALKY WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 266</p>
<p>MRVKEYQHLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTFENFMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRFVVTIGKIGNMRQAHCNIS RAKWNNTLQKIDSKLREQFGNNKTIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTEGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGGDMRDNRSELYKYKVVKIEPLGVAPTKAKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL</p>	<p>SEQ ID NO : 267</p>

<p>LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERR FKDQQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMT WMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWASL WNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVRQ GYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSLA LIWDDLRSCLFSYHRLRDLLIVTRIVELLGRRGWEALKYW WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQGAY RAIRHIPRRIRQGLERILL</p>	
<p>MRVKEYQHLLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTENFNMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERR GKDQQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRSCLFSYHRLRDLLIVTRIVELLGRRGWEALKY WNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL</p>	<p>SEQ ID NO : 268</p>
<p>MRVKEYQHLLWRWGWRWGTMLLGMMLICSATEKLWVTV YYGVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTD PNPQEVVLVNVTENFNMWKNDMVEQMHEDIISLWDQSLKP CVKLTPLCVSLKCTDLKNDTNTNSSSGRMIMEKGEIKNCSFNI STSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQAC PKVSFEPIPIHYCAPAGFAILKCNKTFNGTGPCTNVSTVQCT HGIRPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQS VEINCTRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNIS RAKWNNTLKQIDSKLREQFGNNKTIIFKQSSGGDPEIVTHSFN CGGEFFYCNSTQLFNSTWFNSTWSTKGSNNTGSDTITLPCRI KQIINMWQEVGKAMYAPPISGQIRCSSNITGLLLTRDGGNSN NESEIFRPGGDMRDNRSELYKYKVVKIEPLGVAPTAKKR RVVQREKRAVGIGALFLGFLGAAGSTMGAASMTLTVQARQL LSGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERR RKDQQQLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNM TWMEWDREINNYTSLIHSLIEESQNQQEKNEQELLELDKWAS LWNWFNITNWLWYIKLFIMIVGGLVGLRIVFAVLSVVNRVR QGYSPLSFQTHLPIPRGPDRPEGIEEEGGERDRDRSIRLVNGSL ALIWDDLRSCLFSYHRLRDLLIVTRIVELLGRRGWEALKY</p>	<p>SEQ ID NO : 269</p>

WWNLLQYWSQELKNSAVSLLNATAIAVAEGTDRVIEVVQG AYRAIRHIPRRIRQGLERILL	
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The invention also encompasses the variants of the above sequences, harbouring the above mentioned mutations, and conferring to said variant a lack of immunosuppressive properties.

In particular, the invention relates to a pharmaceutical composition as defined above, wherein said isolated mutated human or simian lentiviral ENV protein consists of one of the amino acid sequences : SEQ ID NO : 212, 214, 216, 218 to 224, 226 to 232, 234 to 241, 243, 245, 247 to 253, 255 to 261, 263 to 269.

The invention also relates to a pharmaceutical composition as defined above, wherein said mutated protein comprises additional mutations either downstream the C-terminal end of the sequence SEQ ID NO : 416 or upstream the N-terminal end of SEQ ID NO : 416.

These additional mutations can be advantageous to maintain the three-dimensional structure of the immunosuppressive domain and of the ENV protein (see details concerning the membrane expression of the ENV protein [see figure 4 and example] and concerning infectivity [see figure 5 and example]).

In a particular embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said mutated protein comprises an additional mutation in one at least of the amino acids at positions 29, 36 and 37 of SEQ ID NO: 426 :

A[I/V]E[K/R]X_aX_bX₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPX_cX₁₀Z₁Z₂Z₃Z₄Z₅
X_dX_e[S/T] (SEQ ID NO: 426)

wherein

X_a and X_b are as defined above,

X₁ to X₁₀ represent any amino acid,

Z₁ to Z₅ represent no amino acid or any amino acid, independently from each other

such that

- X_c is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, Y, or deleted, preferably A, D, or N,

- X_d is A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, Y, W, or deleted, preferably A, G, S or Y,
- X_e is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, T, V, Y, W, or deleted, preferably A, D or N.

This sequence contains the following amino acid sequence A[I/V]E[K/R]X_aX_bX₁DQ (SEQ ID NO : 416) elongated on its C-terminal end in which additional mutations are present.

It is to be noted that “X₁” in the above mentioned sequence SEQ ID NO : 426 has the same meaning as “X” in SEQ ID NO : 416.

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, said mutated protein comprising the amino acid sequences of the group consisting of SEQ ID NO: 271 to SEQ ID NO: 283.

SEQ ID NO: 271, contains the mutations Y41R, K72A:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPWNASWSNASLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 272, contains the mutations Y41R, K72G:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPWNASWSNGSLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 273, contains the mutations Y41R, K72S:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPWNASWSNSSLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 274, contains the mutations Y41R, W65D:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPDNASWSNKSLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 275, contains the mutations Y41R, W65A:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPANASWSNKSLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 276, contains the mutations Y41R, W65N:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPNNASWSNKSLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 277, contains the mutations Y41R, S73D:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPWNASWSNKDLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 278, contains the mutations Y41R, S73A:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPWNASWSNKALEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 279, contains the mutations Y41R, S73N:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
KLICTTAVPWNASWSNKNLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
NEQEL.

SEQ ID NO: 280, contains the mutations Y41R, K72Y:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
 KLICTTAVPWNASWSNYSLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
 NEQEL.

SEQ ID NO: 281, contains the mutations R40A, Y41R, K72A:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVEARRLKDQQLLGIWGCSG
 KLICTTAVPWNASWSNASLEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
 NEQEL.

SEQ ID NO: 282, contains the mutations Y41R, K72A, S73A:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
 KLICTTAVPWNASWSNAALEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
 NEQEL.

SEQ ID NO: 283, contains the mutations Y41R, W65A, K72A, S73A:

SGIVQQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRRLKDQQLLGIWGCSG
 KLICTTAVPANASWSNAALEQIWNNMTWMEWDREINNYTSLIHSLIEESQNQQEK
 NEQEL.

The mutations in the above mutated peptides are indicated by the amino acid residues which are bolded and underlined.

In another aspect, the invention relates to a pharmaceutical composition as defined above, comprising an additional mutation of one at least of the amino acids X₁₁, X₁₂, X₁₃ and X₁₄ in the following sequence (containing the above defined SEQ ID NO: 416 sequence) :

X₁₁X₁₂ X₁₃X₁₄I LA[I/V]E[K/R]X_aX_bX₁DQ (SEQ ID NO: 428),

wherein

X₁ represents any amino acid,

X_a and X_b are as defined above,

X₁₁ is :

- either deleted,
- or A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, Y or W, in particular A, G or R,

and/or

X₁₂ is :

- either deleted,
 - or A, C, D, E, F, G, H, I, K, L, M, N, P, R, S, T, V, Y or W, in particular A, G or R,
- and/or

X₁₃ is :

- either deleted,
 - or C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, Y or W, in particular R or G,
- and/or

X₁₄ is :

- either deleted,
- or A, C, D, E, F, G, H, I, K, L, M, N, P, Q, S, T, V, Y or W, in particular A or G.

This sequence contains the following amino acid sequence A[I/V]E[K/R]X_aX_bX₁DQ (SEQ ID NO : 416) elongated on its N-terminal end in which additional mutations are present.

It is to be noted that “X₁” in SEQ ID NO : 428 has exactly the same meaning as “X” in SEQ ID NO : 416.

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said mutated lentiviral protein, said variant or said fragment harbour a three-dimensional structure similar to the structure of the natural non mutated lentiviral ENV protein, non mutated lentiviral ENV variant or non mutated lentiviral ENV fragment thereof.

The skilled person knows how to measure the antigenicity, by using standard proceedings.

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said mutated lentiviral protein, said variant or said fragment are expressed at the plasma membrane at a level substantially identical to the expression at the plasma membrane of the natural non mutated lentiviral ENV protein, non mutated lentiviral ENV variant or non mutated lentiviral ENV fragment thereof.

The membrane expression of the lentiviral ENV protein according to the invention can be measured by any techniques allowing determination of a plasma membrane protein. For instance, cells can be transfected with an expression vector allowing the expression of the

mutated lentiviral ENV protein according to the invention. Cells are then incubated with an antibody recognizing specifically the extracellular part of said lentiviral mutated ENV protein. The complex (antibody/ENV protein) is detected by another antibody, and the complex can be quantified by flow cytometry (see for instance figure 4, and example).

If no complex is detected, the mutated ENV protein is not expressed at the plasma membrane. On the contrary, if the protein is expressed, this means that the mutated ENV protein is expressed at the plasma membrane.

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said mutated lentiviral ENV protein is such that a lentivirus, or a pseudotype, expressing said mutated lentiviral ENV protein, instead of the non mutated ENV protein, has a viral titer similar to the viral titer of said lentivirus, or pseudotype, expressing the non mutated lentiviral protein.

Viral titer is measured according to a commonly used protocol, and as described in the example (see figure 5).

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said mutated lentiviral ENV protein is a HIV-1 lentiviral protein consisting of the amino acid sequences SEQ ID NO: 284 to SEQ ID NO: 292.

SEQ ID NO: 284 Y41R	MRVKEKYQHLWRWGWKWTMLLGILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDNP QEVVLNVNVTENFNMWKNMVEQMHEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNTEGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNMTWMEWDREINNYT SLIHSLEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSFQTHLPTPRGP
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	<p>DRPEGIEEEGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
<p>SEQ ID NO: 285 Y41R, K72A</p>	<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVYY GVPVWKEATTLFCASDAKAYDTEVHNWVATHACVPTDNP QEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLNGLSLAEEVIRSANFTDNAKTIVQLNQSVEINC TRPNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNTEGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLLGIWGC SGKLICTTAVPWNASWSNASLEQIWNNMTWMEWDREINNYT SLIHSLEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSFQTHLPTPRGP DRPEGIEEEGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
<p>SEQ ID NO: 286 Y41R, K72G</p>	<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVYY GVPVWKEATTLFCASDAKAYDTEVHNWVATHACVPTDNP QEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLNGLSLAEEVIRSANFTDNAKTIVQLNQSVEINC TRPNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNTEGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLLGIWGC SGKLICTTAVPWNASWSNGSLEQIWNNMTWMEWDREINNYT SLIHSLEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSFQTHLPTPRGP DRPEGIEEEGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
<p>SEQ ID NO: 287 Y41R, K72S</p>	<p>MRVKEKYQHLWRWGKWTMLLGILMICSATEKLWVTVYY GVPVWKEATTLFCASDAKAYDTEVHNWVATHACVPTDNP QEVVLVNVTENFNMWKNMVEQMHEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP</p>

	KVSFEPPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLLNGSLAEEEEVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNNTEGSDTITLPCRICKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMMLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARIL AVERRLKDQQLLGIWGC SGKLICTTAVPWNASWSNSSLEQIWNNMTWMEWDREINNYTS LIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYIK IFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSQTHLPTPRGPD RPEGIEEEGGERDRDRSIRLVNGSLALIWDDLRSCLFSYHRLR DLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVSL LNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL
SEQ ID NO: 288 Y41G	MRVKEKYQHLWRWGWKWTMLL GILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNWVWATHACVPTDNP QEVVLVNVTENFNMWKNDMVEQM HEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLLNGSLAEEEEVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNNTEGSDTITLPCRICKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMMLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARIL AVERGLKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYT SLIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSQTHLPTPRGPD DRPEGIEEEGGERDRDRSIRLVNGSLALIWDDLRSCLFSYHRL RDLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL
SEQ ID NO: 289 Y41L	MRVKEKYQHLWRWGWKWTMLL GILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNWVWATHACVPTDNP QEVVLVNVTENFNMWKNDMVEQM HEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPPIHYCAPAGFAILKCNNKTFNGTGPCTNVSTVQCTHGI RPVVSTQLLLNGSLAEEEEVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNNTEGSDTITLPCRICKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMMLTVQARQLLSGIVQQQNNLL

	<p>RAIEAQQHLLQLTVWGIKQLQARILAVERLLKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYT SLIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSQTHLPTPRGP DRPEGIEEEGGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
SEQ ID NO: 290 Y41A	<p>MRVKEKYQHLWRWGKWKGTMLLGILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNWVATHACVPTDNP QEVVVLNVNVTENFNMWKNDMVEQMHEHDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCNTVSTVQCTHGI RPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNTEGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNNNGSEIFRPGG GDMRDNWRSELYKYKVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERALKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYT SLIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSQTHLPTPRGP DRPEGIEEEGGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
SEQ ID NO: 291 Y41F	<p>MRVKEKYQHLWRWGKWKGTMLLGILMICSATEKLWVTVYY GVPVWKEATTTLFCASDAKAYDTEVHNWVATHACVPTDNP QEVVVLNVNVTENFNMWKNDMVEQMHEHDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCNTVSTVQCTHGI RPVVSTQLLNGLSLAEEVVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNTEGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNNNGSEIFRPGG GDMRDNWRSELYKYKVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERFLKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYT SLIHSLIEESQNQQEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSQTHLPTPRGP DRPEGIEEEGGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQGACRAIRHIPRRIRQGLERILL</p>
SEQ ID NO: 292	<p>MRVKEKYQHLWRWGKWKGTMLLGILMICSATEKLWVTVYY</p>

L42R	<p>GVPVWKEATTTLFCASDAKAYDTEVHNVWATHACVPTDPNP QEVVLVNV TENFNMWKNMVEQM HEDIISLWDQSLKPCVKL TPLCVSLKCTDLGNATNTNSSNTNSSSGEMMMEKGEIKNCSFN ISTSIRGKVQKEYAFFYKLDIIPIDNDTTSYTLTSCNTSVITQACP KVSFEPIPIHYCAPAGFAILKCNNKTFNGTGPCNTVSTVQCTHGI RPVVSTQLLNGSLAEEEVVIRSANFTDNAKTIIVQLNQSVEINC TRPNNNTRKSIRIQRGPGRAFVTIGKIGNMRQAHCNISRAKWN ATLKQIASKLREQFGNNKTIIFKQSSGGDPEIVTHSFNCGGEFFY CNSTQLFNSTWFNSTWSTEGSNTEGSDTITLPCRKQFINMWQ EVGKAMYAPPISGQIRCSSNITGLLLTRDGGNNNGSEIFRPGG GDMRDNRSELYKYKVVKIEPLGVAPTKAKRRVVQREKRAV GIGALFLGFLGAAGSTMGARSMTLTVQARQLLSGIVQQQNNLL RAIEAQQHLLQLTVWGIKQLQARILAVERYRKDQQLLGIWGC SGKLICTTAVPWNASWSNKSLEQIWNNMTWMEWDREINNYT SLIHSLEESQNQOEKNEQELLELDKWASLWNWFNITNWLWYI KIFIMIVGGLVGLRIVFAVLSIVNRVRQGYSPFSQTHLPTPRGP DRPEGIEEEGGERDRDRSIRLVNGSLALIWDLLRSLCLFSYHRL RDLLLIVTRIVELLGRRGWEALKYWWNLLQYWSQELKNSAVS LLNATAIAVAEGTDRVIEVVQACRAIRHIPRRIRQGLERILL</p>
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Also, the following proteins are advantageous in the invention:

SEQ ID NO: 293 SIV MM251 L42R	<p>MGCLGNQLLIAILLSSVYGIYCTQYVTVFYGVPAWRNATIPLFC ATKNRDWTGTTQCLPDNGDYSELALNVTESFDAWENTVTEQA IEDVWQLFETSIIKPCVKLSPLCITMRCNKSETDRWGLTKSSTTIT TAAPTSAPVSEKIDMVNETSSCIAQNNCTGLEQEQMISCKFTMT GLKRDKTKEYNETWYSTDLVCEQGNSTDNESRCYMNH CNTS VIQESCDKHYWDTIRFRYCAPPGYALLRCNDTNYSGFMPKCSK VVVSSCTRM METQTSTWFGFNGTRAENRTYIYWHGRDNRTIIS LNKYYNLTMKCRPGNKTVLPVTIMSGLVFHSQPINDRPKQA WCWFGGKWKDAIKEVKQTIVKHPRYTGTNNTDKINLTAPGGG DPEVTFMWTNCRGEFLYCKMNWFLNWVEDRDVTTQRPKERH RRNYVPCHIRQIINTWHKVGKNVYLPPREGDLTCNSTVTS LIAN IDWTDGNQTSITMSAEVAELYRLELGDYKLVEITPIGLAPTDVK RYTTGGTSRNKRGVFLGFLGFLATAGSAMGAASLTTLTAQSR TLLAGIVQQQQQLLDVVKRQOELLRLTVWGTKNLQTRVTAIE KYRKDQAQLNAWGCAFRQVCHTTVPWPNASLTPDWNNDTW QEWERKVD FLEENITALLEEAQIQQEKNMYELQKLSWDVFG NWFDLASWIKYIQYGIYVVVGVILLRIVYIVQMLAKLRQGYRP VFSSPPSYFQXHTHTQQDPALPTREGKEGDGGEGGNSWPWQI EYIHFLIRQLIRLLTWLFSNCR TLLSRAYQILQPILQRLSATLRRV REVLRELTYLQYGWSYFHEAVQAGWRSATETLAGAWRDLW ETLRRGGRWILAIPRRIRQGLELTLL</p>
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SEQ ID NO: 294 HIV2 L42R	<p>MEPGRNQLFVVILLTSACL VYCSQYVTVFYGIPAWKNASIP LFC ATKNRDWTG TIQCLPDND D YQE IILNVTEAFDAWNNTVTEQA VEDVWHLFETSIIKPCVKL TPLCVAMNCSR VQGNTTTPNPRTSS STTSRPPTSAASIINETSNCIENNTCAGLGYEEMMQCEFNMKGL</p>
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	<p>EQDKKRRYKDTWYLEDVVCDNTTAGTCYMRHCNTSIIKESCD KHYWDAMRFRYCAPPGFALLRCNDTNYSGFEPKCTKVVAASC TRMMETQTSTWFGFNGTRAENRTYIYWHGRDNRTIISLNKYY NLTMRCKRPGNKTVLPITLMSGLVFHSQPINTRPRQAWCRFGG RWREAMQEVKQTLVQHPRYKGINDTGKINFTKPGAGSDPEVA FMWTNCRGEFLYCNMTWFLNWVEDKNQTRRNYCHIKQIINT WHKVGKNVYLPREGELACESTVTSIIANIDIDKNRTHTNITFS AEVAELYRLELGDYKLIETPIGFAPTDQRRYSSTPVRNKRGVF VLGFLGFLATAGSAMGARSLTLSAQSRLLAGIVQQQQQLLDV VKRQQEMLRLTVWGTKNLQARVTAIEKYRKHQAQLNSWGC AFRQVCHTTVPWVNDLSLSPDWKNMTWQEWKQVRYLEANIS QSLEEAQIQQEKMYELQKLNSWDILGNWFDLTSWVKYIQYG VHIVVGIIALRIAYVVQLLSRFRKGYRPVFSPPGYLQQIHIHK DRGQPANEGTEEDVGGDSGYDLWPWPINYPVQFLIHLRLLLIG LYNICRDLLSKNSPTRRLISQSLTAIRDWLRLKAAQLQYGCEWI QEAFQAFARTTRETLAGAWGWLWEAARRIGRILAVPRRIRQ GAELALL</p>
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Advantageously: the mutated lentiviral proteins are

1 - SEQ ID NO: 284 to 294; these mutated ENV proteins having substantially no immunosuppressive properties

advantageously,

2- SEQ ID NO: 284 to SEQ ID NO: 292; these mutated ENV proteins having substantially no immunosuppressive properties and being highly expressed at the plasma membrane, more advantageously,

3- SEQ ID NO: 284 to SEQ ID NO: 291; these mutated ENV proteins having substantially no immunosuppressive properties, being highly expressed at the plasma membrane and conferring to a virus expressing them a medium or high viral titer,

in particular

4- Y41F (SEQ ID NO: 291), Y41L (SEQ ID NO: 289), Y41A (SEQ ID NO: 290); these mutated ENV proteins having substantially no immunosuppressive properties, being highly expressed at the plasma membrane and confers to a virus expressing them a high viral titer.

As mentioned above “conferring to a virus expressing them a medium or high viral titer” means that, when the sequence of the wild type ENV protein is substituted by the sequence of the mutated ENV in the lentiviral or pseudotype retrovirus, the virus thus expresses a mutated lentiviral ENV protein, along with other wild type viral proteins (GAG, PRO, POL), said mutated ENV protein conferring to the virus the ability to enter in its target cell:

- either at a level comparable or higher to the ability of the wild type virus, i.e. high ability
- or at a lower level compared to the ability of the wild type virus, i.e. medium or low ability.

The ability to enter in the target cell can be measured by the viral load. Viral load is a measure of the amount of target cells that can be infected by a given amount (1 mL) of virus (see figure 5 and example).

The invention also relates to a pharmaceutical composition comprising a nucleic acid molecule coding for a mutated lentiviral ENV protein, or variant of said protein, or fragments thereof, as defined above, in association with a pharmaceutically acceptable carrier.

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said nucleic acid is comprised in a vector, said vector comprising means allowing the expression of said mutated lentiviral ENV protein, or variant of said protein, or fragments thereof.

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said nucleic acid is comprised in a vector, said nucleic acid being placed under the control of sequences that allow the expression of said mutated lentiviral ENV protein, or variant of said protein, or fragments thereof.

In another embodiment, the invention relates to a pharmaceutical composition, as defined above, in particular as a vaccine, comprising a DNA molecule coding for said mutated lentiviral ENV protein, or variant of said protein, or fragments thereof.

DNA vaccines expressing ENV proteins can be produced as described in Bellier et al., (DNA vaccines expressing retrovirus-like particles are efficient immunogens to induce neutralizing antibodies, *Vaccine*, 27(42):5772-80, 2009).

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, said vector being chosen among an eukaryotic or

prokaryotic expression vector, in particular an eukaryotic vector which is a viral vector, in particular a pox vector, such as a fowlpox, a canarypox, or a MVA (modified vaccinia virus Ankara) vector, an adenoviral vector, a lentiviral vector, a measles vector, a Sendai virus or a CMV (cytomegalovirus) vector.

In another advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, further comprising at least one nucleotide molecule coding for a GAG and/or a PRO and/or a POL protein and/or a mutated NEF protein substantially devoid of immunosuppressive properties, of a lentivirus, preferably a lentivirus of the same origin as the mutated lentiviral ENV protein.

The advantageous mutated Nef protein contains a substitution at position 93, as described in the international application WO 2006/018289 (Inventors : Renard M., Mangeney M. and Heidmann T.) and a deletion of the N-terminus of the protein involved in its myristoylation.

GAG expression will produce virus like particles (VLPs) which are particularly advantageous for a vaccine, in particular if the ENV protein is associated with the VLP (Guerbois et al., *Virology*, 388:191-203, 2009).

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains said at least one nucleic acid molecule coding for a GAG protein and/or a PRO protein and/or a POL protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains all said at least one nucleic acid molecule coding for a GAG protein and/or a PRO protein and/or a POL

protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in a vector which is different from the at least one vector containing said at least one nucleic acid molecule coding for a GAG protein and/or a PRO protein and/or a POL protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, said nucleic acid molecule coding for a GAG protein, said nucleic acid molecule coding for a PRO protein, said nucleic acid molecule coding for a POL protein and said nucleic acid molecule coding for a mutated NEF protein substantially devoid of immunosuppressive properties, are all contained in vectors which are different from each other.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, comprising at least one nucleic acid molecule coding for a GAG protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties,

of a human or simian lentivirus, said lentivirus being preferably of the same origin as the mutated lentiviral ENV protein.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains said at least one nucleic acid molecule coding for a GAG protein and/or a mutated NEF protein,

wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains a nucleic acid molecule coding for a GAG protein and for a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains a nucleic acid molecule coding for a GAG protein and for a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties, said vector being preferably a measles vector (Guerbois et al., *Virology*, 388:191-203, 2009) or a canary pox vector (Poulet et al., *Veterinary Record*, 153(5):141-145, 2003 ; Vaccari et al., *Expert Review of Vaccines*, Vol. 9, No 9, pages 997-1005, 2010).

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in a vector which is different from the at least one vector containing said at least one nucleic acid molecule coding for a GAG protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

In another particular embodiment, the invention relates to a pharmaceutical composition, as defined above, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, said nucleic acid molecule coding for a GAG protein and said nucleic acid

molecule coding for a mutated NEF protein substantially devoid of immunosuppressive properties, are all contained in vectors which are different from each other.

The invention also relates to a pharmaceutical composition according the above definition, for its use as vaccine, for the treatment or the prevention of lentiviral infection, in particular HIV-1 infection, HIV-2 infection and SIV infection.

As mentioned above the pharmaceutical composition according to the invention encompasses a vaccine, comprising as active substance, a protein as defined above, or a nucleic acid as defined above, or a vector as defined above, or a combination thereof.

In a particular embodiment, the pharmaceutical composition according to the invention encompasses a vaccine, comprising a nucleic acid coding for a GAG protein as defined above, a nucleic acid coding for a mutated ENV protein as defined above and a nucleic acid coding for a mutated NEF protein as defined above.

In a particular embodiment, the pharmaceutical composition according to the invention encompasses a vaccine as defined above, comprising a nucleic acid coding for a GAG protein as defined above, a nucleic acid coding for a mutated ENV protein as defined above and a nucleic acid coding for a mutated NEF protein as defined above, which are contained in a same vector, said same vector being preferably a canary pox vector or a measles vector, more preferably a measles vector.

In a particular embodiment, the pharmaceutical composition according to the invention encompasses a vaccine, comprising a GAG protein as defined above, a mutated ENV protein as defined above and a mutated NEF protein as defined above.

In a particular embodiment, the pharmaceutical composition according to the invention encompasses a vaccine, comprising a GAG protein as defined above, a mutated ENV protein as defined above and a mutated NEF protein as defined above which are associated to at least one adjuvant.

A comprehensive list of adjuvants that can be used in HIV vaccines is indicated in the FRANKLIN PIERCE LAW CENTER EDUCATIONAL REPORT: PATENT LANDSCAPE OF ADJUVANT FOR HIV VACCINES which is incorporated herein by reference. This list includes:

- Aluminum-based compounds such as aluminum phosphate and aluminum hydroxide.
- Immunostimulatory adjuvants like CpG, MPL and QS21 or MF59 which is a squalene oil-in water emulsions.
- Freund's incomplete adjuvant (FIA) or Freund's complete adjuvant (FCA), can also be used.
- Calcium phosphate in particular orthophosphates, metaphosphates or pyrophosphates and occasionally hydrogen or hydroxide ions.
- Muramyl dipeptide (*N*-Acetyl muramyl-L-alanine-D-isoglutamine, MDP) and its synthetic analogs such as muramyl tripeptide phosphatidylethanolamine (MTPPTdEtn),
- Trehalose-6,6'-dimycolate,
- Saponins (Triterpenoid glycosides) like QS-21
- Stearyl Tyrosine (octadecyl ester hydrochloride salt of tyrosine)

Immunostimulatory adjuvants can include the following classes of compounds:

- Polysaccharides like:
 - *Chitosan*:
 - *Inulin*:

- *Beta – Glucans*
- *Lipo-Polysaccharides or endotoxin*
- *MGN-3Actinidia eriantha (AEPS):*
- *Eldexomer:*
- *CpG ODN*

- Liposomes like
 - *Dehydration-rehydration liposome vesicles (DRVs)*
 - *Cytotoxic T lymphocyte (CTL)*
 - *CAF01*
 - *Liposomes containing lipid A (LA)*

- Lipid Polysine Core Peptides (LCP)

- Cytokine like GM-CSF, IL-2,IL-12 or IL-15

- Lipid A and Monophosphoryl Lipid A (MPL)

- Lipopeptide which are molecules consisting of lipid connected to a peptide. They are derived from the lipoprotein of bacterial cell wall.
- Exogenous Immunostimulatory Adjuvants which are compounds, or proteins that are not found in the human body like:
 - Proteosomes in particular Multiple Antigenic Peptides (MAP) or Exogenous Toxins

Suitably, the total amount mutated lentiviral ENV protein in a single dose of the immunogenic composition is 10µg and/or the total amount of unfused polypeptide in a single dose of the immunogenic composition is 20µg. In one embodiment, the total amount of all antigens in a single dose of the immunogenic composition is 0.5-50µg, 2-40µg, 5-30µg, 10µ-20µg or around 30µg, around 20µg or around 10µg.

The amount of mutated lentiviral ENV protein in a dose of the immunogenic composition is selected as an amount which induces an immune response without significant, adverse side effects in typical recipients. Such amount will vary depending upon which specific immunogen is employed and the dosing or vaccination regimen that is selected. An optimal amount for a particular immunogenic composition can be ascertained by standard studies involving observation of relevant immune responses in subjects.

Administration of the pharmaceutical composition can take the form of one or of more than one individual dose, for example as repeat doses of the same polypeptide containing composition, or in a heterologous "prime-boost" vaccination regime, including proteins and vectors. In one embodiment, the immunogenic composition of the invention is initially administered to a subject as two or three doses, wherein the doses are separated by a period of two weeks to three months, preferably one month.

Conveniently, the composition is administered to a subject (for instance as a booster) every 6-24, or 9-18 months, for instance annually. For instance, the composition is administered to a subject (for instance as a booster) at six month or 1 year intervals. Suitably in this respect, subsequent administrations of the composition to the subject boost the immune response of earlier administrations of the composition to the same subject.

In an embodiment, the immunogenic composition of the invention is used as part of a prime -boost regimen for use in the treatment or prevention of disease or infection by HIV-1 strains from one or more clades different from the one or more HIV-1 clades in the immunogenic composition. Conveniently, the composition is the priming dose. Alternatively, the composition is the boosting dose.

Suitably, two or more priming and/ or boosting doses are administered. A heterologous prime-boost regime uses administration of different forms of immunogenic composition or vaccine in the prime and the boost, each of which can itself include two or more administrations. The priming composition and the boosting composition will have at least one antigen in common, although it is not necessarily an identical form of the antigen, it can be a different form of the same antigen.

Prime boost immunisations according to the invention can be homologous prime-boost regimes or heterologous prime-boost regimes. Homologous prime-boost regimes utilize the same composition for prime and boost, for instance the immunogenic composition of the

invention. Heterologous prime-boost regimes can be performed with a combination of protein and DNA- based formulations. Such a strategy is considered to be effective in inducing broad immune responses. Adjuvanted protein vaccines induce mainly antibodies and CD4+ T cell immune responses, while delivery of DNA as a plasmid or a recombinant vector induces strong CD8+ T cell responses. Thus, the combination of protein and DNA vaccination can provide for a wide variety of immune responses. This is particularly relevant in the context of HIV, since neutralizing antibodies, CD4+ T cells and CD8+ T cells are thought to be important for the immune defense against HIV-1.

The invention also relates to a pharmaceutical composition according the above definition, for its use for the treatment of lentiviral infection.

The invention also relates to a method for treating patient afflicted by pathologies related to lentiviral infection, comprising the administration to a patient in a need thereof of a pharmaceutically efficient amount of the pharmaceutical composition as defined above.

The invention also relates to a method for inducing an immune response against a lentivirus that has infected an individual, said method comprising the administration to a patient in a need thereof of a pharmaceutically efficient amount of the pharmaceutical composition as defined above.

The invention also relates to a pharmaceutical composition as defined above, further comprising an antiviral agent.

The composition according to the invention can also be used in combination with the antiviral compositions listed in Table 1 below:

Intence [®] (TMC 125/etravirine) Tibotec -	Non-nucleoside reverse transcriptase inhibitor
Agenerase [®] (APV/amprenavir) GSK -	Protease inhibitor
Aptivus [®] (TPV/tipranavir) Boehringer -	Protease inhibitor
Crixivan [®] (IDV/indinavir) MSD -	Protease inhibitor
Invirase [®] (SQV/saquinavir) Roche -	Protease inhibitor
Kaletra [®] (LPV.r/lopinavir + ritonavir) Abbott	Protease inhibitor
Norvir [®] (ritonavir) Abbott -	Protease inhibitor
Prezista [®] (TMC 114/darunavir)	Protease inhibitor

Tibotec/Janssen-Cilag -	
Reyataz [®] (ATZ/atazanavir) BMS -	Protease inhibitor
Telzir [®] (APV/fosamprenavir) GSK -	Protease inhibitor
Viracept [®] (nelfinavir) Roche -	Protease inhibitor
Fuzeon [®] (T20/enfuvirtide) Roche -	Fusion inhibitor
Celsentri [®] (maraviroc) Pfizer -	Entry inhibitor
Isentress [®] (MK 0518/raltegravir) Merck -	Integrase inhibitor
Rescriptor [®] (delavirdine) Agouron -	Non-nucleoside reverse transcriptase inhibitor
Sustiva [®] (EFV/efavirenz) BMS -	Non-nucleoside reverse transcriptase inhibitor
Viramune [®] (nevirapine) Boehringer -	Non-nucleoside reverse transcriptase inhibitor
Combivir [®] (Retrovir [®] +Epivir [®]) GSK -	Nucleoside reverse transcriptase inhibitor
Emtriva [®] (FTC, emtricitabine) Gilead -	Nucleoside reverse transcriptase inhibitor
Epivir [®] (3TC, lamivudine) GSK -	Nucleoside reverse transcriptase inhibitor
Kivexa [®] (Ziagen [®] + Epivir [®]) GSK -	Nucleoside reverse transcriptase inhibitor
Retrovir [®] (AZT/zidovudine) QSK -	Nucleoside reverse transcriptase inhibitor
Trizivir [®] (Retrovir [®] + Epivir [®] + Ziagen [®]) GSK -	Nucleoside reverse transcriptase inhibitor
Videx [®] (ddI/didanosine) BMS -	Nucleoside reverse transcriptase inhibitor
Viread [®] (TDF/tenofovir) Gilead -	Nucleoside reverse transcriptase inhibitor
Zerit [®] (d4T/stavudine) BMS -	Nucleoside reverse transcriptase inhibitor
Ziagen [®] (ABC/abacavir) GSK -	Nucleoside reverse transcriptase inhibitor
Truvada [®] (Emtriva [®] + Viread [®]) Gilead -	Nucleoside reverse transcriptase inhibitor
Atripla [®] (Sustiva [®] + Emtriva [®] + Viread [®]) BMS / GILEAD	Nucleoside and non-nucleoside reverse transcriptase inhibitor

Table 1: Trade names of the different antivirals used for HIV treatment, as well as their specificity of action.

The invention also relates to a pharmaceutical composition as defined above, for simultaneous, separate or sequential use.

The invention relates to a pharmaceutical composition as defined above for its use for stimulating an immune response in a host organism.

The invention relates to a pharmaceutical composition as defined above for its use for inducing a specific immune response against the HIV ENV protein.

The invention relates to a pharmaceutical composition as defined above for its use for the prevention or the treatment of HIV infection, or pathologies related to AIDS.

The invention relates to a pharmaceutical composition as defined above, as a vaccine, for its use for inducing a specific immune response against the HIV ENV protein.

The invention relates to a pharmaceutical composition as defined above, as a vaccine, for its use for the prevention or the treatment of HIV infection, or pathologies related to AIDS.

In another aspect, the invention also relates to a method to obtain the active substance of a pharmaceutical composition, as defined above, consisting of modifying the immunosuppressive property of:

a wild-type human or simian lentiviral ENV protein,

or a fragment of said wild-type human or simian lentiviral ENV protein, said fragment comprising at least 40 amino acids,

said ENV protein or fragment thereof presenting a transmembrane subunit (TM) comprising an immunosuppressive domain (ISU) containing the following amino acid sequence :

A-[I/V]-E-[K/R]-X'_a-X'_b-X-D-Q (SEQ ID NO: 427),

wherein

X represents any amino acid,

X'_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and

X'_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

said method comprising a step of introduction of at least one mutation of X'_a and/or X'_b,

to obtain:

an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein having at least 70% identity, preferably at least 80% identity, to one sequence chosen from the group consisting of SEQ ID NO : 216, SEQ ID NO : 420 and SEQ ID NO : 421,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L, R or deleted, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G, R or deleted, or

X_a is A, F, G, L, R or deleted, and X_b is A, F, G, R or deleted,

said substantial absence of immunosuppressive activity of the above mentioned mutated human or simian lentiviral ENV protein or of the above defined fragment being liable to be assessed by the fact that in an *in vivo* assay involving engrafted tumor cells rejection, said tumor cells being transduced either so as to express said mutated ENV protein or said fragment (mutated ENV tumor cells),

or said tumor cells being transduced so as to express said wild type ENV protein or a fragment thereof (wild type ENV tumor cells),

or said tumor cells being not transduced (normal tumor cells),

the following ratio :

immunosuppression index of said mutated ENV protein or of said fragment ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV protein ($i_{\text{wild type env}}$) is less than 0.5,

$i_{\text{mutated env}}$ being defined by : (maximum area reached by mutated ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells), and

$i_{\text{wild type env}}$ being defined by : (maximum area reached by wild type ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells).

The invention also relates to a method to obtain the active substance of a pharmaceutical composition, according to the above defined method, consisting of modifying the immunosuppressive property of :

a wild-type human or simian lentiviral ENV protein,

or a fragment of said wild-type human or simian lentiviral ENV protein, said fragment comprising at least 40 amino acids,

said ENV protein or fragment thereof presenting a transmembrane subunit (TM) comprising an immunosuppressive domain (ISU) containing the following amino acid sequence :

A-[I/V]-E-[K/R]-Y-L-X-D-Q (SEQ ID NO : 1),

wherein

X represents any amino acid,

said method comprising a step of introduction of at least one mutation of Y in position 5 and/or L in position 6,

to obtain:

an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L, R or deleted, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G, R or deleted, or

X_a is A, F, G, L, R or deleted, and X_b is A, F, G, R or deleted.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition, ,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R, or

X_a is F or L, and X_b is F, G or R, or

X_a is A, G or R, and X_b is A.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is F, G or R, or

X_a is F or L, and X_b is F, G or R, or

X_a is A, G or R, and X_b is A, or

X_a is A, G or R, and X_b is F, G or R.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is F, G or R.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is R, or

X_a is R, and X_b is R.

The invention also relates to a method as defined above to obtain the active substance of a pharmaceutical composition,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutation in one at least of the amino acids at positions, 29, 36 and 37 of SEQ ID NO: 426 :

A[I/V]E[K/R] X_a X_b X_1 DQ X_2 X_3 L X_4 X_5 WGC[A/S][F/G] X_6 X_7 CV X_8 TX $_9$ VP X_c X_{10} Z $_1$ Z $_2$ Z $_3$ Z $_4$ Z $_5$
 X_d X_e [S/T] (SEQ ID NO: 426)

wherein

X_a and X_b are as defined in anyone of claims 1 to 15,

X_1 to X_{10} represent any amino acid,

Z_1 to Z_5 represent no amino acid or any amino acid, independently from each other such that

- X_c is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, Y, or deleted, preferably A, D, or N,
- X_d is A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, Y, W, or deleted, preferably A, G, S or Y,
- X_e is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, T, V, Y, W, or deleted, preferably A, D or N.

The present invention relates to a pharmaceutical composition comprising as active substance :

a) an isolated non naturally occurring mutated human or simian lentiviral ENV having substantially no immunosuppressive activity, said mutated human or simian lentiviral ENV resulting from mutation of the transmembrane subunit (TM) of a wild type human or simian lentiviral ENV protein, the transmembrane subunit comprising an immunosuppressive domain (ISU) comprising the following amino acid sequence:

A-[I/V]-E-[K/R]-Y-L-X-D-Q (SEQ ID NO: 1),

said sequence encompassing

AIEKYLXDQ (SEQ ID NO: 2), AIERYLXDQ (SEQ ID NO: 3), AVEKYLXDQ (SEQ ID NO: 4) and AVERYLXDQ (SEQ ID NO: 5), wherein X represents any natural amino acid,

wherein the amino acids at the positions chosen among

- the position 4 of SEQ ID NO: 1,*
- the position 5 of SEQ ID NO: 1,*
- the position 6 of SEQ ID NO: 1,*
- the positions 4 and 5 of SEQ ID NO: 1,*
- the position 4 and 6 of SEQ ID NO: 1,*
- the position 5 and 6 of SEQ ID NO: 1, and*

- the position 4, 5 and 6 of SEQ ID NO: 1,

are :

- either deleted,
- or substituted by another amino acid such that
 - when the immunosuppressive domain comprises SEQ ID NO : 2 or 4
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, W or Y, and/or
 - the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or
 - the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,
 - when the immunosuppressive domain comprises SEQ ID NO : 3 or 5,
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, L, M, N, P, Q, S, T, V, W or Y, and/or
 - the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or
 - the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,

or

b) a fragment of said mutated human or simian lentiviral ENV, said fragment comprising at least the immunosuppressive domain (ISU) of said wild type human or simian lentiviral ENV comprising the following amino acid sequence:

A-[I/V]-E-[K/R]-Y-L-X-D-Q (SEQ ID NO: 1),

wherein the amino acids at the positions chosen among

- the position 4 of SEQ ID NO: 1,
- the position 5 of SEQ ID NO: 1,
- the position 6 of SEQ ID NO: 1,
- the positions 4 and 5 of SEQ ID NO: 1,
- the position 4 and 6 of SEQ ID NO: 1,
- the position 5 and 6 of SEQ ID NO: 1, and
- the position 4, 5 and 6 of SEQ ID NO: 1,

are :

- either deleted,
- or substituted by another amino acid such that
 - when the immunosuppressive domain comprises SEQ ID NO : 2 or 4
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, W or Y, and/or
 - the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or
 - the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,
 - when the immunosuppressive domain comprises SEQ ID NO : 3 or 5,
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, L, M, N, P, Q, S, T, V, W or Y, and/or
 - the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or
 - the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,

said fragment having substantially no immunosuppressive properties,
in association of a pharmaceutically acceptable carrier.

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein the amino acids at the positions of the group consisting of

- the position 4 of SEQ ID NO: 1,
- the position 5 of SEQ ID NO: 1,
- the position 6 of SEQ ID NO: 1,
- the positions 4 and 5 of SEQ ID NO: 1,
- the position 4 and 6 of SEQ ID NO: 1,
- the position 5 and 6 of SEQ ID NO: 1, and
- the position 4, 5 and 6 of SEQ ID NO: 1,
- are substituted by another amino acid such that
 - when the immunosuppressive domain comprises SEQ ID NO : 2 or 4
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, W or Y, and/or

- *the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or*
- *the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,*
- *when the immunosuppressive domain comprises SEQ ID NO : 3 or 5,*
 - *the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, L, M, N, P, Q, S, T, V, W or Y, and/or*
 - *the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or*
 - *the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y.*

Advantageously, the mutated lentiviral protein according to the invention, comprised in the above defined composition, comprises the amino acid sequence of the list consisting of: SEQ ID NO: 407 to SEQ ID NO: 414.

The correspondence is the following one:

<i>AVEAALKD</i>	<i>SEQ ID NO : 407</i>	<i>AIEAALKD</i>	<i>SEQ ID NO : 411</i>
<i>AVEEALKD</i>	<i>SEQ ID NO : 408</i>	<i>AIEEALKD</i>	<i>SEQ ID NO : 412</i>
<i>AVESALKD</i>	<i>SEQ ID NO : 409</i>	<i>AIESALKD</i>	<i>SEQ ID NO : 413</i>
<i>AVETALKD</i>	<i>SEQ ID NO : 410</i>	<i>AIETALKD</i>	<i>SEQ ID NO : 414</i>

These proteins have been mutated at position 4 of SEQ ID NO: 1, by substitution, and the amino acid at position 5 has been substituted by A.

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein the amino acids at the positions of the group consisting of

- *the position 5 of SEQ ID NO: 1,*
- *the position 6 of SEQ ID NO: 1, and*
- *the position 5 and 6 of SEQ ID NO: 1,*
- *are substituted by another amino acid such that*
 - *the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or*

- *the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y.*

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said immunosuppressive domain (ISU) comprises the amino acid sequence of the group consisting of: SEQ ID NO : 4 or 5.

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said immunosuppressive domain (ISU) comprises the amino acid sequence SEQ ID NO : 5, and wherein

- *the amino acid residue at position 5 of SEQ ID NO: 5 is substituted by A, F, G, L or R, or*
- *the amino acid residue at position 6 of SEQ ID NO: 5 is substituted by A, F, G or R, or*
- *the amino acid residues at position 5 and 6 of SEQ ID NO: 5 are substituted by A, F, G, or R.*

This is the case when said ENV protein is the ENV protein of HIV 1 virus.

The invention also relates to a pharmaceutical composition as defined above, wherein said immunosuppressive domain (ISU) comprises the amino acid sequence SEQ ID NO : 2, and wherein

- *the amino acid residue at position 5 of SEQ ID NO: 2 is substituted by A, F, G, L or R, or*
- *the amino acid residue at position 6 of SEQ ID NO: 2 is substituted by A, F, G or R, or*
- *the amino acid residues at position 5 and 6 of SEQ ID NO: 2 are substituted by A, F, G, or R.*

The invention also relates to a pharmaceutical composition as defined above, wherein said immunosuppressive domain (ISU) comprises the amino acid sequence SEQ ID NO : 3, and wherein

- *the amino acid residue at position 5 of SEQ ID NO: 3 is substituted by A, F, G, L or R, or*
- *the amino acid residue at position 6 of SEQ ID NO: 3 is substituted by A, F, G or R, or*
- *the amino acid residues at position 5 and 6 of SEQ ID NO: 3 are substituted by A, F, G, or R.*

In one advantageous embodiment, the invention relates to a pharmaceutical composition as defined above, wherein said mutated protein further comprises the wild type amino acid sequence SEQ ID NO: 270,

X₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPWX₁₀Z₁Z₂Z₃Z₄Z₅[N/S][E/D/N/A][S/T]
(SEQ ID NO: 270)

wherein X₁-X₁₀ represents any amino acid,

wherein Z₁ to Z₅ represent no amino acid or any natural amino acid, independently from each other,

said mutated lentiviral protein further comprising a mutation in one at least of the amino acids at positions 23, 30 and 31 of SEQ ID NO: 270

said mutation being :

- *either a deletion,*
- *or a substitution such that*
 - *the amino acid at position 25 of SEQ ID NO: 270 is substituted by A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, Y,*
 - *the amino acid at position 32 of SEQ ID NO: 270 is substituted by A, C, D, E, F, G, H, I, K, L, M, P, Q, R, T, V, Y, W,*
 - *the amino acid at position 33 of SEQ ID NO: 270 is substituted by C, F, G, H, I, K, L, M, P, Q, R, S, T, V, Y, W.*

In one advantageous embodiment, the invention relates to a pharmaceutical composition comprising as active substance :

an isolated non naturally occurring mutated human or simian lentiviral ENV having substantially no immunosuppressive activity, said mutated human or simian lentiviral ENV resulting from mutation of a wild type human or simian lentiviral ENV protein, said wild

type human or simian lentiviral ENV protein comprising the following amino acid sequence:

A[I/V]E[K/R]Y₁LX₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPWX₁₀Z₁Z₂Z₃Z₄Z₅[N/S][E/D/N/A][S/T] (SEQ ID NO: 415),

wherein X₁-X₁₀ represents any amino acid,

wherein Z₁ to Z₅ represent no amino acid or any natural amino acid, independently from each other,

wherein the amino acids at the positions chosen among

- the position 4 of SEQ ID NO: 1,
- the position 5 of SEQ ID NO: 1,
- the position 6 of SEQ ID NO: 1,
- the positions 4 and 5 of SEQ ID NO: 1,
- the position 4 and 6 of SEQ ID NO: 1,
- the position 5 and 6 of SEQ ID NO: 1, and
- the position 4, 5 and 6 of SEQ ID NO: 1,

are :

- either deleted,
- or substituted by another amino acid such that
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, L, M, N, P, Q, S, T, V, W or Y, and/or
 - the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or
 - the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,

and further wherein

one at least of the amino acids at positions 29, 36 and 37 of the amino acid sequence SEQ ID NO : 415 are mutated, by

- either a deletion,
- or a substitution such that
 - the amino acid at position 29 of SEQ ID NO: 415 is substituted by A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, Y,

- *the amino acid at position 36 of SEQ ID NO: 415 is substituted by A, C, D, E, F, G, H, I, K, L, M, P, Q, R, T, V, Y, W,*
- *the amino acid at position 37 of SEQ ID NO: 415 is substituted by C, F, G, H, I, K, L, M, P, Q, R, S, T, V, Y, W.*

In another aspect, the invention relates to a pharmaceutical composition as defined above, further comprising a mutation at the position X₄

X₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPWX₁₀Z₁Z₂Z₃Z₄Z₅[N/S][E/D/N/A][S/T]
(SEQ ID NO: 270)

wherein X₁-X₃ and X₅-X₁₀ represents any amino acid,

Z₁ to Z₅ represent no amino acid or any amino acid, independently from each other

said mutation being :

- *either a deletion,*
- *or a substitution by A, C, D, E, F, H, I, K, L, M, P, Q, R, S, T, V, Y or W.*

In one advantageous embodiment, the invention relates to a pharmaceutical composition comprising as active substance :

an isolated non naturally occurring mutated human or simian lentiviral ENV having substantially no immunosuppressive activity, said mutated human or simian lentiviral ENV resulting from mutation of a wild type human or simian lentiviral ENV protein, said wild type human or simian lentiviral ENV protein comprising the following amino acid sequence:

A[I/V]E[K/R]Y LX₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPWX₁₀Z₁Z₂Z₃Z₄Z₅[N/S][E/D/N/A][S/T] *(SEQ ID NO: 415),*

wherein X₁-X₁₀ represents any amino acid,

wherein Z₁ to Z₅ represent no amino acid or any natural amino acid, independently from each other,

wherein the amino acids at the positions chosen among

- *the position 4 of SEQ ID NO: 1,*
- *the position 5 of SEQ ID NO: 1,*
- *the position 6 of SEQ ID NO: 1,*
- *the positions 4 and 5 of SEQ ID NO: 1,*

- the position 4 and 6 of SEQ ID NO: 1,
 - the position 5 and 6 of SEQ ID NO: 1, and
 - the position 4, 5 and 6 of SEQ ID NO: 1,
- are :

- either deleted,
- or substituted by another amino acid such that
 - the amino acid residue at position 4 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, L, M, N, P, Q, S, T, V, W or Y, and/or
 - the amino acid residue at position 5 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, I, L, M, N, P, Q, R, S, T, V or W, and/or
 - the amino acid residue at position 6 of SEQ ID NO: 1 is substituted by A, C, D, E, F, G, H, I, K, M, N, P, Q, R, S, T, V, W or Y,

and possibly, further wherein

one at least of the amino acids at positions 29, 36 and 37 of the amino acid sequence SEQ ID NO : 415 are mutated, by

- either a deletion,
- or a substitution such that
 - the amino acid at position 29 of SEQ ID NO: 415 is substituted by A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V, Y,
 - the amino acid at position 36 of SEQ ID NO: 415 is substituted by A, C, D, E, F, G, H, I, K, L, M, P, Q, R, T, V, Y, W,
 - the amino acid at position 37 of SEQ ID NO: 415 is substituted by C, F, G, H, I, K, L, M, P, Q, R, S, T, V, Y, W,

and/or possibly further wherein the amino acid at position X_4 is either deleted or substituted by A, C, D, E, F, H, I, K, L, M, P, Q, R, S, T, V, Y or W, preferably by R or H.

LEGEND TO THE FIGURES

Figure 1 :

Amino acid sequences of the HIV ectodomain deletants, with the indicated length, and of single mutant (substitution are underlined) ectodomains.

Figure 2:

Functional delineation of the immunosuppressive domain of the HIV envelope. The immunosuppressive activity of 115aa-long and truncated HIV envelope ectodomains (see structures on the left) was tested using the MCA205 tumor rejection *in vivo* assay. Immunosuppression indexes are given as histograms on the right (mean values +/- SD).

Figure 3 :

Functional identification of the aminoacid in the HIV envelope ectodomain directly involved in immunosuppressive activity, and search for aminoacid substitutions inhibiting this activity. Immunosuppressive activity was tested *in vivo* as in Figure 2.

Figure 4 :

Expression profile of mutated HIV envelopes using full-length HIVenv-expressing vectors and an anti-SU HIVenv(110H) monoclonal antibody (FACS). The data are expressed as percentages of the control HIVenv WT.

Figure 5 :

Infectivity of HIVenv-pseudotyped viral particles showing functionality of the full-length WT and of specifically mutated HIV envelopes. The results are expressed as viral titers (number of infected target cells/mL of virus supernatant).

Figure 6 :

Functional identification of the amino acid in the SIV envelope ectodomain directly involved in immunosuppressive activity, and search for aminoacid substitutions inhibiting this activity. Immunosuppressive activity was tested using the *in vivo* MCA205 tumor rejection assay. Immunosuppression indexes are represented as histograms (mean values +/- SD).

EXAMPLE

Immunosuppressive domains have been identified on the envelope proteins of oncoretroviruses, of either the gamma- (murine MLV) or the delta (e.g. human HTLV) type, as well as of some endogenous retroviruses (e.g. HERV-FRD). These domains have a

highly conserved crystallographic structure, although their primary sequences are quite diverse, and an amino-acid (either Q, E, or K) at a definite position has previously been demonstrated to be essential for the immunosuppressive activity of the corresponding envelope protein. Mutation of this amino acid to an Arginine (R) was further demonstrated to result in inhibition of the immunosuppressive activity of the mutated envelope protein, with in some cases complete conservation of the other functional properties of the envelope protein, including its ability to be normally expressed at the cell membrane, to be captured by a nascent retroviral particle, and finally to confer infectivity of the mutant virus in vitro. Such mutants allowed an unambiguous demonstration of the essential role of the immunosuppressive activity for viremia in vivo, with the mutant IS- virus being unable to escape the host immune system and propagate in an immunocompetent animal [Schlecht-Louf et al., *Proc Natl Acad Sci U S A.* 2010;107(8):3782-7].

On the basis of the identification of an IS domain and on the ability of the IS function to be inhibited by specific mutations within the IS domain that do not disrupt the antigenic structure of the corresponding viral protein, vaccinal approaches are being developed with vaccines containing mutated « optimized » IS-negative viral antigens, which demonstrate an increased immunogenicity as compared to those using the native viral antigens. Search for similar IS domains and appropriate mutations in the case of non-oncogenic—but still pathogenic- retroviruses such as HIV is therefore of interest to tentatively develop improved vaccines.

Actually, in the case of the other major class of retroviruses, namely the lentiviruses (among which the human HIV1 and HIV2, and the SIV simian homologues), the crystallographic structure of the envelope protein discloses some similarities but also very important differences, especially in the domain corresponding to the IS domain of oncoretroviruses, with evidence in HIV and SIV for a severely extended helix-loop-helix domain, and no evidence for sequence similarities with the IS domain of oncoretroviruses. Furthermore, an IS domain with strong amino-acid similarities with the IS domain of oncoretroviruses was identified within an accessory protein of HIV and SIV, namely within the Nef protein. This protein is specifically produced by these complex retroviruses and is not encoded by oncoretroviruses. The Nef protein is essential for viremia in vivo, and it possesses several domains responsible for immune escape, among which the identified IS domain, thus strongly suggesting that lentiviruses had transferred the

immunosuppressive activity found in oncoretroviruses within their envelope protein, to the accessory Nef protein. In agreement with this hypothesis, Nef-deleted or Nef-mutated retroviruses have a severely attenuated pathogenicity in *in vivo* macaque animal models. Here, the Inventors report on the identification of an IS activity carried by the HIV and SIV envelope proteins, and on specific mutations that inhibit this activity without major disruption of the overall structure of the corresponding envelope proteins.

Results

Delineation of the immunosuppressive domain of the HIV Envelope

Delineation of the immunosuppressive domain of the HIV envelope was achieved using an *in vivo* tumor rejection assay, that the Inventors had previously used to demonstrate the immunosuppressive activity of the Env protein of oncoretroviruses (murine MoMLV and simian MPMV). The rationale of the assay can be summarized as follows: while injection of MCA205 tumor cells (H-2^b) into allogeneic Balb/c mice (H-2^d) leads to the formation of no tumor or transient tumors that are rapidly rejected, injection of the same cells, but stably expressing an immunosuppressive retroviral Env protein, leads to the growth of larger tumors that persist for a longer time –in spite of the expression of the new exogenous antigen. This difference is not associated with a difference in intrinsic cell growth rate since it is not observed in syngeneic C57BL/6 mice, and is immune system-dependent. The extent of "immunosuppression" can be quantified by an index based on tumor size: $(A_{env} - A_{none})/A_{none}$, where A_{env} and A_{none} are the mean areas at the peak of growth of tumors from Balb/c mice injected with *env*-expressing or control cells, respectively. A positive index indicates that *env* expression facilitates tumor growth, as a consequence of its immunosuppressive activity; a null or negative index points to no effect or even an inhibitory effect, respectively. The latter may be explained by a stimulation of the immune response of the host against the new foreign antigen, represented by a non-immunosuppressive Env protein, expressed at the surface of tumor cells.

To first delineate a minimal immunosuppressive domain (ISD) active *in vivo*, the Inventors analyzed the effect of a series of truncations/deletions within the HIV *env* gene. Assay of the series of C-term truncations in Fig. 1&2 identified a 49 residue-long domain with IS activity, with the 6 aa shorter HIV43 fragment and subsequent C-term truncated HIV37 and HIV30 fragments being IS-negative. HIV49 is embedded into the so-called

ectodomain, which corresponds to a soluble part of the extracellular domain of the TM subunit, and consists in the α -helical domain involved in HIV TM trimerization, and the N-term part of the loop containing the 2 well-conserved, 6 aa-distant, cysteine residues found in most retroviral envelopes. Refine delineation of the amino acids responsible for IS activity was then performed by arginine-scanning between aa 36 and 51, i.e. in the domain associated with the transition from IS positive to negative Env subdomains. As illustrated in Fig. 3, all the X-to-Arg substitutions resulted in no significant change in the IS activity of HIV115, with one exception for Y41 and L42, which resulted in loss of IS activity of the mutant peptide.

Mutations at position 41

The Inventors then checked that the Y41R substitution did not alter the overall capacity of the HIV envelope to be expressed by an eucaryotic cell and to be exported at the cell membrane, by introducing the Y41R substitution into an expression vector for the HIV envelope. A FACS analysis of cell transfected with both the wild-type and the Y41R mutant using an anti-SU specific monoclonal antibody actually demonstrated quantitative expression of the mutant envelope at the cell surface, thus indicating that the Y41R mutation does not significantly alter the HIV Env structure and SU-TM interaction (figure 4).

A series of distinct substitutions at the Y41 position were finally performed to tentatively identify whether other amino acids could be substituted to generate IS-negative variants: the amino acid assayed included the positively charged K and H (in addition to R), the small A and G, and the hydrophobic L and F residues. Again it was checked that these substitutions did not alter the overall structure of the HIV Env with the corresponding mutations.

Among them, substitution of Y41 by A, G, L and F resulted in loss of IS (figure 3), and did not alter the overall structure of the HIV Env with the corresponding mutations (figure 4 and 5).

Further mutation outside of the ISU domain

The effects of a set of second mutations at positions that are structurally close to Y41 position within the ENV ectodomain three-dimensional structure were also tested. Two of them (Y41R-K72A and Y41R-K78G) maintain a high level expression at cell membrane and confer infectivity (figure 5).

Mutations at position 42

Interestingly, the adjacent position to the position Y41, position L42, also resulted in loss of IS activity of the mutant peptide when the mutation L42R is introduced. This mutation maintains a high level expression at cell membrane of the mutated HIV ENV protein (figure 4).

Mutations within the SIV ENV protein

Interestingly, mutation at the homologous position in SIV ENV (L42) of the position L42 in HIV-1 ENV ectodomain also resulted in specific loss of IS activities (Figure 6).

Accordingly, the present investigation has clearly identified a definite location and a definite substitution(s) within the HIV env resulting in the loss of its IS activity. Being compatible with the conservation of the overall structure of the human and simian lentiviral Env proteins, these substitutions should be introduced in all pharmaceutical preparations which include the Env protein as a vaccine antigen.

Materials and Methods

Mice and cell lines: C57Bl/6 and Balb/c mice, 6-10 weeks old, were obtained from CER Janvier (Laval, France). Mice were maintained in the animal facility of the Gustave Roussy Institute in accordance with institutional regulations. 293T (ATCC CRL11268), and MCA205 cells were cultured in DMEM supplemented with 10% fetal calf serum (Invitrogen), streptomycin (100 µg/ml) and penicillin (100 units/ml).

Plasmid construction: The PCEL/E160 encoding the envelope protein of the BRU/LAI HIV-1 isolate is a gift from Dr Marc Sitbon. To generate the pDFG plasmids encoding the various fragments of HIV-1 envelope ectodomain, PCR fragments generated using PCEL/E160 as a template and primer pairs 1-2 (pDFG-HIV115), 1-3 (pDFG-HIV81), 1-4 (pDFG-HIV67), 1-5 (pDFG-HIV55), 1-6 (pDFG-HIV49), 1-7 (pDFG-HIV43), 1-8 (pDFG-HIV37), 1-9 (pDFG-HIV30) were digested with SfiI and MluI and inserted into pDFG-ectoSyncytin-1 (Mangency et al, 2007) opened with the same enzymes.

Mutated pDFG-HIV115 were obtained by successive PCR using appropriate primers. A first series of PCR was performed with pDFG HIV115 as template using primers 1-11 and 10-2, 1-13 and 12-2, 1-15 and 14-2 or 1-17 and 16-2 to introduce the E39R, Y41R, K43R or D44R mutations respectively, and primers 1-18 and 19-2, 1-20 and 21-2, 1-22 and 23-2,

1-24 and 25-2, 1-26 and 27-2, 1-28 and 29-2, 1-30 and 31-2, 1-32 and 33-2, 1-34 and 35-2, or 1-36 and 37-2 to introduce the L36R, A37R, V38R, L42R, Q45R, Q46R, L47R, L48R, G49R, or I50R mutations respectively. The PCR products were then used as templates in subsequent PCR using primers 1-2. These PCR fragments were digested with SfiI and MluI and inserted into pDFG-ectoSyncytin-1 (Mangeny et al, 2007) opened with the same enzymes.

All the constructions were sequenced before use.

Table 1. Primer list

N°	Name	Primer sequence (5'-3')	SEQ ID
1	TM HIV Sfi-Sens	ACATggcccagccggccTCTGGTATAGTGCAGCAGC	SEQ ID NO: 295
2	TM HIV115 Mlu-AS	GTATacgcgtTTATAATTCTGTTCATTCTTTTC	SEQ ID NO: 296
3	TM HIV81 Mlu AS	GTATacgcgtTTACATGTTATTCCAAATCTGTTCC	SEQ ID NO: 297
4	TM HIV67 Mlu AS	GTATacgcgtTTAAGCATTCCAAGGCACAGC	SEQ ID NO: 298
5	TM HIV55 Mlu AS	GTATacgcgtTTATCCAGAGCAACCCCAAATCC	SEQ ID NO: 299
6	TM HIV49 Mlu AS	GTATacgcgtTTACCCCAGGAGCTGTTGATCC	SEQ ID NO: 300
7	TM HIV43 Mlu AS	GTATacgcgtTTACTTTAGGTATCTTTCCACAGC	SEQ ID NO: 301
8	TM HIV37 Mlu AS	GTATacgcgtTTAAGCCAGGATTCTTGCCCTGGAG	SEQ ID NO: 302
9	TM HIV30 Mlu AS	GTATacgcgtTTACTGCTTGATGCCCCAGAC	SEQ ID NO: 303
18	TM HIV115 L36R as	CTTTCCACAGCCcGGATTCTTGCCCTG	SEQ ID NO: 304
19	TM HIV115 L36R s	CAGGCAAGAATCCgGGCTGTGGAAAG	SEQ ID NO: 305
20	TM HIV115 A37R as	GTATCTTTCCACAcgCAGGATTCTTGCC	SEQ ID NO: 306
21	TM HIV115 A37R s	GGCAAGAATCCTGcgTGTGGAAAGATAC	SEQ ID NO: 307
22	TM HIV115 V38R as	CTTTAGGTATCTTTCCctAGCCAGGATTCTTGCC	SEQ ID NO: 308
23	TM HIV115 V38R s	GGCAAGAATCCTGGCTagGGAAAGATACCTAAAG	SEQ ID NO: 309
11	TM HIV115 E39R AS	CCTTTAGGTATCTTctCACAGCCAGGATTC	SEQ ID NO: 310
10	TM HIV115 E39R S	GAATCCTGGCTGTGagAAGATACCTAAAGG	SEQ ID NO: 311
13	TM HIV115 Y41R AS	GTTGATCCTTTAGGcgTCTTTCCACAGCCAG	SEQ ID NO: 312
12	TM HIV115 Y41R S	CTGGCTGTGGAAAGAcgCCTAAAGGATCAAC	SEQ ID NO: 313
24	TM HIV115 L42R as	GGAGCTGTTGATCCTTTcGGTATCTTTCCACAGCC	SEQ ID NO: 314
25	TM HIV115 L42R s	GGCTGTGGAAAGATACCGAAAGGATCAACAGCTCC	SEQ ID NO: 315
15	TM HIV115 K43R AS	GAGCTGTTGATCCcTTAGGTATCTTTCCAC	SEQ ID NO: 316
14	TM HIV115 K43R S	GTGGAAAGATACCTAAgGGATCAACAGCTC	SEQ ID NO: 317
17	TM HIV115 D44R AS	AGGAGCTGTTGAcgCTTTAGGTATCTTT	SEQ ID NO: 318
16	TM HIV115 D44R S	AAAGATACCTAAAGcgTCAACAGCTCCT	SEQ ID NO: 319
26	TM HIV115 Q45R as	CCCAGGAGCTGTcGATCCTTTAGGTATC	SEQ ID NO: 320
27	TM HIV115 Q45R s	GATACCTAAAGGATCgACAGCTCCTGGG	SEQ ID NO: 321

N°	Name	Primer sequence (5'-3')	SEQ ID
28	TMHIV115 Q46R as	CAAATCCCCAGGAGCcGTTGATCCTTTAG	SEQ ID NO: 322
29	TMHIV115 Q46R s	CTAAAGGATCAACgGCTCCTGGGGATTTG	SEQ ID NO: 323
30	TMHIV115 L47R as	CAAATCCCCAGGcGCTGTTGATCCTTTAG	SEQ ID NO: 324
31	TMHIV115 L47R s	CTAAAGGATCAACAGCgCCTGGGGATTTG	SEQ ID NO: 325
32	TMHIV115 L48R as	CCCAAATCCCCcGGAGCTGTTG	SEQ ID NO: 326
33	TMHIV115 L48R s	CAACAGCTCCgGGGGATTTGGG	SEQ ID NO: 327
34	TMHIV115 G49R as	CCCAAATCCcCAGGAGCTGTTG	SEQ ID NO: 328
35	TMHIV115 G49R s	CAACAGCTCCTGcGGATTTGGGG	SEQ ID NO: 329
36	TMHIV115 I50R as	GCAACCCCAtcTCCCCAGGAGCTG	SEQ ID NO: 330
37	TMHIV115 I50R s	CAGCTCCTGGGGAgTGGGGTTGC	SEQ ID NO: 331
38	TMHIV115 W51R as	GCAACCCCAtcTCCCCAGGAGCTG	SEQ ID NO: 332
39	TMHIV115 W51R s	CAGCTCCTGGGGAgTGGGGTTGC	SEQ ID NO: 333

HIV115 Y41 mutant construction: To explore the amino acid sequence necessary for the loss of immunosuppression, HIV115 Y41 mutants were produced by PCR using the pDFG HIV115 WT plasmid as template and pairs of primers 1-42 and 43-2, 1-44 and 45-2, 1-46 and 47-2, 1-48 and 49-2, 1-50 and 51-2, or 1-52 and 53-2, to introduce the Y41K, Y41H, Y41A, Y41G, Y41F, or Y41L mutations respectively. The PCR products were then used as templates in subsequent PCR using primers 1-2. These PCR fragments were digested with SfiI and MluI and inserted into pDFG-ectoSyncytin-1 (Mangeny et al, 2007) opened with the same enzymes

All the constructions were sequenced before use.

N°	Name	Primer sequence (5'-3')	SEQ ID NO
42	TMHIV115 Y41K as	GTTGATCCTTTAGtTfTCTTTCCACAGCCAG	SEQ ID NO: 334
43	TMHIV115 Y41K s	CTGGCTGTGGAAAGAAaAaCTAAAGGATCAAC	SEQ ID NO: 335
44	TMHIV115 Y41H as	GTTGATCCTTTAGGTgTCTTTCCACAGCCAG	SEQ ID NO: 336
45	TMHIV115 Y41H s	CTGGCTGTGGAAAGAcACCTAAAGGATCAAC	SEQ ID NO: 337
46	TMHIV115 Y41A as	GTTGATCCTTTAGGgcTCTTTCCACAGCCAG	SEQ ID NO: 338
47	TMHIV115 Y41A s	CTGGCTGTGGAAAGAgcCCTAAAGGATCAAC	SEQ ID NO: 339
48	TMHIV115 Y41G as	GTTGATCCTTTAGGccTCTTTCCACAGCCAG	SEQ ID NO: 340
49	TMHIV115 Y41G s	CTGGCTGTGGAAAGAggCCTAAAGGATCAAC	SEQ ID NO: 341
50	TMHIV115 Y41F as	GTTGATCCTTTAGGaATCTTTCCACAGCCAG	SEQ ID NO: 342

51	TMHIV115 Y41F s	CTGGCTGTGGAAAGATcCCTAAAGGATCAAC	SEQ ID NO: 343
52	TMHIV115 Y41L as	GTTGATCCTTTAGGagTCTTTCCACAGCCAG	SEQ ID NO: 344
53	TMHIV115 Y41L s	CTGGCTGTGGAAAGAcCCTAAAGGATCAAC	SEQ ID NO: 345

HIV115 Y41 double mutants construction: HIV115 Y41R K72A, G or S double mutants were produced by PCR using the pDFG HIV115 Y41R plasmid as template and pairs of primers 1-94 and 93-2, 1-96 and 95-2, or 1-98 and 97-2. The PCR products were then used as templates in subsequent PCR using primers 1-2. These PCR fragments were digested with SfiI and MluI and inserted into pDFG-ectoSyncytin-1 (Mangeny et al, 2007) opened with the same enzymes. HIV115 R40X Y41R double-mutants were produced by PCR using the pDFG HIV115 Y41R as template and pairs of primers 1-54 and 55-2, 1-56 and 57-2 to introduce the R40A and R40E mutations respectively. HIV115 R40X Y41A double-mutants were produced by PCR using the pDFG HIV115 Y41A as template and pairs of primers 1-58 and 59-2, 1-60 and 61-2, 1-62 and 63-2, 1-64 and 65-2 to introduce the R40A, R40E, R40S, R40T mutations respectively. HIV115 R40X simple mutants were also generated using pDFG HIV115 WT as template and pairs of primers 1-66 and 67-2, 1-68 and 69-2, 1-70 and 71-2, 1-72 and 73-2.

The PCR products were then used as templates in subsequent PCR using primers 1-2. These PCR fragments were digested with SfiI and MluI and inserted into pDFG-ectoSyncytin-1 (Mangeny et al, 2007) opened with the same enzyme.

N°	Name	Primer sequence (5'-3')	SEQ ID
54	TM HIV115 R40AY41R- S	CTGGCTGTGGAAgcAcgCCTAAAGGATCAAC	SEQ ID NO: 378
55	TM HIV115 R40AY41R- AS	GTTGATCCTTTAGGcgTgcTTCCACAGCCAG	SEQ ID NO: 379
56	TM HIV115 R40EY41R- S	CTGGCTGTGGAAgaAcgCCTAAAGGATCAAC	SEQ ID NO: 380

N°	Name	Primer sequence (5'-3')	SEQ ID
57	TM HIV115 R40EY41R- AS	GTTGATCCTTTAGGcgTtcTTCCACAGCCAG	SEQ ID NO: 381
58	TM HIV115 R40AY41A- S	CTGGCTGTGGAAgctgcCCTAAAGGATCAAC	SEQ ID NO: 382
59	TM HIV115 R40AY41A- AS	GTTGATCCTTTAGGgcagcTTCCACAGCCAG	SEQ ID NO: 383
60	TM HIV115 R40EY41A- S	CTGGCTGTGGAAgaagcCCTAAAGGATCAAC	SEQ ID NO: 384
61	TM HIV115 R40EY41A- AS	GTTGATCCTTTAGGgcttcTTCCACAGCCAG	SEQ ID NO: 385
62	TM HIV115 R40SY41A- S	CTGGCTGTGGAAagcgcCCTAAAGGATCAAC	SEQ ID NO: 386
63	TM HIV115 R40SY41A- AS	GTTGATCCTTTAGGgcgctTTCCACAGCCAG	SEQ ID NO: 387
64	TM HIV115 R40TY41A- S	CTGGCTGTGGAAacagcCCTAAAGGATCAAC	SEQ ID NO: 388
65	TM HIV115 R40TY41A- AS	GTTGATCCTTTAGGgctgfTTCCACAGCCAG	SEQ ID NO: 389
66	TM HIV115 R40A-S	CTGGCTGTGGAAgctTACCTAAAGGATCAAC	SEQ ID NO: 390
67	TM HIV115 R40A-AS	GTTGATCCTTTAGGTAagcTTCCACAGCCAG	SEQ ID NO: 391

N°	Name	Primer sequence (5'-3')	SEQ ID
68	TM HIV115 R40E-S	CTGGCTGTGGAAgaaTACCTAAAGGATCAAC	SEQ ID NO: 392
69	TM HIV115 R40E-AS	GTTGATCCTTTAGGTAttcTTCCACAGCCAG	SEQ ID NO: 393
70	TM HIV115 R40S-S	CTGGCTGTGGAAagcTACCTAAAGGATCAAC	SEQ ID NO: 394
71	TM HIV115 R40S-AS	GTTGATCCTTTAGGTAgctTTCCACAGCCAG	SEQ ID NO: 395
72	TM HIV115 R40T-S	CTGGCTGTGGAAacaTACCTAAAGGATCAAC	SEQ ID NO: 396
73	TM HIV115 R40T-AS	GTTGATCCTTTAGGTAtgtTTCCACAGCCAG	SEQ ID NO: 397
93	TMHIV115 K72A s	GCTAGTTGGAGTAATgcATCTCTGGAACAGATTTGG	SEQ ID NO: 398
94	TMHIV115 K72A as	CCAAATCTGTTCCAGAGATgcATTACTCCAACACTAGC	SEQ ID NO: 399
95	TMHIV115 K72G s	GCTAGTTGGAGTAATggATCTCTGGAACAGATTTGG	SEQ ID NO: 400
96	TMHIV115 K72G as	CCAAATCTGTTCCAGAGATccATTACTCCAACACTAGC	SEQ ID NO: 401
97	TMHIV115 K72S s	GCTAGTTGGAGTAATtcATCTCTGGAACAGATTTGG	SEQ ID NO: 402
98	TMHIV115 K72S as	CCAAATCTGTTCCAGAGATgaATTACTCCAACACTAGC	SEQ ID NO: 403

Introduction of HIV115 mutations into an HIV Env expression vector: To introduce these HIV envelope mutations into the HIVenv pTr712 expression vector (Schnierle et al PNAS 1997), a silent mutation was first generated to create an SfiI insertion site. PCR fragments were generated using pTr712as a template and primer pairs 38-39 and 40-41. The PCR products were then used as templates in subsequent PCR using primers 38-41.

The resulting PCR fragment was then digested with BsaBI and HindIII and inserted into the pTr712 plasmid opened with the same enzymes resulting in the pTr712-Sfi plasmid.

PCR fragments of each HIV115 mutation were then generated by using primers 40-41 and the pDFG HIV115 mutant plasmids as templates, digestion with SfiI and HindIII and then insertion into the pTr712-Sfi plasmid opened with the same enzymes.

All the constructions were sequenced before use.

N°	Name	Primer sequence (5'-3')	SEQ ID NO
38	HIV BH10 BsaBI s	acaaattagatgttcatcaaatattacaggg	SEQ ID NO: 346
39	HIV BH10 SfiI mutsil as	GCAACAGATGCTGTTGgGCCTCA ATgGCCCTCAGCAAATTGTTC	SEQ ID NO: 347
40	HIV BH10 SfiI mutsil s	GAACAATTTGCTGAGGGCcATTG AGGCcCAACAGCATCTGTTGC	SEQ ID NO: 348
41	HIV BH10 HindIII as	gtgtattaagcttgtgtaattgtaattctc	SEQ ID NO: 349
74	EnvHIVTr712-Xho- Age-Kozak-S	NNNNNCTCGAGACCGGTccaactaga accATGAGAGTGAAGGAGAAATA TCAGC	SEQ ID NO : 404
75	EnvHIVTr712-Mlu-AS	NNNNNACGCGTTCAATATCCCTG CCTAACTC	SEQ ID NO : 405
76	EnvHIVWT-Mlu-AS	NNNNNACGCGTTTATAGCAAAAT CCTTTCCAAGC	SEQ ID NO : 406

Establishment of *env*-expressing tumor cells and MCA205 tumor-rejection assay: 7.5×10^5 293T cells were cotransfected with the *env*-expressing pDFG retroviral vector to be tested (1.75 μ g) and expression vectors for the MLV proteins (0.55 μ g for the amphotropic MLV *env* vector and 1.75 μ g for the MLV *gag* and *pol* vector). 36 hours post-transfection, supernatants were harvested for infection of MCA205 tumor cells (2.5 ml of supernatant per 5×10^5 cells with 8 μ g/ml polybrene). Cells were maintained in selective medium (400 units/ml hygromycin) for 3 weeks, and then washed with PBS, trypsinized and inoculated subcutaneously in the shaved area of each mouse right flank as in Mangeney et al (1998, 2007). Tumor growth was monitored by palpation twice or thrice weekly and tumor area (mm^2) determined by measuring perpendicular tumor diameters. The extent of

"immunosuppression" was quantified by an index based on tumor size: $(A_{env} - A_{none}) / A_{none}$, where A_{env} and A_{none} are the mean areas at the peak of growth of tumors from Balb/c mice injected with *env*-expressing or control cells, respectively.

Analysis of HIVenv expression : 293T cells were transfected with 4mg of the expression vector for the HIV-1 envelope (pTr712) either wild-type or mutated at the indicated positions, by Calcium Phosphate precipitation. Cells are washed 16h later and then harvested 2 days post-transfection using PBS-EDTA 5mM. The 110-H monoclonal antibody (anti-V3 loop, gift from Hybridolab, Pasteur Institute) was used (1/200 dilution) to stain the HIV envelope. As a secondary antibody, the Inventors used the goat anti mouse IgG Alexa 488 (1/400) (Invitrogen). For intracellular HIVenv staining, 293T cells were fixed with a formaldehyde buffer and then permeabilized (BD cytofix /cytoperm, BD Biosciences). The isotype mouse anti IgG1Kappa (BD Biosciences) was used to control non-specific staining. Fluorescence was acquired by flow cytometry using a FACS Calibur (BD Biosciences), and data analysed by the CellQuest software (BD Biosciences).

Mutated Env pseudotyping and measure of viral titer: 293T cells are triple transfected with 3mg of a reporter MLV vector carrying GFP (CNCG), 1.75mg Mo-MLV gag-pol vector and 0.55mg phCMV vector encoding HIV-1 envelope wt or mutated at the indicated positions. The infectivity of Mo-MLV virions pseudotyped with HIV-1 Env, harvested 48 hours post-transfection, is measured using U87 cells (CD4⁺, CXCR4⁺) as target cells. The infectivity of the envelopes is analysed after 72h exposure diluted 0.45mm-filtered supernatant in presence of 4mg/mL polybrene. The fluorescence (GFP) is acquired by flow cytometry using a FACS Calibur (BD Biosciences). The results are analysed by the CellQuest software (BD Biosciences). The resulting titers (number of infected cells/mL) are calculated as the following : $(\% \text{ GFP}^+ \text{ cells (infected)} \times \text{plated cell number}) / \text{volume of supernatant} \times 1000$.

Env SIV mutants construction : PCR fragments were generated using p239 SPE3' (the plasmid encoding the SIV half virus containing the envelope protein) as a template and primer pairs 77-79 and 78-80, 77-81 and 78-82, 77-83 and 78-84, 77-85 and 78-86, 77-87 and 78-88, 77-89 and 78-90 to introduce the E39R, K40R, Y41R, L42R, K43R, D44R

mutations respectively. The PCR products were the used as templates in subsequent PCR using primers 77-78. The resulting PCR fragment was then digested with BmgBI and NheI and inserted into the p239 SPE3' plasmid opened with the same enzymes.

All the constructions were sequenced before use.

Introduction of SIV mutants ectodomain into pDFG : To generate the pDFG plasmids encoding the fragment of SIV envelope ectodomain-55, PCR fragments generated using p239 SPE3' WT and mutants as a template and primer pairs 91-92, were digested with SfiI and MluI and inserted into pDFG opened with the same enzymes.

77	Env SIV S	ccgctcagtcgccgaactttattggc	SEQ ID NO: 350
78	Env SIV AS	ggtggggaagagaacactggcc	SEQ ID NO: 351
79	SIV env E39R s	cagactagggtcactgccatcCGCaagtacttaaaggaccaggcgcg	SEQ ID NO: 352
80	SIV env E39R as	cgcttggtcctttaagtacttGCGgatggcagtgaccctagtctg	SEQ ID NO: 353
81	SIV env K40R s	actagggtcactgccatcgagCGCtacttaaaggaccaggcgcag	SEQ ID NO: 354
82	SIV env K40R as	ctgcgcttggtcctttaagtaGCGctcgatggcagtgaccctagt	SEQ ID NO: 355
83	SIV env Y41R s	GCCATCGAGAAGcgCTTAAAGGACCAGGCG	SEQ ID NO: 356
84	SIV env Y41R as	CGCCTGGTCCTTTAAGcgCTTCTCGATGGC	SEQ ID NO: 357
85	SIV env L42R s	gtcactgccatcgagaagtacCGCaaggaccaggcgcagctg	SEQ ID NO: 358
86	SIV env L42R as	cagctgcgcttggtccttGCGgtacttctcgatggcagtgac	SEQ ID NO: 359
87	SIV env K43R s	gccatcgagaagtacttaCGCgaccaggcgcagctgaatgcttgg	SEQ ID NO: 360
88	SIV env K43R as	ccaagcattcagctgcgcttggtcGCGtaagtacttctcgatggc	SEQ ID NO: 361
89	SIV env D44R s	gagaagtacttaaagCGCcaggcgcagctgaatgcttgg	SEQ ID NO: 362
90	SIV env D44R as	attcagctgcgctgGCGctttaagtacttctcgatggc	SEQ ID NO: 363
91	TMSIV55 Sfi S	ACATggcccagccggccgctgggatagtgcagcaac	SEQ ID NO: 364
92	TMSIV55 Mlu AS	GTATacgcgtTTAaaacgcacatccccaagcattc	SEQ ID NO: 365

Comparative example: test of the *in vivo* effect of the mutation G49R, which corresponds to the mutation “G19R” in the international application WO 2010/022,740.

WO 2010/022,740 discloses a consensus sequence of 50 amino acid of the HIV ENV protein. In this sequence, it is suggested that substitution of amino acids in positions 10, 19, 24, 34 and 40 affect the immunosuppressive properties of the HIV ENV protein.

These mutations are a transposition in lentivirus of the teaching of WO 2005/095,442 limited to endogenous or onco retroviruses. The authors of WO 2005/095,442 are also the

authors of the present application and they early observed that such a transposition is not effective.

Despite the fact that any amino acids can be assigned to the positions 10, 19, 24, 34 or 40 in the consensus sequence of WO 2010/022,740, only one substitution (one residue for one position) was tested by *ex vivo* experiments in WO 2010/022,740. It is the mutation “G19R”, which corresponds to the mutation G49R in the present figures 1 and 3.

Because the immune response of an individual involves different organs and different cellular and non-cellular components, *ex vivo* results have no predictive value concerning the *in vivo* immunosuppressive properties of a viral protein.

To determine if the mutations previously disclosed in WO 2010/022,740 are suitable for a medical use, the mutation G49R has been tested using the MCA205 tumor rejection *in vivo* assay, as defined in the section “*Establishment of env-expressing tumor cells and MCA205 tumor-rejection assay*” of the Example.

As shown in Figure 3, the peptide G49R remains almost as immunosuppressive *in vivo* as the wild type HIV 115 peptide, with an immunosuppression index which is higher than 0,5. Thus, the mutation G49R does not induce a significant decrease of the *in vivo* immunosuppressive properties of the HIV ENV protein, with a ratio of immunosuppression index of G49R mutated ENV HIV 115 peptide ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV HIV 115 peptide ($i_{\text{wild type env}}$), which is 0,7 (*i.e.* higher than 0,5).

This result demonstrates the insufficiently described teaching of WO 2010/022740 since the only one mutation tested in WO 2010/022,740, using an *ex vivo* test, does not significantly affect the *in vivo* immunosuppressive properties of the HIV ENV protein.

As a consequence, WO 2010/022,740 raises the same technical problem as the present invention but does not offer a technical solution.

CLAIMS

1. Pharmaceutical composition comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein having at least 90% identity, to one sequence chosen from the group consisting of SEQ ID NO : 216, SEQ ID NO : 420 and SEQ ID NO : 421,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

in association with a pharmaceutically acceptable carrier,

said substantial absence of immunosuppressive activity of the above mentioned mutated human or simian lentiviral ENV protein or of the above defined fragment being liable to be assessed by the fact that in an in vivo assay involving engrafted tumor cells rejection, said tumor cells being transduced either so as to express said mutated ENV protein or said fragment (mutated ENV tumor cells),

or said tumor cells being transduced so as to express said wild type ENV protein or a fragment thereof (wild type ENV tumor cells),

or said tumor cells being not transduced (normal tumor cells),

the following ratio :

immunosuppression index of said mutated ENV protein or of said fragment ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV protein ($i_{\text{wild type env}}$) is less than 0.5,

$i_{\text{mutated env}}$ being defined by : (maximum area reached by mutated ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells), and

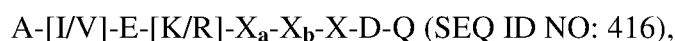
$i_{\text{wild type env}}$ being defined by : (maximum area reached by wild type ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells).

2. Pharmaceutical composition according to claim 1 comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:



wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

in association with a pharmaceutically acceptable carrier.

3. Pharmaceutical composition according to claim 1 comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,
or

X_a is F or L, and X_b is F, G or R,

or

X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids, said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

or

X_a is F or L, and X_b is F, G or R,

or

X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G, R,

in association with a pharmaceutically acceptable carrier.

4. Pharmaceutical composition according to any one of claims 1 to 3 comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

or

X_a is F or L, and X_b is F, G or R,

or

X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

or

X_a is F or L, and X_b is F, G or R,

or

X_a is A, G or R, and X_b is A,

or

X_a is A, G or R, and X_b is F, G or R,

in association with a pharmaceutically acceptable carrier.

5. Pharmaceutical composition according to claim 1 or claim 3, comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R,

in association with a pharmaceutically acceptable carrier.

6. Pharmaceutical composition according to any one of claims 1 to 5 comprising as active substance :

a) an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein resulting from mutation of the transmembrane (TM) subunit of a wild type human or simian lentiviral ENV protein,

said mutated human or simian lentiviral ENV protein comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

or

b) at least one fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said fragment comprising at least 40 amino acids,

said fragment comprising a mutated immunosuppressive domain (ISU) containing the following amino acid sequence:

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P,

or

X_a is Y, I, H, C or T, and X_b is F, G or R,

in association with a pharmaceutically acceptable carrier.

7. Pharmaceutical composition according to any one of claims 1, 3 or 5,

wherein

X_a is R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is R, or

X_a is R, and X_b is R.

8. Pharmaceutical composition according to any one of claims 1 to 7, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the following amino acid sequences :

A-I-E-K-X_a-X_b-X-DQ (SEQ ID NO: 422),

A-I-E-R-X_a-X_b-X-DQ (SEQ ID NO: 423),

A-V-E-K-X_a-X_b-X-DQ (SEQ ID NO: 424),

A-V-E-R-X_a-X_b-X-DQ (SEQ ID NO: 425).

9. Pharmaceutical composition according to any one of claims 1 to 8, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences :

SEQ ID NO : 13, SEQ ID NO : 42, SEQ ID NO : 71,

SEQ ID NO : 9 to 12,

SEQ ID NO : 14 to 41,

SEQ ID NO : 43 to 70, and
SEQ ID NO : 72 to 95.

- 10.** Pharmaceutical composition according to any one of claims 1 to 8, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences :

SEQ ID NO : 13, SEQ ID NO : 42, SEQ ID NO : 71,

SEQ ID NO : 9, 11, 15 to 21, 23 to 29, 31 to 38, 40, 44 to 50, 52 to 58, 60 to 67, 69, 73 to 79, 81 to 87, 89 to 95.

- 11.** Pharmaceutical composition according to any one of claims 1 to 8, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences : SEQ ID NO : 96 to 211.

- 12.** Pharmaceutical composition according to any one of claims 1 to 8, wherein said isolated mutated human or simian lentiviral ENV protein or said fragment of said isolated mutated human or simian lentiviral ENV protein comprises one of the amino acid sequences : SEQ ID NO : 96, 98, 100, 102 to 108, 110 to 116, 118 to 125, 127, 129, 131 to 137, 139 to 145, 147 to 154, 156, 158, 160 to 166, 168 to 174, 176 to 183, 185, 187, 189 to 195, 197 to 203, 205 to 211.

- 13.** Pharmaceutical composition according to any one of claims 1 to 8, wherein said isolated mutated human or simian lentiviral ENV protein consists of one of the amino acid sequences : SEQ ID NO : 212 to 269.

- 14.** Pharmaceutical composition according to any one of claims 1 to 8, wherein said isolated mutated human or simian lentiviral ENV protein consists of one of the amino acid sequences : SEQ ID NO : 212, 214, 216, 218 to 224, 226 to 232, 234 to 241, 243, 245, 247 to 253, 255 to 261, 263 to 269.

- 15.** Pharmaceutical composition according to any one of claims 1 to 14, wherein said isolated mutated human or simian lentiviral ENV protein comprises an additional mutation in one at least of the amino acids at positions 29, 36 and 37 of SEQ ID NO: 426 :

A[I/V]E[K/R]X_aX_bX₁DQX₂X₃LX₄X₅WGC[A/S][F/G]X₆X₇CVX₈TX₉VPX_cX₁₀Z₁Z₂Z₃Z₄Z₅X_dX_e[S/T] (SEQ ID NO: 426)

wherein

X_a and X_b are as defined in any one of claims 1 to 14,

X₁ to X₁₀ represent any amino acid,

Z₁ to Z₅ represent no amino acid or any amino acid, independently from each other

such that

- X_c is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V or Y, preferably A, D, or N,
- X_d is A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, Y or W, preferably A, G, S or Y,
- X_e is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, T, V, Y or W, preferably A, D or N.

- 16.** Pharmaceutical composition according to any one of claims 1 to 15, wherein said mutated protein consists of one of the amino acid sequences of the group consisting of SEQ ID NO : 271 to 283.

- 17.** Pharmaceutical composition comprising as active substance a nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, as defined in any one of claims 1 to 16.

- 18.** Pharmaceutical composition according to claim 18, wherein said nucleic acid molecule being contained in a vector, said vector comprising means allowing the expression of the mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, as defined in any one of claims 1 to 16.

- 19.** Pharmaceutical composition according to claim 18, wherein said vector being chosen among a measles vector, a canarypox vector, a pox vector, a fowlpox, an adenoviral vector, a lentiviral vector, a Sendai virus, a Cytomegalovirus vector or a Modified Vaccinia Ankara vector.
- 20.** Pharmaceutical composition, according to any one of claims 17 to 19, comprising at least one nucleic acid molecule coding for a GAG protein and/or a PRO protein and/or a POL protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties, of a human or simian lentivirus, said lentivirus being preferably of the same origin as the mutated lentiviral ENV protein.
- 21.** Pharmaceutical composition, according to any one of claims 18 to 20, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains said at least one nucleic acid molecule coding for a GAG protein and/or a PRO protein and/or a POL protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.
- 22.** Pharmaceutical composition, according to claim 20 or claim 21, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains all said at least one nucleic acid molecule coding for a GAG protein and/or a PRO protein and/or a POL protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.
- 23.** Pharmaceutical composition, according to claim 20 or claim 21, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in a vector which is different from the at least one vector containing said at least one nucleic

acid molecule coding for a GAG protein and/or a PRO protein and/or a POL protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

- 24.** Pharmaceutical composition, according to claim 20 or claim 23, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, said nucleic acid molecule coding for a GAG protein, said nucleic acid molecule coding for a PRO protein, said nucleic acid molecule coding for a POL protein and said nucleic acid molecule coding for a mutated NEF protein substantially devoid of immunosuppressive properties, are all contained in vectors which are different from each other.
- 25.** Pharmaceutical composition, according to any one of claims 17 to 20, comprising at least one nucleic acid molecule coding for a GAG protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties,
of a human or simian lentivirus, said lentivirus being preferably of the same origin as the mutated lentiviral ENV protein.
- 26.** Pharmaceutical composition, according to claims 20 or claim 25, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains said at least one nucleic acid molecule coding for a GAG protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.
- 27.** Pharmaceutical composition, according to any one of claims 20, 25 or 26, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains a nucleic acid molecule coding for a GAG protein and for a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.

- 28.** Pharmaceutical composition, according to any one of claims 20, or 25 to 27, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in the same vector as the one which also contains a nucleic acid molecule coding for a GAG protein and for a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties, said vector being a measles vector or a canary pox vector.
- 29.** Pharmaceutical composition, according to claim 20 or claim 25, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, is contained in a vector which is different from the at least one vector containing said at least one nucleic acid molecule coding for a GAG protein and/or a mutated NEF protein, wherein said mutated NEF protein is substantially devoid of immunosuppressive properties.
- 30.** Pharmaceutical composition, according to any one of claims 20, 25 or 29, wherein said nucleic acid molecule coding for a mutated human or simian lentiviral ENV protein, or a fragment of said mutated human or simian lentiviral ENV protein, said nucleic acid molecule coding for a GAG protein and said nucleic acid molecule coding for a mutated NEF protein substantially devoid of immunosuppressive properties, are all contained in vectors which are different from each other.
- 31.** Pharmaceutical composition according to any one of claims 1 to 30, in association with at least one antiviral compound, preferably for a simultaneous, separated or sequential use.
- 32.** Pharmaceutical composition according to any one of claims 1 to 31, for its use for stimulating an immune response in a host organism.

33. Pharmaceutical composition according to any one of claims 1 to 32, for its use for the prevention or the treatment of lentiviral infection, preferably HIV-1 infection, HIV-2 infection or SIV infection.
34. Pharmaceutical composition according to any one of claims 1 to 33, for its use as a vaccine, in particular against HIV-1 infection, HIV-2 infection or SIV infection.
35. A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, consisting of modifying the immunosuppressive property of:
 a wild-type human or simian lentiviral ENV protein,
 or a fragment of said wild-type human or simian lentiviral ENV protein, said fragment comprising at least 40 amino acids,
 said ENV protein or fragment thereof presenting a transmembrane subunit (TM) comprising an immunosuppressive domain (ISU) containing the following amino acid sequence :

$$A-[I/V]-E-[K/R]-X'_a-X'_b-X-D-Q \text{ (SEQ ID NO: 427)},$$

wherein

X represents any amino acid,

X'_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and

X'_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y,

said method comprising a step of introduction of at least one mutation of X'_a and/or X'_b,

to obtain:

an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

said mutated human or simian lentiviral ENV protein having at least 90% identity, to one sequence chosen from the group consisting of SEQ ID NO : 216, SEQ ID NO : 420 and SEQ ID NO : 421,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R,

said substantial absence of immunosuppressive activity of the above mentioned mutated human or simian lentiviral ENV protein or of the above defined fragment being liable to be assessed by the fact that in an in vivo assay involving engrafted tumor cells rejection, said tumor cells being transduced either so as to express said mutated ENV protein or said fragment (mutated ENV tumor cells),

or said tumor cells being transduced so as to express said wild type ENV protein or a fragment thereof (wild type ENV tumor cells),

or said tumor cells being not transduced (normal tumor cells),

the following ratio :

immunosuppression index of said mutated ENV protein or of said fragment ($i_{\text{mutated env}}$) / immunosuppression index of wild type ENV protein ($i_{\text{wild type env}}$) is less than 0.5,

$i_{\text{mutated env}}$ being defined by : (maximum area reached by mutated ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells), and

$i_{\text{wild type env}}$ being defined by : (maximum area reached by wild type ENV tumor cells - maximum area reached by normal tumor cells) / (maximum area reached by normal tumor cells).

36. A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35, consisting of modifying the immunosuppressive property of :

a wild-type human or simian lentiviral ENV protein,

or a fragment of said wild-type human or simian lentiviral ENV protein, said fragment comprising at least 40 amino acids,

said ENV protein or fragment thereof presenting a transmembrane subunit (TM) comprising an immunosuppressive domain (ISU) containing the following amino acid sequence :

A-[I/V]-E-[K/R]-Y-L-X-D-Q (SEQ ID NO : 1),

wherein

X represents any amino acid,

said method comprising a step of introduction of at least one mutation of Y in position 5 and/or L in position 6,

to obtain:

an isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,

or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,

said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R.

- 37.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35,

wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
 or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
 said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, F, G, L or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is A, F, G or R, or

X_a is A, F, G, L or R, and X_b is A, F, G or R.

- 38.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35,
 wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
 or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
 said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R, or

X_a is F or L, and X_b is F, G or R, or

X_a is A, G or R, and X_b is A.

- 39.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35,
 wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
 or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
 said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is F, G or R, or

X_a is F or L, and X_b is F, G or R, or

X_a is A, G or R, and X_b is A, or

X_a is A, G or R, and X_b is F, G or R.

- 40.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35,
 wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity,
 or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids,
 said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]-X_a-X_b-X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is F, G or R.

- 41.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35, wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids, said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is A, G, or R, and X_b is L, I, V, M or P, or

X_a is Y, I, H, C or T, and X_b is F, G or R.

- 42.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to any one of claim 35, wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids, said mutated ENV protein and fragment thereof comprising a mutated immunosuppressive domain (ISU) containing the following amino sequence :

A-[I/V]-E-[K/R]- X_a - X_b -X-D-Q (SEQ ID NO: 416),

wherein,

X represents any amino acid,

and either

X_a is R, and X_b is C, D, E, H, I, K, L, M, N, P, Q, S, T, V, W or Y, or

X_a is C, D, E, H, I, K, M, N, P, Q, S, T, V, W or Y, and X_b is R, or

X_a is R, and X_b is R.

- 43.** A method to obtain the active substance of a pharmaceutical composition, as defined in any one of claims 1 to 31, according to claim 35, wherein said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, or a fragment of said isolated mutated human or simian lentiviral ENV protein having substantially no immunosuppressive activity, said fragment comprising at least 40 amino acids, said mutated ENV protein and fragment thereof comprising an additional mutation in one at least of the amino acids at positions, 29, 36 and 37 of SEQ ID NO: X :

A[I/V]E[K/R] X_a X_b X_1 DQ X_2 X_3 L X_4 X_5 WGC[A/S][F/G] X_6 X_7 CV X_8 TX $_9$ VP X_c X_{10} Z $_1$ Z $_2$ Z $_3$ Z $_4$ Z $_5$ X_d X_e [S/T] (SEQ ID NO: 426)

wherein

X_a and X_b are as defined in any one of claims 1 to 14,

X_1 to X_{10} represent any amino acid,

Z $_1$ to Z $_5$ represent no amino acid or any amino acid, independently from each other

such that

- X_c is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, S, T, V or Y, preferably A, D, or N,
- X_d is A, C, D, E, F, G, H, I, L, M, N, P, Q, R, S, T, V, Y or W, preferably A, G, S or Y,
- X_e is A, C, D, E, F, G, H, I, K, L, M, N, P, Q, R, T, V, Y or W, preferably A, D or N.

- 44.** The active substance when prepared according to method of any one of claims 35 to 43.

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQ **HIV30**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILA **HIV37**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLK **HIV43**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLG **HIV49**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSG **HIV55**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPwana **HIV67**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNM **HIV81**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **HIV115**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 L36R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 A37R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILARERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 V38R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVRRYLKQDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 E39R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 Y41R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 L42R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLRDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 K43R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKQDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 D44R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDRQLLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 Q45R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDRQLLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 Q46R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 L47R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 L48R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLRIWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 G49R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGRWGCSSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNE
 QEL **115 I50R**
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIRGCSGKLLICTTAVPWNASWSNKSLEQIWNNMTWMEWDRREINNYTSLIHSLIEESQOQEKNEQ
 EL **115 W51R**

Figure 1 (first part)

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **HIV 115**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41R**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERHLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41H**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41K**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERGLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41G**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERRLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41L**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERALKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41A**

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERFLKDQQLGIWGCSGKLLICTTAVPWNASWSNKSLEQIWNMMTWMEWDREINNYTSLIHSLEESQNQQEKNEQ
 EL **115 Y41F**

Figure 1 (end)

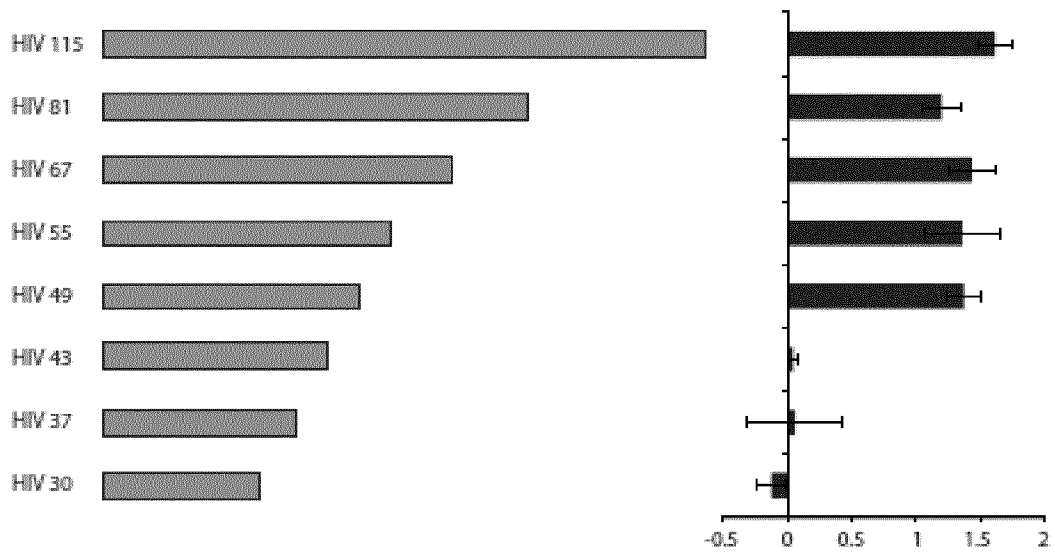


Figure 2

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQ - HIV 30
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILA - HIV 37
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLK - HIV 43
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLG - HIV 49
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC - HIV 55
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC - HIV 67
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC - HIV 81
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC - HIV 81
 EL - HIV 115

SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 L36R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 A37R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 V38R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 E39R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41R K72A
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41R K72G
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41R K72S
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41H
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41K
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41G
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41L
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41A
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Y41F
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 L42R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 K43R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 D44R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Q45R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 Q46R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 L47R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 L48R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 G49R
 SGIVQQNNLLRAIEAQQHLLQLTVWGIKQLQARILAVERYLKDQQLGIWGC...KNEQEL - HIV 115 I50R
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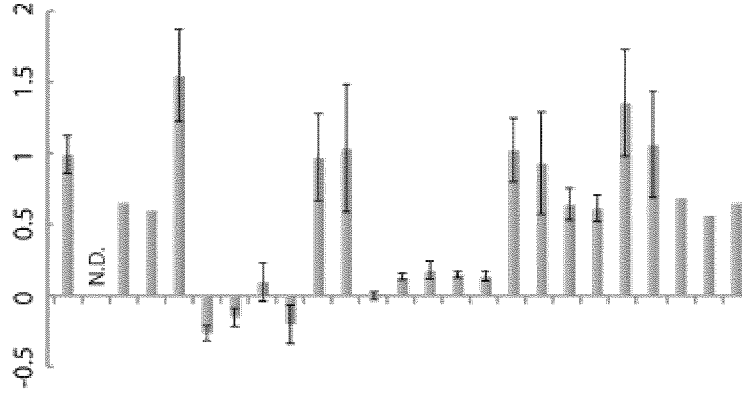


Figure 3

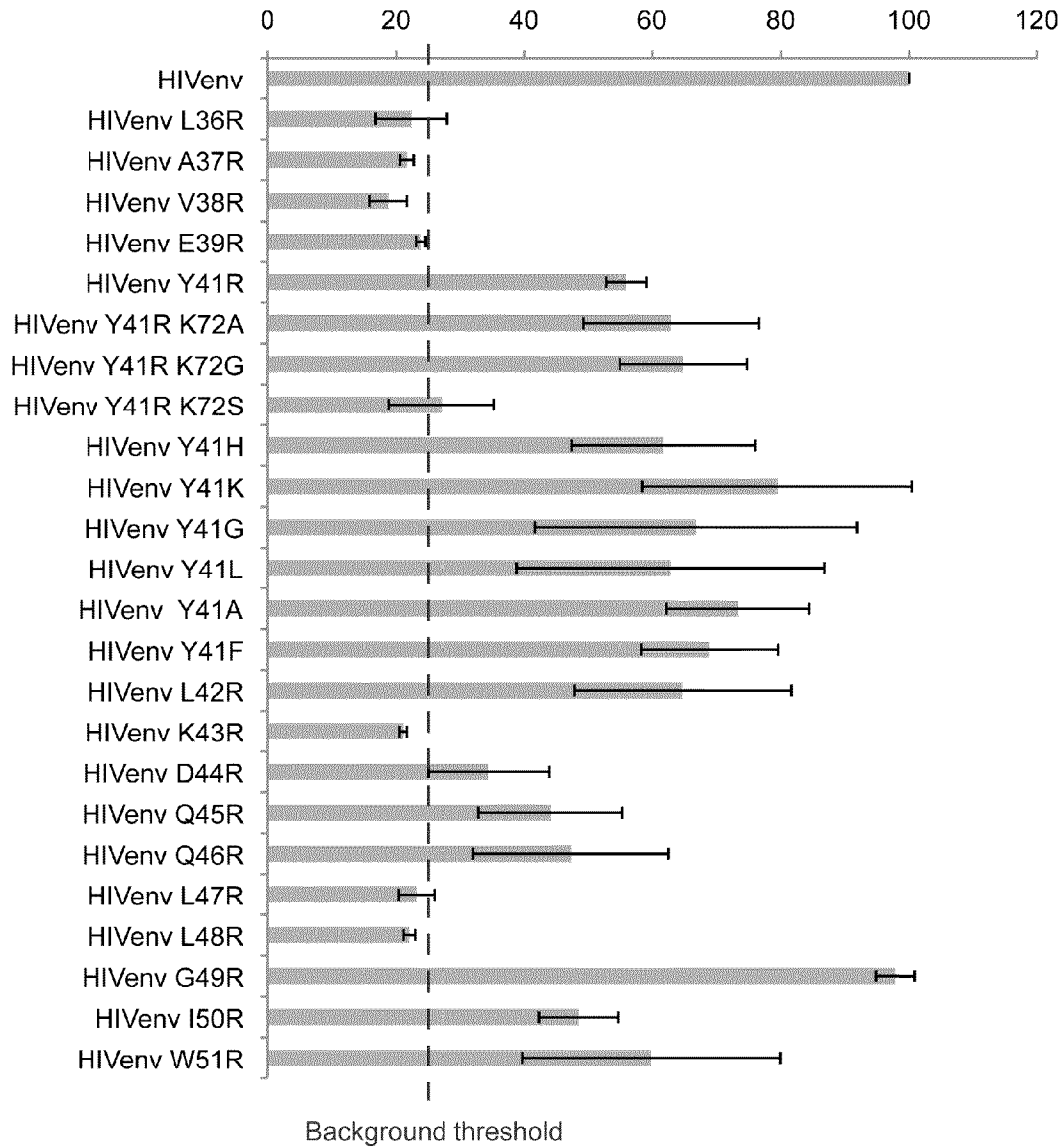


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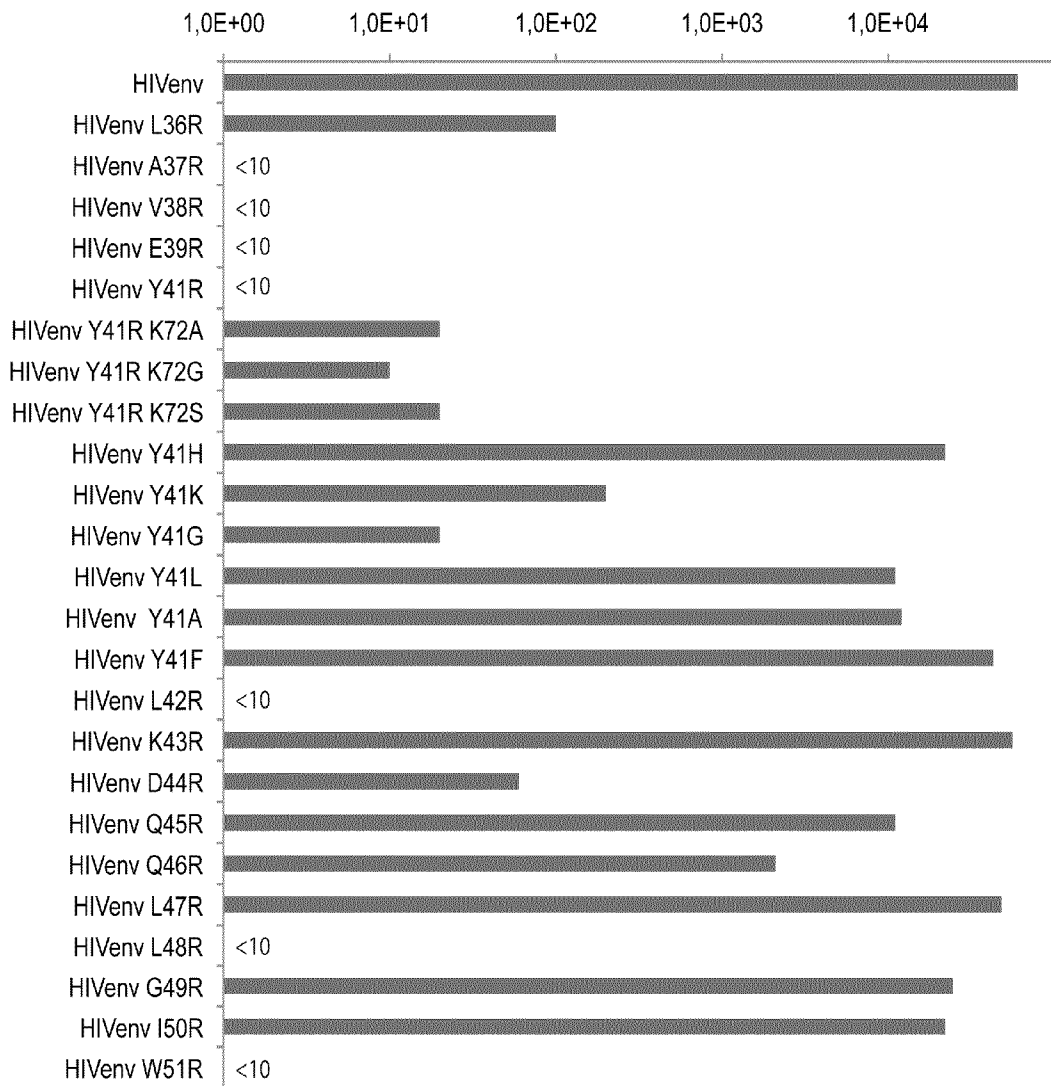
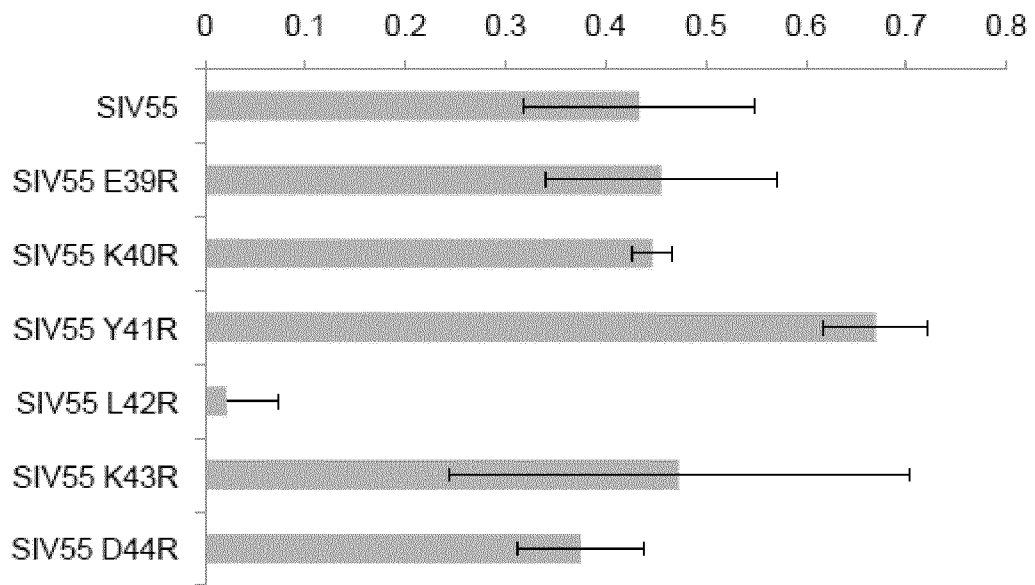


Figure 5

7/7

**Figure 6**

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INSTITUT GUSTAVE ROUSSY
UNIVERSITE PARIS SUD XI
VIROXIS SAS

<120> MUTATED LENTIVIRAL ENV PROTEINS AND THEIR USE AS DRUGS

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<170> PatentIn version 3.5

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Al a I l e Gl u Arg Arg Phe Lys Asp
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<400> 94

Al a I l e Gl u Arg Arg Gly Lys Asp
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<220>
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<400> 95

Al a I l e Gl u Arg Arg Arg Lys Asp
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<210> 96
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1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly I l e Lys Gln Leu Gln
20 25 30

Al a Arg Val Leu Al a Val Gl u Arg Al a Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 97
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<220>
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<400> 97

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Phe Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 98
<211> 49
<212> PRT
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<400> 98

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Gly Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 99
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<220>
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<400> 99

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

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Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Leu Leu Lys Asp Gln Gln Leu Leu
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Gly

<210> 100
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<400> 100

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Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly

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<212> PRT
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<400> 101

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1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Tyr Ala Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 102
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<220>

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Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Tyr Phe Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 1 0 3

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Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Tyr Gly Lys Asp Gln Gln Leu Leu
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Gly

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Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
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 1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

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 35 40 45

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<210> 106
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Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
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Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Ala Phe Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 107
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Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Ala Gly Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 108
<211> 49
<212> PRT
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<220>
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<400> 108

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Ala Arg Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 109
<211> 49
<212> PRT
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<220>
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<400> 109

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Phe Ala Lys Asp Gln Gln Leu Leu
35 40 45

Gly

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<210> 110
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<212> PRT
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Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
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Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Phe Phe Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 111
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<212> PRT
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<220>
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<400> 111

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1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Phe Gly Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 112
<211> 49
<212> PRT
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<220>
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<400> 112

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
Page 28

Al a Arg Val Leu Al a Val Gl u Arg Phe Arg Lys Asp Gl n Gl n Leu Leu
35 40 45

Gly

<210> 113
<211> 49
<212> PRT
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<400> 113

Ser Gly Ile Val Gl n Gl n Gl n Ser Asn Leu Leu Arg Al a Ile Gl n Al a
1 5 10 15

Arg Gl n His Met Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu Gl n
20 25 30

Al a Arg Val Leu Al a Val Gl u Arg Gly Al a Lys Asp Gl n Gl n Leu Leu
35 40 45

Gly

<210> 114
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Ser Gly Ile Val Gl n Gl n Gl n Ser Asn Leu Leu Arg Al a Ile Gl n Al a
1 5 10 15

Arg Gl n His Met Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu Gl n
20 25 30

Al a Arg Val Leu Al a Val Gl u Arg Gly Phe Lys Asp Gl n Gl n Leu Leu
35 40 45

Gly

<210> 115
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<400> 115

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Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Gly Gly Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 116

<211> 49

<212> PRT

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<220>

<223> fragment of HIV1 ENV ISU

<400> 116

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1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Gly Arg Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 117

<211> 49

<212> PRT

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<220>

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<400> 117

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Leu Ala Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 118
<211> 49
<212> PRT
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<220>
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<400> 118

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1 5 10 15
Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30
Ala Arg Val Leu Ala Val Glu Arg Leu Phe Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 119
<211> 49
<212> PRT
<213> Arti f i c i a l s e q u e n c e

<220>
<223> f r a g m e n t o f H I V 1 E N V I S U
<400> 119

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15
Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30
Ala Arg Val Leu Ala Val Glu Arg Leu Gly Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 120
<211> 49
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1

5

10

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Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Leu Arg Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 121
 <211> 49
 <212> PRT
 <213> Artificial sequence

<220>
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<400> 121

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
 1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Arg Ala Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 122
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 <212> PRT
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<220>
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<400> 122

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 1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Arg Phe Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 123

<211> 49
<212> PRT
<213> Arti f i c i a l s e q u e n c e

<220>
<223> f r a g m e n t o f H I V 1 E N V I S U

<400> 123

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Arg Gly Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 124
<211> 49
<212> PRT
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<220>
<223> f r a g m e n t o f H I V 1 E N V I S U

<400> 124

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Arg Arg Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 125
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<220>
<223> f r a g m e n t o f H I V 2 E N V I S U

<400> 125

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

eol f-seq1 . txt

Al a Arg Val Thr Al a Ile Gl u Lys Al a Leu Lys Asp Gl n Al a Gl n Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 126
<211> 57
<212> PRT
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<220>
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<400> 126

Al a Gly Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gl n Gl n Gl u Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Gl u Lys Phe Leu Lys Asp Gl n Al a Gl n Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

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<211> 57
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<220>
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<400> 127

Al a Gly Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gl n Gl n Gl u Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Gl u Lys Gly Leu Lys Asp Gl n Al a Gl n Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 128
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eol f-seql . txt

<400> 128

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

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Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Arg Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
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Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Tyr Ala Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln

eol f-seql . txt

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Tyr Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
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1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Ala Ala Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

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Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Ala Phe Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 136
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35

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Asn Ser Trp Gly Cys Ala Phe Arg Gln
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Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Phe Phe Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
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<210> 140
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 1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Phe Gly Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

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1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Phe Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 142
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<400> 142

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Gly Ala Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
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<400> 143

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1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Gly Phe Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

eol f-seq1 . txt

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1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Gly Gly Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
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1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Gly Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
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Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
Page 41

Al a Arg Val Thr Al a Ile Gl u Lys Leu Al a Lys Asp Gl n Al a Gl n Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

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1 5 10 15

Gl n Gl n Gl u Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Gl u Lys Leu Phe Lys Asp Gl n Al a Gl n Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
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<400> 148

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1 5 10 15

Gl n Gl n Gl u Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Gl u Lys Leu Gly Lys Asp Gl n Al a Gl n Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

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<400> 149

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20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 150

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of HIV2 ENV ISU

<400> 150

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Arg Ala Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 151

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of HIV2 ENV ISU

<400> 151

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Arg Phe Lys Asp Gln Ala Gln Leu
35 40 45

eol f-seq1 . txt

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 152
<211> 57
<212> PRT
<213> Arti f i c i a l s e q u e n c e

<220>
<223> f r a g m e n t o f H I V 2 E N V I S U
<400> 152

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Arg Gly Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 153
<211> 57
<212> PRT
<213> Arti f i c i a l S e q u e n c e

<220>
<223> f r a g m e n t o f H I V 2 E N V I S U
<400> 153

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Arg Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 154
<211> 57
<212> PRT
<213> Arti f i c i a l s e q u e n c e

<220>
<223> f r a g m e n t o f H I V 2 E N V I S U
<400> 154

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg

1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Ala Leu Gln Asp Gln Ala Arg Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 155
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of HIV2 ENV ISU

<400> 155

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Phe Leu Gln Asp Gln Ala Arg Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 156
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of HIV2 ENV ISU

<400> 156

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Gly Leu Gln Asp Gln Ala Arg Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 157

eol f-seq1 . txt

<211> 57
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<213> Arti f i c i a l sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 157

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Leu Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 158
<211> 57
<212> PRT
<213> Arti f i c i a l sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 158

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Arg Leu Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 159
<211> 57
<212> PRT
<213> Arti f i c i a l sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 159

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

eol f-seql . txt

Al a Arg Val Thr Al a Ile Gl u Lys Tyr Al a Gl n Asp Gl n Al a Arg Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 160
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 160

Al a Gly Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gl n Gl n Gl u Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Gl u Lys Tyr Phe Gl n Asp Gl n Al a Arg Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 161
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 161

Al a Gly Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gl n Gl n Gl u Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Gl u Lys Tyr Gly Gl n Asp Gl n Al a Arg Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 162
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV ISU

eol f-seql . txt

<400> 162

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Tyr Arg Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 163

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of HIV2 ENV ISU

<400> 163

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Ala Ala Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 164

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of HIV2 ENV ISU

<400> 164

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Ala Phe Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln

50

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<210> 165
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 <212> PRT
 <213> Arti f i c i a l s e q u e n c e

<220>
 <223> f r a g m e n t o f H I V 2 E N V I S U

<400> 165

Al a G l y I l e V a l G l n G l n G l n G l n G l n L e u L e u A s p V a l V a l L y s A r g
 1 5 10 15

G l n G l n G l u L e u L e u A r g L e u T h r V a l T r p G l y T h r L y s A s n L e u G l n
 20 25 30

Al a A r g V a l T h r A l a I l e G l u L y s A l a G l y G l n A s p G l n A l a A r g L e u
 35 40 45

A s n S e r T r p G l y C y s A l a P h e A r g G l n
 50 55

<210> 166
 <211> 57
 <212> PRT
 <213> Arti f i c i a l s e q u e n c e

<220>
 <223> f r a g m e n t o f H I V 2 E N V I S U

<400> 166

Al a G l y I l e V a l G l n G l n G l n G l n G l n L e u L e u A s p V a l V a l L y s A r g
 1 5 10 15

G l n G l n G l u L e u L e u A r g L e u T h r V a l T r p G l y T h r L y s A s n L e u G l n
 20 25 30

Al a A r g V a l T h r A l a I l e G l u L y s A l a A r g G l n A s p G l n A l a A r g L e u
 35 40 45

A s n S e r T r p G l y C y s A l a P h e A r g G l n
 50 55

<210> 167
 <211> 57
 <212> PRT
 <213> Arti f i c i a l s e q u e n c e

<220>
 <223> f r a g m e n t o f H I V 2 E N V I S U

<400> 167

Al a G l y I l e V a l G l n G l n G l n G l n G l n L e u L e u A s p V a l V a l L y s A r g
 1 5 10 15

eol f-seql . txt

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Phe Ala Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 168
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 168

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Phe Phe Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 169
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 169

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Phe Gly Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 170
<211> 57
<212> PRT

35

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Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 173
 <211> 57
 <212> PRT
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<220>
 <223> fragment of HIV2 ENV ISU
 <400> 173

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Gly Gly Gln Asp Gln Ala Arg Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 174
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of HIV2 ENV ISU
 <400> 174

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Ala Arg Val Thr Ala Ile Glu Lys Gly Arg Gln Asp Gln Ala Arg Leu
 35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 175
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of HIV2 ENV ISU
 <400> 175

eol f-seq1 . txt

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Ala Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 176
<211> 57
<212> PRT
<213> Arti ficial sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 176

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Phe Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 177
<211> 57
<212> PRT
<213> Arti ficial sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 177

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Gly Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

eol f-seq1 . txt

<210> 178
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV I SU

<400> 178

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Leu Arg Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 179
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV I SU

<400> 179

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Al a Arg Val Thr Ala Ile Glu Lys Arg Ala Gln Asp Gln Ala Arg Leu
35 40 45

Asn Ser Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 180
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV I SU

<400> 180

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln

Al a Arg Val Thr Al a Ile Glu Lys Arg Phe Gl n Asp Gl n Al a Arg Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 181
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 181

Al a Gly Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gl n Gl n Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Glu Lys Arg Gly Gl n Asp Gl n Al a Arg Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 182
<211> 57
<212> PRT
<213> Arti fici al Sequence

<220>
<223> fragment of HIV2 ENV ISU

<400> 182

Al a Gly Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gl n Gl n Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n
20 25 30

Al a Arg Val Thr Al a Ile Glu Lys Arg Arg Gl n Asp Gl n Al a Arg Leu
35 40 45

Asn Ser Trp Gly Cys Al a Phe Arg Gl n
50 55

<210> 183
<211> 57
<212> PRT
<213> Arti fici al sequence

<220>

<223> fragment of SIV ENV ISU

<400> 183

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Ala Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 184

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of SIV ENV ISU

<400> 184

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Phe Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 185

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of SIV ENV ISU

<400> 185

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Gly Leu Lys Asp Gln Ala Gln Leu
35 40 45

eol f-seq1 . txt

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 186
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 186

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Leu Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 187
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 187

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Arg Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 188
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 188

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg

1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Tyr Ala Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 189
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 189

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Tyr Phe Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 190
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 190

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Tyr Gly Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 191

eol f-seq1 . txt

<211> 57
 <212> PRT
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<220>
 <223> fragment of SIV ENV ISU

<400> 191

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Tyr Arg Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 192
 <211> 57
 <212> PRT
 <213> Arti fici al sequence

<220>
 <223> fragment of SIV ENV ISU

<400> 192

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Ala Ala Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 193
 <211> 57
 <212> PRT
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<220>
 <223> fragment of SIV ENV ISU

<400> 193

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

eol f-seq1 . txt

Thr Arg Val Thr Ala Ile Glu Lys Ala Phe Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 194
<211> 57
<212> PRT
<213> Arti ficial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 194

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Ala Gly Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 195
<211> 57
<212> PRT
<213> Arti ficial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 195

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Ala Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 196
<211> 57
<212> PRT
<213> Arti ficial sequence

<220>
<223> fragment of SIV ENV ISU

eol f-seql . txt

<400> 196

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Phe Ala Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 197

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of SIV ENV ISU

<400> 197

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Phe Phe Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 198

<211> 57

<212> PRT

<213> Artificial sequence

<220>

<223> fragment of SIV ENV ISU

<400> 198

Al a Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Phe Gly Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln

50

55

<210> 199
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of SIV ENV ISU

<400> 199

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Phe Arg Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 200
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of SIV ENV ISU

<400> 200

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Gly Ala Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 201
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of SIV ENV ISU

<400> 201

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

eol f-seql . txt

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Gly Phe Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 202
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 202

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Gly Gly Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 203
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 203

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Gly Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 204
<211> 57
<212> PRT

<213> Arti f i c i a l s e q u e n c e

<220>

<223> f r a g m e n t o f S I V E N V I S U

<400> 204

A l a G l y I l e V a l G l n G l n G l n G l n G l n L e u L e u A s p V a l V a l L y s A r g
1 5 10 15

G l n G l n G l u L e u L e u A r g L e u T h r V a l T r p G l y T h r L y s A s n L e u G l n
20 25 30

T h r A r g V a l T h r A l a I l e G l u L y s L e u A l a L y s A s p G l n A l a G l n L e u
35 40 45

A s n A l a T r p G l y C y s A l a P h e A r g G l n
50 55

<210> 205

<211> 57

<212> P R T

<213> A r t i f i c i a l s e q u e n c e

<220>

<223> f r a g m e n t o f S I V E N V I S U

<400> 205

A l a G l y I l e V a l G l n G l n G l n G l n G l n L e u L e u A s p V a l V a l L y s A r g
1 5 10 15

G l n G l n G l u L e u L e u A r g L e u T h r V a l T r p G l y T h r L y s A s n L e u G l n
20 25 30

T h r A r g V a l T h r A l a I l e G l u L y s L e u P h e L y s A s p G l n A l a G l n L e u
35 40 45

A s n A l a T r p G l y C y s A l a P h e A r g G l n
50 55

<210> 206

<211> 57

<212> P R T

<213> A r t i f i c i a l s e q u e n c e

<220>

<223> f r a g m e n t o f S I V E N V I S U

<400> 206

A l a G l y I l e V a l G l n G l n G l n G l n G l n L e u L e u A s p V a l V a l L y s A r g
1 5 10 15

G l n G l n G l u L e u L e u A r g L e u T h r V a l T r p G l y T h r L y s A s n L e u G l n
20 25 30

T h r A r g V a l T h r A l a I l e G l u L y s L e u G l y L y s A s p G l n A l a G l n L e u
Page 64

35

40

45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 207
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of SIV ENV ISU
 <400> 207

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Leu Arg Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 208
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of SIV ENV ISU
 <400> 208

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Arg Ala Lys Asp Gln Ala Gln Leu
 35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
 50 55

<210> 209
 <211> 57
 <212> PRT
 <213> Artificial sequence

<220>
 <223> fragment of SIV ENV ISU
 <400> 209

eol f-seq1 . txt

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Arg Phe Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 210
<211> 57
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of SIV ENV ISU

<400> 210

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Arg Gly Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

<210> 211
<211> 57
<212> PRT
<213> Artificial Sequence

<220>
<223> fragment of SIV ENV ISU

<400> 211

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Arg Arg Lys Asp Gln Ala Gln Leu
35 40 45

Asn Ala Trp Gly Cys Ala Phe Arg Gln
50 55

eol f-seql . txt

<210> 212
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 212
 Met Arg Val Lys Glu Lys Tyr Gl n His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15
 Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30
 Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gl n Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gl n Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380
 Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430
 Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460
 Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495
 Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540 545

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560 565

Leu Arg Ala Ile Gl u Ala Gl n Gl n Hi s Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Ala Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640 645

Asn Asn Tyr Thr Ser Leu Ile Hi s Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gl n Thr Hi s Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Gl u
 725 730 735

Gly Ile Gl u Gl u Gl u Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr Hi s Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 213

<211> 861

<212> PRT

<213> Human immunodeficiency virus type 1

<400> 213

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160 165

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys

Gl u Tyr Al a Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205
 Al a Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Al a
 210 215 220
 Pro Al a Gly Phe Al a Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Al a Gl u
 260 265 270
 Gl u Gl u Val Val Ile Arg Ser Al a Asn Phe Thr Asp Asn Al a Lys Thr
 275 280 285
 Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320
 Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
 325 330 335
 Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
 340 345 350
 Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365
 Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380
 Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430
 Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile

435

440

445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Phe Leu
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu

705 710 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 214
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 214

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

eol f-seql . txt

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

eol f-seql . txt

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Glu Arg Gly Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

eol f-seql . txt

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 215
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 215

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

eol f-seql . txt

Trp Gly Thr Met 20 Leu Leu Gly Ile 25 Leu Met Ile Cys Ser Ala 30 Thr Glu

Lys Leu Trp 35 Val Thr Val Tyr Tyr 40 Gly Val Pro Val Trp 45 Lys Glu Ala

Thr Thr 50 Thr Leu Phe Cys Ala 55 Ser Asp Ala Lys Ala 60 Tyr Asp Thr Glu

Val 65 His Asn Val Trp Ala 70 Thr His Ala Cys Val 75 Pro Thr Asp Pro Asn 80

Pro Gln Glu Val Val 85 Leu Val Asn Val Thr 90 Glu Asn Phe Asn Met Trp 95

Lys Asn Asp 100 Met Val Glu Gln Met His 105 Glu Asp Ile Ile Ser 110 Leu Trp

Asp Gln Ser 115 Leu Lys Pro Cys Val 120 Lys Leu Thr Pro Leu 125 Cys Val Ser

Leu Lys 130 Cys Thr Asp Leu Gly 135 Asn Ala Thr Asn Thr 140 Asn Ser Ser Asn

Thr 145 Asn Ser Ser Ser Gly 150 Glu Met Met Met Glu 155 Lys Gly Glu Ile Lys 160

Asn Cys Ser Phe 165 Asn Ile Ser Thr Ser Ile 170 Arg Gly Lys Val Gln Lys 175

Glu Tyr Ala 180 Phe Phe Tyr Lys Leu Asp 185 Ile Ile Pro Ile Asp 190 Asn Asp

Thr Thr 195 Ser Tyr Thr Leu Thr Ser 200 Cys Asn Thr Ser Val 205 Ile Thr Gln

Ala Cys 210 Pro Lys Val Ser Phe 215 Glu Pro Ile Pro Ile His Tyr Cys Ala

Pro 225 Ala Gly Phe Ala Ile 230 Leu Lys Cys Asn 235 Asn Lys Thr Phe Asn Gly 240

Thr Gly Pro Cys Thr 245 Asn Val Ser Thr Val 250 Gln Cys Thr His Gly 255 Ile

Arg Pro Val 260 Val Ser Thr Gln Leu Leu 265 Leu Asn Gly Ser Leu Ala Glu 270

Glu Glu Val 275 Val Ile Arg Ser Ala 280 Asn Phe Thr Asp Asn 285 Ala Lys Thr

eol f-seq1 . txt

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

eol f-seq1 . txt

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Leu Leu
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

eol f-seq1 . txt

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
 835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
 850 855 860

<210> 216
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 216

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr His Gly Ile
245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Glu
260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Glu Ile Asn Cys Thr Arg Pro
290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Ala His Cys
325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
340 345 350

Lys Leu Arg Glu Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
420 425 430

Gl n Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 560

Leu Arg Ala Ile Glu Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Glu Arg Arg Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gl n Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gl n Asn
 645 650 655

Gl n Gl n Glu Lys Asn Glu Gl n Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gl n Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 217
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 217

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn

130

135

Thr 145 Asn Ser Ser Ser Gly 150 Glu Met Met Met Glu 155 Lys Gly Glu Ile Lys 160

Asn Cys Ser Phe Asn 165 Ile Ser Thr Ser Ile 170 Arg Gly Lys Val Gln Lys 175

Glu Tyr Ala Phe 180 Phe Tyr Lys Leu Asp 185 Ile Ile Pro Ile Asp 190 Asn Asp

Thr Thr Ser 195 Tyr Thr Leu Thr Ser 200 Cys Asn Thr Ser Val 205 Ile Thr Gln

Ala Cys Pro Lys Val Ser Phe 215 Glu Pro Ile Pro Ile His Tyr Cys Ala 220

Pro 225 Ala Gly Phe Ala Ile 230 Leu Lys Cys Asn Asn 235 Lys Thr Phe Asn Gly 240

Thr Gly Pro Cys Thr 245 Asn Val Ser Thr Val 250 Gln Cys Thr His Gly 255 Ile

Arg Pro Val Val 260 Ser Thr Gln Leu Leu 265 Leu Asn Gly Ser Leu Ala Glu 270

Glu Glu Val 275 Val Ile Arg Ser Ala 280 Asn Phe Thr Asp Asn 285 Ala Lys Thr

Ile Ile Val Gln Leu Asn Gln 295 Ser Val Glu Ile Asn 300 Cys Thr Arg Pro

Asn 305 Asn Asn Thr Arg Lys 310 Ser Ile Arg Ile Gln 315 Arg Gly Pro Gly Arg 320

Ala Phe Val Thr Ile 325 Gly Lys Ile Gly Asn 330 Met Arg Gln Ala His Cys 335

Asn Ile Ser Arg 340 Ala Lys Trp Asn Ala 345 Thr Leu Lys Gln Ile Ala Ser 350

Lys Leu Arg 355 Glu Gln Phe Gly Asn 360 Asn Lys Thr Ile Ile 365 Phe Lys Gln

Ser Ser Gly Gly Asp Pro Glu 375 Ile Val Thr His Ser 380 Phe Asn Cys Gly

Gly 385 Glu Phe Phe Tyr Cys 390 Asn Ser Thr Gln Leu 395 Phe Asn Ser Thr Trp 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser

675

680

685

I l e P h e I l e M e t I l e V a l G l y G l y L e u V a l G l y L e u A r g I l e V a l P h e
 690 695 700

A l a V a l L e u S e r I l e V a l A s n A r g V a l A r g G l n G l y T y r S e r P r o L e u
 705 710 715 720

S e r P h e G l n T h r H i s L e u P r o T h r P r o A r g G l y P r o A s p A r g P r o G l u
 725 730 735

G l y I l e G l u G l u G l u G l y G l y G l u A r g A s p A r g A s p A r g S e r I l e A r g
 740 745 750

L e u V a l A s n G l y S e r L e u A l a L e u I l e T r p A s p A s p L e u A r g S e r L e u
 755 760 765

C y s L e u P h e S e r T y r H i s A r g L e u A r g A s p L e u L e u L e u I l e V a l T h r
 770 775 780

A r g I l e V a l G l u L e u L e u G l y A r g A r g G l y T r p G l u A l a L e u L y s T y r
 785 790 795 800

T r p T r p A s n L e u L e u G l n T y r T r p S e r G l n G l u L e u L y s A s n S e r A l a
 805 810 815

V a l S e r L e u L e u A s n A l a T h r A l a I l e A l a V a l A l a G l u G l y T h r A s p
 820 825 830

A r g V a l I l e G l u V a l V a l G l n G l y A l a C y s A r g A l a I l e A r g H i s I l e
 835 840 845

P r o A r g A r g I l e A r g G l n G l y L e u G l u A r g I l e L e u L e u
 850 855 860

<210> 218
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 218

M e t A r g V a l L y s G l u L y s T y r G l n H i s L e u T r p A r g T r p G l y T r p L y s
 1 5 10 15

T r p G l y T h r M e t L e u L e u G l y I l e L e u M e t I l e C y s S e r A l a T h r G l u
 20 25 30

L y s L e u T r p V a l T h r V a l T y r T y r G l y V a l P r o V a l T r p L y s G l u A l a
 35 40 45

T h r T h r T h r L e u P h e C y s A l a S e r A s p A l a L y s A l a T y r A s p T h r G l u
 50 55 60

eol f-seql . txt

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

eol f-seql . txt

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510 515

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Phe
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

eol f-seql . txt

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 219

<211> 861

<212> PRT

<213> Human immunodeficiency virus type 1

<400> 219

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

eol f-seql . txt

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Al a Gl u
260 265 270

Gl u Gl u Val Val Ile Arg Ser Al a Asn Phe Thr Asp Asn Al a Lys Thr
275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
305 310 315 320

Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
325 330 335

Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
420 425 430

Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Gl u Pro Leu Gly Val Al a Pro Thr Lys Al a Lys Arg Arg Val Val Gl n
500 505 510

Arg Gl u Lys Arg Al a Val Gly Ile Gly Al a Leu Phe Leu Gly Phe Leu
515 520 525

eol f-seq1 . txt

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Gly
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

eol f-seql . txt

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 220
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 220

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
165 170 175

Gl u Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205

Ala Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Gl u
 260 265 270

Gl u Gl u Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Ala Hi s Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
 340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Glu
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Glu Ala Arg Glu Leu Leu Ser Gly Ile Val Glu Glu Glu Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Glu Glu His Leu Leu Glu Leu Thr Val Trp
 565 570 575

Gly Ile Lys Glu Leu Glu Ala Arg Ile Leu Ala Val Glu Arg Tyr Arg
 580 585 590

Lys Asp Glu Glu Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Glu Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Glu Asn
 645 650 655

Glu Glu Glu Lys Asn Glu Glu Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Glu Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His 725 Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg 745 Asp Arg Asp Arg Ser Ile Arg 750

Leu Val Asn Gly Ser Leu Ala Leu 760 Ile Trp Asp Asp Leu Arg Ser Leu 765

Cys Leu Phe Ser Tyr His 775 Arg Leu Arg Asp Leu Leu 780 Ile Val Thr

Arg Ile Val Glu Leu Leu 790 Gly Arg Arg Gly Trp 795 Glu Ala Leu Lys Tyr 800

Trp Trp Asn Leu Leu 805 Gln Tyr Trp Ser Gln 810 Glu Leu Lys Asn Ser Ala 815

Val Ser Leu Leu 820 Asn Ala Thr Ala 825 Ile Ala Val Ala Glu Gly Thr Asp 830

Arg Val Ile 835 Glu Val Val Gln Gly Ala 840 Cys Arg Ala 845 Ile Arg His Ile

Pro Arg 850 Arg Ile Arg Gln Gly 855 Leu Glu Arg Ile Leu Leu 860

<210> 221

<211> 861

<212> PRT

<213> Human immunodeficiency virus type 1

<400> 221

Met Arg Val Lys 5 Glu Lys Tyr Gln His 10 Leu Trp Arg Trp Gly Trp Lys 15

Trp Gly Thr 20 Met Leu Leu Gly Ile 25 Leu Met Ile Cys Ser Ala Thr Glu 30

Lys Leu Trp 35 Val Thr Val Tyr Tyr 40 Gly Val Pro Val Trp Lys Glu Ala 45

Thr Thr Thr 50 Leu Phe Cys Ala 55 Ser Asp Ala Lys Ala Tyr Asp Thr Glu 60

Val His Asn Val Trp 70 Ala Thr His Ala Cys Val 75 Pro Thr Asp Pro Asn 80

Pro Gln Glu Val 85 Val Leu Val Asn Val Thr 90 Glu Asn Phe Asn Met Trp 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp

100

105

110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly

370

375

Gly 385 Glu Phe Phe Tyr Cys 390 Asn Ser Thr Gl n Leu Phe 395 Asn Ser Thr Trp 400
Phe Asn Ser Thr Trp 405 Ser Thr Gl u Gly Ser 410 Asn Asn Thr Gl u Gly 415 Ser
Asp Thr Ile Thr 420 Leu Pro Cys Arg Ile 425 Lys Gl n Phe Ile Asn 430 Met Trp
Gl n Gl u Val 435 Gly Lys Ala Met Tyr 440 Ala Pro Pro Ile Ser Gly Gl n Ile
Arg Cys 450 Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr 460 Arg Asp Gly Gly
Asn 465 Asn Asn Asn Gly Ser 470 Gl u Ile Phe Arg Pro 475 Gly Gly Gly Asp Met 480
Arg Asp Asn Trp Arg 485 Ser Gl u Leu Tyr Lys 490 Tyr Lys Val Val Lys 495 Ile
Glu Pro Leu Gly 500 Val Ala Pro Thr Lys 505 Ala Lys Arg Arg Val 510 Val Gl n
Arg Gl u Lys 515 Arg Ala Val Gly Ile 520 Gly Ala Leu Phe Leu 525 Gly Phe Leu
Gly Ala 530 Ala Gly Ser Thr Met 535 Gly Ala Arg Ser Met 540 Thr Leu Thr Val
Gl n Ala Arg Gl n Leu Leu 550 Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu 560
Leu Arg Ala Ile Gl u 565 Ala Gl n Gl n His Leu 570 Leu Gl n Leu Thr Val 575 Trp
Gly Ile Lys Gl n 580 Leu Gl n Ala Arg Ile 585 Leu Ala Val Gl u Arg 590 Ala Ala
Lys Asp Gl n 595 Gl n Leu Leu Gly Ile 600 Trp Gly Cys Ser Gly 605 Lys Leu Ile
Cys Thr Thr Ala Val Pro Trp 615 Asn Ala Ser Trp Ser 620 Asn Lys Ser Leu
Glu Gl n Ile Trp Asn 630 Asn Met Thr Trp Met Gl u 635 Trp Asp Arg Gl u Ile 640
Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn

Gln Gln Glu Lys 660 Asn Glu Gln Glu Leu 665 Leu Glu Leu Asp Lys 670 Trp Ala

Ser Leu Trp 675 Asn Trp Phe Asn Ile 680 Thr Asn Trp Leu Trp 685 Tyr Ile Lys

Ile Phe 690 Ile Met Ile Val Gly 695 Gly Leu Val Gly Leu 700 Arg Ile Val Phe

Ala Val Leu Ser Ile Val 710 Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu 720

Ser Phe Gln Thr His 725 Leu Pro Thr Pro Arg 730 Gly Pro Asp Arg Pro Glu 735

Gly Ile Glu Glu 740 Glu Gly Gly Glu Arg 745 Asp Arg Asp Arg Ser Ile Arg 750

Leu Val Asn Gly Ser Leu Ala Leu 760 Ile Trp Asp Asp Leu Arg Ser Leu 765

Cys Leu Phe Ser Tyr His Arg 775 Leu Arg Asp Leu Leu 780 Leu Ile Val Thr

Arg Ile Val Glu Leu Leu 790 Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr 800

Trp Trp Asn Leu Leu 805 Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala 815

Val Ser Leu Leu 820 Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile 845

Pro Arg Arg Ile Arg Gln Gly 855 Leu Glu Arg Ile Leu 860 Leu

<210> 222
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 222

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu 20 25 30

eol f-seql . txt

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

eol f-seql . txt

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
305 310 315 320

Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
325 330 335

Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
420 425 430

Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Gl u Pro Leu Gly Val Al a Pro Thr Lys Al a Lys Arg Arg Val Val Gl n
500 505 510

Arg Gl u Lys Arg Al a Val Gly Ile Gly Al a Leu Phe Leu Gly Phe Leu
515 520 525

Gly Al a Al a Gly Ser Thr Met Gly Al a Arg Ser Met Thr Leu Thr Val
530 535 540

Gl n Al a Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
545 550 555 560

Leu Arg Al a Ile Gl u Al a Gl n Gl n Hi s Leu Leu Gl n Leu Thr Val Trp
565 570 575

eol f-seql . txt

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Ala Phe
 580 585 590
 Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605
 Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620
 Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640
 Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655
 Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670
 Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685
 Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 700
 Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720
 Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735
 Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750
 Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765
 Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780
 Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
 785 790 795 800
 Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
 805 810 815
 Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
 820 825 830
 Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
 835 840 845

eol f-seql . txt

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
 850 855 860

<210> 223
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 223

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

eol f-seq1 . txt

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380
 Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430
 Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460
 Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

eol f-seql . txt

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555

Leu Arg Ala Ile Gl u Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Ala Gly
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gl n Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Gl u
 725 730 735

Gly Ile Gl u Gl u Gl u Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

eol f-seq1 . txt

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 224
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 224

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile Hi s Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Ala Hi s Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Ala Arg
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

I l e P h e I l e M e t I l e V a l G l y G l y L e u V a l G l y L e u A r g I l e V a l P h e
 690 695 700

A l a V a l L e u S e r I l e V a l A s n A r g V a l A r g G l n G l y T y r S e r P r o L e u
 705 710 715 720

S e r P h e G l n T h r H i s L e u P r o T h r P r o A r g G l y P r o A s p A r g P r o G l u
 725 730 735

G l y I l e G l u G l u G l u G l y G l y G l u A r g A s p A r g A s p A r g S e r I l e A r g
 740 745 750

L e u V a l A s n G l y S e r L e u A l a L e u I l e T r p A s p A s p L e u A r g S e r L e u
 755 760 765

C y s L e u P h e S e r T y r H i s A r g L e u A r g A s p L e u L e u L e u I l e V a l T h r
 770 775 780

A r g I l e V a l G l u L e u L e u G l y A r g A r g G l y T r p G l u A l a L e u L y s T y r
 785 790 800

T r p T r p A s n L e u L e u G l n T y r T r p S e r G l n G l u L e u L y s A s n S e r A l a
 805 810 815

V a l S e r L e u L e u A s n A l a T h r A l a I l e A l a V a l A l a G l u G l y T h r A s p
 820 825 830

A r g V a l I l e G l u V a l V a l G l n G l y A l a C y s A r g A l a I l e A r g H i s I l e
 835 840 845

P r o A r g A r g I l e A r g G l n G l y L e u G l u A r g I l e L e u L e u
 850 855 860

<210> 225
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 225

M e t A r g V a l L y s G l u L y s T y r G l n H i s L e u T r p A r g T r p G l y T r p L y s
 1 5 10 15

T r p G l y T h r M e t L e u L e u G l y I l e L e u M e t I l e C y s S e r A l a T h r G l u
 20 25 30

L y s L e u T r p V a l T h r V a l T y r T y r G l y V a l P r o V a l T r p L y s G l u A l a
 35 40 45

T h r T h r T h r L e u P h e C y s A l a S e r A s p A l a L y s A l a T y r A s p T h r G l u
 50 55 60

V a l H i s A s n V a l T r p A l a T h r H i s A l a C y s V a l P r o T h r A s p P r o A s n
 Page 109

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Phe Ala
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu

610

615

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 226
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 226

Met Arg Val Lys Gl u Lys Tyr Gl n Hi s Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15
 Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Gl u
 20 25 30
 Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Gl u Ala
 35 40 45
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gl u
 50 55 60
 Val Hi s Asn Val Trp Ala Thr Hi s Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gl n Gl u Val Val Leu Val Asn Val Thr Gl u Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Gl u Gl n Met Hi s Gl u Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Gl u Met Met Met Gl u Lys Gly Gl u Ile Lys
 145 150 155 160
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys
 165 170 175
 Gl u Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Gl u
 260 265 270

eol f-seql . txt

Gl u Gl u Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Gl u Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380
 Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430
 Gln Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460
 Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495
 Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510
 Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525
 Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

eol f-seql . txt

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Phe Phe
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
 785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
 805 810 815

eol f-seql . txt

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
 820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
 835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
 850 855 860

<210> 227
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 227

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

eol f-seq1 . txt

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205

Al a Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Al a
 210 215 220

Pro Al a Gly Phe Al a Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Al a Gl u
 260 265 270

Gl u Gl u Val Val Ile Arg Ser Al a Asn Phe Thr Asp Asn Al a Lys Thr
 275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320

Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
 325 330 335

Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
 340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

eol f-seql . txt

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495
 Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510
 Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525
 Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540
 Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560
 Leu Arg Ala Ile Glu Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575
 Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Glu Arg Phe Gly
 580 585 590
 Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605
 Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620
 Glu Gl n Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640
 Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gl n Asn
 645 650 655
 Gl n Gl n Glu Lys Asn Glu Gl n Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670
 Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685
 Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700
 Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
 705 710 715 720
 Ser Phe Gl n Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

eol f-seq1 . txt

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 228
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 228

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly 385 Gl u Phe Phe Tyr Cys 390 Asn Ser Thr Gl n Leu 395 Phe Asn Ser Thr Trp 400
 Phe Asn Ser Thr Trp 405 Ser Thr Gl u Gly Ser 410 Asn Asn Thr Gl u Gly Ser 415
 Asp Thr Ile Thr 420 Leu Pro Cys Arg Ile 425 Lys Gl n Phe Ile Asn Met Trp 430
 Gl n Gl u Val 435 Gly Lys Ala Met Tyr 440 Ala Pro Pro Ile Ser Gly Gl n Ile 445
 Arg Cys 450 Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr 460 Arg Asp Gly Gly
 Asn 465 Asn Asn Asn Gly Ser 470 Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met 480
 Arg Asp Asn Trp Arg 485 Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile 495
 Gl u Pro Leu Gly 500 Val Ala Pro Thr Lys 505 Ala Lys Arg Arg Val Val Gl n 510
 Arg Gl u Lys 515 Arg Ala Val Gly Ile 520 Gly Ala Leu Phe Leu Gly Phe Leu 525
 Gly Ala 530 Ala Gly Ser Thr Met 535 Gly Ala Arg Ser Met Thr Leu Thr Val 540
 Gl n Ala Arg Gl n Leu 550 Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu 560
 Leu Arg Ala Ile Gl u 565 Ala Gl n Gl n His Leu 570 Leu Gl n Leu Thr Val Trp 575
 Gly Ile Lys Gl n 580 Leu Gl n Ala Arg Ile 585 Leu Ala Val Gl u Arg Phe Arg 590
 Lys Asp Gl n 595 Gl n Leu Leu Gly Ile 600 Trp Gly Cys Ser Gly Lys Leu Ile 605
 Cys Thr Thr Ala Val Pro Trp 610 Asn Ala Ser Trp Ser Asn Lys Ser Leu 620
 Gl u Gl n Ile Trp Asn 630 Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile 640
 Asn Asn Tyr Thr Ser 645 Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn 655

Gln Gln Glu Lys₆₆₀ Asn Glu Gln Glu Leu₆₆₅ Leu Glu Leu Asp Lys₆₇₀ Trp Ala

Ser Leu Trp₆₇₅ Asn Trp Phe Asn Ile₆₈₀ Thr Asn Trp Leu Trp₆₈₅ Tyr Ile Lys

Ile Phe₆₉₀ Ile Met Ile Val Gly₆₉₅ Gly Leu Val Gly₇₀₀ Leu Arg Ile Val Phe

Ala Val Leu Ser Ile₇₁₀ Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu₇₂₀

Ser Phe Gln Thr His₇₂₅ Leu Pro Thr Pro Arg₇₃₀ Gly Pro Asp Arg Pro₇₃₅ Glu

Gly Ile Glu Glu₇₄₀ Glu Gly Gly Glu Arg₇₄₅ Asp Arg Asp Arg Ser Ile Arg

Leu Val Asn₇₅₅ Gly Ser Leu Ala Leu₇₆₀ Ile Trp Asp Asp Leu₇₆₅ Arg Ser Leu

Cys Leu₇₇₀ Phe Ser Tyr His Arg₇₇₅ Leu Arg Asp Leu Leu₇₈₀ Leu Ile Val Thr

Arg Ile Val Glu Leu Leu₇₉₀ Gly Arg Arg Gly Trp₇₉₅ Glu Ala Leu Lys Tyr₈₀₀

Trp Trp Asn Leu₈₀₅ Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala₈₁₅

Val Ser Leu Leu₈₂₀ Asn Ala Thr Ala Ile₈₂₅ Ala Val Ala Glu Gly Thr Asp

Arg Val Ile₈₃₅ Glu Val Val Gln Gly Ala Cys Arg Ala Ile₈₄₅ Arg His Ile

Pro Arg Arg Ile Arg Gln Gly₈₅₅ Leu Glu Arg Ile Leu₈₆₀ Leu

<210> 229
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 229

Met Arg Val Lys₅ Glu Lys Tyr Gln His₁₀ Leu Trp Arg Trp Gly Trp₁₅ Lys

Trp Gly Thr Met₂₀ Leu Leu Gly Ile₂₅ Leu Met Ile Cys Ser Ala Thr Glu

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala

35

40

45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg

305

310

320

Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
420 425 430

Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Gl u Ala Gl n Gl n Hi s Leu Leu Gl n Leu Thr Val Trp
565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Gly Ala

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu

850

855

<210> 230
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 230

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160 165

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
225 230 235 240

eol f-seql . txt

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

eol f-seq1 . txt

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
545 550 555 560 565

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Gly Phe
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

eol f-seql . txt

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 231
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 231

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160

eol f-seq1 . txt

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

eol f-seq1 . txt

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Gly Gly
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

eol f-seql . txt

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 232
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 232

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
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Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gl n Gl u Val Val Leu Val Asn Val Thr Gl u Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Gl u Gl n Met Hi s Gl u Asp Il e Il e Ser Leu Trp
 100 105 110

Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gl y Asn Al a Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gl y Gl u Met Met Met Gl u Lys Gl y Gl u Il e Lys
 145 150 155 160

Asn Cys Ser Phe Asn Il e Ser Thr Ser Il e Arg Gl y Lys Val Gl n Lys
 165 170 175

Gl u Tyr Al a Phe Phe Tyr Lys Leu Asp Il e Il e Pro Il e Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Il e Thr Gl n
 195 200 205

Al a Cys Pro Lys Val Ser Phe Gl u Pro Il e Pro Il e Hi s Tyr Cys Al a
 210 215 220

Pro Al a Gl y Phe Al a Il e Leu Lys Cys Asn Asn Lys Thr Phe Asn Gl y
 225 230 235 240

Thr Gl y Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gl y Il e
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gl y Ser Leu Al a Gl u
 260 265 270

Gl u Gl u Val Val Il e Arg Ser Al a Asn Phe Thr Asp Asn Al a Lys Thr
 275 280 285

Il e Il e Val Gl n Leu Asn Gl n Ser Val Gl u Il e Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Il e Arg Il e Gl n Arg Gl y Pro Gl y Arg
 305 310 315 320

Al a Phe Val Thr Il e Gl y Lys Il e Gl y Asn Met Arg Gl n Al a Hi s Cys
 325 330 335

Asn Il e Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Il e Al a Ser
 340 345 350

Lys Leu Arg₃₅₅ Glu Gln Phe Gly Asn₃₆₀ Asn Lys Thr Ile Ile₃₆₅ Phe Lys Gln
 Ser Ser₃₇₀ Gly Gly Asp Pro Glu₃₇₅ Ile Val Thr His Ser₃₈₀ Phe Asn Cys Gly
 Gly₃₈₅ Glu Phe Phe Tyr Cys₃₉₀ Asn Ser Thr Gln Leu₃₉₅ Phe Asn Ser Thr Trp₄₀₀
 Phe Asn Ser Thr Trp₄₀₅ Ser Thr Glu Gly Ser₄₁₀ Asn Asn Thr Glu Gly₄₁₅ Ser
 Asp Thr Ile Thr₄₂₀ Leu Pro Cys Arg Ile₄₂₅ Lys Gln Phe Ile Asn₄₃₀ Met Trp
 Gln Glu Val₄₃₅ Gly Lys Ala Met Tyr₄₄₀ Ala Pro Pro Ile Ser₄₄₅ Gly Gln Ile
 Arg Cys₄₅₀ Ser Ser Asn Ile Thr₄₅₅ Gly Leu Leu Leu Thr₄₆₀ Arg Asp Gly Gly
 Asn Asn Asn Asn Gly Ser₄₇₀ Glu Ile Phe Arg Pro₄₇₅ Gly Gly Gly Asp Met₄₈₀
 Arg Asp Asn Trp Arg₄₈₅ Ser Glu Leu Tyr Lys₄₉₀ Tyr Lys Val Val Lys₄₉₅ Ile
 Glu Pro Leu Gly₅₀₀ Val Ala Pro Thr Lys₅₀₅ Ala Lys Arg Arg Val₅₁₀ Val Gln
 Arg Glu Lys₅₁₅ Arg Ala Val Gly Ile₅₂₀ Gly Ala Leu Phe Leu₅₂₅ Gly Phe Leu
 Gly Ala₅₃₀ Ala Gly Ser Thr Met₅₃₅ Gly Ala Arg Ser Met₅₄₀ Thr Leu Thr Val
 Gln Ala Arg Gln Leu Leu₅₅₀ Ser Gly Ile Val Gln₅₅₅ Gln Gln Asn Asn Leu₅₆₀
 Leu Arg Ala Ile Glu₅₆₅ Ala Gln Gln His Leu₅₇₀ Leu Gln Leu Thr Val₅₇₅ Trp
 Gly Ile Lys Gln Leu Gln Ala Arg Ile₅₈₅ Leu Ala Val Glu Arg₅₉₀ Gly Arg
 Lys Asp Gln Gln Leu Leu Gly Ile₆₀₀ Trp Gly Cys Ser Gly₆₀₅ Lys Leu Ile
 Cys Thr₆₁₀ Thr Ala Val Pro Trp₆₁₅ Asn Ala Ser Trp Ser₆₂₀ Asn Lys Ser Leu

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 233
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 233

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
Page 135

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 Trp Gly Thr Met₂₀ Leu Leu Gly Ile Leu₂₅ Met Ile Cys Ser Ala₃₀ Thr Glu
 Lys Leu Trp₃₅ Val Thr Val Tyr₄₀ Tyr Gly Val Pro Val Trp₄₅ Lys Glu Ala
 Thr Thr₅₀ Thr Leu Phe Cys₅₅ Ala Ser Asp Ala Lys₆₀ Ala Tyr Asp Thr Glu
 Val His₆₅ Asn Val Trp Ala₇₀ Thr His Ala Cys₇₅ Val Pro Thr Asp Pro Asn₈₀
 Pro Gln Glu Val Val₈₅ Leu Val Asn Val Thr₉₀ Glu Asn Phe Asn₉₅ Met Trp
 Lys Asn Asp Met₁₀₀ Val Glu Gln Met His₁₀₅ Glu Asp Ile Ile Ser₁₁₀ Leu Trp
 Asp Gln Ser₁₁₅ Leu Lys Pro Cys₁₂₀ Val Lys Leu Thr Pro Leu₁₂₅ Cys Val Ser
 Leu Lys₁₃₀ Cys Thr Asp Leu Gly₁₃₅ Asn Ala Thr Asn Thr₁₄₀ Asn Ser Ser Asn
 Thr₁₄₅ Asn Ser Ser Ser Gly₁₅₀ Glu Met Met Met Glu₁₅₅ Lys Gly Glu Ile Lys₁₆₀
 Asn Cys Ser Phe Asn₁₆₅ Ile Ser Thr Ser Ile₁₇₀ Arg Gly Lys Val Gln₁₇₅ Lys
 Glu Tyr Ala Phe₁₈₀ Phe Tyr Lys Leu Asp₁₈₅ Ile Ile Pro Ile Asp₁₉₀ Asn Asp
 Thr Thr₁₉₅ Ser Tyr Thr Leu Thr Ser₂₀₀ Cys Asn Thr Ser Val₂₀₅ Ile Thr Gln
 Ala Cys₂₁₀ Pro Lys Val Ser Phe₂₁₅ Glu Pro Ile Pro Ile₂₂₀ His Tyr Cys Ala
 Pro₂₂₅ Ala Gly Phe Ala Ile₂₃₀ Leu Lys Cys Asn Asn₂₃₅ Lys Thr Phe Asn Gly₂₄₀
 Thr Gly Pro Cys Thr₂₄₅ Asn Val Ser Thr Val₂₅₀ Gln Cys Thr His Gly₂₅₅ Ile
 Arg Pro Val Val₂₆₀ Ser Thr Gln Leu Leu₂₆₅ Leu Asn Gly Ser Leu₂₇₀ Ala Glu
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 Page 136

275

280

285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380
 Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430
 Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460
 Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495
 Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510
 Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525
 Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540
 Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 234
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 234

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
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Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
195 200 205

eol f-seql . txt

Al a Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Al a
 210 215 220

Pro Al a Gly Phe Al a Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Al a Gl u
 260 265 270

Gl u Gl u Val Val Ile Arg Ser Al a Asn Phe Thr Asp Asn Al a Lys Thr
 275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320

Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
 325 330 335

Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
 340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

eol f-seql . txt

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Gl u Ala Gl n Gl n Hi s Leu Leu Gl n Leu Thr Val Trp
565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Leu Phe
580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile Hi s Ser Leu Ile Gl u Gl u Ser Gl n Asn
645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gl n Thr Hi s Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Gl u
725 730 735

Gly Ile Gl u Gl u Gl u Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

eol f-seql . txt

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 235
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 235

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
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Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

eol f-seq1 . txt

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

eol f-seq1 . txt

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Gl u Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Leu Gly
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

eol f-seql . txt

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 236
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 236

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Leu Arg
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640 645

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
 785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
 805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
 820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
 835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
 850 855 860

<210> 237
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 237

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
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Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Gl u
 260 265 270
 Gl u Gl u Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
 340 345 350
 Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365
 Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380
 Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430
 Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460
 Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495
 Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510
 Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 Page 150

515

520

525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Ala
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr

785 790 800

Trp Trp Asn Leu Leu 805 Gl n Tyr Trp Ser Gl n Gl u Leu Lys Asn Ser Ala 815

Val Ser Leu Leu 820 Asn Ala Thr Ala Ile 825 Ala Val Ala Gl u Gly 830 Thr Asp

Arg Val Ile 835 Gl u Val Val Gl n Gly 840 Ala Cys Arg Ala Ile 845 Arg His Ile

Pro Arg 850 Arg Ile Arg Gl n Gly 855 Leu Gl u Arg Ile Leu Leu 860

<210> 238
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 238

Met Arg Val Lys Gl u Lys Tyr Gl n His Leu Trp Arg Trp Gly Trp Lys 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Gl u 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gl y Val Pro Val Trp Lys Gl u Ala 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gl u 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn 65 70 75 80

Pro Gl n Gl u Val Val Leu Val Asn Val Thr Gl u Asn Phe Asn Met Trp 85 90 95

Lys Asn Asp Met Val Gl u Gl n Met His Gl u Asp Ile Ile Ser Leu Trp 100 105 110

Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn 130 135 140

Thr Asn Ser Ser Ser Gly Gl u Met Met Met Gl u Lys Gly Gl u Ile Lys 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys 165 170 175

eol f-seql . txt

Gl u Tyr Al a Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205
 Al a Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Al a
 210 215 220
 Pro Al a Gly Phe Al a Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Al a Gl u
 260 265 270
 Gl u Gl u Val Val Ile Arg Ser Al a Asn Phe Thr Asp Asn Al a Lys Thr
 275 280 285
 Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320
 Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
 325 330 335
 Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
 340 345 350
 Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365
 Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380
 Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430
 Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

eol f-seql . txt

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Phe
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720

eol f-seql . txt

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 239
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 239

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
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Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

eol f-seq1 . txt

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
355 360 365

eol f-seq1 . txt

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Glu Arg Arg Gly
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

eol f-seql . txt

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 240
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 240

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
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 Lys Leu Trp₃₅ Val Thr Val Tyr Tyr Gly Val Pro Val Trp₄₅ Lys Glu Ala
 Thr Thr Thr Leu Phe Cys Ala₅₅ Ser Asp Ala Lys Ala₆₀ Tyr Asp Thr Glu
 Val His Asn Val Trp Ala₇₀ Thr His Ala Cys Val₇₅ Pro Thr Asp Pro Asn₈₀
 Pro Gln Glu Val Val₈₅ Leu Val Asn Val Thr Glu Asn Phe Asn Met₉₅ Trp
 Lys Asn Asp Met₁₀₀ Val Glu Gln Met His₁₀₅ Glu Asp Ile Ile Ser Leu Trp
 Asp Gln Ser₁₁₅ Leu Lys Pro Cys Val₁₂₀ Lys Leu Thr Pro Leu₁₂₅ Cys Val Ser
 Leu Lys₁₃₀ Cys Thr Asp Leu Gly₁₃₅ Asn Ala Thr Asn Thr₁₄₀ Asn Ser Ser Asn
 Thr Asn Ser Ser Ser Gly₁₅₀ Glu Met Met Met Glu₁₅₅ Lys Gly Glu Ile Lys₁₆₀
 Asn Cys Ser Phe Asn₁₆₅ Ile Ser Thr Ser Ile₁₇₀ Arg Gly Lys Val Gln Lys₁₇₅
 Glu Tyr Ala Phe₁₈₀ Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp₁₉₀ Asn Asp
 Thr Thr Ser₁₉₅ Tyr Thr Leu Thr Ser₂₀₀ Cys Asn Thr Ser Val₂₀₅ Ile Thr Gln
 Ala Cys Pro Lys Val Ser Phe₂₁₅ Glu Pro Ile Pro Ile₂₂₀ His Tyr Cys Ala
 Pro Ala Gly Phe Ala Ile₂₃₀ Leu Lys Cys Asn Asn₂₃₅ Lys Thr Phe Asn Gly₂₄₀
 Thr Gly Pro Cys Thr₂₄₅ Asn Val Ser Thr Val₂₅₀ Gln Cys Thr His Gly₂₅₅ Ile
 Arg Pro Val Val₂₆₀ Ser Thr Gln Leu Leu₂₆₅ Leu Asn Gly Ser Leu Ala Glu₂₇₀
 Glu Glu Val₂₇₅ Val Ile Arg Ser Ala₂₈₀ Asn Phe Thr Asp Asn₂₈₅ Ala Lys Thr

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Arg
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640 645

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
 785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
 805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
 820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 241
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 241

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
Page 162

210

215

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
290 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val

Al a Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Al a
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
545 550 555 560

Al a Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Ala Leu Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
725 730 735

Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr

755

760

765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 242
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 242

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

eol f-seql . txt

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160
 Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175
 Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300
 Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320
 Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335
 Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350
 Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365
 Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380
 Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

eol f-seql . txt

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560 565

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Phe Leu Lys Asp Gln Gln Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685

eol f-seql . txt

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 243
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 243

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

eol f-seql . txt

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140
 Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160
 Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175
 Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300
 Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320
 Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

eol f-seq1 . txt

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Gl u Gl n
340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
355 360 365

Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly Gly Gl u Phe Phe Tyr
370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Gl u Gly Ser Asp Thr Ile Thr Leu
405 410 415

Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gly Lys
420 425 430

Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile Arg Cys Ser Ser Asn
435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
450 455 460

Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
485 490 495

Al a Pro Thr Lys Al a Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Al a
500 505 510

Val Gly Ile Gly Al a Leu Phe Leu Gly Phe Leu Gly Al a Al a Gly Ser
515 520 525

Thr Met Gly Al a Al a Ser Met Thr Leu Thr Val Gl n Al a Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Al a Ile Gl u
545 550 555 560

Al a Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Al a Arg Ile Leu Al a Val Gl u Arg Gly Leu Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Al a Val
595 600 605

eol f-seq1 . txt

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 244
<211> 856
<212> PRT

<213> Human immunodeficiency virus type 1

<400> 244

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Gl u Gl u Gl u Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gl n Leu
 275 280 285

Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gl n Ala Hi s Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Gl u Gl n
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365

Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly Gly Gl u Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Gl u Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
 450 455 460

Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Leu Leu Lys Asp Gln Gln Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 245
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 245

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
Page 175

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Ala Pro Ala Gl y Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gl y Thr Gl y Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gl n Cys Thr Hi s Gl y Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gl n Leu Leu Leu Asn Gl y Ser Leu Ala Gl u Gl u Gl u Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gl n Leu
 275 280 285
 Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300
 Lys Ser Ile Arg Ile Gl n Arg Gl y Pro Gl y Arg Ala Phe Val Thr Ile
 305 310 315 320
 Gl y Lys Ile Gl y Asn Met Arg Gl n Ala Hi s Cys Asn Ile Ser Arg Ala
 325 330 335
 Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Gl u Gl n
 340 345 350
 Phe Gl y Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gl y Gl y Asp
 355 360 365
 Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gl y Gl y Gl u Phe Phe Tyr
 370 375 380
 Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gl y Ser Asn Asn Thr Gl u Gl y Ser Asp Thr Ile Thr Leu
 405 410 415
 Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gl y Lys
 420 425 430
 Ala Met Tyr Ala Pro Pro Ile Ser Gl y Gl n Ile Arg Cys Ser Ser Asn
 435 440 445
 Ile Thr Gl y Leu Leu Leu Thr Arg Asp Gl y Gl y Asn Ser Asn Asn Gl u

450

455

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
545 550 555 560 565

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 246
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 246

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

eol f-seql . txt

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
370 375 380

eol f-seql . txt

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415
 Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430
 Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445
 Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460
 Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510
 Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525
 Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540
 Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560
 Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575
 Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Ala Lys Asp Gln Gln Leu
 580 585 590
 Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605
 Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620
 Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640
 Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655

eol f-seql . txt

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 247
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 247

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

eol f-seq1 . txt

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140
 Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160 165
 Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175
 Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

eol f-seq1 . txt

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575

eol f-seql . txt

Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Phe Lys Asp Gln Gln Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

eol f-seql . txt

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 248
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 248

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Al a Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
545 550 555 560

Al a Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Tyr Gly Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
725 730 735

Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 249
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 249

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
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Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
Page 188

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
545 550 555 560 565

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Arg Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val

690

695

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 250
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 250

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

eol f-seql . txt

Pro Gl n Gl u Val Val 85 Leu Val Asn Val Thr 90 Gl u Asn Phe Asn Met 95 Trp

Lys Asn Asp Met 100 Val Gl u Gl n Met Hi s 105 Gl u Asp Ile Ile Ser 110 Leu Trp

Asp Gl n Ser 115 Leu Lys Pro Cys Val 120 Lys Leu Thr Pro Leu 125 Cys Val Ser

Leu Lys 130 Cys Thr Asp Leu Lys 135 Asn Asp Thr Asn Thr 140 Asn Ser Ser Ser

Gly 145 Arg Met Ile Met Gl u 150 Lys Gly Gl u Ile Lys 155 Asn Cys Ser Phe Asn 160

Ile Ser Thr Ser Ile 165 Arg Gly Lys Val Gl n 170 Lys Gl u Tyr Ala Phe 175 Phe

Tyr Lys Leu Asp 180 Ile Ile Pro Ile Asp 185 Asn Asp Thr Thr Ser 190 Tyr Thr

Leu Thr Ser 195 Cys Asn Thr Ser Val 200 Ile Thr Gl n Ala Cys 205 Pro Lys Val

Ser Phe 210 Gl u Pro Ile Pro Ile 215 Hi s Tyr Cys Ala Pro 220 Ala Gly Phe Ala

Ile 225 Leu Lys Cys Asn Asn 230 Lys Thr Phe Asn Gly 235 Thr Gly Pro Cys Thr 240

Asn Val Ser Thr Val 245 Gl n Cys Thr Hi s Gly 250 Ile Arg Pro Val Val 255 Ser

Thr Gl n Leu Leu 260 Leu Asn Gly Ser Leu 265 Ala Gl u Gl u Gl u Val 270 Val Ile

Arg Ser Ala 275 Asn Phe Thr Asp Asn 280 Ala Lys Thr Ile Ile Val Gl n Leu

Asn Gl n Ser Val Gl u Ile Asn 295 Cys Thr Arg Pro Asn 300 Asn Asn Thr Arg

Lys 305 Ser Ile Arg Ile Gl n 310 Arg Gly Pro Gly Arg 315 Ala Phe Val Thr Ile 320

Gly Lys Ile Gly Asn 325 Met Arg Gl n Ala Hi s 330 Cys Asn Ile Ser Arg 335 Ala

Lys Trp Asn Asn 340 Thr Leu Lys Gl n Ile 345 Asp Ser Lys Leu Arg 350 Gl u Gl n

eol f-seql . txt

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365
 Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly Gly Gl u Phe Phe Tyr
 370 375 380
 Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gly Ser Asn Asn Thr Gl u Gly Ser Asp Thr Ile Thr Leu
 405 410 415
 Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gly Lys
 420 425 430
 Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile Arg Cys Ser Ser Asn
 435 440 445
 Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
 450 455
 Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480
 Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
 485 490 495
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
 500 505 510
 Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525
 Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
 530 535 540
 Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
 545 550 555 560
 Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
 565 570 575
 Gl n Ala Arg Ile Leu Ala Val Gl u Arg Ala Ala Lys Asp Gl n Gl n Leu
 580 585 590
 Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605
 Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
 610 615 620

eol f-seql . txt

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Glu Asn Glu Glu Glu Lys Asn
645 650 655

Glu Glu Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Glu Gly Tyr Ser Pro Leu Ser Phe Glu Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Glu Tyr Trp Ser Glu Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Glu Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Glu Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 251
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 251

eol f-seq1 . txt

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
260 265 270

eol f-seq1 . txt

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

eol f-seq1 . txt

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
545 550 555 560

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Ala Phe Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

eol f-seql . txt

Al a Thr Al a Ile Al a Val Al a Gl u Gly Thr Asp Arg Val Ile Gl u Val
820 825 830

Val Gl n Gly Al a Tyr Arg Al a Ile Arg Hi s Ile Pro Arg Arg Ile Arg
835 840 845

Gl n Gly Leu Gl u Arg Ile Leu Leu
850 855

<210> 252
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 252

Met Arg Val Lys Gl u Lys Tyr Gl n Hi s Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Al a Thr Gl u
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Gl u Al a
35 40 45

Thr Thr Thr Leu Phe Cys Al a Ser Asp Al a Lys Al a Tyr Asp Thr Gl u
50 55 60

Val Hi s Asn Val Trp Al a Thr Hi s Al a Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gl n Gl u Val Val Leu Val Asn Val Thr Gl u Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Gl u Gl n Met Hi s Gl u Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Gl u Lys Gly Gl u Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys Gl u Tyr Al a Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
500 505 510 515

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
545 550 555 560

Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Ala Gly Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

- <210> 253
- <211> 856
- <212> PRT
- <213> Human immunodeficiency virus type 1
- <400> 253

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 Page 201

115

120

125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys Glu Tyr Ala Phe Phe
 165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gl n Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gl n Leu
 275 280 285

Asn Gl n Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gl n Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Glu Gl n
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 Page 202

660

665

670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 254
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 254

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

eol f-seql . txt

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
305 310 315 320

eol f-seql . txt

Gly Lys Ile Gly Asn 325 Met Arg Gln Ala His 330 Cys Asn Ile Ser Arg Ala
 Lys Trp Asn 340 Asn Thr Leu Lys Gln Ile 345 Asp Ser Lys Leu Arg 350 Glu Gln
 Phe Gly Asn 355 Asn Lys Thr Ile Ile 360 Phe Lys Gln Ser Ser 365 Gly Gly Asp
 Pro Glu 370 Ile Val Thr His Ser 375 Phe Asn Cys Gly Gly 380 Glu Phe Phe Tyr
 Cys 385 Asn Ser Thr Gln Leu 390 Phe Asn Ser Thr Trp 395 Phe Asn Ser Thr Trp 400
 Ser Thr Lys Gly Ser 405 Asn Asn Thr Glu Gly 410 Ser Asp Thr Ile Thr 415 Leu
 Pro Cys Arg Ile 420 Lys Gln Ile Ile Asn 425 Met Trp Gln Glu Val 430 Gly Lys
 Ala Met Tyr 435 Ala Pro Pro Ile Ser 440 Gly Gln Ile Arg Cys 445 Ser Ser Asn
 Ile Thr Gly Leu Leu Leu Thr 455 Arg Asp Gly Gly Asn 460 Ser Asn Asn Glu
 Ser Glu Ile Phe Arg Pro 470 Gly Gly Gly Asp Met 475 Arg Asp Asn Trp Arg 480
 Ser Glu Leu Tyr Lys 485 Tyr Lys Val Val Lys 490 Ile Glu Pro Leu Gly Val 495
 Ala Pro Thr Lys 500 Ala Lys Arg Arg Val 505 Val Gln Arg Glu Lys 510 Arg Ala
 Val Gly Ile Gly Ala Leu Phe Leu 520 Gly Phe Leu Gly Ala 525 Ala Gly Ser
 Thr Met Gly Ala Ala Ser 535 Met Thr Leu Thr Val Gln 540 Ala Arg Gln Leu
 Leu Ser Gly Ile Val Gln 550 Gln Gln Asn Asn 555 Leu Leu Arg Ala Ile Glu 560
 Ala Gln Gln His Leu 565 Leu Gln Leu Thr Val 570 Trp Gly Ile Lys Gln 575 Leu
 Gln Ala Arg Ile 580 Leu Ala Val Glu Arg 585 Phe Ala Lys Asp Gln 590 Gln Leu

eol f-seql . txt

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605
 Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620
 Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640
 Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655
 Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670
 Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685
 Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700
 Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720
 Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735
 Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750
 Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765
 His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780
 Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800
 Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815
 Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830
 Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845
 Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

eol f-seql . txt

<210> 255
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 255
 Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15
 Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30
 Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140
 Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160
 Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175
 Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

eol f-seq1 . txt

Asn Val Ser Thr Val 245 Gln Cys Thr His Gly 250 Ile Arg Pro Val Val 255 Ser

Thr Gln Leu Leu 260 Leu Asn Gly Ser Leu 265 Ala Glu Glu Glu Val 270 Val Ile

Arg Ser Ala 275 Asn Phe Thr Asp Asn 280 Ala Lys Thr Ile Ile 285 Val Gln Leu

Asn Gln 290 Ser Val Glu Ile Asn 295 Cys Thr Arg Pro Asn 300 Asn Asn Thr Arg

Lys 305 Ser Ile Arg Ile Gln 310 Arg Gly Pro Gly Arg 315 Ala Phe Val Thr Ile 320

Gly Lys Ile Gly Asn 325 Met Arg Gln Ala His 330 Cys Asn Ile Ser Arg Ala 335

Lys Trp Asn Asn 340 Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg 350 Glu Gln

Phe Gly Asn 355 Asn Lys Thr Ile Ile 360 Phe Lys Gln Ser Ser 365 Gly Gly Asp

Pro Glu Ile Val Thr His Ser 375 Phe Asn Cys Gly Gly 380 Glu Phe Phe Tyr

Cys 385 Asn Ser Thr Gln Leu 390 Phe Asn Ser Thr Trp 395 Phe Asn Ser Thr Trp 400

Ser Thr Lys Gly Ser 405 Asn Asn Thr Glu Gly 410 Ser Asp Thr Ile Thr 415 Leu

Pro Cys Arg Ile 420 Lys Gln Ile Ile Asn 425 Met Trp Gln Glu Val 430 Gly Lys

Ala Met Tyr 435 Ala Pro Pro Ile Ser 440 Gly Gln Ile Arg Cys 445 Ser Ser Asn

Ile Thr Gly Leu Leu Leu Thr 455 Arg Asp Gly Gly Asn 460 Ser Asn Asn Glu

Ser Glu Ile Phe Arg Pro 470 Gly Gly Gly Asp Met 475 Arg Asp Asn Trp Arg 480

Ser Glu Leu Tyr Lys 485 Tyr Lys Val Val Lys 490 Ile Glu Pro Leu Gly Val 495

Ala Pro Thr Lys 500 Ala Lys Arg Arg Val 505 Val Gln Arg Glu Lys 510 Arg Ala

eol f-seq1 . txt

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
 530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
 545 550 555 560

Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
 565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Phe Phe Lys Asp Gl n Gl n Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
 625 630 635 640

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
 645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
 705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
 725 730 735

Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Gl u Leu
 770 775 780

eol f-seql . txt

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 256
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 256

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175
 Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300
 Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320
 Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335
 Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350
 Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365
 Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380
 Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415
 Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
545 550 555 560 565

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Phe Gly Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 257
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 257

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
Page 214

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
Page 215

355

360

365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Phe Arg Lys Asp Gln Gln Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
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eol f-seq1 . txt

Trp Gly Thr Met 20 Leu Leu Gly Met 25 Leu Met Ile Cys Ser Ala Thr Glu
Lys Leu Trp 35 Val Thr Val Tyr 40 Gly Val Pro Val 45 Trp Lys Glu Ala
Thr Thr Thr 50 Leu Phe Cys 55 Ala Ser Asp Ala Lys 60 Ala Tyr Asp Thr Glu
Val His Asn 65 Val Trp 70 Ala Thr His Ala Cys 75 Val Pro Thr Asp Pro Asn 80
Pro Gln Glu Val 85 Val Leu Val Asn Val 90 Thr Glu Asn Phe Asn Met 95 Trp
Lys Asn Asp 100 Met Val Glu Gln Met 105 His Glu Asp Ile Ile Ser 110 Leu Trp
Asp Gln Ser 115 Leu Lys Pro Cys 120 Val Lys Leu Thr Pro Leu Cys Val Ser
Leu Lys 130 Cys Thr Asp Leu Lys 135 Asn Asp Thr Asn Thr 140 Asn Ser Ser Ser
Gly Arg Met 145 Ile Met Glu Lys Gly Glu Ile Lys 155 Asn Cys Ser Phe Asn 160
Ile Ser Thr Ser 165 Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe 175
Tyr Lys Leu Asp 180 Ile Ile Pro Ile Asp 185 Asn Asp Thr Thr Ser 190 Tyr Thr
Leu Thr Ser 195 Cys Asn Thr Ser Val 200 Ile Thr Gln Ala Cys 205 Pro Lys Val
Ser Phe Glu 210 Pro Ile Pro Ile His Tyr Cys Ala 220 Pro Ala Gly Phe Ala
Ile Leu Lys Cys 225 Asn Asn Lys Thr Phe Asn Gly 235 Thr Gly Pro Cys Thr 240
Asn Val Ser Thr 245 Val Gln Cys Thr His Gly 250 Ile Arg Pro Val Val Ser 255
Thr Gln Leu 260 Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val 270 Val Ile
Arg Ser Ala 275 Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile 285 Val Gln Leu

eol f-seql . txt

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560

eol f-seql . txt

Al a Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Gly Ala Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

eol f-seql . txt

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 259
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 259

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
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 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205

eol f-seq1 . txt

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

eol f-seq1 . txt

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
485 490 495

Al a Pro Thr Lys Al a Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Al a
500 505 510

Val Gly Ile Gly Al a Leu Phe Leu Gly Phe Leu Gly Al a Al a Gly Ser
515 520 525

Thr Met Gly Al a Al a Ser Met Thr Leu Thr Val Gl n Al a Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Al a Ile Gl u
545 550 555 560 565

Al a Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Al a Arg Ile Leu Al a Val Gl u Arg Gly Phe Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Al a Val
595 600 605

Pro Trp Asn Al a Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
625 630 635 640 645

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Al a Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Al a Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
725 730 735

Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

eol f-seq1 . txt

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 260
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 260

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
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Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys Glu Tyr Ala Phe Phe
 165 170

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gl n Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gl n Leu
 275 280 285

Asn Gl n Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gl n Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Glu Gl n
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser 405 Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu 415
 Pro Cys Arg Ile 420 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys 430
 Ala Met Tyr 435 Ala Pro Pro Ile Ser 440 Gly Gln Ile Arg Cys Ser Ser Asn 445
 Ile Thr Gly Leu Leu Leu Thr 455 Arg Asp Gly Gly Asn Ser Asn Asn Glu 460
 Ser Glu Ile Phe Arg Pro 470 Gly Gly Gly Asp Met 475 Arg Asp Asn Trp Arg 480
 Ser Glu Leu Tyr Lys 485 Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val 495
 Ala Pro Thr Lys 500 Ala Lys Arg Arg Val 505 Val Gln Arg Glu Lys Arg Ala 510
 Val Gly Ile Gly Ala Leu Phe Leu 520 Gly Phe Leu Gly Ala Ala Gly Ser 525
 Thr Met Gly Ala Ala Ser Met 535 Thr Leu Thr Val Gln Ala Arg Gln Leu 540
 Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu 555 560
 Ala Gln Gln His Leu 565 Leu Gln Leu Thr Val 570 Trp Gly Ile Lys Gln Leu 575
 Gln Ala Arg Ile 580 Leu Ala Val Glu Arg 585 Gly Gly Lys Asp Gln Gln Leu 590
 Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val 605
 Pro Trp Asn Ala Ser Trp Ser 615 Asn Lys Ser Leu Glu Gln Ile Trp Asn 620
 Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser 635 640
 Leu Ile His Ser Leu 645 Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn 655
 Glu Gln Glu Leu 660 Leu Glu Leu Asp Lys 665 Trp Ala Ser Leu Trp Asn Trp 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 261
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 261

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
Page 227

50

55

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Gl u Gl n
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365

Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly Gly Gl u Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Gl u Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
 450 455 460

Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
 530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
 545 550 555 560

Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
 565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Gly Arg Lys Asp Gl n Gl n Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val

595

600

605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620
 Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640
 Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655
 Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670
 Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685
 Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700
 Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720
 Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735
 Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750
 Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765
 His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780
 Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800
 Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815
 Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830
 Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845
 Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 262

<211> 856

<212> PRT

<213> Human immunodeficiency virus type 1

<400> 262

Met Arg Val Lys Glu Lys Tyr Gl n Hi s Leu Trp Arg Trp Gly Trp Arg
1 5 10 15Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Gl u
20 25 30Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Gl u Ala
35 40 45Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gl u
50 55 60Val Hi s Asn Val Trp Ala Thr Hi s Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80Pro Gl n Gl u Val Val Leu Val Asn Val Thr Gl u Asn Phe Asn Met Trp
85 90 95Lys Asn Asp Met Val Gl u Gl n Met Hi s Gl u Asp Ile Ile Ser Leu Trp
100 105 110Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys Glu Tyr Ala Phe Phe
165 170 175Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n Ala Cys Pro Lys Val
195 200 205Ser Phe Glu Pro Ile Pro Ile Hi s Tyr Cys Ala Pro Ala Gly Phe Ala
210 215 220Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
225 230 235 240Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile Arg Pro Val Val Ser
245 250 255

eol f-seql . txt

Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Gl u Gl u Gl u Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gl n Leu
 275 280 285

Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gl n Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Gl u Gl n
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365

Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly Gly Gl u Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Gl u Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
 450 455 460

Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

eol f-seql . txt

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
545 550 555 560

Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Leu Ala Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
725 730 735

Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Gl u Leu
770 775 780

Leu Gly Arg Arg Gly Trp Gl u Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

eol f-seql . txt

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 263
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 263

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
165 170 175

eol f-seql . txt

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300
 Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320
 Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335
 Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350
 Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365
 Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380
 Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415
 Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430
 Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

eol f-seq1 . txt

I l e Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
450 455 460

Ser Gl u I l e Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys I l e Gl u Pro Leu Gly Val
485 490 495

Al a Pro Thr Lys Al a Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Al a
500 505 510

Val Gly I l e Gly Al a Leu Phe Leu Gly Phe Leu Gly Al a Al a Gly Ser
515 520 525

Thr Met Gly Al a Al a Ser Met Thr Leu Thr Val Gl n Al a Arg Gl n Leu
530 535 540

Leu Ser Gly I l e Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Al a I l e Gl u
545 550 555 560

Al a Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly I l e Lys Gl n Leu
565 570 575

Gl n Al a Arg I l e Leu Al a Val Gl u Arg Leu Phe Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly I l e Trp Gly Cys Ser Gly Lys Leu I l e Cys Thr Thr Al a Val
595 600 605

Pro Trp Asn Al a Ser Trp Ser Asn Lys Ser Leu Gl u Gl n I l e Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u I l e Asn Asn Tyr Thr Ser
625 630 635 640

Leu I l e His Ser Leu I l e Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Al a Ser Leu Trp Asn Trp
660 665 670

Phe Asn I l e Thr Asn Trp Leu Trp Tyr I l e Lys Leu Phe I l e Met I l e
675 680 685

Val Gly Gly Leu Val Gly Leu Arg I l e Val Phe Al a Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr Hi s
705 710 715 720

eol f-seql . txt

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 264
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 264

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
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Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Leu Gly Lys Asp Gln Gln Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640

eol f-seq1 . txt

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 265
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 265

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
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Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
Page 240

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140
 Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160
 Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175
 Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190
 Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205
 Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220
 Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 Page 241

290

295

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
545 550 555 560

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu

Gln Ala Arg Ile Leu Ala Val Glu Arg Leu Arg Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg

835

840

845

Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 266
 <211> 856
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 266

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
 130 135 140

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

eol f-seql . txt

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255
 Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270
 Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285
 Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300
 Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320
 Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335
 Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350
 Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365
 Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380
 Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400
 Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415
 Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430
 Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445
 Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460
 Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

eol f-seql . txt

Al a Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Al a
500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
545 550 555 560

Al a Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Arg Ala Lys Asp Gl n Gl n Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Gl u Gl n Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn Gl n Gl n Gl u Lys Asn
645 650 655

Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu Ser Phe Gl n Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Gl u Gly Ile Gl u Gl u Gl u
725 730 735

Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

eol f-seql . txt

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 267
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 267

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140

eol f-seql . txt

Gly Arg Met Ile Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe Asn
 145 150 155 160

Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys Glu Tyr Ala Phe Phe
 165 170 175

Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp Thr Thr Ser Tyr Thr
 180 185 190

Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Val
 195 200 205

Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala
 210 215 220

Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly Thr Gly Pro Cys Thr
 225 230 235 240

Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser
 245 250 255

Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile
 260 265 270

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

eol f-seql . txt

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 530 535 540

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
 545 550 555 560 565

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
 565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Phe Lys Asp Gln Gln Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685

eol f-seq1 . txt

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
805 810 815

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 268
<211> 856
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 268

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Arg
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Met Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val 65 His Asn Val Trp Ala 70 Thr His Ala Cys 75 Val Pro Thr Asp Pro Asn 80
 Pro Gln Glu Val Val 85 Leu Val Asn Val Thr 90 Glu Asn Phe Asn Met Trp 95
 Lys Asn Asp Met 100 Val Glu Gln Met His 105 Glu Asp Ile Ile Ser Leu Trp 110
 Asp Gln Ser 115 Leu Lys Pro Cys Val 120 Lys Leu Thr Pro Leu Cys Val Ser 125
 Leu Lys 130 Cys Thr Asp Leu Lys 135 Asn Asp Thr Asn Thr 140 Asn Ser Ser Ser
 Gly 145 Arg Met Ile Met Glu 150 Lys Gly Glu Ile Lys 155 Asn Cys Ser Phe Asn 160
 Ile Ser Thr Ser Ile 165 Arg Gly Lys Val Gln 170 Lys Glu Tyr Ala Phe Phe 175
 Tyr Lys Leu Asp 180 Ile Ile Pro Ile Asp 185 Asn Asp Thr Thr Ser Tyr Thr 190
 Leu Thr Ser 195 Cys Asn Thr Ser Val 200 Ile Thr Gln Ala Cys 205 Pro Lys Val
 Ser Phe 210 Glu Pro Ile Pro Ile 215 His Tyr Cys Ala Pro 220 Ala Gly Phe Ala
 Ile 225 Leu Lys Cys Asn Asn 230 Lys Thr Phe Asn Gly 235 Thr Gly Pro Cys Thr 240
 Asn Val Ser Thr Val 245 Gln Cys Thr His Gly 250 Ile Arg Pro Val Val Ser 255
 Thr Gln Leu Leu 260 Leu Asn Gly Ser Leu 265 Ala Glu Glu Glu Val Val Ile 270
 Arg Ser Ala 275 Asn Phe Thr Asp Asn 280 Ala Lys Thr Ile Ile Val Gln Leu 285
 Asn Gln Ser 290 Val Glu Ile Asn 295 Cys Thr Arg Pro Asn 300 Asn Asn Thr Arg
 Lys 305 Ser Ile Arg Ile Gln 310 Arg Gly Pro Gly Arg 315 Ala Phe Val Thr Ile 320
 Gly Lys Ile Gly Asn 325 Met Arg Gln Ala His 330 Cys Asn Ile Ser Arg Ala 335

Lys Trp Asn Asn Thr Leu Lys Gl n Ile Asp Ser Lys Leu Arg Gl u Gl n
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n Ser Ser Gly Gly Asp
 355 360 365

Pro Gl u Ile Val Thr His Ser Phe Asn Cys Gly Gly Gl u Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Gl u Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gl n Ile Ile Asn Met Trp Gl n Gl u Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Gl u
 450 455 460

Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile Gl u Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n Arg Gl u Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gl n Ala Arg Gl n Leu
 530 535 540

Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu Leu Arg Ala Ile Gl u
 545 550 555 560

Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp Gly Ile Lys Gl n Leu
 565 570 575

Gl n Ala Arg Ile Leu Ala Val Gl u Arg Arg Gly Lys Asp Gl n Gl n Leu
 580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
 595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
 610 615 620
 Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
 625 630 635
 Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
 645 650
 Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
 660 665 670
 Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
 675 680 685
 Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
 690 695 700
 Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
 705 710 715 720
 Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
 725 730 735
 Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
 740 745 750
 Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
 755 760 765
 His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
 770 775 780
 Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
 785 790 795 800
 Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
 805 810 815
 Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
 820 825 830
 Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
 835 840 845
 Gln Gly Leu Glu Arg Ile Leu Leu
 850 855

<210> 269

<211> 856

<212> PRT

<213> Human immunodeficiency virus type 1

eol f-seq1 . txt

<400> 269

Met Arg Val Lys Gl u Lys Tyr Gl n Hi s Leu Trp Arg Trp Gl y Trp Arg
1 5 10 15Trp Gl y Thr Met Leu Leu Gl y Met Leu Met Il e Cys Ser Al a Thr Gl u
20 25 30Lys Leu Trp Val Thr Val Tyr Tyr Gl y Val Pro Val Trp Lys Gl u Al a
35 40 45Thr Thr Thr Leu Phe Cys Al a Ser Asp Al a Lys Al a Tyr Asp Thr Gl u
50 55 60Val Hi s Asn Val Trp Al a Thr Hi s Al a Cys Val Pro Thr Asp Pro Asn
65 70 75 80Pro Gl n Gl u Val Val Leu Val Asn Val Thr Gl u Asn Phe Asn Met Trp
85 90 95Lys Asn Asp Met Val Gl u Gl n Met Hi s Gl u Asp Il e Il e Ser Leu Trp
100 105 110Asp Gl n Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125Leu Lys Cys Thr Asp Leu Lys Asn Asp Thr Asn Thr Asn Ser Ser Ser
130 135 140Gl y Arg Met Il e Met Gl u Lys Gl y Gl u Il e Lys Asn Cys Ser Phe Asn
145 150 155 160Il e Ser Thr Ser Il e Arg Gl y Lys Val Gl n Lys Gl u Tyr Al a Phe Phe
165 170 175Tyr Lys Leu Asp Il e Il e Pro Il e Asp Asn Asp Thr Thr Ser Tyr Thr
180 185 190Leu Thr Ser Cys Asn Thr Ser Val Il e Thr Gl n Al a Cys Pro Lys Val
195 200 205Ser Phe Gl u Pro Il e Pro Il e Hi s Tyr Cys Al a Pro Al a Gl y Phe Al a
210 215 220Il e Leu Lys Cys Asn Asn Lys Thr Phe Asn Gl y Thr Gl y Pro Cys Thr
225 230 235 240Asn Val Ser Thr Val Gl n Cys Thr Hi s Gl y Il e Arg Pro Val Val Ser
245 250 255Thr Gl n Leu Leu Leu Asn Gl y Ser Leu Al a Gl u Gl u Gl u Val Val Il e
Page 254

Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr Ile Ile Val Gln Leu
 275 280 285

Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg
 290 295 300

Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg Ala Phe Val Thr Ile
 305 310 315 320

Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 325 330 335

Lys Trp Asn Asn Thr Leu Lys Gln Ile Asp Ser Lys Leu Arg Glu Gln
 340 345 350

Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln Ser Ser Gly Gly Asp
 355 360 365

Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr
 370 375 380

Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Phe Asn Ser Thr Trp
 385 390 395 400

Ser Thr Lys Gly Ser Asn Asn Thr Glu Gly Ser Asp Thr Ile Thr Leu
 405 410 415

Pro Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 420 425 430

Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445

Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Ser Asn Asn Glu
 450 455 460

Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 465 470 475 480

Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val
 485 490 495

Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 500 505 510

Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala Ala Gly Ser
 515 520 525

Thr Met Gly Ala Ala Ser Met Thr Leu Thr Val Gln Ala Arg Gln Leu
 Page 255

530

535

Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu
545 550 555 560

Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu
565 570 575

Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Arg Lys Asp Gln Gln Leu
580 585 590

Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val
595 600 605

Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn
610 615 620

Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser
625 630 635 640

Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn
645 650 655

Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala Ser Leu Trp Asn Trp
660 665 670

Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys Leu Phe Ile Met Ile
675 680 685

Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe Ala Val Leu Ser Val
690 695 700

Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu Ser Phe Gln Thr His
705 710 715 720

Leu Pro Ile Pro Arg Gly Pro Asp Arg Pro Glu Gly Ile Glu Glu Glu
725 730 735

Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg Leu Val Asn Gly Ser
740 745 750

Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu Cys Leu Phe Ser Tyr
755 760 765

His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr Arg Ile Val Glu Leu
770 775 780

Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr Trp Trp Asn Leu Leu
785 790 795 800

Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala Val Ser Leu Leu Asn
Page 256

Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp Arg Val Ile Glu Val
820 825 830

Val Gln Gly Ala Tyr Arg Ala Ile Arg His Ile Pro Arg Arg Ile Arg
835 840 845

Gln Gly Leu Glu Arg Ile Leu Leu
850 855

<210> 270
<211> 32
<212> PRT
<213> Artificial Sequence

<220>
<223> fragemnt of Lenti viral ENV protein

<220>
<221> mi sc_feature
<222> (1)..(1)
<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
<221> mi sc_feature
<222> (4)..(5)
<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
<221> mi sc_feature
<222> (7)..(8)
<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
<221> MI SC_FEATURE
<222> (12)..(12)
<223> Xaa = A or S

<220>
<221> mi sc_feature
<222> (13)..(15)
<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
<221> mi sc_feature
<222> (18)..(18)
<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
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<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
<221> mi sc_feature
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<223> Xaa can be any natural ly occurri ng ami no aci d

<220>
<221> MI SC_FEATURE
<222> (25)..(29)
<223> Xaa = no or any ami no aci d

<220>
 <221> MI SC_FEATURE
 <222> (30).. (30)
 <223> Xaa = N or S

<220>
 <221> MI SC_FEATURE
 <222> (31).. (31)
 <223> Xaa = E, D, N or A

<220>
 <221> MI SC_FEATURE
 <222> (32).. (32)
 <223> Xaa = S or T

<400> 270

Xaa Asp Gln Xaa Xaa Leu Xaa Xaa Trp Gly Cys Xaa Xaa Xaa Xaa Cys
 1 5 10 15

Val Xaa Thr Xaa Val Pro Trp Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
 20 25 30

<210> 271
 <211> 115
 <212> PRT
 <213> Arti ficial sequence

<220>
 <223> derived from lenti viral ENV ISU

<400> 271

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
 35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
 50 55 60

Trp Asn Ala Ser Trp Ser Asn Ala Ser Leu Glu Gln Ile Trp Asn Asn
 65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
 85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
 100 105 110

Gln Glu Leu
 115

<210> 272
 <211> 115
 <212> PRT
 <213> Artificial sequence

<220>
 <223> derived from lenti viral ENV ISU

<400> 272

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
 35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
 50 55 60

Trp Asn Ala Ser Trp Ser Asn Gly Ser Leu Glu Gln Ile Trp Asn Asn
 65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
 85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
 100 105 110

Gln Glu Leu
 115

<210> 273
 <211> 115
 <212> PRT
 <213> Artificial sequence

<220>
 <223> derived from lenti viral ENV ISU

<400> 273

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
 35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
 50 55 60

eol f-seq1 . txt

Trp Asn Ala Ser Trp Ser Asn Ser Ser Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 274
<211> 115
<212> PRT
<213> Artificial sequence

<220>
<223> derived from lenti viral ENV ISU

<400> 274

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Asp Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 275
<211> 115
<212> PRT
<213> Artificial sequence

<220>
<223> derived from lenti viral ENV ISU

<400> 275

eol f-seq1 . txt

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Ala Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 276
<211> 115
<212> PRT
<213> Artificial sequence

<220>
<223> derived from lenti viral ENV ISU

<400> 276

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Asn Asn Ala Ser Trp Ser Asn Lys Ser Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 277
<211> 115
<212> PRT
<213> Artificial sequence

<220>
<223> derived from lenti viral ENV ISU
<400> 277

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Trp Asn Ala Ser Trp Ser Asn Lys Asp Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 278
<211> 115
<212> PRT
<213> Artificial sequence

<220>
<223> derived from lenti viral ENV ISU
<400> 278

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

eol f-seql . txt

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Trp Asn Ala Ser Trp Ser Asn Lys Ala Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 279
<211> 115
<212> PRT
<213> Artificial sequence

<220>
<223> derived from lentiviral ENV ISU

<400> 279

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Trp Asn Ala Ser Trp Ser Asn Lys Asn Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 280
<211> 115
<212> PRT
<213> Artificial sequence

eol f-seql . txt

<220>

<223> derived from lenti viral ENV ISU

<400> 280

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Trp Asn Ala Ser Trp Ser Asn Tyr Ser Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 281

<211> 115

<212> PRT

<213> Artificial sequence

<220>

<223> derived from lenti viral ENV ISU

<400> 281

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Ala Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Trp Asn Ala Ser Trp Ser Asn Ala Ser Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

eol f-seq1 . txt

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
 85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
 100 105 110

Gln Glu Leu
 115

<210> 282
 <211> 115
 <212> PRT
 <213> Artificial sequence

<220>
 <223> derived from lenti viral ENV ISU

<400> 282

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
 35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
 50 55 60

Trp Asn Ala Ser Trp Ser Asn Ala Ala Leu Glu Gln Ile Trp Asn Asn
 65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
 85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
 100 105 110

Gln Glu Leu
 115

<210> 283
 <211> 115
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> derived from lenti viral ENV ISU

<400> 283

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Arg Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile Cys Thr Thr Ala Val Pro
50 55 60

Ala Asn Ala Ser Trp Ser Asn Ala Ala Leu Glu Gln Ile Trp Asn Asn
65 70 75 80

Met Thr Trp Met Glu Trp Asp Arg Glu Ile Asn Asn Tyr Thr Ser Leu
85 90 95

Ile His Ser Leu Ile Glu Glu Ser Gln Asn Gln Gln Glu Lys Asn Glu
100 105 110

Gln Glu Leu
115

<210> 284
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 284

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
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130

135

Thr 145 Asn Ser Ser Ser Gly 150 Glu Met Met Met Glu 155 Lys Gly Glu Ile Lys 160

Asn Cys Ser Phe Asn 165 Ile Ser Thr Ser Ile 170 Arg Gly Lys Val Gln Lys 175

Glu Tyr Ala Phe 180 Phe Tyr Lys Leu Asp 185 Ile Ile Pro Ile Asp 190 Asn Asp

Thr Thr Ser 195 Tyr Thr Leu Thr Ser 200 Cys Asn Thr Ser Val 205 Ile Thr Gln

Ala Cys Pro Lys Val Ser Phe 215 Glu Pro Ile Pro Ile His Tyr Cys Ala 210

Pro 225 Ala Gly Phe Ala Ile 230 Leu Lys Cys Asn Asn 235 Lys Thr Phe Asn Gly 240

Thr Gly Pro Cys Thr 245 Asn Val Ser Thr Val 250 Gln Cys Thr His Gly 255 Ile

Arg Pro Val Val 260 Ser Thr Gln Leu Leu 265 Leu Asn Gly Ser Leu Ala Glu 270

Glu Glu Val 275 Val Ile Arg Ser Ala 280 Asn Phe Thr Asp Asn 285 Ala Lys Thr

Ile Ile Val Gln Leu Asn Gln 295 Ser Val Glu Ile Asn 300 Cys Thr Arg Pro

Asn 305 Asn Asn Thr Arg Lys 310 Ser Ile Arg Ile Gln 315 Arg Gly Pro Gly Arg 320

Ala Phe Val Thr Ile 325 Gly Lys Ile Gly Asn 330 Met Arg Gln Ala His Cys 335

Asn Ile Ser Arg 340 Ala Lys Trp Asn Ala 345 Thr Leu Lys Gln Ile Ala Ser 350

Lys Leu Arg 355 Glu Gln Phe Gly Asn 360 Asn Lys Thr Ile Ile 365 Phe Lys Gln

Ser Ser Gly Gly Asp Pro Glu 375 Ile Val Thr His Ser 380 Phe Asn Cys Gly

Gly 385 Glu Phe Phe Tyr Cys 390 Asn Ser Thr Gln Leu 395 Phe Asn Ser Thr Trp 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Gl u Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Arg Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 Page 268

675

680

685

I l e P h e I l e M e t I l e V a l G l y G l y L e u V a l G l y L e u A r g I l e V a l P h e
 690 695 700

A l a V a l L e u S e r I l e V a l A s n A r g V a l A r g G l n G l y T y r S e r P r o L e u
 705 710 715 720

S e r P h e G l n T h r H i s L e u P r o T h r P r o A r g G l y P r o A s p A r g P r o G l u
 725 730 735

G l y I l e G l u G l u G l u G l y G l y G l u A r g A s p A r g A s p A r g S e r I l e A r g
 740 745 750

L e u V a l A s n G l y S e r L e u A l a L e u I l e T r p A s p A s p L e u A r g S e r L e u
 755 760 765

C y s L e u P h e S e r T y r H i s A r g L e u A r g A s p L e u L e u L e u I l e V a l T h r
 770 775 780

A r g I l e V a l G l u L e u L e u G l y A r g A r g G l y T r p G l u A l a L e u L y s T y r
 785 790 795 800

T r p T r p A s n L e u L e u G l n T y r T r p S e r G l n G l u L e u L y s A s n S e r A l a
 805 810 815

V a l S e r L e u L e u A s n A l a T h r A l a I l e A l a V a l A l a G l u G l y T h r A s p
 820 825 830

A r g V a l I l e G l u V a l V a l G l n G l y A l a C y s A r g A l a I l e A r g H i s I l e
 835 840 845

P r o A r g A r g I l e A r g G l n G l y L e u G l u A r g I l e L e u L e u
 850 855 860

<210> 285
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1

<400> 285

M e t A r g V a l L y s G l u L y s T y r G l n H i s L e u T r p A r g T r p G l y T r p L y s
 1 5 10 15

T r p G l y T h r M e t L e u L e u G l y I l e L e u M e t I l e C y s S e r A l a T h r G l u
 20 25 30

L y s L e u T r p V a l T h r V a l T y r T y r G l y V a l P r o V a l T r p L y s G l u A l a
 35 40 45

T h r T h r T h r L e u P h e C y s A l a S e r A s p A l a L y s A l a T y r A s p T h r G l u
 50 55 60

eol f-seql . txt

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80
 Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95
 Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110
 Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

eol f-seql . txt

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Arg Leu
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

eol f-seql . txt

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Ala Ser Leu
610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 286

<211> 861

<212> PRT

<213> Human immunodeficiency virus type 1

<400> 286

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

eol f-seql . txt

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

eol f-seq1 . txt

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560 565

Leu Arg Ala Ile Gl u Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Arg Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Gly Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gl n Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Gl u
 725 730 735

Gly Ile Gl u Gl u Gl u Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Gl u Leu Leu Gly Arg Arg Gly Trp Gl u Ala Leu Lys Tyr
 785 790 795 800

eol f-seq1 . txt

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 287
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1

<400> 287

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
165 170 175

Gl u Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205

Ala Cys Pro Lys Val Ser Phe Gl u Pro Ile Pro Ile Hi s Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Gl u
 260 265 270

Gl u Gl u Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Gl u Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Ala Hi s Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
 340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Glu
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Glu Ala Arg Glu Leu Leu Ser Gly Ile Val Glu Glu Glu Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Glu Ala Glu Glu His Leu Leu Glu Leu Thr Val Trp
 565 570 575

Gly Ile Lys Glu Leu Glu Ala Arg Ile Leu Ala Val Glu Arg Arg Leu
 580 585 590

Lys Asp Glu Glu Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Ser Ser Leu
 610 615 620

Glu Glu Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Glu Asn
 645 650 655

Glu Glu Glu Lys Asn Glu Glu Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Glu Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His 725 Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg 745 Asp Arg Asp Arg Ser Ile Arg 750

Leu Val Asn Gly Ser Leu Ala Leu 760 Ile Trp Asp Asp Leu Arg Ser Leu 765

Cys Leu Phe Ser Tyr His 775 Arg Leu Arg Asp Leu Leu 780 Leu Ile Val Thr

Arg Ile Val Glu Leu Leu 790 Gly Arg Arg Gly Trp 795 Glu Ala Leu Lys Tyr 800

Trp Trp Asn Leu Leu 805 Gln Tyr Trp Ser Gln 810 Glu Leu Lys Asn Ser Ala 815

Val Ser Leu Leu 820 Asn Ala Thr Ala 825 Ile Ala Val Ala Glu Gly Thr Asp 830

Arg Val Ile 835 Glu Val Val Gln Gly Ala 840 Cys Arg Ala 845 Ile Arg His Ile

Pro Arg 850 Arg Ile Arg Gln Gly 855 Leu Glu Arg Ile Leu Leu 860

<210> 288

<211> 861

<212> PRT

<213> Human immunodeficiency virus type 1

<400> 288

Met Arg Val Lys 5 Glu Lys Tyr Gln His 10 Leu Trp Arg Trp Gly Trp Lys 15

Trp Gly Thr 20 Met Leu Leu Gly Ile 25 Leu Met Ile Cys Ser Ala Thr Glu 30

Lys Leu Trp 35 Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala 45

Thr Thr Thr 50 Leu Phe Cys Ala 55 Ser Asp Ala Lys Ala Tyr Asp Thr Glu 60

Val His Asn Val Trp 70 Ala Thr His Ala Cys Val 75 Pro Thr Asp Pro Asn 80

Pro Gln Glu Val 85 Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp

100

105

110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125
 Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140
 Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165
 Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175
 Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190
 Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205
 Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly

370

375

Gly 385 Glu Phe Phe Tyr Cys 390 Asn Ser Thr Gl n Leu Phe 395 Asn Ser Thr Trp 400
Phe Asn Ser Thr Trp 405 Ser Thr Gl u Gly Ser 410 Asn Asn Thr Gl u Gly Ser 415
Asp Thr Ile Thr 420 Leu Pro Cys Arg Ile 425 Lys Gl n Phe Ile Asn Met Trp 430
Gl n Gl u Val 435 Gly Lys Ala Met Tyr 440 Ala Pro Pro Ile Ser Gly Gl n Ile
Arg Cys 450 Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly 460
Asn 465 Asn Asn Asn Gly Ser 470 Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met 480
Arg Asp Asn Trp Arg 485 Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile 495
Glu Pro Leu Gly 500 Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n 510
Arg Gl u Lys 515 Arg Ala Val Gly Ile 520 Gly Ala Leu Phe Leu Gly Phe Leu 525
Gly Ala Ala Gly Ser Thr Met 535 Gly Ala Arg Ser Met Thr Leu Thr Val 540
Gl n Ala Arg Gl n Leu Leu 550 Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu 560
Leu Arg Ala Ile Gl u Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp 570 575
Gly Ile Lys Gl n Leu Gl n Ala Arg Ile 585 Leu Ala Val Gl u Arg Gly Leu 590
Lys Asp Gl n Gl n Leu Leu Gly Ile 600 Trp Gly Cys Ser Gly Lys Leu Ile 605
Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu 610 620
Glu Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile 625 630 635 640
Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn

Gln Gln Glu Lys 660 Asn Glu Gln Glu Leu 665 Leu Glu Leu Asp Lys 670 Trp Ala

Ser Leu Trp 675 Asn Trp Phe Asn Ile 680 Thr Asn Trp Leu Trp 685 Tyr Ile Lys

Ile Phe 690 Ile Met Ile Val Gly 695 Gly Leu Val Gly Leu 700 Arg Ile Val Phe

Ala Val 705 Leu Ser Ile Val 710 Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu 720

Ser Phe Gln Thr His 725 Leu Pro Thr Pro Arg 730 Gly Pro Asp Arg Pro Glu 735

Gly Ile Glu 740 Glu Glu Gly Gly Glu Arg 745 Asp Arg Asp Arg Ser Ile Arg 750

Leu Val 755 Asn Gly Ser Leu Ala Leu 760 Ile Trp Asp Asp Leu Arg Ser Leu 765

Cys Leu 770 Phe Ser Tyr His Arg 775 Leu Arg Asp Leu Leu 780 Leu Ile Val Thr

Arg Ile Val 785 Glu Leu Leu 790 Gly Arg Arg Gly Trp 795 Glu Ala Leu Lys Tyr 800

Trp Trp Asn Leu 805 Leu Gln Tyr Trp Ser Gln 810 Glu Leu Lys Asn Ser Ala 815

Val Ser Leu 820 Leu Asn Ala Thr Ala Ile 825 Ala Val Ala Glu Gly Thr Asp 830

Arg Val 835 Ile Glu Val Val Gln Gly 840 Ala Cys Arg Ala Ile 845 Arg His Ile

Pro Arg 850 Arg Ile Arg Gln Gly 855 Leu Glu Arg Ile Leu 860 Leu

<210> 289
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 289

Met Arg Val 1 Lys 5 Glu Lys Tyr Gln His 10 Leu Trp Arg Trp Gly Trp Lys 15

Trp Gly Thr 20 Met 25 Leu Leu Gly Ile Leu 25 Met Ile Cys Ser Ala Thr Glu 30

eol f-seql . txt

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
290 295 300

eol f-seql . txt

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
305 310 315 320

Al a Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Al a Hi s Cys
325 330 335

Asn Ile Ser Arg Al a Lys Trp Asn Al a Thr Leu Lys Gl n Ile Al a Ser
340 345 350

Lys Leu Arg Gl u Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
355 360 365

Ser Ser Gly Gly Asp Pro Gl u Ile Val Thr Hi s Ser Phe Asn Cys Gly
370 375 380

Gly Gl u Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Gl u Gly Ser Asn Asn Thr Gl u Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
420 425 430

Gl n Gl u Val Gly Lys Al a Met Tyr Al a Pro Pro Ile Ser Gly Gl n Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Gl u Pro Leu Gly Val Al a Pro Thr Lys Al a Lys Arg Arg Val Val Gl n
500 505 510

Arg Gl u Lys Arg Al a Val Gly Ile Gly Al a Leu Phe Leu Gly Phe Leu
515 520 525

Gly Al a Al a Gly Ser Thr Met Gly Al a Arg Ser Met Thr Leu Thr Val
530 535 540

Gl n Al a Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
545 550 555 560

Leu Arg Al a Ile Gl u Al a Gl n Gl n Hi s Leu Leu Gl n Leu Thr Val Trp
565 570 575

eol f-seql . txt

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Leu Leu
 580 585 590
 Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605
 Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620
 Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640
 Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655
 Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670
 Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685
 Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 700
 Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720
 Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735
 Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750
 Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765
 Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780
 Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
 785 790 795 800
 Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
 805 810 815
 Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
 820 825 830
 Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
 835 840 845

eol f-seql . txt

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
 850 855 860

<210> 290
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 290

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

eol f-seq1 . txt

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240
 Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255
 Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 270
 Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285
 Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300
 Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320
 Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335
 Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350
 Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365
 Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380
 Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400
 Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415
 Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430
 Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460
 Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480
 Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

eol f-seql . txt

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510 515

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540 545

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555

Leu Arg Ala Ile Gl u Ala Gl n Gl n His Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Ala Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gl n Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gl n Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Gl u
 725 730 735

Gly Ile Gl u Gl u Gl u Gly Gly Gl u Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

eol f-seq1 . txt

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 291
<211> 861
<212> PRT
<213> Human immunodeficiency virus type 1
<400> 291

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gl n Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gl n
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile Hi s Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gl n Cys Thr Hi s Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gl n Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gl n Leu Asn Gl n Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gl n Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gl n Ala Hi s Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gl n Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gl n Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gl n
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr Hi s Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gl n Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gl n Phe Ile Asn Met Trp
 420 425 430

Gl n Gl u Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gl n Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Gl u Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Gl u Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Gl u Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gl n
 500 505 510

Arg Gl u Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gl n Ala Arg Gl n Leu Leu Ser Gly Ile Val Gl n Gl n Gl n Asn Asn Leu
 545 550 555 560

Leu Arg Ala Ile Gl u Ala Gl n Gl n Hi s Leu Leu Gl n Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gl n Leu Gl n Ala Arg Ile Leu Ala Val Gl u Arg Phe Leu
 580 585 590

Lys Asp Gl n Gl n Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Gl u Gl n Ile Trp Asn Asn Met Thr Trp Met Gl u Trp Asp Arg Gl u Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile Hi s Ser Leu Ile Gl u Gl u Ser Gl n Asn
 645 650 655

Gl n Gl n Gl u Lys Asn Gl u Gl n Gl u Leu Leu Gl u Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

I l e P h e I l e M e t I l e V a l G l y G l y L e u V a l G l y L e u A r g I l e V a l P h e
 690 695 700

A l a V a l L e u S e r I l e V a l A s n A r g V a l A r g G l n G l y T y r S e r P r o L e u
 705 710 715 720

S e r P h e G l n T h r H i s L e u P r o T h r P r o A r g G l y P r o A s p A r g P r o G l u
 725 730 735

G l y I l e G l u G l u G l u G l y G l y G l u A r g A s p A r g A s p A r g S e r I l e A r g
 740 745 750

L e u V a l A s n G l y S e r L e u A l a L e u I l e T r p A s p A s p L e u A r g S e r L e u
 755 760 765

C y s L e u P h e S e r T y r H i s A r g L e u A r g A s p L e u L e u L e u I l e V a l T h r
 770 775 780

A r g I l e V a l G l u L e u L e u G l y A r g A r g G l y T r p G l u A l a L e u L y s T y r
 785 790 800

T r p T r p A s n L e u L e u G l n T y r T r p S e r G l n G l u L e u L y s A s n S e r A l a
 805 810 815

V a l S e r L e u L e u A s n A l a T h r A l a I l e A l a V a l A l a G l u G l y T h r A s p
 820 825 830

A r g V a l I l e G l u V a l V a l G l n G l y A l a C y s A r g A l a I l e A r g H i s I l e
 835 840 845

P r o A r g A r g I l e A r g G l n G l y L e u G l u A r g I l e L e u L e u
 850 855 860

<210> 292
 <211> 861
 <212> PRT
 <213> Human immunodeficiency virus type 1
 <400> 292

M e t A r g V a l L y s G l u L y s T y r G l n H i s L e u T r p A r g T r p G l y T r p L y s
 1 5 10 15

T r p G l y T h r M e t L e u L e u G l y I l e L e u M e t I l e C y s S e r A l a T h r G l u
 20 25 30

L y s L e u T r p V a l T h r V a l T y r T y r G l y V a l P r o V a l T r p L y s G l u A l a
 35 40 45

T h r T h r T h r L e u P h e C y s A l a S e r A s p A l a L y s A l a T y r A s p T h r G l u
 50 55 60

V a l H i s A s n V a l T r p A l a T h r H i s A l a C y s V a l P r o T h r A s p P r o A s n
 Page 292

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
545 550 555 560

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Arg
580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu

610

615

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
785 790 795 800

Trp Trp Asn Leu Leu Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala
805 810 815

Val Ser Leu Leu Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp
820 825 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile
835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu
850 855 860

<210> 293
<211> 881
<212> PRT
<213> Simian immunodeficiency virus

eol f-seql . txt

<220>

<221> mi sc_feature

<222> (736)..(736)

<223> Xaa can be any naturally occurring amino acid

<400> 293

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Val Tyr Gly Ile Tyr Cys Thr Gl n Tyr Val Thr Val Phe Tyr Gly Val
20 25 30

Pro Ala Trp Arg Asn Ala Thr Ile Pro Leu Phe Cys Ala Thr Lys Asn
35 40 45

Arg Asp Thr Trp Gly Thr Thr Gl n Cys Leu Pro Asp Asn Gly Asp Tyr
50 55 60

Ser Gl u Leu Ala Leu Asn Val Thr Gl u Ser Phe Asp Ala Trp Gl u Asn
65 70 75 80

Thr Val Thr Gl u Gl n Ala Ile Gl u Asp Val Trp Gl n Leu Phe Gl u Thr
85 90 95

Ser Ile Lys Pro Cys Val Lys Leu Ser Pro Leu Cys Ile Thr Met Arg
100 105 110

Cys Asn Lys Ser Gl u Thr Asp Arg Trp Gly Leu Thr Lys Ser Ser Thr
115 120 125

Thr Ile Thr Thr Ala Ala Pro Thr Ser Ala Pro Val Ser Gl u Lys Ile
130 135 140

Asp Met Val Asn Gl u Thr Ser Ser Cys Ile Ala Gl n Asn Asn Cys Thr
145 150 155 160

Gly Leu Gl u Gl n Gl u Gl n Met Ile Ser Cys Lys Phe Thr Met Thr Gly
165 170 175

Leu Lys Arg Asp Lys Thr Lys Gl u Tyr Asn Gl u Thr Trp Tyr Ser Thr
180 185 190

Asp Leu Val Cys Gl u Gl n Gly Asn Ser Thr Asp Asn Gl u Ser Arg Cys
195 200 205

Tyr Met Asn His Cys Asn Thr Ser Val Ile Gl n Gl u Ser Cys Asp Lys
210 215 220

His Tyr Trp Asp Thr Ile Arg Phe Arg Tyr Cys Ala Pro Pro Gly Tyr
225 230 235 240

Ala Leu Leu Arg Cys₂₄₅ Asn Asp Thr Asn Tyr₂₅₀ Ser Gly Phe Met Pro Lys₂₅₅

Cys Ser Lys Val₂₆₀ Val Val Ser Ser Cys₂₆₅ Thr Arg Met Met Glu₂₇₀ Thr Gln

Thr Ser Thr₂₇₅ Trp Phe Gly Phe Asn₂₈₀ Gly Thr Arg Ala Glu₂₈₅ Asn Arg Thr

Tyr Ile₂₉₀ Tyr Trp His Gly Arg₂₉₅ Asp Asn Arg Thr Ile₃₀₀ Ile Ser Leu Asn

Lys Tyr Tyr Asn Leu Thr₃₁₀ Met Lys Cys Arg Arg₃₁₅ Pro Gly Asn Lys Thr₃₂₀

Val Leu Pro Val Thr₃₂₅ Ile Met Ser Gly Leu₃₃₀ Val Phe His Ser Gln₃₃₅ Pro

Ile Asn Asp Arg₃₄₀ Pro Lys Gln Ala Trp₃₄₅ Cys Trp Phe Gly Gly₃₅₀ Lys Trp

Lys Asp Ala₃₅₅ Ile Lys Glu Val Lys₃₆₀ Gln Thr Ile Val Lys₃₆₅ His Pro Arg

Tyr Thr Gly Thr Asn Asn Thr₃₇₅ Asp Lys Ile Asn Leu Thr Ala Pro Gly

Gly Gly Asp Pro Glu Val₃₉₀ Thr Phe Met Trp Thr₃₉₅ Asn Cys Arg Gly Glu₄₀₀

Phe Leu Tyr Cys Lys₄₀₅ Met Asn Trp Phe Leu₄₁₀ Asn Trp Val Glu Asp Arg₄₁₅

Asp Val Thr Thr₄₂₀ Gln Arg Pro Lys Glu₄₂₅ Arg His Arg Arg Asn Tyr Val₄₃₀

Pro Cys His Ile Arg Gln Ile Ile₄₄₀ Asn Thr Trp His Lys₄₄₅ Val Gly Lys

Asn Val Tyr Leu Pro Pro Arg₄₅₅ Glu Gly Asp Leu Thr₄₆₀ Cys Asn Ser Thr

Val Thr Ser Leu Ile Ala Asn Ile Asp Trp Thr₄₇₅ Asp Gly Asn Gln Thr₄₈₀

Ser Ile Thr Met Ser₄₈₅ Ala Glu Val Ala Glu₄₉₀ Leu Tyr Arg Leu Glu₄₉₅ Leu

Gly Asp Tyr Lys₅₀₀ Leu Val Glu Ile Thr₅₀₅ Pro Ile Gly Leu Ala₅₁₀ Pro Thr

Asp Val Lys Arg Tyr Thr Thr Gly Gly Thr Ser Arg Asn Lys Arg Gly
 515 520 525

Val Phe Val Leu Gly Phe Leu Gly Phe Leu Ala Thr Ala Gly Ser Ala
 530 535 540 545

Met Gly Ala Ala Ser Leu Thr Leu Thr Ala Gln Ser Arg Thr Leu Leu
 545 550 555 560 565

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
 565 570 575

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
 580 585 590

Thr Arg Val Thr Ala Ile Glu Lys Tyr Arg Lys Asp Gln Ala Gln Leu
 595 600 605

Asn Ala Trp Gly Cys Ala Phe Arg Gln Val Cys His Thr Thr Val Pro
 610 615 620

Trp Pro Asn Ala Ser Leu Thr Pro Asp Trp Asn Asn Asp Thr Trp Gln
 625 630 635 640 645

Glu Trp Glu Arg Lys Val Asp Phe Leu Glu Glu Asn Ile Thr Ala Leu
 645 650 655

Leu Glu Glu Ala Gln Ile Gln Gln Glu Lys Asn Met Tyr Glu Leu Gln
 660 665 670

Lys Leu Asn Ser Trp Asp Val Phe Gly Asn Trp Phe Asp Leu Ala Ser
 675 680 685

Trp Ile Lys Tyr Ile Gln Tyr Gly Ile Tyr Val Val Val Gly Val Ile
 690 695 700

Leu Leu Arg Ile Val Ile Tyr Ile Val Gln Met Leu Ala Lys Leu Arg
 705 710 715 720

Gln Gly Tyr Arg Pro Val Phe Ser Ser Pro Pro Ser Tyr Phe Gln Xaa
 725 730 735

Thr His Thr Gln Gln Asp Pro Ala Leu Pro Thr Arg Glu Gly Lys Glu
 740 745 750

Gly Asp Gly Gly Glu Gly Gly Gly Asn Ser Ser Trp Pro Trp Gln Ile
 755 760 765

Glu Tyr Ile His Phe Leu Ile Arg Gln Leu Ile Arg Leu Leu Thr Trp
 770 775 780

Leu Phe Ser Asn Cys Arg Thr Leu Leu Ser Arg Ala Tyr Gln Ile Leu
785 790 795 800

Gln Pro Ile Leu Gln Arg Leu Ser Ala Thr Leu Arg Arg Val Arg Glu
805 810 815

Val Leu Arg Thr Glu Leu Thr Tyr Leu Gln Tyr Gly Trp Ser Tyr Phe
820 825 830

His Glu Ala Val Gln Ala Gly Trp Arg Ser Ala Thr Glu Thr Leu Ala
835 840 845

Gly Ala Trp Arg Asp Leu Trp Glu Thr Leu Arg Arg Gly Gly Arg Trp
850 855 860

Ile Leu Ala Ile Pro Arg Arg Ile Arg Gln Gly Leu Glu Leu Thr Leu
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Leu

<210> 294
<211> 860
<212> PRT
<213> Human immunodeficiency virus type 2

<400> 294

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Ala Cys Leu Val Tyr Cys Ser Gln Tyr Val Thr Val Phe Tyr Gly Ile
20 25 30

Pro Ala Trp Lys Asn Ala Ser Ile Pro Leu Phe Cys Ala Thr Lys Asn
35 40 45

Arg Asp Thr Trp Gly Thr Ile Gln Cys Leu Pro Asp Asn Asp Asp Tyr
50 55 60

Gln Glu Ile Ile Leu Asn Val Thr Glu Ala Phe Asp Ala Trp Asn Asn
65 70 75 80

Thr Val Thr Glu Gln Ala Val Glu Asp Val Trp His Leu Phe Glu Thr
85 90 95

Ser Ile Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ala Met Asn
100 105 110

Cys Ser Arg Val Gln Gly Asn Thr Thr Thr Pro Asn Pro Arg Thr Ser
115 120 125

Ser Ser Thr Thr Ser Arg Pro Pro Thr Ser Ala Ala Ser Ile Ile Asn
Page 299

130

135

Gl u Thr Ser Asn Cys Ile Gl u Asn Asn Thr Cys Ala Gly Leu Gly Tyr
145 150 155 160

Gl u Gl u Met Met Gl n Cys Gl u Phe Asn Met Lys Gly Leu Gl u Gl n Asp
165 170 175

Lys Lys Arg Arg Tyr Lys Asp Thr Trp Tyr Leu Gl u Asp Val Val Cys
180 185 190

Asp Asn Thr Thr Ala Gly Thr Cys Tyr Met Arg His Cys Asn Thr Ser
195 200 205

Ile Ile Lys Gl u Ser Cys Asp Lys His Tyr Trp Asp Ala Met Arg Phe
210 215 220

Arg Tyr Cys Ala Pro Pro Gly Phe Ala Leu Leu Arg Cys Asn Asp Thr
225 230 235 240

Asn Tyr Ser Gly Phe Gl u Pro Lys Cys Thr Lys Val Val Ala Ala Ser
245 250 255

Cys Thr Arg Met Met Gl u Thr Gl n Thr Ser Thr Trp Phe Gly Phe Asn
260 265 270

Gly Thr Arg Ala Gl u Asn Arg Thr Tyr Ile Tyr Trp His Gly Arg Asp
275 280 285

Asn Arg Thr Ile Ile Ser Leu Asn Lys Tyr Tyr Asn Leu Thr Met Arg
290 295 300

Cys Lys Arg Pro Gly Asn Lys Thr Val Leu Pro Ile Thr Leu Met Ser
305 310 315 320

Gly Leu Val Phe His Ser Gl n Pro Ile Asn Thr Arg Pro Arg Gl n Ala
325 330 335

Trp Cys Arg Phe Gly Gly Arg Trp Arg Gl u Ala Met Gl n Gl u Val Lys
340 345 350

Gl n Thr Leu Val Gl n His Pro Arg Tyr Lys Gly Ile Asn Asp Thr Gly
355 360 365

Lys Ile Asn Phe Thr Lys Pro Gly Ala Gly Ser Asp Pro Gl u Val Ala
370 375 380

Phe Met Trp Thr Asn Cys Arg Gly Gl u Phe Leu Tyr Cys Asn Met Thr
385 390 395 400

Trp Phe Leu Asn Trp Val Gl u Asp Lys Asn Gl n Thr Arg Arg Asn Tyr

Cys His Ile Lys Gln Ile Ile Asn Thr Trp His Lys Val Gly Lys Asn
 420 425 430
 Val Tyr Leu Pro Pro Arg Glu Gly Glu Leu Ala Cys Glu Ser Thr Val
 435 440 445
 Thr Ser Ile Ile Ala Asn Ile Asp Ile Asp Lys Asn Arg Thr His Thr
 450 455 460
 Asn Ile Thr Phe Ser Ala Glu Val Ala Glu Leu Tyr Arg Leu Glu Leu
 465 470 475 480
 Gly Asp Tyr Lys Leu Ile Glu Ile Thr Pro Ile Gly Phe Ala Pro Thr
 485 490 495
 Asp Gln Arg Arg Tyr Ser Ser Thr Pro Val Arg Asn Lys Arg Gly Val
 500 505 510
 Phe Val Leu Gly Phe Leu Gly Phe Leu Ala Thr Ala Gly Ser Ala Met
 515 520 525
 Gly Ala Arg Ser Leu Thr Leu Ser Ala Gln Ser Arg Thr Leu Leu Ala
 530 535 540
 Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg Gln
 545 550 555 560
 Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln Ala
 565 570
 Arg Val Thr Ala Ile Glu Lys Tyr Arg Lys His Gln Ala Gln Leu Asn
 580 585 590
 Ser Trp Gly Cys Ala Phe Arg Gln Val Cys His Thr Thr Val Pro Trp
 595 600 605
 Val Asn Asp Ser Leu Ser Pro Asp Trp Lys Asn Met Thr Trp Gln Glu
 610 615 620
 Trp Glu Lys Gln Val Arg Tyr Leu Glu Ala Asn Ile Ser Gln Ser Leu
 625 630 635 640
 Glu Glu Ala Gln Ile Gln Gln Glu Lys Asn Met Tyr Glu Leu Gln Lys
 645 650 655
 Leu Asn Ser Trp Asp Ile Leu Gly Asn Trp Phe Asp Leu Thr Ser Trp
 660 665 670
 Val Lys Tyr Ile Gln Tyr Gly Val His Ile Val Val Gly Ile Ile Ala

675

680

685

Leu Arg Ile Ala Ile Tyr Val Val Gln Leu Leu Ser Arg Phe Arg Lys
 690 695 700

Gly Tyr Arg Pro Val Phe Ser Ser Pro Pro Gly Tyr Leu Gln Gln Ile
 705 710 715 720

His Ile His Lys Asp Arg Gly Gln Pro Ala Asn Glu Gly Thr Glu Glu
 725 730 735

Asp Val Gly Gly Asp Ser Gly Tyr Asp Leu Trp Pro Trp Pro Ile Asn
 740 745 750

Tyr Val Gln Phe Leu Ile His Leu Leu Thr Arg Leu Leu Ile Gly Leu
 755 760 765

Tyr Asn Ile Cys Arg Asp Leu Leu Ser Lys Asn Ser Pro Thr Arg Arg
 770 775 780

Leu Ile Ser Gln Ser Leu Thr Ala Ile Arg Asp Trp Leu Arg Leu Lys
 785 790 795 800

Ala Ala Gln Leu Gln Tyr Gly Cys Glu Trp Ile Gln Glu Ala Phe Gln
 805 810 815

Ala Phe Ala Arg Thr Thr Arg Glu Thr Leu Ala Gly Ala Trp Gly Trp
 820 825 830

Leu Trp Glu Ala Ala Arg Arg Ile Gly Arg Gly Ile Leu Ala Val Pro
 835 840 845

Arg Arg Ile Arg Gln Gly Ala Glu Leu Ala Leu Leu
 850 855 860

<210> 295
 <211> 36
 <212> DNA
 <213> Artificial sequence

<220>
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<400> 295
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36

<210> 296
 <211> 34
 <212> DNA
 <213> Artificial sequence

<220>
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 lentiviral ENV protein

<400> 296
gtatacgcgt ttataattct tgttcattct tttc 34

<210> 297
<211> 35
<212> DNA
<213> Arti f i c i a l s e q u e n c e

<220>
<223> o l i g o n u c l e o t i d e f o r i n t r o d u c i n g m u t a t i o n w i t h i n t h e I S U o f
l e n t i v i r a l E N V p r o t e i n

<400> 297
gtatacgcgt ttacatgta ttccaaatct gttcc 35

<210> 298
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<220>
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l e n t i v i r a l E N V p r o t e i n

<400> 298
gtatacgcgt ttaagcattc caaggcacag c 31

<210> 299
<211> 33
<212> DNA
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<220>
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l e n t i v i r a l E N V p r o t e i n

<400> 299
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<210> 300
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<220>
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l e n t i v i r a l E N V p r o t e i n

<400> 300
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<210> 301
<211> 34
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l e n t i v i r a l E N V p r o t e i n

<400> 301
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 l e n t i v i r a l E N V p r o t e i n

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 l e n t i v i r a l E N V p r o t e i n

 <400> 303
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 <211> 26
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 <220>
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 l e n t i v i r a l E N V p r o t e i n

 <400> 304
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 <211> 26
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 l e n t i v i r a l E N V p r o t e i n

 <400> 305
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<210> 306
 <211> 28
 <212> DNA
 <213> Arti f i c i a l s e q u e n c e

 <220>
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 l e n t i v i r a l E N V p r o t e i n

 <400> 306
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<210> 307

<211> 28
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 l e n t i v i r a l E N V p r o t e i n

 <400> 307
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<210> 308
 <211> 34
 <212> DNA
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 <223> o l i g o n u c l e o t i d e f o r i n t r o d u c i n g m u t a t i o n w i t h i n t h e I S U o f
 l e n t i v i r a l E N V p r o t e i n

 <400> 308
 ctttaggtat ctttcctag ccaggattct tgcc 34

<210> 309
 <211> 34
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 l e n t i v i r a l E N V p r o t e i n

 <400> 309
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<210> 310
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 l e n t i v i r a l E N V p r o t e i n

 <400> 310
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<210> 311
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 <220>
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 l e n t i v i r a l E N V p r o t e i n

 <400> 311
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<210> 312
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 <213> Arti f i c i a l s e q u e n c e

<220>
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<400> 312
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<210> 313
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 <212> DNA
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<220>
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 lentiviral ENV protein

<400> 313
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<210> 314
 <211> 35
 <212> DNA
 <213> Artificial sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 lentiviral ENV protein

<400> 314
 ggagctgttg atcctttcgg tatctttcca cagcc 35

<210> 315
 <211> 35
 <212> DNA
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<220>
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 lentiviral ENV protein

<400> 315
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<210> 316
 <211> 30
 <212> DNA
 <213> Artificial sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 lentiviral ENV protein

<400> 316
 gagctgttga tcccttaggt atctttccac 30

<210> 317
 <211> 30
 <212> DNA
 <213> Artificial sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of

Lentiviral ENV protein

<400> 317
gtggaagat acctaaggga tcaacagctc 30

<210> 318
<211> 28
<212> DNA
<213> Artificial sequence

<220>
<223> oligonucleotide for introducing mutation within the ISU of
Lentiviral ENV protein

<400> 318
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<210> 319
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Lentiviral ENV protein

<400> 319
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Lentiviral ENV protein

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Lentiviral ENV protein

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Lentiviral ENV protein

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<400> 333
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<210> 334
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<400> 334
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lenti viral ENV protei n

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<210> 336
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<223> oligonucleotide for introducing mutation within the ISU of
lenti viral ENV protei n

<400> 336
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<210> 337
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lenti viral ENV protei n

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<210> 338
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<400> 338
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<210> 339
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<210> 340
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lentiviral ENV protein

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<210> 341
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lentiviral ENV protein

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<210> 342
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lentiviral ENV protein

<400> 342
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<210> 343
<211> 31
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lentiviral ENV protein

<400> 343
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<210> 346
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<220>
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<400> 346
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<210> 347
 <211> 44
 <212> DNA
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<220>
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<400> 347
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<210> 348
 <211> 44
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 <213> Arti fici al sequence

<220>
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<400> 348
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<210> 349
<211> 32
<212> DNA
<213> Artificial sequence

<220>
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lentiviral ENV protein

<400> 349
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<210> 350
<211> 25
<212> DNA
<213> Artificial sequence

<220>
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lentiviral ENV protein

<400> 350
ccgctcagtc ccgaacttta ttggc 25

<210> 351
<211> 22
<212> DNA
<213> Artificial sequence

<220>
<223> oligonucleotide for introducing mutation within the ISU of
lentiviral ENV protein

<400> 351
ggtggggaag agaactgg cc 22

<210> 352
<211> 45
<212> DNA
<213> Artificial sequence

<220>
<223> oligonucleotide for introducing mutation within the ISU of
lentiviral ENV protein

<400> 352
cagactaggg tcaactgcat ccgcaagtac ttaaaggacc aggcg 45

<210> 353
<211> 45
<212> DNA
<213> Artificial sequence

<220>
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lentiviral ENV protein

<400> 353
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<210> 354
<211> 45

<220>
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<400> 359
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<210> 360
 <211> 45
 <212> DNA
 <213> Artificial sequence

<220>
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 lentiviral ENV protein

<400> 360
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<210> 361
 <211> 45
 <212> DNA
 <213> Artificial sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 lentiviral ENV protein

<400> 361
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<210> 362
 <211> 39
 <212> DNA
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<220>
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 lentiviral ENV protein

<400> 362
 gagaagtact taaagcgcca ggcgagctg aatgcttg 39

<210> 363
 <211> 39
 <212> DNA
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<220>
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 lentiviral ENV protein

<400> 363
 attcagctgc gcctggcgct ttaagtactt ctcgatggc 39

<210> 364
 <211> 36
 <212> DNA
 <213> Artificial sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 lentiviral ENV protein

<400> 364
 acatggccca gccggccgct gggatagtg c agcaac 36

<210> 365
 <211> 35
 <212> DNA
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<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 Lentiviral ENV protein

<400> 365
 gtatacgcgt ttaaacgca catccccaag cattc 35

<210> 366
 <211> 49
 <212> PRT
 <213> Artificial sequence

<220>
 <223> derived from the ISU domain of HIV1

<400> 366
 Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15
 Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30
 Ala Arg Ile Leu Ala Val Glu Arg Tyr Leu Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 367
 <211> 49
 <212> PRT
 <213> Artificial sequence

<220>
 <223> derived from the ISU domain of HIV1

<400> 367
 Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
 1 5 10 15
 Gln Gln His Leu Leu Lys Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30
 Ala Arg Ile Leu Ala Val Glu Arg Tyr Leu Lys Asp Gln Gln Leu Leu
 35 40 45

Gly

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<210> 368
<211> 49
<212> PRT
<213> Arti ficial sequence

<220>
<223> derived from the ISU domain of HIV1
<400> 368

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Val Glu Arg Tyr Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 369
<211> 49
<212> PRT
<213> Arti ficial sequence

<220>
<223> derived from the ISU domain of HIV1
<400> 369

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Ile Leu Ala Val Glu Arg Tyr Leu Lys Asp Gln Gln Leu Leu
35 40 45

Gly

<210> 370
<211> 49
<212> PRT
<213> Arti ficial sequence

<220>
<223> derived from the ISU domain of HIV1
<400> 370

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln Gln Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Arg
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Tyr Leu Arg Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 371
 <211> 49
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> derived from the ISU domain of HIV1

<400> 371

Ser Gly Ile Val Gln Gln Gln Ser Asn Leu Leu Arg Ala Ile Gln Ala
 1 5 10 15

Arg Gln His Met Leu Gln Leu Thr Val Trp Gly Ile Lys Gln Leu Gln
 20 25 30

Ala Arg Val Leu Ala Val Glu Arg Tyr Leu Arg Asp Gln Gln Leu Leu
 35 40 45

Gly

<210> 372
 <211> 49
 <212> PRT
 <213> Artificial sequence

<220>
 <223> derived from the ISU domain of SIV

<400> 372

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Lys Ala Ile Glu Ala
 1 5 10 15

Gln Gln His Leu Leu Gln Leu Ser Ile Trp Gly Val Lys Gln Leu Gln
 20 25 30

Ala Arg Leu Leu Ala Val Glu Arg Tyr Leu Gln Asp Gln Gln Ile Leu
 35 40 45

Gly

<210> 373
 <211> 49
 <212> PRT
 <213> Artificial sequence

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<220>

<223> derived from the ISU domain of SIV

<400> 373

Ser Gly Ile Val Gln Gln Gln Asn Asn Leu Leu Arg Ala Ile Glu Ala
1 5 10 15

Gln Gln His Leu Leu Gln Leu Ser Val Trp Gly Ile Lys Gln Leu Gln
20 25 30

Ala Arg Val Leu Ala Ile Glu Arg Tyr Leu Arg Asp Gln Gln Ile Leu
35 40 45

Gly

<210> 374

<211> 49

<212> PRT

<213> Artificial sequence

<220>

<223> derived from the ISU domain of SIV

<400> 374

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Ser Ala Ile Glu Lys Tyr Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn

<210> 375

<211> 49

<212> PRT

<213> Artificial Sequence

<220>

<223> derived from the ISU domain of SIV

<400> 375

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15

Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30

Thr Arg Val Thr Ala Ile Glu Lys Tyr Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn

<210> 376
<211> 49
<212> PRT
<213> Arti ficial sequence

<220>
<223> derived from the ISU domain of HIV2

<400> 376

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15
Gln Gln Glu Met Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30
Ala Arg Val Thr Ala Ile Glu Lys Tyr Leu Lys Asp Gln Ala Gln Leu
35 40 45

Asn

<210> 377
<211> 49
<212> PRT
<213> Arti ficial Sequence

<220>
<223> derived from the ISU domain of HIV2

<400> 377

Ala Gly Ile Val Gln Gln Gln Gln Gln Leu Leu Asp Val Val Lys Arg
1 5 10 15
Gln Gln Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gln
20 25 30
Thr Arg Val Thr Ala Ile Glu Lys Tyr Leu Lys Asp Gln Ala Leu Leu
35 40 45

Asn

<210> 378
<211> 31
<212> DNA
<213> Arti ficial sequence

<220>
<223> Oligonucleotide used for introducing a second mutation

<400> 378
ctggctgtgg aagcacgcct aaaggatcaa c

<210> 379
<211> 31
<212> DNA
<213> Arti f i c i a l s e q u e n c e

<220>
<223> O l i g o n u c l e o t i d e u s e d f o r i n t r o d u c i n g a s e c o n d m u t a t i o n

<400> 379
g t t g a t c c t t t a g g c g t g c t t c c a c a g c c a g 31

<210> 380
<211> 31
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<220>
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<400> 380
c t g g c t g t g g a a g a a c g c c t a a g g a t c a a c 31

<210> 381
<211> 31
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<220>
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<400> 381
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<210> 382
<211> 31
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<213> Arti f i c i a l s e q u e n c e

<220>
<223> O l i g o n u c l e o t i d e u s e d f o r i n t r o d u c i n g a s e c o n d m u t a t i o n

<400> 382
c t g g c t g t g g a a g c t g c c c t a a g g a t c a a c 31

<210> 383
<211> 31
<212> DNA
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<220>
<223> O l i g o n u c l e o t i d e u s e d f o r i n t r o d u c i n g a s e c o n d m u t a t i o n

<400> 383
g t t g a t c c t t t a g g g c a g c t t c c a c a g c c a g 31

<210> 384
<211> 31
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<213> Arti f i c i a l s e q u e n c e

<220>

<223> Oligonucleotide used for introducing a second mutation

<400> 384
ctggctgtgg aagaagccct aaaggatcaa c 31

<210> 385
<211> 31
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<213> Artificial sequence

<220>
<223> Oligonucleotide used for introducing a second mutation

<400> 385
gttgatcctt tagggcttct tccacagcca g 31

<210> 386
<211> 31
<212> DNA
<213> Artificial sequence

<220>
<223> Oligonucleotide used for introducing a second mutation

<400> 386
ctggctgtgg aaagcgcct aaaggatcaa c 31

<210> 387
<211> 31
<212> DNA
<213> Artificial sequence

<220>
<223> Oligonucleotide used for introducing a second mutation

<400> 387
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<210> 388
<211> 31
<212> DNA
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<220>
<223> Oligonucleotide used for introducing a second mutation

<400> 388
ctggctgtgg aacagccct aaaggatcaa c 31

<210> 389
<211> 31
<212> DNA
<213> Artificial sequence

<220>
<223> Oligonucleotide used for introducing a second mutation

<400> 389
gttgatcctt tagggctggt tccacagcca g 31

<210> 390
<211> 31

<212> DNA
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 <220>
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 <400> 390
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<210> 391
 <211> 31
 <212> DNA
 <213> Artificial sequence

 <220>
 <223> Oligonucleotide used for introducing a second mutation

 <400> 391
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<210> 392
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 <400> 392
 ctggctgtgg aagaatacct aaaggatcaa c 31

<210> 393
 <211> 31
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 <400> 393
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<210> 394
 <211> 31
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 <220>
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 <400> 394
 ctggctgtgg aaagctacct aaaggatcaa c 31

<210> 395
 <211> 31
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 <400> 395
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<210> 396
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 <400> 396
 ctggctgtgg aaacatacct aaaggatcaa c 31

<210> 397
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<210> 398
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 <400> 398
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<210> 399
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<210> 400
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 <213> Arti f i c i a l s e q u e n c e
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 <223> O l i g o n u c l e o t i d e u s e d f o r i n t r o d u c i n g a s e c o n d m u t a t i o n
 <400> 400
 gctagttgga gtaatggatc tctggaacag atttgg 36

<210> 401
 <211> 36
 <212> DNA
 <213> Arti f i c i a l s e q u e n c e
 <220>

<223> Oligonucleotide used for introducing a second mutation
 <400> 401
 ccaaatctgt tccagagatc cattactcca actagc 36

<210> 402
 <211> 36
 <212> DNA
 <213> Artificial sequence

<220>
 <223> Oligonucleotide used for introducing a second mutation
 <400> 402
 gctagttgga gtaattcatc tctggaacag atttgg 36

<210> 403
 <211> 36
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Oligonucleotide used for introducing a second mutation
 <400> 403
 ccaaatctgt tccagagatg aattactcca actagc 36

<210> 404
 <211> 54
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 lentiviral ENV protein

<220>
 <221> misc_feature
 <222> (1)..(5)
 <223> n i s a , c , g , o r t

<400> 404
 nnnnctcga gaccggtcca actagaacca tgagagtgaa ggagaaatat cagc 54

<210> 405
 <211> 31
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> oligonucleotide for introducing mutation within the ISU of
 lentiviral ENV protein

<220>
 <221> misc_feature
 <222> (1)..(5)
 <223> n i s a , c , g , o r t

<400> 405
 nnnnacgcg ttcaatatcc ctgcctaact c 31

<210> 406
<211> 34
<212> DNA
<213> Artificial Sequence

<220>
<223> oligonucleotide for introducing mutation within the ISU of
Lentiviral ENV protein

<220>
<221> misc_feature
<222> (1)..(5)
<223> n i s a , c , g , o r t

<400> 406
nnnnnacgcg tttatagcaa aatcctttcc aagc

34

<210> 407
<211> 8
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of Lentiviral ENV ISU

<400> 407

Ala Val Glu Ala Ala Leu Lys Asp
1 5

<210> 408
<211> 8
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of Lentiviral ENV ISU

<400> 408

Ala Val Glu Glu Ala Leu Lys Asp
1 5

<210> 409
<211> 8
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of Lentiviral ENV ISU

<400> 409

Ala Val Glu Ser Ala Leu Lys Asp
1 5

<210> 410
<211> 8
<212> PRT
<213> Artificial sequence

<220>
<223> fragment of Lentiviral ENV ISU

<400> 410

Al a Val Gl u Thr Al a Leu Lys Asp
1 5

<210> 411

<211> 8

<212> PRT

<213> Arti fici al sequence

<220>

<223> fragment of Lenti vi ral ENV ISU

<400> 411

Al a Ile Gl u Al a Al a Leu Lys Asp
1 5

<210> 412

<211> 8

<212> PRT

<213> Arti fici al sequence

<220>

<223> fragment of Lenti vi ral ENV ISU

<400> 412

Al a Ile Gl u Gl u Al a Leu Lys Asp
1 5

<210> 413

<211> 8

<212> PRT

<213> Arti fici al sequence

<220>

<223> fragment of Lenti vi ral ENV ISU

<400> 413

Al a Ile Gl u Ser Al a Leu Lys Asp
1 5

<210> 414

<211> 8

<212> PRT

<213> Arti fici al sequence

<220>

<223> fragment of Lenti vi ral ENV ISU

<400> 414

Al a Ile Gl u Thr Al a Leu Lys Asp
1 5

<210> 415

<211> 38

<212> PRT

<213> Arti fici al Sequence

<220>
 <223> derived from lenti viral ENV protein

<220>
 <221> MI SC_FEATURE
 <222> (2)..(2)
 <223> Xaa = I or V

<220>
 <221> MI SC_FEATURE
 <222> (4)..(4)
 <223> Xaa = K or R

<220>
 <221> mi sc_feature
 <222> (7)..(7)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> mi sc_feature
 <222> (10)..(11)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> mi sc_feature
 <222> (13)..(14)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> MI SC_FEATURE
 <222> (18)..(18)
 <223> Xaa = A or S

<220>
 <221> MI SC_FEATURE
 <222> (19)..(19)
 <223> Xaa = F or G

<220>
 <221> mi sc_feature
 <222> (20)..(21)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> mi sc_feature
 <222> (24)..(24)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> mi sc_feature
 <222> (26)..(26)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> mi sc_feature
 <222> (30)..(30)
 <223> Xaa can be any naturally occurring amino acid

<220>
 <221> MI SC_FEATURE
 <222> (31)..(35)
 <223> Xaa = no or any amino acid

<220>
 <221> MI SC_FEATURE
 <222> (36)..(36)
 <223> Xaa = N or S

eol f-seql . txt

<220>
<221> MI SC_FEATURE
<222> (37).. (37)
<223> Xaa = E, D, N or A

<220>
<221> MI SC_FEATURE
<222> (38).. (38)
<223> Xaa = S or T

<400> 415

Al a Xaa Gl u Xaa Tyr Leu Xaa Asp Gl n Xaa Xaa Leu Xaa Xaa Trp Gl y
1 5 10 15

Cys Xaa Xaa Xaa Xaa Cys Val Xaa Thr Xaa Val Pro Trp Xaa Xaa Xaa
20 25 30

Xaa Xaa Xaa Xaa Xaa Xaa
35

<210> 416
<211> 9
<212> PRT
<213> Arti fici al sequence

<220>
<223> fragment of lenti viral ENV I SU

<220>
<221> MI SC_FEATURE
<222> (2).. (2)
<223> Xaa = I or V

<220>
<221> MI SC_FEATURE
<222> (4).. (4)
<223> Xaa = K or R

<220>
<221> MI SC_FEATURE
<222> (5).. (5)
<223> Xaa is any natural ly occuring ami no acid or is mutated

<220>
<221> MI SC_FEATURE
<222> (6).. (6)
<223> Xaa is any natural ly occuring ami no acid or is mutated

<220>
<221> MI SC_FEATURE
<222> (7).. (7)
<223> Xaa can be any natural ly occuring ami no aci d

<400> 416

Al a Xaa Gl u Xaa Xaa Xaa Xaa Asp Gl n
1 5

<210> 417
<211> 861
<212> PRT

<213> Human immunodeficiency virus type 1

<400> 417

Met Arg Val Lys Glu Lys Tyr Gln His Leu Trp Arg Trp Gly Trp Lys
 1 5 10 15

Trp Gly Thr Met Leu Leu Gly Ile Leu Met Ile Cys Ser Ala Thr Glu
 20 25 30

Lys Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala
 35 40 45

Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 50 55 60

Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn
 65 70 75 80

Pro Gln Glu Val Val Leu Val Asn Val Thr Glu Asn Phe Asn Met Trp
 85 90 95

Lys Asn Asp Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp
 100 105 110

Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ser
 115 120 125

Leu Lys Cys Thr Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser Asn
 130 135 140

Thr Asn Ser Ser Ser Gly Glu Met Met Met Glu Lys Gly Glu Ile Lys
 145 150 155 160 165

Asn Cys Ser Phe Asn Ile Ser Thr Ser Ile Arg Gly Lys Val Gln Lys
 165 170 175

Glu Tyr Ala Phe Phe Tyr Lys Leu Asp Ile Ile Pro Ile Asp Asn Asp
 180 185 190

Thr Thr Ser Tyr Thr Leu Thr Ser Cys Asn Thr Ser Val Ile Thr Gln
 195 200 205

Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala
 210 215 220

Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asn Lys Thr Phe Asn Gly
 225 230 235 240

Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile
 245 250 255

Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 260 265 270

Glu Glu Val Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 275 280 285

Ile Ile Val Gln Leu Asn Gln Ser Val Glu Ile Asn Cys Thr Arg Pro
 290 295 300

Asn Asn Asn Thr Arg Lys Ser Ile Arg Ile Gln Arg Gly Pro Gly Arg
 305 310 315 320

Ala Phe Val Thr Ile Gly Lys Ile Gly Asn Met Arg Gln Ala His Cys
 325 330 335

Asn Ile Ser Arg Ala Lys Trp Asn Ala Thr Leu Lys Gln Ile Ala Ser
 340 345 350

Lys Leu Arg Glu Gln Phe Gly Asn Asn Lys Thr Ile Ile Phe Lys Gln
 355 360 365

Ser Ser Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys Gly
 370 375 380

Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp
 385 390 395 400

Phe Asn Ser Thr Trp Ser Thr Glu Gly Ser Asn Asn Thr Glu Gly Ser
 405 410 415

Asp Thr Ile Thr Leu Pro Cys Arg Ile Lys Gln Phe Ile Asn Met Trp
 420 425 430

Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 435 440 445

Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 450 455 460

Asn Asn Asn Asn Gly Ser Glu Ile Phe Arg Pro Gly Gly Gly Asp Met
 465 470 475 480

Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 485 490 495

Glu Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 500 505 510

Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Leu Phe Leu Gly Phe Leu
 515 520 525

Gly Ala Ala Gly Ser Thr Met Gly Ala Arg Ser Met Thr Leu Thr Val
 530 535 540

Gln Ala Arg Gln Leu Leu Ser Gly Ile Val Gln Gln Gln Asn Asn Leu
 545 550 555

Leu Arg Ala Ile Glu Ala Gln Gln His Leu Leu Gln Leu Thr Val Trp
 565 570 575

Gly Ile Lys Gln Leu Gln Ala Arg Ile Leu Ala Val Glu Arg Tyr Leu
 580 585 590

Lys Asp Gln Gln Leu Leu Gly Ile Trp Gly Cys Ser Gly Lys Leu Ile
 595 600 605

Cys Thr Thr Ala Val Pro Trp Asn Ala Ser Trp Ser Asn Lys Ser Leu
 610 615 620

Glu Gln Ile Trp Asn Asn Met Thr Trp Met Glu Trp Asp Arg Glu Ile
 625 630 635 640

Asn Asn Tyr Thr Ser Leu Ile His Ser Leu Ile Glu Glu Ser Gln Asn
 645 650 655

Gln Gln Glu Lys Asn Glu Gln Glu Leu Leu Glu Leu Asp Lys Trp Ala
 660 665 670

Ser Leu Trp Asn Trp Phe Asn Ile Thr Asn Trp Leu Trp Tyr Ile Lys
 675 680 685

Ile Phe Ile Met Ile Val Gly Gly Leu Val Gly Leu Arg Ile Val Phe
 690 695 700

Ala Val Leu Ser Ile Val Asn Arg Val Arg Gln Gly Tyr Ser Pro Leu
 705 710 715 720

Ser Phe Gln Thr His Leu Pro Thr Pro Arg Gly Pro Asp Arg Pro Glu
 725 730 735

Gly Ile Glu Glu Glu Gly Gly Glu Arg Asp Arg Asp Arg Ser Ile Arg
 740 745 750

Leu Val Asn Gly Ser Leu Ala Leu Ile Trp Asp Asp Leu Arg Ser Leu
 755 760 765

Cys Leu Phe Ser Tyr His Arg Leu Arg Asp Leu Leu Leu Ile Val Thr
 770 775 780

Arg Ile Val Glu Leu Leu Gly Arg Arg Gly Trp Glu Ala Leu Lys Tyr
 785 790 795 800

Trp Trp Asn Leu Leu 805 Gln Tyr Trp Ser Gln Glu Leu Lys Asn Ser Ala 815

Val Ser Leu Leu 820 Asn Ala Thr Ala Ile Ala Val Ala Glu Gly Thr Asp 830

Arg Val Ile Glu Val Val Gln Gly Ala Cys Arg Ala Ile Arg His Ile 835 840 845

Pro Arg Arg Ile Arg Gln Gly Leu Glu Arg Ile Leu Leu 850 855 860

<210> 418
 <211> 859
 <212> PRT
 <213> Human immunodeficiency virus type 2
 <400> 418

Met Cys Gly Arg Asn Gln Leu Phe Val Ala Ser Leu Leu Ala Ser Ala 1 5 10 15

Cys Leu Ile Tyr Cys Val Gln Tyr Val Thr Val Phe Tyr Gly Val Pro 20 25 30

Val Trp Arg Asn Ala Ser Ile Pro Leu Phe Cys Ala Thr Lys Asn Arg 35 40 45

Asp Thr Trp Gly Thr Ile Gln Cys Leu Pro Asp Asn Asp Asp Tyr Gln 50 55 60

Glu Ile Ala Leu Asn Val Thr Glu Ala Phe Asp Ala Trp Asn Asn Thr 65 70 75 80

Val Thr Glu Gln Ala Val Glu Asp Val Trp Ser Leu Phe Glu Thr Ser 85 90 95

Ile Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ala Met Arg Cys 100 105 110

Asn Ser Thr Thr Ala Lys Asn Thr Thr Ser Thr Pro Thr Thr Thr 115 120 125

Thr Ala Asn Thr Thr Ile Gly Glu Asn Ser Ser Cys Ile Arg Thr Asp 130 135 140

Asn Cys Thr Gly Leu Gly Glu Glu Glu Met Val Asp Cys Gln Phe Asn 145 150 155 160

Met Thr Gly Leu Glu Arg Asp Lys Lys Lys Leu Tyr Asn Glu Thr Trp 165 170 175

Tyr Ser Lys Asp Val Val Cys Glu Ser Asn Asp Thr Lys Lys Glu Lys

180

185

190

Thr Cys Tyr Met Asn His Cys Asn Thr Ser Val Ile Thr Glu Ser Cys
 195 200 205
 Asp Lys His Tyr Trp Asp Thr Met Arg Phe Arg Tyr Cys Ala Pro Pro
 210 215 220
 Gly Phe Ala Leu Leu Arg Cys Asn Asp Thr Asn Tyr Ser Gly Phe Glu
 225 230 235 240
 Pro Asn Cys Ser Lys Val Val Ala Ala Thr Cys Thr Arg Met Met Glu
 245 250 255
 Thr Gln Thr Ser Thr Trp Phe Gly Phe Asn Gly Thr Arg Ala Glu Asn
 260 265 270
 Arg Thr Tyr Ile Tyr Trp His Gly Arg Asp Asn Arg Thr Ile Ile Ser
 275 280 285
 Leu Asn Lys Phe Tyr Asn Leu Thr Val His Cys Lys Arg Pro Gly Asn
 290 295 300
 Lys Thr Val Val Pro Ile Thr Leu Met Ser Gly Leu Val Phe His Ser
 305 310 315 320
 Gln Pro Ile Asn Arg Arg Pro Arg Gln Ala Trp Cys Trp Phe Lys Gly
 325 330 335
 Glu Trp Lys Glu Ala Met Lys Glu Val Lys Leu Thr Leu Ala Lys His
 340 345 350
 Pro Arg Tyr Lys Gly Thr Asn Asp Thr Glu Lys Ile Arg Phe Ile Ala
 355 360 365
 Pro Gly Glu Arg Ser Asp Pro Glu Val Ala Tyr Met Trp Thr Asn Cys
 370 375 380
 Arg Gly Glu Phe Leu Tyr Cys Asn Met Thr Trp Phe Leu Asn Trp Val
 385 390 395 400
 Glu Asn Arg Thr Asn Gln Thr Gln His Asn Tyr Val Pro Cys His Ile
 405 410 415
 Lys Gln Ile Ile Asn Thr Trp His Lys Val Gly Lys Asn Val Tyr Leu
 420 425 430
 Pro Pro Arg Glu Gly Gln Leu Thr Cys Asn Ser Thr Val Thr Ser Ile
 435 440 445
 Ile Ala Asn Ile Asp Gly Gly Glu Asn Gln Thr Asn Ile Thr Phe Ser

450

455

Ala Glu Val Ala Glu Leu Tyr Arg Leu Glu Leu Gly Asp Tyr Lys Leu
465 470 475 480

Ile Glu Val Thr Pro Ile Gly Phe Ala Pro Thr Pro Val Lys Arg Tyr
485 490 495

Ser Ser Ala Pro Val Arg Asn Lys Arg Gly Val Phe Val Leu Gly Phe
500 505 510

Leu Gly Phe Leu Thr Thr Ala Gly Ala Ala Met Gly Ala Ala Ser Leu
515 520 525

Thr Leu Ser Ala Gln Ser Arg Thr Leu Leu Ala Gly Ile Val Gln Gln
530 535 540

Gln Gln Gln Leu Leu Asp Val Val Lys Arg Gln Gln Glu Met Leu Arg
545 550 555 560 565

Leu Thr Val Trp Gly Thr Lys Asn Leu Gln Ala Arg Val Thr Ala Ile
565 570 575

Glu Lys Tyr Leu Lys Asp Gln Ala Gln Leu Asn Ser Trp Gly Cys Ala
580 585 590

Phe Arg Gln Val Cys His Thr Thr Val Pro Trp Val Asn Asp Thr Leu
595 600 605

Thr Pro Asp Trp Asn Asn Met Thr Trp Gln Glu Trp Glu Gln Arg Ile
610 615 620

Arg Asn Leu Glu Ala Asn Ile Ser Glu Ser Leu Glu Gln Ala Gln Ile
625 630 635 640

Gln Gln Glu Lys Asn Met Tyr Glu Leu Gln Lys Leu Asn Ser Trp Asp
645 650 655

Val Phe Gly Asn Trp Phe Asp Leu Thr Ser Trp Ile Lys Tyr Ile Gln
660 665 670

Tyr Gly Val Tyr Ile Val Val Gly Ile Ile Val Leu Arg Ile Val Ile
675 680 685

Tyr Val Val Gln Met Leu Ser Arg Leu Arg Lys Gly Tyr Arg Pro Val
690 695 700

Phe Ser Ser Pro Pro Ala Tyr Phe Gln Gln Ile His Ile His Lys Asp
705 710 715 720

Arg Glu Gln Pro Ala Arg Glu Glu Thr Glu Glu Asp Val Gly Asn Ser

Val Gly Asp Asn Trp Trp Pro Trp Pro Ile Arg Tyr Ile His Phe Leu
740 745 750

Ile Arg Gln Leu Ile Arg Leu Leu Asn Arg Leu Tyr Asn Ile Cys Arg
755 760 765

Asp Leu Leu Ser Arg Ser Phe Gln Thr Leu Gln Leu Ile Ser Gln Ser
770 775 780

Leu Arg Arg Ala Leu Thr Ala Val Arg Asp Trp Leu Arg Phe Asn Thr
785 790 795 800

Ala Tyr Leu Gln Tyr Gly Gly Glu Trp Ile Gln Glu Ala Phe Arg Ala
805 810 815

Phe Ala Arg Ala Thr Gly Glu Thr Leu Thr Asn Ala Trp Arg Gly Phe
820 825 830

Trp Gly Thr Leu Gly Gln Ile Gly Arg Gly Ile Leu Ala Val Pro Arg
835 840 845

Arg Ile Arg Gln Gly Ala Glu Ile Ala Leu Leu
850 855

<210> 419
<211> 879
<212> PRT
<213> Simian immunodeficiency virus

<400> 419

Met Gly Cys Leu Gly Asn Gln Leu Leu Ile Ala Ile Leu Leu Leu Ser
1 5 10 15

Val Tyr Gly Ile Tyr Cys Thr Leu Tyr Val Thr Val Phe Tyr Gly Val
20 25 30

Pro Ala Trp Arg Asn Ala Thr Ile Pro Leu Phe Cys Ala Thr Lys Asn
35 40 45

Arg Asp Thr Trp Gly Thr Thr Gln Cys Leu Pro Asp Asn Gly Asp Tyr
50 55 60

Ser Glu Val Ala Leu Asn Val Thr Glu Ser Phe Asp Ala Trp Asn Asn
65 70 75 80

Thr Val Thr Glu Gln Ala Ile Glu Asp Val Trp Gln Leu Phe Glu Thr
85 90 95

Ser Ile Lys Pro Cys Val Lys Leu Ser Pro Leu Cys Ile Thr Met Arg
100 105 110

eol f-seql . txt

Cys Asn Lys Ser Glu Thr Asp Arg Trp Gly Leu Thr Lys Ser Ile Thr
 115 120 125
 Thr Thr Ala Ser Thr Thr Ser Thr Thr Ala Ser Ala Lys Val Asp Met
 130 135 140
 Val Asn Glu Thr Ser Ser Cys Ile Ala Gln Asp Asn Cys Thr Gly Leu
 145 150 155 160
 Glu Gln Glu Gln Met Ile Ser Cys Lys Phe Asn Met Thr Gly Leu Lys
 165 170
 Arg Asp Lys Lys Lys Glu Tyr Asn Glu Thr Trp Tyr Ser Ala Asp Leu
 180 185 190
 Val Cys Glu Gln Gly Asn Asn Thr Gly Asn Glu Ser Arg Cys Tyr Met
 195 200 205
 Asn His Cys Asn Thr Ser Val Ile Gln Glu Ser Cys Asp Lys His Tyr
 210 215 220
 Trp Asp Ala Ile Arg Phe Arg Tyr Cys Ala Pro Pro Gly Tyr Ala Leu
 225 230 235 240
 Leu Arg Cys Asn Asp Thr Asn Tyr Ser Gly Phe Met Pro Lys Cys Ser
 245 250 255
 Lys Val Val Val Ser Ser Cys Thr Arg Met Met Glu Thr Gln Thr Ser
 260 265 270
 Thr Trp Phe Gly Phe Asn Gly Thr Arg Ala Glu Asn Arg Thr Tyr Ile
 275 280 285
 Tyr Trp His Gly Arg Asp Asn Arg Thr Ile Ile Ser Leu Asn Lys Tyr
 290 295 300
 Tyr Asn Leu Thr Met Lys Cys Arg Arg Pro Gly Asn Lys Thr Val Leu
 305 310 315 320
 Pro Val Thr Ile Met Ser Gly Leu Val Phe His Ser Gln Pro Ile Asn
 325 330 335
 Asp Arg Pro Lys Gln Ala Trp Cys Trp Phe Gly Gly Lys Trp Lys Asp
 340 345 350
 Ala Ile Lys Glu Val Lys Gln Thr Ile Val Lys His Pro Arg Tyr Thr
 355 360 365
 Gly Thr Asn Asn Thr Asp Lys Ile Asn Leu Thr Ala Pro Gly Gly Gly
 370 375 380

eol f-seql . txt

Asp 385 Pro Glu Val Thr Phe 390 Met Trp Thr Asn Cys 395 Arg Gly Glu Phe Leu 400
 Tyr Cys Lys Met Asn 405 Trp Phe Leu Asn Trp 410 Val Glu Asp Arg Asn Thr 415
 Ala Asn Gln Lys 420 Pro Lys Glu Gln His 425 Lys Arg Asn Tyr Val 430 Pro Cys
 His Ile Arg 435 Gln Ile Ile Asn Thr 440 Trp His Lys Val Gly 445 Lys Asn Val
 Tyr Leu 450 Pro Pro Arg Glu Gly 455 Asp Leu Thr Cys Asn 460 Ser Thr Val Thr
 Ser Leu Ile Ala Asn 470 Ile Asp Trp Ile Asp Gly 475 Asn Gln Thr Asn Ile 480
 Thr Met Ser Ala Glu 485 Val Ala Glu Leu Tyr Arg 490 Leu Glu Leu Gly Asp 495
 Tyr Lys Leu Val 500 Glu Ile Thr Pro Ile 505 Gly Leu Ala Pro Thr Asp Val
 Lys Arg Tyr 515 Thr Thr Gly Gly Thr 520 Ser Arg Asn Lys Arg 525 Gly Val Phe
 Val Leu 530 Gly Phe Leu Gly Phe 535 Leu Ala Thr Ala Gly 540 Ser Ala Met Gly
 Ala Ala Ser Leu Thr 550 Leu Thr Ala Gln Ser Arg 555 Thr Leu Leu Ala Gly 560
 Ile Val Gln Gln Gln 565 Gln Gln Leu Leu Asp 570 Val Val Lys Arg Gln Gln 575
 Glu Leu Leu Arg 580 Leu Thr Val Trp Gly 585 Thr Lys Asn Leu Gln Thr Arg
 Val Thr Ala 595 Ile Glu Lys Tyr Leu 600 Lys Asp Gln Ala Gln 605 Leu Asn Ala
 Trp Gly 610 Cys Ala Phe Arg Gln 615 Val Cys His Thr Thr 620 Val Pro Trp Pro
 Asn Ala Ser Leu Thr 630 Pro Lys Trp Asn Asn Glu 635 Thr Trp Gln Glu Trp 640
 Glu Arg Lys Val Asp 645 Phe Leu Glu Glu Asn Ile Thr Ala Leu Leu Glu 655

eol f-seql . txt

Glu Ala Gln Ile Gln Gln Glu Lys Asn Met Tyr Glu Leu Gln Lys Leu
660 665 670

Asn Ser Trp Asp Val Phe Gly Asn Trp Phe Asp Leu Ala Ser Trp Ile
675 680 685

Lys Tyr Ile Gln Tyr Gly Val Tyr Ile Val Val Gly Val Ile Leu Leu
690 695 700

Arg Ile Val Ile Tyr Ile Val Gln Met Leu Ala Lys Leu Arg Gln Gly
705 710 715 720

Tyr Arg Pro Val Phe Ser Ser Pro Pro Ser Tyr Phe Gln Gln Thr His
725 730 735

Ile Gln Gln Asp Pro Ala Leu Pro Thr Arg Glu Gly Lys Glu Arg Asp
740 745 750

Gly Gly Glu Gly Gly Gly Asn Ser Ser Trp Pro Trp Gln Ile Glu Tyr
755 760 765

Ile His Phe Leu Ile Arg Gln Leu Ile Arg Leu Leu Thr Trp Leu Phe
770 775 780

Ser Asn Cys Arg Thr Leu Leu Ser Arg Val Tyr Gln Ile Leu Gln Pro
785 790 795 800

Ile Leu Gln Arg Leu Ser Ala Thr Leu Gln Arg Ile Arg Glu Val Leu
805 810 815

Arg Thr Glu Leu Thr Tyr Leu Gln Tyr Gly Trp Ser Tyr Phe His Glu
820 825 830

Ala Val Gln Ala Val Trp Arg Ser Ala Thr Glu Thr Leu Ala Gly Ala
835 840 845

Trp Gly Asp Leu Trp Glu Thr Leu Arg Arg Gly Gly Arg Trp Ile Leu
850 855 860

Ala Ile Pro Arg Arg Ile Arg Gln Gly Leu Glu Leu Thr Leu Leu
865 870 875

<210> 420
<211> 859
<212> PRT
<213> Human immunodeficiency virus type 2

<400> 420

Met Cys Gly Arg Asn Gln Leu Phe Val Ala Ser Leu Leu Ala Ser Ala
1 5 10 15

eol f-seq1 . txt

Cys Leu Ile Tyr Cys Val Gln Tyr Val Thr Val Phe Tyr Gly Val Pro
 20 25 30
 Val Trp Arg Asn Ala Ser Ile Pro Leu Phe Cys Ala Thr Lys Asn Arg
 35 40 45
 Asp Thr Trp Gly Thr Ile Gln Cys Leu Pro Asp Asn Asp Asp Tyr Gln
 50 55 60
 Glu Ile Ala Leu Asn Val Thr Glu Ala Phe Asp Ala Trp Asn Asn Thr
 65 70 75 80
 Val Thr Glu Gln Ala Val Glu Asp Val Trp Ser Leu Phe Glu Thr Ser
 85 90 95
 Ile Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Ala Met Arg Cys
 100 105 110
 Asn Ser Thr Thr Ala Lys Asn Thr Thr Ser Thr Pro Thr Thr Thr Thr
 115 120 125
 Thr Ala Asn Thr Thr Ile Gly Glu Asn Ser Ser Cys Ile Arg Thr Asp
 130 135 140
 Asn Cys Thr Gly Leu Gly Glu Glu Glu Met Val Asp Cys Gln Phe Asn
 145 150 155 160
 Met Thr Gly Leu Glu Arg Asp Lys Lys Lys Leu Tyr Asn Glu Thr Trp
 165 170 175
 Tyr Ser Lys Asp Val Val Cys Glu Ser Asn Asp Thr Lys Lys Glu Lys
 180 185 190
 Thr Cys Tyr Met Asn His Cys Asn Thr Ser Val Ile Thr Glu Ser Cys
 195 200 205
 Asp Lys His Tyr Trp Asp Thr Met Arg Phe Arg Tyr Cys Ala Pro Pro
 210 215 220
 Gly Phe Ala Leu Leu Arg Cys Asn Asp Thr Asn Tyr Ser Gly Phe Glu
 225 230 235 240
 Pro Asn Cys Ser Lys Val Val Ala Ala Thr Cys Thr Arg Met Met Glu
 245 250 255
 Thr Gln Thr Ser Thr Trp Phe Gly Phe Asn Gly Thr Arg Ala Glu Asn
 260 265 270
 Arg Thr Tyr Ile Tyr Trp His Gly Arg Asp Asn Arg Thr Ile Ile Ser
 275 280 285

eol f-seql . txt

Leu Asn Lys Phe Tyr Asn Leu Thr Val Hi s Cys Lys Arg Pro Gly Asn
 290 295 300
 Lys Thr Val Val Pro Ile Thr Leu Met Ser Gly Leu Val Phe Hi s Ser
 305 310 315 320
 Gl n Pro Ile Asn Arg Arg Pro Arg Gl n Ala Trp Cys Trp Phe Lys Gly
 325 330 335
 Gl u Trp Lys Gl u Ala Met Lys Gl u Val Lys Leu Thr Leu Ala Lys Hi s
 340 345 350
 Pro Arg Tyr Lys Gly Thr Asn Asp Thr Gl u Lys Ile Arg Phe Ile Ala
 355 360 365
 Pro Gly Gl u Arg Ser Asp Pro Gl u Val Ala Tyr Met Trp Thr Asn Cys
 370 375 380
 Arg Gly Gl u Phe Leu Tyr Cys Asn Met Thr Trp Phe Leu Asn Trp Val
 385 390 395 400
 Gl u Asn Arg Thr Asn Gl n Thr Gl n Hi s Asn Tyr Val Pro Cys Hi s Ile
 405 410 415
 Lys Gl n Ile Ile Asn Thr Trp Hi s Lys Val Gly Lys Asn Val Tyr Leu
 420 425 430
 Pro Pro Arg Gl u Gly Gl n Leu Thr Cys Asn Ser Thr Val Thr Ser Ile
 435 440 445
 Ile Ala Asn Ile Asp Gly Gly Gl u Asn Gl n Thr Asn Ile Thr Phe Ser
 450 455 460
 Ala Gl u Val Ala Gl u Leu Tyr Arg Leu Gl u Leu Gly Asp Tyr Lys Leu
 465 470 475 480
 Ile Gl u Val Thr Pro Ile Gly Phe Ala Pro Thr Pro Val Lys Arg Tyr
 485 490 495
 Ser Ser Ala Pro Val Arg Asn Lys Arg Gly Val Phe Val Leu Gly Phe
 500 505 510
 Leu Gly Phe Leu Thr Thr Ala Gly Ala Ala Met Gly Ala Ala Ser Leu
 515 520 525
 Thr Leu Ser Ala Gl n Ser Arg Thr Leu Leu Ala Gly Ile Val Gl n Gl n
 530 535 540
 Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg Gl n Gl n Gl u Met Leu Arg
 545 550 555 560

eol f-seq1 . txt

Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n Ala Arg Val Thr Ala Ile
 565 570 575

Gl u Lys Arg Leu Lys Asp Gl n Ala Gl n Leu Asn Ser Trp Gly Cys Ala
 580 585 590

Phe Arg Gl n Val Cys Hi s Thr Thr Val Pro Trp Val Asn Asp Thr Leu
 595 600 605

Thr Pro Asp Trp Asn Asn Met Thr Trp Gl n Gl u Trp Gl u Gl n Arg Ile
 610 615 620

Arg Asn Leu Gl u Ala Asn Ile Ser Gl u Ser Leu Gl u Gl n Ala Gl n Ile
 625 630 635 640

Gl n Gl n Gl u Lys Asn Met Tyr Gl u Leu Gl n Lys Leu Asn Ser Trp Asp
 645 650 655

Val Phe Gly Asn Trp Phe Asp Leu Thr Ser Trp Ile Lys Tyr Ile Gl n
 660 665 670

Tyr Gly Val Tyr Ile Val Val Gly Ile Ile Val Leu Arg Ile Val Ile
 675 680 685

Tyr Val Val Gl n Met Leu Ser Arg Leu Arg Lys Gly Tyr Arg Pro Val
 690 695 700

Phe Ser Ser Pro Pro Ala Tyr Phe Gl n Gl n Ile Hi s Ile Hi s Lys Asp
 705 710 715 720

Arg Gl u Gl n Pro Ala Arg Gl u Gl u Thr Gl u Gl u Asp Val Gly Asn Ser
 725 730 735

Val Gly Asp Asn Trp Trp Pro Trp Pro Ile Arg Tyr Ile Hi s Phe Leu
 740 745 750

Ile Arg Gl n Leu Ile Arg Leu Leu Asn Arg Leu Tyr Asn Ile Cys Arg
 755 760 765

Asp Leu Leu Ser Arg Ser Phe Gl n Thr Leu Gl n Leu Ile Ser Gl n Ser
 770 775 780

Leu Arg Arg Ala Leu Thr Ala Val Arg Asp Trp Leu Arg Phe Asn Thr
 785 790 795 800

Ala Tyr Leu Gl n Tyr Gly Gly Gl u Trp Ile Gl n Gl u Ala Phe Arg Ala
 805 810 815

Phe Ala Arg Ala Thr Gly Gl u Thr Leu Thr Asn Ala Trp Arg Gly Phe
 820 825 830

eol f-seql . txt

Trp Gly Thr Leu Gly Gln Ile Gly Arg Gly Ile Leu Ala Val Pro Arg
835 840 845

Arg Ile Arg Gln Gly Ala Glu Ile Ala Leu Leu
850 855

<210> 421
<211> 879
<212> PRT
<213> Simian immunodeficiency virus
<400> 421

Met Gly Cys Leu Gly Asn Gln Leu Leu Ile Ala Ile Leu Leu Leu Ser
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Val Tyr Gly Ile Tyr Cys Thr Leu Tyr Val Thr Val Phe Tyr Gly Val
20 25 30

Pro Ala Trp Arg Asn Ala Thr Ile Pro Leu Phe Cys Ala Thr Lys Asn
35 40 45

Arg Asp Thr Trp Gly Thr Thr Gln Cys Leu Pro Asp Asn Gly Asp Tyr
50 55 60

Ser Glu Val Ala Leu Asn Val Thr Glu Ser Phe Asp Ala Trp Asn Asn
65 70 75 80

Thr Val Thr Glu Gln Ala Ile Glu Asp Val Trp Gln Leu Phe Glu Thr
85 90 95

Ser Ile Lys Pro Cys Val Lys Leu Ser Pro Leu Cys Ile Thr Met Arg
100 105 110

Cys Asn Lys Ser Glu Thr Asp Arg Trp Gly Leu Thr Lys Ser Ile Thr
115 120 125

Thr Thr Ala Ser Thr Thr Ser Thr Thr Ala Ser Ala Lys Val Asp Met
130 135 140

Val Asn Glu Thr Ser Ser Cys Ile Ala Gln Asp Asn Cys Thr Gly Leu
145 150 155 160

Glu Gln Glu Gln Met Ile Ser Cys Lys Phe Asn Met Thr Gly Leu Lys
165 170 175

Arg Asp Lys Lys Lys Glu Tyr Asn Glu Thr Trp Tyr Ser Ala Asp Leu
180 185 190

Val Cys Glu Gln Gly Asn Asn Thr Gly Asn Glu Ser Arg Cys Tyr Met
195 200 205

Asn His Cys Asn Thr Ser Val Ile Gln Glu Ser Cys Asp Lys His Tyr
 210 215 220

Trp Asp Ala Ile Arg Phe Arg Tyr Cys Ala Pro Pro Gly Tyr Ala Leu
 225 230 235

Leu Arg Cys Asn Asp Thr Asn Tyr Ser Gly Phe Met Pro Lys Cys Ser
 245 250 255

Lys Val Val Val Ser Ser Cys Thr Arg Met Met Glu Thr Gln Thr Ser
 260 265 270

Thr Trp Phe Gly Phe Asn Gly Thr Arg Ala Glu Asn Arg Thr Tyr Ile
 275 280 285

Tyr Trp His Gly Arg Asp Asn Arg Thr Ile Ile Ser Leu Asn Lys Tyr
 290 300

Tyr Asn Leu Thr Met Lys Cys Arg Arg Pro Gly Asn Lys Thr Val Leu
 305 310 315 320

Pro Val Thr Ile Met Ser Gly Leu Val Phe His Ser Gln Pro Ile Asn
 325 330 335

Asp Arg Pro Lys Gln Ala Trp Cys Trp Phe Gly Gly Lys Trp Lys Asp
 340 345 350

Ala Ile Lys Glu Val Lys Gln Thr Ile Val Lys His Pro Arg Tyr Thr
 355 360 365

Gly Thr Asn Asn Thr Asp Lys Ile Asn Leu Thr Ala Pro Gly Gly Gly
 370 375 380

Asp Pro Glu Val Thr Phe Met Trp Thr Asn Cys Arg Gly Glu Phe Leu
 385 390 395 400

Tyr Cys Lys Met Asn Trp Phe Leu Asn Trp Val Glu Asp Arg Asn Thr
 405 410 415

Ala Asn Gln Lys Pro Lys Glu Gln His Lys Arg Asn Tyr Val Pro Cys
 420 425 430

His Ile Arg Gln Ile Ile Asn Thr Trp His Lys Val Gly Lys Asn Val
 435 440 445

Tyr Leu Pro Pro Arg Glu Gly Asp Leu Thr Cys Asn Ser Thr Val Thr
 450 455 460

Ser Leu Ile Ala Asn Ile Asp Trp Ile Asp Gly Asn Gln Thr Asn Ile
 465 470 475 480

Thr Met Ser Ala Glu Val Ala Glu Leu Tyr Arg Leu Glu Leu Gly Asp
 485 490 495

Tyr Lys Leu Val Glu Ile Thr Pro Ile Gly Leu Ala Pro Thr Asp Val
 500 505 510 515

Lys Arg Tyr Thr Thr Gly Gly Thr Ser Arg Asn Lys Arg Gly Val Phe
 515 520 525

Val Leu Gly Phe Leu Gly Phe Leu Ala Thr Ala Gly Ser Ala Met Gly
 530 535 540

Ala Ala Ser Leu Thr Leu Thr Ala Gl n Ser Arg Thr Leu Leu Ala Gly
 545 550 555 560 565

Ile Val Gl n Gl n Gl n Gl n Gl n Leu Leu Asp Val Val Lys Arg Gl n Gl n
 565 570 575

Glu Leu Leu Arg Leu Thr Val Trp Gly Thr Lys Asn Leu Gl n Thr Arg
 580 585 590

Val Thr Ala Ile Glu Lys Arg Leu Lys Asp Gl n Ala Gl n Leu Asn Ala
 595 600 605

Trp Gly Cys Ala Phe Arg Gl n Val Cys His Thr Thr Val Pro Trp Pro
 610 615 620

Asn Ala Ser Leu Thr Pro Lys Trp Asn Asn Glu Thr Trp Gl n Glu Trp
 625 630 635 640

Glu Arg Lys Val Asp Phe Leu Glu Glu Asn Ile Thr Ala Leu Leu Glu
 645 650 655

Glu Ala Gl n Ile Gl n Gl n Glu Lys Asn Met Tyr Glu Leu Gl n Lys Leu
 660 665 670

Asn Ser Trp Asp Val Phe Gly Asn Trp Phe Asp Leu Ala Ser Trp Ile
 675 680 685

Lys Tyr Ile Gl n Tyr Gly Val Tyr Ile Val Val Gly Val Ile Leu Leu
 690 695 700

Arg Ile Val Ile Tyr Ile Val Gl n Met Leu Ala Lys Leu Arg Gl n Gly
 705 710 715 720

Tyr Arg Pro Val Phe Ser Ser Pro Pro Ser Tyr Phe Gl n Gl n Thr His
 725 730 735

Ile Gl n Gl n Asp Pro Ala Leu Pro Thr Arg Glu Gly Lys Glu Arg Asp
 740 745 750

eol f-seq1 . txt

Gly Gly Glu Gly Gly Gly Asn Ser Ser Trp Pro Trp Gln Ile Glu Tyr
755 760 765

Ile His Phe Leu Ile Arg Gln Leu Ile Arg Leu Leu Thr Trp Leu Phe
770 775 780

Ser Asn Cys Arg Thr Leu Leu Ser Arg Val Tyr Gln Ile Leu Gln Pro
785 790 795 800

Ile Leu Gln Arg Leu Ser Ala Thr Leu Gln Arg Ile Arg Glu Val Leu
805 810 815

Arg Thr Glu Leu Thr Tyr Leu Gln Tyr Gly Trp Ser Tyr Phe His Glu
820 825 830

Ala Val Gln Ala Val Trp Arg Ser Ala Thr Glu Thr Leu Ala Gly Ala
835 840 845

Trp Gly Asp Leu Trp Glu Thr Leu Arg Arg Gly Gly Arg Trp Ile Leu
850 855 860

Ala Ile Pro Arg Arg Ile Arg Gln Gly Leu Glu Leu Thr Leu Leu
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<213> Artificial sequence

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<223> Xaa is any naturally occurring amino acid or is mutated

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<220>
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<400> 422

Ala Ile Glu Lys Xaa Xaa Xaa Asp Gln
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<210> 423
<211> 9
<212> PRT
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Ala Ile Glu Arg Xaa Xaa Xaa Asp Gln

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<210> 424

<211> 9

<212> PRT

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<400> 424

Ala Val Glu Lys Xaa Xaa Xaa Asp Gln

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5

<210> 425

<211> 9

<212> PRT

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<223> Xaa is any naturally occurring amino acid or is mutated

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<220>
<221> MI SC_FEATURE
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<223> Xaa can be any natural ly occuring ami no aci d

<400> 425

Al a Val Gl u Arg Xaa Xaa Xaa Asp Gl n
1 5

<210> 426
<211> 38
<212> PRT
<213> Arti ficial sequence

<220>
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<222> (2)..(2)
<223> Xaa = I or V

<220>
<221> MI SC_FEATURE
<222> (4)..(4)
<223> Xaa = K or R

<220>
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<223> Xaa is any natural ly occuring ami no aci d or is mutated

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<223> Xaa can be any natural ly occuring ami no aci d

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 <223> Xaa = S or T

<400> 426

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Cys Xaa Xaa Xaa Xaa Cys Val Xaa Thr Xaa Val Pro Xaa Xaa Xaa Xaa
 20 25 30

Xaa Xaa Xaa Xaa Xaa Xaa
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<210> 427
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 <223> Xaa = I or V

<220>
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 <223> Xaa = K or R

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<400> 427

Al a Xaa Gl u Xaa Xaa Xaa Xaa Asp Gl n
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<210> 428
 <211> 15
 <212> PRT
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<220>
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<223> Xaa is any naturally occurring amino acid or is mutated

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<223> Xaa = K or R

<220>
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<222> (11)..(11)
<223> Xaa is any naturally occurring amino acid or is mutated

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<223> Xaa is any naturally occurring amino acid or is mutated

<220>
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<223> Xaa = any naturally occurring amino acid

<400> 428

Xaa Xaa Xaa Xaa Ile Leu Ala Xaa Glu Xaa Xaa Xaa Xaa Asp Gln
1 5 10 15