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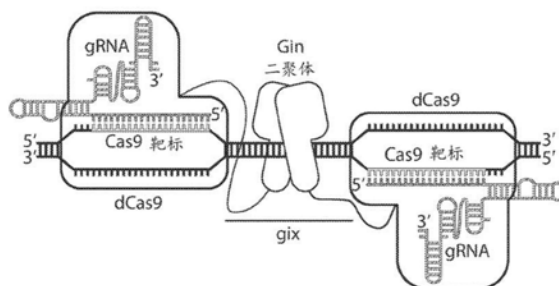
权利要求书7页 说明书128页
序列表285页 附图20页

(54)发明名称

可编程CAS9-重组酶融合蛋白及其用途

(57)摘要

本公开的一些方面提供了融合蛋白,其包含引导核苷酸序列-可编程DNA结合蛋白域(例如,Cas9的核酸酶无活性变体,例如dCas9)、任选的接头和重组酶催化域(例如,酪氨酸重组酶催化域或丝氨酸重组酶催化域,例如Gin重组酶催化域)。该融合蛋白可以重组含有侧翼为引导RNA指定序列的最小重组酶核心位点的DNA位点。本公开代表朝向不依赖于内源性细胞机制或细胞状态的在未修饰的细胞中的可编程无痕基因组编辑的步骤。



1. 融合蛋白,其包含:
 - (i) 引导核苷酸序列-可编程DNA结合蛋白域;
 - (ii) 接头;和
 - (iii) 重组酶催化域。
2. 权利要求1的融合蛋白,其中所述引导核苷酸序列-可编程DNA结合蛋白域选自下组:核酸酶无活性的Cas9 (dCas9) 域、核酸酶无活性的Cpf1域、核酸酶无活性的Argonaute域,及其变体。
3. 权利要求1的融合蛋白,其中所述引导核苷酸序列-可编程DNA结合蛋白域是核酸酶无活性的Cas9 (dCas9) 域。
4. 权利要求2或3的融合蛋白,其中所述dCas9域的氨基酸序列包含对应于SEQ ID NO:1中D10A或H840A突变的突变。
5. 权利要求2-4中任一项的融合蛋白,其中所述dCas9域的氨基酸序列包含对应于SEQ ID NO:1中D10A突变的突变和对应于SEQ ID NO:1中H840A突变的突变。
6. 权利要求2-5中任一项的融合蛋白,其中所述dCas9域的氨基酸序列进一步包含对应于SEQ ID NO:1中缺少的N-末端甲硫氨酸的突变。
7. 权利要求2或3的融合蛋白,其中所述dCas9域的氨基酸序列包含SEQ ID NO:712。
8. 权利要求2或3的融合蛋白,其中所述dCas9域的氨基酸序列与SEQ ID NO:712具有95%或更高的序列一致性。
9. 权利要求2或3的融合蛋白,其中所述dCas9域的氨基酸序列与SEQ ID NO:712具有96%、97%、98%、99%或更高的序列一致性。
10. 前述权利要求中任一项的融合蛋白,其中所述重组酶催化域是丝氨酸重组酶催化域或酪氨酸重组酶催化域。
11. 权利要求10的融合蛋白,其中所述丝氨酸重组酶催化域或酪氨酸重组酶催化域选自Gin、Sin、Tn3、Hin、 β 、 γ δ 、PhiC31、Cre或FLP重组酶催化域。
12. 权利要求10或权利要求11的融合蛋白,其中所述Gin重组酶催化域的氨基酸序列包含选自下组的一个或多个突变:SEQ ID NO:713中的H106Y、I127L、I136R或G137F突变。
13. 权利要求11或权利要求12的融合蛋白,其中所述Gin重组酶催化域的氨基酸序列包含SEQ ID NO:713。
14. 权利要求10或权利要求11的融合蛋白,其中所述Cre重组酶的氨基酸序列是截短的。
15. 权利要求11或权利要求14的融合蛋白,其中所述酪氨酸重组酶催化域是所述Cre重组酶的25kDa羧基-末端域。
16. 权利要求14的融合蛋白,其中所述Cre重组酶以氨基酸R118、A127、E138或R154开始。
17. 权利要求1-16中任一项的融合蛋白,其中所述重组酶的氨基酸序列已得以进一步突变。
18. 前述权利要求中任一项的融合蛋白,其中所述重组酶催化域是经演化的重组酶催化域。
19. 权利要求1-18中任一项的融合蛋白,其中所述接头具有约33埃至约81埃的长度。

20. 权利要求1-19中任一项的融合蛋白,其中所述接头是肽接头。

21. 权利要求20的融合蛋白,其中所述肽接头包含XTEN接头(SGSETPGTSESATPES (SEQ ID NO:7)、SGSETPGTSESA (SEQ ID NO:8)或SGSETPGTSESATPEGGSGGS (SEQ ID NO:9))、包含三肽GGG的一个或多个重复的氨基酸序列,或任何以下氨基酸序列:VPFLLEPDNINGKTC (SEQ ID NO:10)、GSAGSAAGSGEF (SEQ ID NO:11)、SIVAQLSRPDPA (SEQ ID NO:12)、MKIIEQLPSA (SEQ ID NO:13)、VRHKLKRVGS (SEQ ID NO:14)、GHGTGSTGSGSS (SEQ ID NO:15)、MSRPDPA (SEQ ID NO:16)或GGSM (SEQ ID NO:17)。

22. 权利要求20或权利要求21的融合蛋白,其中所述肽接头包含所述三肽GGG的一个或多个重复。

23. 权利要求20-22中任一项的融合蛋白,其中所述肽接头包含所述三肽GGG的六至十个重复。

24. 权利要求20-23中任一项的融合蛋白,其中所述肽接头包含所述三肽GGG的八个重复。

25. 权利要求20-24中任一项的融合蛋白,其中所述肽接头长约18至约27个氨基酸。

26. 权利要求20-25中任一项的融合蛋白,其中所述肽接头长24个氨基酸。

27. 权利要求20-26中任一项的融合蛋白,其中所述肽接头具有氨基酸序列GGSGSGSGSGSGSGSGSGSGGS (SEQ ID NO:183)。

28. 权利要求1-19中任一项的融合蛋白,其中所述接头是非肽接头。

29. 权利要求28的融合蛋白,其中所述非肽接头包含聚乙二醇(PEG)、聚丙二醇(PPG)、共聚(乙烯/丙烯)二醇、聚氧乙烯(POE)、聚氨酯、聚磷腈、多糖、右旋糖酐、聚乙烯醇、聚乙烯吡咯烷酮、聚乙烯乙醚、聚丙烯酰胺、聚丙烯酸酯、聚氰基丙烯酸酯、脂质聚合物、甲壳质、透明质酸、肝素或烷基接头。

30. 权利要求29的融合蛋白,其中所述烷基接头具有式—NH—(CH₂)_s—C(O)—,其中s可以是1-100的任何整数,包括端点。

31. 权利要求30的融合蛋白,其中s是1-20的整数,包括端点。

32. 权利要求1-31中任一项的融合蛋白,其进一步包含核定位信号(NLS)域。

33. 权利要求32的融合蛋白,其中所述NLS域经由一个或多个第二接头与所述引导核苷酸序列-可编程DNA结合蛋白域或所述Gin重组酶催化域结合。

34. 权利要求33的融合蛋白,其中所述第二接头是肽接头。

35. 权利要求33或权利要求34的融合蛋白,其中所述第二接头包含XTEN接头(SGSETPGTSESATPES (SEQ ID NO:7)、SGSETPGTSESA (SEQ ID NO:8)或SGSETPGTSESATPEGGSGGS (SEQ ID NO:9))、包含一个或多个三肽GGG的重复的氨基酸序列,或任何以下氨基酸序列:VPFLLEPDNINGKTC (SEQ ID NO:10)、GSAGSAAGSGEF (SEQ ID NO:11)、SIVAQLSRPDPA (SEQ ID NO:12)、MKIIEQLPSA (SEQ ID NO:13)、VRHKLKRVGS (SEQ ID NO:14)、GHGTGSTGSGSS (SEQ ID NO:15)、MSRPDPA (SEQ ID NO:16)或GGSM (SEQ ID NO:17)。

36. 权利要求33-35中任一项的融合蛋白,其中所述第二接头包含所述三肽GGG的一个或多个重复。

37. 权利要求33-36中任一项的融合蛋白,其中所述第二接头包含所述三肽GGG的一至五个重复。

38. 权利要求33-37中任一项的融合蛋白,其中所述第二接头包含所述三肽GGs的一个重复。

39. 权利要求33-38中任一项的融合蛋白,其中所述第二接头具有序列GGs。

40. 权利要求33的融合蛋白,其中所述第二接头是非肽接头。

41. 权利要求40的融合蛋白,其中所述非肽接头包含聚乙二醇(PEG)、聚丙二醇(PPG)、共聚(乙烯/丙烯)二醇、聚氧乙烯(POE)、聚氨酯、聚磷腈、多糖、右旋糖酐、聚乙烯醇、聚乙烯吡咯烷酮、聚乙烯乙醚、聚丙烯酰胺、聚丙烯酸酯、聚氰基丙烯酸酯、脂质聚合物、甲壳质、透明质酸、肝素或烷基接头。

42. 权利要求41的融合蛋白,其中所述烷基接头具有式—NH—(CH₂)_s—C(O)—,其中s可以是1-100的任何整数,包括端点。

43. 权利要求42的融合蛋白,其中s是1-20的任何整数,包括端点。

44. 权利要求32-43中任一项的融合蛋白,其中所述融合蛋白包含结构NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-COOH。

45. 权利要求32-44中任一项的融合蛋白,其中所述融合蛋白包含结构NH₂-[Gln重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-COOH。

46. 权利要求32-39、44或45中任一项的融合蛋白,其中所述融合蛋白包含SEQ ID NO: 719中所示的氨基酸序列。

47. 前述权利要求中任一项的融合蛋白,其进一步包含一个或多个亲和标签。

48. 权利要求47的融合蛋白,其中所述亲和标签选自下组:FLAG标签、多组氨酸(多His)标签、多精氨酸(多Arg)标签、Myc标签和HA标签。

49. 权利要求47或权利要求48的融合蛋白,其中所述亲和标签是FLAG标签。

50. 权利要求47-49中任一项的融合蛋白,其中所述FLAG标签具有序列PKKKRKV (SEQ ID NO:702)。

51. 权利要求47-50中任一项的融合蛋白,其中所述一个或多个亲和标签经由一个或多个第三接头与所述引导核苷酸序列-可编程DNA结合蛋白域、所述重组酶催化域或所述NLS域结合。

52. 权利要求51的融合蛋白,其中所述第三接头是肽接头。

53. 权利要求51或权利要求52的融合蛋白,其中所述第三接头包含XTEN接头,(SGSETPGTSESATPES (SEQ ID NO:7)、SGSETPGTSESA (SEQ ID NO:8) 或SGSETPGTSESATPEGGSGGS (SEQ ID NO:9))、包含三肽GGs的一个或多个重复的氨基酸序列,或任何以下氨基酸序列:VPFLLEPDNINGKTC (SEQ ID NO:10)、GSAGSAAGSGEF (SEQ ID NO:11)、SIVAQLSRPDPA (SEQ ID NO:12)、MKIIEQLPSA (SEQ ID NO:13)、VRHKLKRVGS (SEQ ID NO:14)、GHGTGSTGSGSS (SEQ ID NO:15)、MSRPDPA (SEQ ID NO:16) 或GGSM (SEQ ID NO:17)。

54. 权利要求51-53中任一项的融合蛋白,其中所述第三接头包含所述三肽GGs的一个或多个重复。

55. 权利要求51-54中任一项的融合蛋白,其中所述第三接头包含所述三肽GGs的一至五个重复。

56. 权利要求51-55中任一项的融合蛋白,其中所述第三接头包含所述三肽GGS的一个重复。

57. 权利要求51-56中任一项的融合蛋白,其中所述第三接头具有序列GGS。

58. 权利要求51的融合蛋白,其中所述第三接头是非肽接头。

59. 权利要求58的融合蛋白,其中所述非肽接头包含聚乙二醇(PEG)、聚丙二醇(PPG)、共聚(乙烯/丙烯)二醇、聚氧乙烯(POE)、聚氨酯、聚磷腈、多糖、右旋糖酐、聚乙烯醇、聚乙烯吡咯烷酮、聚乙烯乙醚、聚丙烯酰胺、聚丙烯酸酯、聚氰基丙烯酸酯、脂质聚合物、甲壳质、透明质酸、肝素或烷基接头。

60. 权利要求59的融合蛋白,其中所述烷基接头具有式—NH—(CH₂)_s—C(O)—,其中s可以是1-100的任何整数,包括端点。

61. 权利要求60的融合蛋白,其中s可以是1-20的任何整数。

62. 权利要求47-61中任一项的融合蛋白,其中所述融合蛋白包含结构NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[亲和标签]-COOH、NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH、NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH、NH₂-[亲和标签]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-COOH、NH₂-[亲和标签]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-COOH或NH₂-[亲和标签]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-COOH。

63. 权利要求47-57中任一项的融合蛋白,其中所述融合蛋白与SEQ ID NO:185中所示的氨基酸序列具有大于99%的序列一致性。

64. 权利要求47-57中任一项的融合蛋白,其中所述融合蛋白与SEQ ID NO:185中所示的氨基酸序列具有大于90%或95%的序列一致性。

65. 权利要求47-57、63或64中任一项的融合蛋白,其中所述融合蛋白具有SEQ ID NO:185中所示的氨基酸序列。

66. 前述权利要求中任一项的融合蛋白,其中所述引导核苷酸序列-可编程DNA结合蛋白域与引导RNA(gRNA)结合。

67. 权利要求66的融合蛋白的二聚体。

68. 权利要求67的融合蛋白的二聚体,其中所述二聚体与DNA分子结合。

69. 权利要求68的融合蛋白的二聚体,其中所述二聚体的每个融合蛋白与所述DNA分子的相同链结合。

70. 权利要求68的融合蛋白的二聚体,其中所述二聚体的每个融合蛋白与所述DNA分子的相对链结合。

71. 权利要求67-70中任一项的融合蛋白的二聚体,其中所述二聚体的gRNA与重组酶位点旁侧的gRNA结合位点杂交。

72. 权利要求71的方法,其中所述重组酶位点包含res、gix、hix、six、resH、LoxP、FTR或att核心或相关核心序列。

73. 权利要求71或权利要求72的融合蛋白的二聚体,其中所述重组酶位点包含gix核心或gix相关核心序列。

74. 权利要求72或权利要求73的二聚体,其中所述gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为3至7个碱基对。

75. 权利要求72-74中任一项的二聚体,其中所述gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为5至6个碱基对。

76. 权利要求67-75中任一项的融合蛋白的二聚体,其中第一二聚体与第二二聚体结合,从而形成所述融合蛋白的四聚体。

77. 权利要求66的融合蛋白的四聚体。

78. 权利要求77的融合蛋白的四聚体,其中所述四聚体与DNA分子结合。

79. 权利要求68的融合物的四聚体,其中每个二聚体与DNA的相对链结合。

80. 权利要求68的融合蛋白的四聚体,其中每个二聚体与DNA的相同链结合。

81. 用于两个DNA分子之间的位点特异性重组的方法,其包括:

(a) 使第一DNA与第一融合蛋白接触,其中所述引导核苷酸序列-可编程DNA结合蛋白域结合第一gRNA,所述第一gRNA与所述第一DNA的第一区域杂交;

(b) 使所述第一DNA与第二融合蛋白接触,其中所述第二融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第二gRNA,所述第二gRNA与所述第一DNA的第二区域杂交;

(c) 使第二DNA与第三融合蛋白接触,其中所述第三融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第三gRNA,所述第三gRNA与所述第二DNA的第一区域杂交;和

(d) 使所述第二DNA与第四融合蛋白接触,其中所述第四融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第四gRNA,所述第四gRNA与所述第二DNA的第二区域杂交;

其中在使得所述DNA重组的条件下,步骤(a)-(d)中所述融合蛋白的结合导致所述融合蛋白的重组酶催化域的四聚化,并且其中所述第一、第二、第三和/或第四融合蛋白是权利要求1-63中任一项的融合蛋白。

82. 权利要求81的方法,其中所述第一和第二DNA分子具有不同的序列。

83. 权利要求81的方法,其中步骤(a)和(b)的所述gRNA与所述第一DNA的相对链杂交,并且步骤(c)和(d)的所述gRNA与所述第二DNA的相对链杂交。

84. 权利要求81-83中任一项的方法,其中步骤(a)和(b)的所述gRNA;和/或步骤(c)和(d)的所述gRNA与它们各自的DNA的区域杂交,所述区域相隔不超过100个碱基对。

85. 权利要求84的方法,其中步骤(a)和(b)的所述gRNA;和/或步骤(c)和(d)的所述gRNA与它们各自的DNA的区域杂交,所述区域相隔不超过10、不超过15、不超过20、不超过25、不超过30、不超过40、不超过50、不超过60、不超过70、不超过80或不超过90个碱基对。

86. 权利要求81-85中任一项的方法,其中步骤(a)和(b)的所述gRNA;和/或步骤(c)和(d)的所述gRNA在重组酶位点旁侧的gRNA结合位点处与它们各自的DNA的区域杂交。

87. 权利要求86的方法,其中所述重组酶位点包含res、gix、hix、six、resH、LoxP、FTR或

att核心或相关核心序列。

88. 权利要求86或权利要求87的方法,其中所述重组酶位点包含gix核心或gix相关核心序列。

89. 权利要求87或权利要求88的方法,其中所述gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为3至7个碱基对。

90. 权利要求87-89中任一项的方法,其中所述gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为5至6个碱基对。

91. 用于单一DNA分子的两个区域之间的位点特异性重组的方法,其包括:

(a) 使所述DNA与第一融合蛋白接触,其中所述引导核苷酸序列-可编程DNA结合蛋白域结合第一gRNA,所述第一gRNA与所述DNA的第一区域杂交;

(b) 使所述DNA与第二融合蛋白接触,其中所述第二融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第二gRNA,所述第二gRNA与所述DNA的第二区域杂交;

(c) 使所述DNA与第三融合蛋白接触,其中所述第三融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第三gRNA,所述第三gRNA与所述DNA的第三区域杂交;和

(d) 使所述DNA与第四融合蛋白接触,其中所述第四融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第四gRNA,所述第四gRNA与所述DNA的第四区域杂交;

其中在使得所述DNA重组的条件下,步骤(a)-(d)中所述融合蛋白的结合导致所述融合蛋白的重组酶催化域的四聚化,并且其中所述第一、第二、第三和/或第四融合蛋白是权利要求1-63中任一项的融合蛋白。

92. 权利要求91的方法,其中重组的所述单一DNA分子的两个区域具有不同的序列。

93. 权利要求91或权利要求92的方法,其中所述重组导致所述DNA分子的区域的缺失。

94. 权利要求93的方法,其中缺失的所述DNA分子的区域易于在减数分裂中发生交换事件。

95. 权利要求91-94中任一项的方法,其中步骤(a)-(d)的所述第一和第二gRNA与所述DNA的相同链杂交,并且步骤(a)-(d)的所述第三和第四gRNA与所述DNA的相对链杂交。

96. 权利要求91-95中任一项的方法,其中步骤(a)和(b)的所述gRNA与所述DNA的区域杂交,所述区域相隔不超过100个碱基对,并且步骤(c)和(d)的所述gRNA与所述DNA的区域杂交,所述区域相隔不超过100个碱基对。

97. 权利要求96的方法,其中步骤(a)和(b)的所述gRNA与所述DNA的区域杂交,所述区域相隔不超过50、不超过60、不超过70、不超过80或不超过90个碱基对,并且步骤(c)和(d)的所述gRNA与所述DNA的区域杂交,所述区域相隔不超过10、不超过15、不超过20、不超过25、不超过30、不超过40、不超过50、不超过60、不超过70、不超过80或不超过90个碱基对。

98. 权利要求91-97中任一项的方法,其中步骤(a)和(b)的所述gRNA;和/或步骤(c)和(d)的所述gRNA与重组酶位点旁的gRNA结合位点杂交。

99. 权利要求98的方法,其中所述重组酶位点包含res、gix、hix、six、resH、LoxP、FTR或att核心或相关核心序列。

100. 权利要求98或权利要求99的方法,其中所述重组酶位点包含gix核心或gix相关核心序列。

101. 权利要求99或权利要求100的方法,其中所述gix核心或gix相关核心序列与至少

一个gRNA结合位点之间的距离为3至7个碱基对。

102. 权利要求99-101中任一项的方法,其中所述gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为5至6个碱基对。

103. 权利要求81-102中任一项的方法,其中所述DNA在细胞中。

104. 权利要求103的方法,其中所述细胞是真核细胞。

105. 权利要求103的方法,其中所述细胞是原核细胞。

106. 权利要求104或权利要求105的方法,其中所述细胞在受试者中。

107. 权利要求106的方法,其中所述受试者是人。

108. 多核苷酸,其编码权利要求1-63中任一项的融合蛋白。

109. 载体,其包含权利要求108的多核苷酸。

110. 用于重组蛋白表达的载体,其包含编码权利要求1-63中任一项的融合蛋白的多核苷酸。

111. 细胞,其包含用于表达权利要求1-63中任一项的融合蛋白的遗传构建体。

112. 试剂盒,其包含权利要求1-63中任一项的融合蛋白。

113. 试剂盒,其包含编码权利要求1-63中任一项的融合蛋白的多核苷酸。

114. 试剂盒,其包含用于重组蛋白表达的载体,其中所述载体包含编码权利要求1-63中任一项的融合蛋白的多核苷酸。

115. 试剂盒,其包含含有用于表达权利要求1-63中任一项的融合蛋白的遗传构建体的细胞。

116. 权利要求112-115中任一项的试剂盒,其进一步包含一个或多个gRNA和/或用于表达一个或多个gRNA的载体。

可编程CAS9-重组酶融合蛋白及其用途

[0001] 相关申请

[0002] 根据35U.S.C.§119(e),本申请要求2016年8月9日提交的美国临时专利申请U.S.S.N.62/372,755和2017年2月7日提交的美国临时专利申请U.S.S.N.62/456,048的优先权,其每一个通过引用并入本文。

[0003] 政府资助

[0004] 本发明是在国立卫生研究院(National Institutes of Health)授予的拨款号R01EB022376和R35GM118062下得到政府支持完成的。政府拥有本发明的某些权利。

[0005] 发明背景

[0006] 高效、可编程且位点特异性同源重组仍然是遗传学和基因组编辑的长期目标。将重组导向感兴趣的基因座的早期尝试依赖于用与靶基因座同源的长侧翼序列转染供体DNA。这种策略受到非常低的效率阻碍,并因此需要严格的选择来识别整合体。最近的努力利用了双链DNA断裂(DSB)诱导同源性定向修复(HDR)的能力。归巢内切核酸酶和后来的可编程内切核酸酶如锌指核酸酶、TALE核酸酶、Cas9和fCas9已用于引入靶定的DSB并在供体DNA存在下诱导HDR。然而,在大多数有丝分裂后细胞中,DSB诱导的HDR强烈下调并且通常效率低。此外,通过易错修复途径(例如非同源末端连接(NHEJ)或单链退火(SSA))修复DSB以比HDR更高的频率导致DSB位点处的核苷酸的随机插入或缺失(插入/缺失)。若细胞经受迫使细胞周期同步的条件或者若NHEJ中涉及的酶得以抑制,则可以提高HDR的效率。但是,此类情况可导致许多随机且不可预测的事件,从而限制了潜在的应用。本公开提供了融合蛋白,其可以重组含有侧翼为引导RNA指定序列的最小重组酶核心位点的DNA位点,并且代表朝向不依赖于内源性细胞机制或细胞状态的在未修饰的细胞中的可编程无痕(scarless)基因组编辑的步骤。

[0007] 发明概述

[0008] 本公开描述了融合蛋白的开发,所述融合蛋白包含引导核苷酸序列-可编程DNA结合蛋白域、任选的接头和重组酶催化域(例如,丝氨酸重组酶催化域如Gin重组酶催化域、酪氨酸重组酶催化域,或任何演化的重组酶催化域)。该融合蛋白在侧翼为两个引导RNA指定的DNA序列的最小gix核心重组酶位点(NNNNAAASSWWSSTTTNNNN,SEQ ID NO:19)上起作用。由所述融合蛋白介导的重组依赖于这两个引导RNA,导致不同引导核苷酸:融合蛋白复合物之间的正交性,并且在培养的人细胞中在与人基因组中发现的DNA序列匹配的DNA序列上有效地发挥功能。本公开的融合蛋白还可以直接在人细胞(例如,培养的人细胞)的基因组上起作用,催化位于相隔约14千碱基之间的两个recCas9假位点(pseudosite)之间的缺失、插入、倒位、易位或重组。这项工作提供了工程化的酶,其可以在未修饰的基因组中以用户定义的单碱基对分辨率催化基因插入、缺失、倒位或染色体易位。

[0009] 在一个方面,本发明提供融合蛋白,其包含:(i)引导核苷酸序列-可编程DNA结合蛋白域;(ii)任选的接头;和(iii)重组酶催化域,如任何丝氨酸重组酶催化域(包括但不限于Gin、Sin、Tn3、Hin、 β 、 γ δ 或PhiC31重组酶催化域)、任何酪氨酸重组酶域(包括但不限于Cre或FLP重组酶催化域),或任何演化的重组酶催化域。

(PEG)、聚丙二醇 (PPG)、共聚(乙烯/丙烯)二醇、聚氧乙烯(POE)、聚氨酯、聚磷腈、多糖、右旋糖酐、聚乙烯醇、聚乙烯吡咯烷酮、聚乙烯醚、聚丙烯酰胺、聚丙烯酸酯、聚氰基丙烯酸酯、脂质聚合物、甲壳质、透明质酸、肝素或烷基接头。在某些实施方案中,烷基接头具有式—NH—(CH₂)_s—C(O)—,其中s是1-100之间的任何整数,包括端点。在某些实施方案中,s是1-20的任何整数,包括端点。

[0016] 在另一个实施方案中,融合蛋白进一步包含核定位信号(NLS)域。在某些实施方案中,NLS域经由一个或多个第二接头与引导核苷酸序列-可编程DNA结合蛋白域或重组酶催化域结合。

[0017] 在一个实施方案中,融合蛋白包含结构NH₂-[重组酶催化域]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的第二接头序列]-[NLS域]-COOH。在某些实施方案中,融合蛋白与SEQ ID NO:719中所示的氨基酸序列具有大于85%、90%、95%、98%或99%的序列一致性。在具体的实施方案中,融合蛋白包含SEQ ID NO:719中所示的氨基酸序列。在一个实施方案中,融合蛋白由SEQ ID NO:719中所示的氨基酸序列组成。

[0018] 在另一个实施方案中,融合蛋白进一步包含一个或多个亲和标签。在一个实施方案中,亲和标签选自下组:FLAG标签、多组氨酸(多His)标签、多精氨酸(多Arg)标签、Myc标签和HA标签。在实施方案中,亲和标签是FLAG标签。在具体的实施方案中,FLAG标签具有序列PKKKRKV (SEQ ID NO:702)。在另一个实施方案中,一个或多个亲和标签经由一个或多个第三接头与引导核苷酸序列-可编程DNA结合蛋白域、重组酶催化域或NLS域结合。在某些实施方案中,第三接头是肽接头。

[0019] 本文描述的融合蛋白的元件可以不做限制以任何顺序。在一些实施方案中,融合蛋白具有结构NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH、NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH或NH₂-[二分叉(bifurcated)或环状置换(circularly permuted)的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。

[0020] 在一些实施方案中,融合蛋白具有结构NH₂-[任选的亲和标签]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-COOH、NH₂-[任选的亲和标签]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-COOH或NH₂-[任选的亲和标签]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-COOH。

[0021] 在某些实施方案中,融合蛋白与SEQ ID NO:185中所示的氨基酸序列具有大于85%、90%、95%、98%或99%的序列一致性。在具体的实施方案中,融合蛋白具有SEQ ID NO:185中所示的氨基酸序列。在某些实施方案中,融合蛋白的重组酶催化域与SEQ ID NO:

185的氨基酸1-142中所示的氨基酸序列(其与SEQ ID NO:713中所示的序列相同)具有大于85%、90%、95%、98%或99%的序列一致性。在某些实施方案中,dCas9域与SEQ ID NO:185的氨基酸167-1533中所示的氨基酸序列(其与SEQ ID NO:712中所示的序列相同)具有大于90%、95%或99%的序列一致性。在某些实施方案中,本公开的融合蛋白与SEQ ID NO:185的氨基酸1-1544中所示的氨基酸序列(其与SEQ ID NO:719中所示的序列相同)具有大于90%、95%或99%的序列一致性。在一个实施方案中,融合蛋白与引导RNA(gRNA)结合。

[0022] 在一个方面,本公开提供了本文所述的融合蛋白的二聚体。在某些实施方案中,二聚体与靶DNA分子结合。在某些实施方案中,二聚体的每个融合蛋白与靶DNA分子的相同链结合。在某些实施方案中,二聚体的每个融合蛋白与靶DNA分子的相对链结合。在某些实施方案中,二聚体的gRNA与靶DNA分子的重组酶位点侧翼的gRNA结合位点杂交。在某些实施方案中,重组酶位点包含res、gix、hix、six、resH、LoxP、FTR或att核心或相关核心序列。在某些实施方案中,重组酶位点包含gix核心或gix相关核心序列。在进一步的实施方案中,gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为3至7个碱基对。在某些实施方案中,gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为5至6个碱基对。

[0023] 在某些实施方案中,第一二聚体与第二二聚体结合,从而形成融合蛋白的四聚体。在一个方面,本公开提供了本文所述的融合蛋白的四聚体。在某些实施方案中,四聚体与靶DNA分子结合。在某些实施方案中,每个二聚体与DNA的相对链结合。在其他实施方案中,每个二聚体与DNA的相同链结合。

[0024] 在另一个方面,本公开提供了用于两个DNA分子之间的位点特异性重组的方法,其包括:(a)使第一DNA与第一融合蛋白接触,其中引导核苷酸序列-可编程DNA结合蛋白域结合第一gRNA,所述第一gRNA与第一DNA的第一区域杂交;(b)使第一DNA与第二融合蛋白接触,其中第二融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第二gRNA,所述第二gRNA与第一DNA的第二区域杂交;(c)使第二DNA与第三融合蛋白接触,其中第三融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第三gRNA,所述第三gRNA与第二DNA的第一区域杂交;和(d)使第二DNA与第四融合蛋白接触,其中第四融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第四gRNA,所述第四gRNA与第二DNA的第二区域杂交;其中在使得DNA重组的条件下,步骤(a)-(d)中融合蛋白的结合导致融合蛋白的重组酶催化域的四聚化,并且其中第一、第二、第三和/或第四融合蛋白是本文所述的任何融合蛋白。

[0025] 在一个实施方案中,第一和第二DNA分子具有不同的序列。在另一个实施方案中,步骤(a)和(b)的gRNA与第一DNA的相对链杂交,并且步骤(c)和(d)的gRNA与第二DNA的相对链杂交。在另一个实施方案中,其中步骤(a)和(b)的gRNA;和/或步骤(c)和(d)的gRNA与它们各自的DNA的区域杂交,所述区域相隔不超过10、不超过15、不超过20、不超过25、不超过30、不超过40、不超过50、不超过60、不超过70、不超过80、不超过90或不超过100个碱基对。在某些实施方案中,步骤(a)和(b)的gRNA;和/或步骤(c)和(d)的gRNA在重组酶位点侧翼的gRNA结合位点处与它们各自的DNA的区域杂交(参见例如图1D)。在某些实施方案中,重组酶位点包含res、gix、hix、six、resH、LoxP、FTR或att核心或相关核心序列。在某些实施方案中,重组酶位点包含gix核心或gix相关核心序列。在某些实施方案中,gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为3至7个碱基对。在某些实施方案中,gix核心

或gix相关核心序列与至少一个gRNA结合位点之间的距离为5至6个碱基对。

[0026] 本文提供的用于位点特异性重组的方法也可以与单一DNA分子一起使用。在一个方面,本公开提供了用于单一DNA分子的两个区域之间的位点特异性重组的方法,其包括:(a)使DNA与第一融合蛋白接触,其中引导核苷酸序列-可编程DNA结合蛋白域结合第一gRNA,所述第一gRNA与DNA的第一区域杂交;(b)使DNA与第二融合蛋白接触,其中第二融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第二gRNA,所述第二gRNA与DNA的第二区域杂交;(c)使DNA与第三融合蛋白接触,其中第三融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第三gRNA,所述第三gRNA与DNA的第三区域杂交;和(d)使DNA与第四融合蛋白接触,其中第四融合蛋白的引导核苷酸序列-可编程DNA结合蛋白域结合第四gRNA,所述第四gRNA与DNA的第四区域杂交;其中在使得DNA重组的条件下,步骤(a)-(d)中融合蛋白的结合导致融合蛋白的重组酶催化域的四聚化,并且其中第一、第二、第三和/或第四融合蛋白是所述的任何融合蛋白。

[0027] 在某些实施方案中,重组的单一DNA分子的两个区域具有不同的序列。在另一个实施方案中,重组导致DNA分子的区域缺失。在具体的实施方案中,缺失的DNA分子的区域易于在减数分裂中发生交换事件。在一个实施方案中,步骤(a)-(d)的第一和第二gRNA与DNA的相同链杂交,并且步骤(a)-(d)的第三和第四gRNA与DNA的相对链杂交。在另一个实施方案中,步骤(a)和(b)的gRNA与DNA的区域杂交,所述区域相隔不超过50、不超过60、不超过70、不超过80、不超过90或不超过100个碱基对,并且步骤(c)和(d)的gRNA与DNA的区域杂交,所述区域相隔不超过10、不超过15、不超过20、不超过25、不超过30、不超过40、不超过50、不超过60、不超过70、不超过80、不超过90或不超过100个碱基对。在某些实施方案中,步骤(a)和(b)的gRNA;和/或步骤(c)和(d)的gRNA与重组酶位点侧翼的gRNA结合位点杂交。在某些实施方案中,重组酶位点包含res、gix、hix、six、resH、LoxP、FTR或att核心或相关核心序列。在一个实施方案中,重组酶位点包含gix核心或gix相关核心序列。在某些实施方案中,gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为3至7个碱基对。在某些实施方案中,gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离为5至6个碱基对。

[0028] 本文所述的DNA可以在细胞中。在某些实施方案中,细胞是真核细胞。在某些实施方案中,细胞是植物细胞。在某些实施方案中,细胞是原核细胞。在一些实施方案中,细胞可以是哺乳动物细胞。在一些实施方案中,细胞可以是人细胞。在某些实施方案中,细胞在受试者中。在一些实施方案中,受试者可以是哺乳动物。在某些实施方案中,受试者是人。在某些实施方案中,细胞可以是植物细胞。

[0029] 在一个方面,本公开提供了编码本文所公开的任何融合蛋白的多核苷酸。在某些实施方案中,本公开提供了包含编码本文所公开的任何融合蛋白的多核苷酸的载体。

[0030] 在另一个方面,本公开提供了包含用于表达本文所公开的任何融合蛋白的遗传构建体的细胞。

[0031] 在一个方面,本公开提供了包含本文所公开的任何融合蛋白的试剂盒。在另一个方面,本公开提供了包含编码本文所公开的任何融合蛋白的多核苷酸的试剂盒。在另一个方面,本公开提供了包含用于重组蛋白表达的载体的试剂盒,其中所述载体包含编码本文所公开的任何融合蛋白的多核苷酸。在另一个方面,本公开提供了包含细胞的试剂盒,所述

细胞包含用于表达本文所公开的任何融合蛋白的遗传构建体。在一个实施方案中,试剂盒进一步包含一个或多个gRNA和/或用于表达一个或多个gRNA的载体。

[0032] 如下所述,在某些实施方案的详细描述中阐述了本发明的某些实施方案的细节。根据定义、实施例、附图和权利要求,本发明的其他特征、目的和优点将显而易见。

[0033] 附图简述

[0034] 图1A-1D。实验设置的概述。用(图1A)在hU6启动子的控制下的引导RNA表达载体、(图1B)在CMV启动子的控制下的recCas9表达载体和(图1C) recCas9报告质粒转染细胞。这些组分的共转染导致在靶位点处重新组装引导RNA编程的recCas9(图1D)。这将介导polyA终止子的缺失,允许GFP的转录。引导RNA表达载体和引导RNA序列缩写为gRNA。

[0035] 图2A-2F。融合接头长度和靶位点间隔区变体的优化。在这些实验中使用单一靶引导RNA表达载体pHU6-NT1或非靶载体pHU6-BC74。序列可以在表6-9中找到。(图2A)显示靶位点的一部分,其中引导RNA靶位点为黑色带有虚线下划线并且gix核心序列位点为黑色。假gix位点的任一侧的5'和3'序列是相同的,但是是反向的,并且被pHU6-NT1识别。将gix假位点与5'和3'结合位点分开的碱基对间隔区的数量分别由X和Y表示。该图分别描绘了SEQ ID NO:700和703。(图2B) Z表示将Ginβ与dCas9连接的GGs重复的数量。当X=Y时,对于连接Gin催化域和dCas9域的接头(图2C) (GGs)₂(SEQ ID NO:182)、(图2D) (GGs)₅(SEQ ID NO:701)和(图2E) (GGs)₈(SEQ ID NO:183),评估recCas9活性。(图2F)测定recCas9在由不均匀碱基对间隔区(X≠Y)组成的靶位点上的活性;包括X=Y=6用于比较。所有实验一式三份进行,并从这些实验中减去背景荧光。eGFP阳性细胞的百分比仅是那些转染的细胞的(即,表达组成型表达的iRFP基因),并且每个实验记录至少6,000个活事件(live event)。引导RNA表达载体和引导RNA序列缩写为“gRNA”。值和误差条分别代表三个独立生物学重复的平均值和标准偏差。

[0036] 图3A-3B。正向和反向引导RNA对recCas9活性的依赖性。(图3A)在PCDH15内发现的序列替换了图1A-1D中测试的靶位点。在假gix核心位点的5'和3'侧两者上,引导RNA可以靶向两个偏移序列。该图分别描绘了SEQ ID NO:704-705。(图3B) recCas9活性通过与所有四个引导RNA表达载体对和个别引导RNA载体与脱靶(O.T.)引导RNA载体共转染recCas9表达载体和报告质粒来测量。脱靶正向物和反向物含有分别靶向CLTA和VEGF的引导RNA序列。还显示了用报告质粒转染但未用靶引导RNA转染的对照实验。还显示了与不同引导RNA表达载体共转染但不与recCas9表达载体共转染的报告质粒的结果。所有实验一式四份进行,并且不从这些实验中减去背景荧光。eGFP阳性细胞的百分比仅是那些转染的细胞的(即,表达组成型表达的iRFP基因),并且每个实验记录至少6,000个活事件。引导RNA表达载体和引导RNA序列缩写为gRNA。值和误差条分别代表四个独立生物学重复的平均值和标准偏差。

[0037] 图4A-4D。recCas9可以靶向与人基因组中的序列相同的多个序列。(图4A)图1A-1D中所示的靶位点被人基因组内发现的序列替换。有关序列参见表6。将recCas9表达载体与引导RNA载体对和报告质粒的所有组合共转化。脱靶引导RNA载体也与recCas9表达载体和报告质粒共转化,并含有靶向CLTA和VEGF的引导RNA序列(参见例如Guilinger et al., Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. Nature biotechnology, (2014),其全部内容在此通过引用并入)。eGFP阳性细胞的百分比反映了经转染的(iRFP阳性)细胞的百分比。每个实

验至少记录6,000个活事件。值和误差条分别代表至少三个独立的生物学重复的平均值和标准偏差。(图4B)再次进行转染实验,用SpecR替换recCas9表达载体和pUC中的抗性标志物。共转染和温育后,提取附加型DNA(episomal DNA),转化到大肠杆菌中并选择羧苄青霉素抗性。然后对菌落进行测序以确定(图4C)重组质粒与完全完整质粒的比率。(图4D)分离自转染细胞的附加型提取物的测序数据。列和行代表转染条件。每个单元格显示重组的质粒的百分比和比率。显示的值反映了两个独立的生物学重复的平均值和标准偏差。平均值和每个重复之间的平均差值显示为误差。引导RNA表达载体和引导RNA序列缩写为gRNA。

[0038] 图5A-5D。recCas9介导培养的人细胞中基因组DNA的引导RNA-和recCas9-依赖性缺失。(图5A)示意图显示位于染色体12的FAM19A2基因座的内含子区域内的预测的recCas9靶位点和用于巢式PCR的引物的位置。该图分别从上到下和从左到右描绘了SEQ ID NO: 706-709。(图5B)来自用指定表达载体转染的细胞的模板的巢式基因组PCR的代表性结果($n=3$ 个生物学重复;NTC=无模板对照)。星号表示1.3-kb预测的初级PCR产物的位置。箭头表示二级PCR后预测的缺失产物。这两个图来自相同的凝胶,但经切割以去除空白泳道。(图5C)对用所有四种gRNA表达载体转染的细胞的巢式基因组PCR产生的PCR产物的Sanger测序,并且recCas9表达载体与预测的重组后产物匹配。该图分别从上到下描绘了SEQ ID NO: 710和711。(图5D)通过有限稀释巢式PCR测定的FAM19A2基因座的估计的最小缺失效率。显示的值反映了三次重复的平均值和标准偏差。

[0039] 图6。报告质粒构建。Golden Gate组装用于构建本工作中描述的报告质粒。所有组装均以共同质粒pCALNL-EGFP-Esp3I开始,该质粒衍生自pCALNL-EGFP并包含Esp3I限制性位点。显示的片段侧翼为Esp3I位点。Esp3I消化产生一系列相容的、独特的4碱基对5'突出端,以便按所示顺序进行组装。为了组装靶位点,将Esp3I(ThermoFisher Scientific, Waltham, MA)和五个片段加入到单个反应管中以允许Esp3I消化和T7连接的迭代循环。然后用Plasmid-Safe-ATP依赖性DNA酶(Epicentre, Madison, WI)消化反应以减少背景。通过菌落PCR分析菌落以鉴定与预期的全长5部分组装产物匹配的PCR产物;然后将来自这些菌落的质粒送去进行sanger测序。对于图4中所示的基因组报告物,将片段1和2以及片段4和5组合成编码整个靶位点的两个gBlock(IDT, Coralville, IA)片段(图中未显示)。然后如上所述完成组装。构建细节可以在支持材料的方法中找到。用于产生片段的寡核苷酸和gBLOCK可以在表2中找到。

[0040] 图7A和7B。演化以靶向人基因组的Rosa基因座中称为“36C6”的位点的Cre重组酶与dCas9融合。然后将该融合物用于以引导RNA依赖性方式重组含有Rosa靶位点的基于质粒的报告物。图7A证明了使用野生型Cre和36C6的接头优化的结果。显示靶向其关联报告物的GinB构建体用于参考。显示的1x 2x、5x和8x接头是接头中GGG重复的数量。图7B显示了回复分析的结果,其证明了对与dCas9融合的36C6进行突变可以影响嵌合融合物的相对引导依赖性。显示靶向其关联报告物的GinB构建体用于参考。GGG-36C6:1x GGS接头;2GGG-36C6(使用接头SEQ ID NO:181):2x GGS接头(使用接头SEQ ID NO:181)。

[0041] 图8。鉴定了人基因组中Rosa26位点侧翼的PAM,其可以支持dCas9结合(参见上图)。设计引导RNA和质粒报告物以测试内源性前间隔区是否可以支持dCas9-36C6活性。显示靶向gix报告物的GinB构建体用于参考。混合:Cas9和36C6之间所有5种接头变体的等份混合物。序列对应于SEQ ID NO:769(核苷酸序列)和770(氨基酸序列)。

[0042] 图9A-9B。Cre重组酶的各种测试的截短的位置显示在图9A中。与dCas9融合的Cre重组酶的截短的变体显示出可察觉的重组酶活性以及对Lox质粒报告物系统中引导RNA的存在的严格依赖性(图9B)。与dCas9融合的野生型Cre显示为阳性对照。

[0043] 定义

[0044] 如本文所用,除非上下文另外明确指出,否则单数形式“一种”、“一个”和“该/所述”包括单数和复数。因此,例如,提及“试剂”包括单一试剂和多个此类试剂。

[0045] 非限制性、示例性RNA-可编程DNA结合蛋白包括Cas9核酸酶、Cas9切口酶、核酸酶无活性的Cas9(dCas9)、CasX、CasY、Cpf1、C2c1、C2c2、C2C3和Argonaute。术语“Cas9”或“Cas9域”是指包含Cas9蛋白或其片段的RNA引导的核酸酶(例如,包含Cas9的活性、无活性或部分活性的DNA切割域,和/或Cas9的gRNA结合域的蛋白质)。Cas9具有两个切割域,其切割特定的DNA链(例如有义链和反义链)。可以产生切割任一条链的Cas9切口酶(包括但不限于spCas9的D10A和H840A)。可以不受限制地在本文所述的融合蛋白和方法中使用Cas9域(例如,核酸酶活性Cas9、核酸酶无活性的Cas9或Cas9切口酶)。此外,本文所述的任何引导核苷酸序列-可编程DNA结合蛋白可以用作切口酶。

[0046] Cas9核酸酶有时也称为casn1核酸酶或CRISPR(聚簇规则间隔短回文重复)相关核酸酶。CRISPR是适应性免疫系统,其提供针对移动遗传元件(病毒、转座元件和接合质粒)的保护。CRISPR簇含有间隔区,与先前的移动元件互补的序列,并靶向侵入核酸。CRISPR簇得以转录并加工成CRISPR RNA(crRNA)。在II型CRISPR系统中,对pre-crRNA的正确加工需要反式编码的小RNA(tracrRNA)、内源性核糖核酸酶3(rnc)和Cas9蛋白。tracrRNA充当用于pre-crRNA的核糖核酸酶3辅助加工的引导。随后,Cas9/crRNA/tracrRNA以内切核水解方式切割与间隔区互补的线性或环状dsDNA靶标。首先以内切核水解方式切割不与crRNA互补的靶链,然后以3'-5'外切核水解方式修剪(trim)。在自然界中,DNA结合和切割通常需要蛋白质和这两种RNA。然而,单一引导RNA(“sgRNA”或简称“gRNA”)可以经工程化以将crRNA和tracrRNA两者的方面并入单一RNA种类中。参见例如Jinek M.,Chylinski K.,Fonfara I.,Hauer M.,Doudna J.A.,Charpentier E.Science337:816-821(2012),其全部内容通过引用并入本文。Cas9识别CRISPR重复序列中的短基序(PAM或前间隔区相邻基序),以帮助区分自我与非自我。Cas9核酸酶序列和结构是本领域技术人员熟知的(参见例如“Complete genome sequence of an M1strain of Streptococcus pyogenes.”Ferretti et al.,J.J.,McShan W.M.,Ajdic D.J.,Savic D.J.,Savic G.,Lyon K.,Primeaux C.,Sezate S.,Suvorov A.N.,Kenton S.,Lai H.S.,Lin S.P.,Qian Y.,Jia H.G.,Najar F.Z.,Ren Q.,Zhu H.,Song L.,White J.,Yuan X.,Clifton S.W.,Roe B.A.,McLaughlin R.E.,Proc.Natl.Acad.Sci.U.S.A.98:4658-4663(2001);“CRISPR RNA maturation by trans-encoded small RNA and host factor RNase III.”Deltcheva E.,Chylinski K.,Sharma C.M.,Gonzales K.,Chao Y.,Pirzada Z.A.,Eckert M.R.,Vogel J.,Charpentier E.,Nature 471:602-607(2011);和“A programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity.”Jinek M.,Chylinski K.,Fonfara I.,Hauer M.,Doudna J.A.,Charpentier E.Science 337:816-821(2012),其各自的全部内容通过引用并入本文)。已经在各种物种中描述了Cas9直系同源物,包括但不限于酿脓链球菌(S.pyogenes)和嗜热链球菌(S.thermophilus)。基于本公开,其他合适的Cas9核酸酶和序

列对于本领域技术人员将是显而易见的,并且此类Cas9核酸酶和序列包括来自Chylinski, Rhun, and Charpentier, “The tracrRNA and Cas9 families of type II CRISPR-Cas immunity systems” (2013) RNA Biology 10:5, 726-737中公开的生物体和基因座的Cas9序列;其全部内容通过引用并入本文。在一些实施方案中,Cas9核酸酶具有无活性的(例如失活的)DNA切割域,也就是说,Cas9是切口的酶。作为一个实例,Cas9核酸酶(例如Cas9切口酶)可以切割与gRNA结合的DNA链。作为另一个实例,Cas9核酸酶(例如Cas9切口酶)可以切割不与gRNA结合的DNA链。在另一个实施方案中,任何引导核苷酸序列-可编程DNA结合蛋白可以具有无活性的(例如失活的)DNA切割域,即,引导核苷酸序列-可编程DNA结合蛋白是切口酶。作为一个实例,引导核苷酸序列-可编程DNA结合蛋白可以切割与gRNA结合的DNA链。作为另一个实例,引导核苷酸序列-可编程DNA结合蛋白可以切割不与gRNA结合的DNA链。

[0047] 另外的示例性Cas9序列可以在2017年4月27日公开的题为“Evolved Cas9 Proteins for Gene Editing”的国际公开号W0/2017/070633中找到。

[0048] 核酸酶失活的Cas9蛋白可以互换地称为“dCas9”蛋白(相当于核酸酶-“死亡的”Cas9)。在一些实施方案中,dCas9对应于,或部分或全部包含如下以SEQ ID NO:1所示的氨基酸。在一些实施方案中,提供了dCas9的变体(例如SEQ ID NO:1的变体)。例如,在一些实施方案中,提供了具有除D10A和H840A之外的突变的dCas9变体,其例如导致核酸酶失活的Cas9(dCas9)。举例来说,此类突变包括D10和H840处的其他氨基酸取代,或Cas9的核酸酶域内的其他取代(例如,HNH核酸酶亚域和/或RuvC1亚域中的取代)。在一些实施方案中,提供了dCas9的变体或同源物(例如SEQ ID NO:1的变体),其与SEQ ID NO:10至少约70%相同、至少约80%相同、至少约90%相同、至少约95%相同、至少约98%相同、至少约99%相同、至少约99.5%相同或至少约99.9%。在一些实施方案中,提供了dCas9的变体(例如SEQ ID NO:1的变体),其具有比SEQ ID NO:1短或长约5个氨基酸、约10个氨基酸、约15个氨基酸、约20个氨基酸、约25个氨基酸、约30个氨基酸、约40个氨基酸、约50个氨基酸、约75个氨基酸、约100个氨基酸或更多的氨基酸序列。

[0049] dCas9 (D10A和H840A) :

[0050] MDKKYSIGLAIGTNSVGVAVITDEYKVPKFKVVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLEESFLVEEDKKHERHPIFGNI VDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRLENLIAQLPGEKKNLFGNLIALSLGLTPNFKSNFDLAEDAQLQLSKD TYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSAMIKRYDEHHQDLTLLKALVRQQLEPKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYFPLKDNREKIEKILTFRI PYYVGPLARGNSRFAMTRKSEETITPWNFEVVDKGSASQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKI IKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPA IKKGILQTVKVVDELVKVMGRHKPENIV IEMARENQTTQKGQKNSRERMKRIE EGIKELGSQILKEHPVENTQLQNEKLYLYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAE RGGLSELKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVI TLKSKLVSDFRKDFQFYKVIKREINNYHHAHDAYLNAVVG TALIKKYPKLESEFVYGDYKVDVRKMI AKSEQEIGKATAKYFFYSNIMNFFKTEITLANGE

IRKRPLIETNGETGEIVWDKGRDFATVRKVLSPQVNIKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYG
GFDSPTVAYSVLVVAKEVEKSKKLSVKELLGITIMERSSEFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLEFLEN
GRKRMLASAGELQKGNELALPSKYVNFYLASHYEKLGKSPEDNEQKQLFVEQHKHYLDEIEQISEFSKRVLAD
ANLDKVL SAYNKHDKPIREQAENI IHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDTL IHQSITGLYETRI
DLSQLGGD (SEQ ID NO:1)

[0051] 用于生成具有无活性的DNA切割域的Cas9蛋白(或其片段)的方法是已知的(参见例如Jinek et al., Science.337:816-821(2012); Qi et al., “Repurposing CRISPR as an RNA-Guided Platform for Sequence-Specific Control of Gene Expression” (2013) Cell.28;152(5):1173-83,其各自的全部内容通过引用并入本文)。例如,已知Cas9的DNA切割域包括两个亚域,即HNH核酸酶亚域和RuvC1亚域。HNH亚域切割与gRNA互补的链,而RuvC1亚域切割非互补链。这些亚域内的突变可以沉默Cas9的核酸酶活性。例如,突变D10A和H840A完全使酿脓链球菌Cas9的核酸酶活性失活(参见例如,Jinek et al., Science.337:816-821(2012); Qi et al., Cell.28;152(5):1173-83(2013))。在一些实施方案中,提供了包含Cas9的片段的蛋白质。例如,在一些实施方案中,蛋白质包含两个Cas9域的一个:(1) Cas9的gRNA结合域;或(2) Cas9的DNA切割域。在一些实施方案中,包含Cas9或其片段的蛋白质称为“Cas9变体”。Cas9变体与Cas9或其片段共享同源性。例如,Cas9变体与野生型Cas9至少约70%相同、至少约80%相同、至少约90%相同、至少约95%相同、至少约96%相同、至少约97%相同、至少约98%相同、至少约99%相同、至少约99.5%相同或至少约99.9%。在一些实施方案中,Cas9变体包含Cas9的片段(例如,gRNA结合域或DNA切割域),使得该片段与野生型Cas9的相应的片段至少约70%相同、至少约80%相同、至少约90%相同、至少约95%相同、至少约96%相同、至少约97%相同、至少约98%相同、至少约99%相同、至少约99.5%相同或至少约99.9%。在一些实施方案中,野生型Cas9对应与来自酿脓链球菌(*Streptococcus pyogenes*)的Cas9(NCBI参考序列:NC_017053.1, SEQ ID NO:2(核苷酸); SEQ ID NO:3(氨基酸))。在一些实施方案中,Cas9域包含与野生型Cas9至少60%、至少65%、至少70%、至少75%、至少80%、至少85%、至少90%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列。在一些实施方案中,与野生型Cas9相比,Cas9域包含具有1、2、3、4、5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、21、24、25、26、27、28、29、30、31、32、33、34、35、36、37、38、39、40、41、42、43、44、45、46、47、48、49、50个或更多个突变的氨基酸序列。在一些实施方案中,与野生型Cas9相比,Cas9域包含具有至少10、至少15、至少20、至少30、至少40、至少50、至少60、至少70、至少80、至少90、至少100、至少150、至少200、至少250、至少300、至少350、至少400、至少500、至少600、至少700、至少800、至少900、至少1000、至少1100或至少1200个相同的连续氨基酸残基的氨基酸序列。在一些实施方案中,Cas9变体包含Cas9的片段(例如,gRNA结合域或DNA切割域),使得该片段与野生型Cas9的相应的片段至少约70%相同、至少约80%相同、至少约90%相同、至少约95%相同、至少约96%相同、至少约97%相同、至少约98%相同、至少约99%相同、至少约99.5%相同或至少约99.9%相同。在一些实施方案中,片段是相应的野生型Cas9的氨基酸长度的至少30%、至少35%、至少40%、至少45%、至少50%、至少55%、至少60%、至少65%、至少70%、至少75%、至少80%、至少85%、至少90%、至少95%相同、至少96%、至少97%、至少98%、至少99%或至少99.5%。

[0052] 在一些实施方案中,片段的长度为至少100个氨基酸。在一些实施方案中,片段的长度为至少100、150、200、250、300、350、400、450、500、550、600、650、700、750、800、850、900、950、1000、1050、1100、1150、1200、1250或1300个氨基酸。

[0053] ATGGATAAGAAATACTCAATAGGCTTAGATATCGGCACAAATAGCGTCGGATGGGCGGTGATCACTGATGATTATAAGGTTCCGCTCTAAAAAGTTCAAGGTTCTGGGAAATACAGACCGCCACAGTATCAAAAAAATCTTATAGGGGCTCTTTTATTTGGCAGTGGAGAGACAGCGGAAGCGACTCGTCTCAAACGGACAGCTCGTAGAAGGTATACACGTCGGAAGAATCGTATTTGTTATCTACAGGAGATTTTTTCAAATGAGATGGCGAAAAGTAGATGATAGTTTCTTTCA TCGACTTGAAGAGTCTTTTTTGGTGGGAAGAAGACAAGAAGCATGAACGTCATCCTATTTTTGGAAATATAGTAGAT GAAGTTGCTTATCATGAGAAAATATCCAATCTATCATCTGCGAAAAAATTTGGCAGATTCTACTGATAAAGCGG ATTTGCGCTTAATCTATTTGGCCTTAGCGCATATGATTAAGTTTTCGTGGTCATTTTTTGATTGAGGGAGATTTAAA TCCTGATAATAGTGATGTGGACAACTATTTATCCAGTTGGTACAAATCTACAATCAATTATTTGAAGAAAACCT ATTAACGCAAGTAGAGTAGATGCTAAAGCGATTCTTTCTGCACGATTGAGTAAATCAAGACGATTAGAAAATCTCA TTGCTCAGCTCCCCGGTGAGAAGAGAAATGGCTTGTTTTGGGAATCTCATTGCTTTGTCATTGGGATTGACCCCTAA TTTTAAATCAAATTTTGATTGGCAGAAGATGCTAAATTACAGCTTTCAAAAAGATACTTACGATGATGATTTAGAT AATTTATTGGCGCAAATTTGGAGATCAATATGCTGATTTGTTTTTGGCAGCTAAGAATTTATCAGATGCTATTTTAC TTTTACAGATATCCTAAGAGTAAATAGTGAATAACTAAGGCTCCCCTATCAGCTTCAATGATTAAGCGCTACGATGA ACATCATCAAGACTTGACTCTTTTAAAAGCTTTAGTTTCGACAACAACCTCCAGAAAAGTATAAAGAAATCTTTTTT GATCAATCAAAAAACGGATATGCAGGTTATATTGATGGGGGAGCTAGCCAAGAAGAATTTTATAAATTTATCAAAC CAATTTTAGAAAAAATGGATGGTACTGAGGAATTATTGGTGAACATAAATCGTGAAGATTGCTGCGCAAGCAACG GACCTTTGACAACGGCTCTATTTCCCATCAAATTCACCTGGGTGAGCTGCATGCTATTTGAGAAGACAAGAAGAC TTTTATCCATTTTTTAAAAGACAATCGTGAGAAGATTGAAAAAATCTTGACTTTTCGAATTCCTTATTATGTTGGTC CATTGGCGCGTGGCAATAGTCGTTTTGCATGGATGACTCGGAAGTCTGAAGAAACAATTACCCCATGGAATTTTGA AGAAGTTGTCGATAAAGGTGCTTCAGCTCAATCATTTATTGAACGCATGACAAACTTTGATAAAAAATCTTCAAAT GAAAAAGTACTACCAAAACATAGTTTGCTTTATGAGTATTTTACGGTTTATAACGAATTGACAAAGGTCAAATATG TTAAGTACTGAGGGAATGCGAAAACCAGCATTCTTTTACAGGTGAACAGAAGAAAGCCATTGTTGATTTACTCTTCAAAC AAATCGAAAAAGTAACCGTTAAGCAATTAAGAAGATTATTTCAAAAAAATAGAATGTTTTGATAGTGTGAAATT TCAGGAGTTGAAGATAGATTTAATGCTTCATTAGGCGCCTACCATGATTTGCTAAAAATTATTAAGATAAAGATT TTTTGGATAATGAAGAAAATGAAGATATCTTAGAGGATATTGTTTTAACATTGACCTTATTTGAAGATAGGGGGAT GATTGAGGAAAGACTTAAAACATATGCTCACCTCTTTGATGATAAGGTGATGAAACAGCTTAAACGTCGCCGTTAT ACTGGTTGGGGACGTTTGTCTCGAAAATTGATTAATGGTATTAGGGATAAGCAATCTGGCAAAAACAATATTAGATT TTTTGAATCAGATGGTTTTGCCAATCGCAATTTTATGCAGCTGATCCATGATGATAGTTTGACATTTAAGAAGA TATTCAAAAAGCACAGGTGTCTGGACAAGGCCATAGTTTACATGAACAGATTGCTAACTTAGCTGGCAGTCCTGCT ATTAAAAAAGGTATTTTACAGACTGTAAAAATTGTTGATGAACTGGTCAAAGTAATGGGGCATAAGCCAGAAAATA TCGTTATTGAAATGGCACGTGAAAATCAGACAACCTCAAAGGGCCAGAAAAATTCGCGAGAGCGTATGAAACGAAT CGAAGAAGGTATCAAAGAATTAGGAAGTCAGATTCTTAAAGAGCATCCTGTTGAAAATACTCAATTGCAAAATGAA AAGCTCTATCTCTATTATCTACAAAATGGAAGAGACATGTATGTGGACCAAGAATTAGATATTAATCGTTTAAAGTG ATTATGATGTGATCACATTGTTCCACAAAGTTTCATTAAGACGATTCAATAGACAATAAGGTTACTAACGCGTTC TGATAAAAAATCGTGGTAAATCGGATAACGTTCCAAGTGAAGAAGTAGTCAAAAAGATGAAAAACTATTGGAGACAA CTTCTAAACGCCAAGTTAATCACTCAACGTAAGTTTGATAATTTAACGAAAGCTGAACGTGGAGGTTTGAGTGAAC

TTGATAAAGCTGGTTTTATCAAACGCCAATTGGTTGAAACTCGCCAAATCACTAAGCATGTGGCACAAATTTTGGA
TAGTCGCATGAATACTAAATACGATGAAAATGATAAACTTATTCGAGAGGTTAAAGTGATTACCTAAAATCTAAA
TTAGTTTCTGACTTCGAAAAGATTTCCAATTCTATAAAGTACGTGAGATTAACAATTACCATCATGCCCATGATG
CGTATCTAAATGCCGTCGTTGGAAGTCTTTGATTAAGAAATATCCAAAACCTGAATCGGAGTTTGTCTATGGTGA
TTATAAGTTTATGATGTTGTAATAATGATTGCTAAGTCTGAGCAAGAAAATAGGCAAAGCAACCGCAAATATTTTC
TTTTACTCTAATATCATGAACTTCTTCAAACAGAAATTACACTTGCAAATGGAGAGATTTCGCAAACGCCCTCTAA
TCGAAACTAATGGGGAACTGGAGAAATTGTCTGGGATAAAGGGCGAGATTTTGCCACAGTGCACAAAGTATTGTC
CATGCCCAAGTCAATATTGTCAAGAAAACAGAAGTACAGACAGGCGGATTCTCCAAGGAGTCAATTTTACAAAA
AGAAATTCGGACAAGCTTATTGCTCGTAAAAAAGACTGGGATCCAAAAAATATGGTGGTTTTGATAGTCCAACGG
TAGCTTATTCAGTCCTAGTGGTTGCTAAGGTGGAAAAAGGAAATCGAAGAAGTTAAAATCCGTTAAAGAGTTACT
AGGGATCACAATTATGGAAAGAAGTTCCTTTGAAAAAATCCGATTGACTTTTTAGAAAGCTAAAGGATATAAGGAA
GTTAAAAAGACTTAATCATTAAACTACCTAAATATAGTCTTTTTGAGTTAGAAAACGGTCGTAAACGGATGCTGG
CTAGTGCCGGAGAATTACAAAAAGGAAATGAGCTGGCTCTGCCAAGCAAATATGTGAATTTTTTATATTTAGCTAG
TCATTATGAAAAGTTGAAGGGTAGTCCAGAAGATAACGAACAAAAACAATTGTTTGTGGAGCAGCATAAGCATTAT
TTAGATGAGATTATTGAGCAAATCAGTGAATTTTCTAAGCGTGTTATTTTAGCAGATGCCAATTTAGATAAAGTTC
TTAGTGCATATAACAAACATAGAGACAAACCAATACGTGAACAAGCAGAAAAATATTATTCATTTATTTACGTTGAC
GAATCTTGGAGCTCCCGCTGCTTTTAAATATTTTGATACAACAATTGATCGTAAACGATATACGTCTACAAAAGAA
GTTTTAGATGCCACTCTTATCCATCAATCCATCACTGGTCTTTATGAAACACGCATTGATTTGAGTCAGCTAGGAG
GTGACTGA (SEQ ID NO:2)

[0054] MDKKYSIGLDIGTNSVGVAVITDDYKVPKFKVLGNTDRHSIKKNLIGALLFGSGETAETRLKRTA
RRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLSEESFLVEEDKKHERHPIFGNI VDEVAYHEKYPTIYHLRKKLAD
STDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQIYNQLFEENPINASRVDAKAILSARLSKSR
RLENLIAQLPGEKRNGLFGNLI ALSLGLTPNFKSNFDLAEDAQLQLSKDYYDDLDNLLAQIGDQYADLFLAAKNL
SDAILLSDILRVNSEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEF
YKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYFPLKDNREKIEKILTFRI
PYVVGPLARGNSRFAMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNEL
TKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGAYHDLLKI
IKDKDFLDNEENEDILEDIVLTLTLFEDRGMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSG
KTILDFLKSDFANRNFMLIHDDSLTFKEDIQKAQVSGQGHSLEHQAANLAGSPA IKKGILQTVKIVDELVKVMG
HKPENIVIEMARENQTTQKGQKNSRERMKRIEEG IKELGSQILKEHPVENTQLQNEKLYLYLQNGRDMYVDQELD
INRLSDYDVDHIVPQSF IKDDS IDNKVLRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAER
GGLSELKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSCLVSDFRKDFQFYKREINNY
HHAHDAYLNAVVGTA LIKKYPKLESEFVYGDYKVYDVRKMIKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEI
RKRPLIETNGETGEIVWDKGRDFATVRKVL SMPQVNI VKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGG
FDSPTVAYSVLVVAKVEKGSKLLKSVKELLGITIMERSSEKNPIDFLEAKGYKEVKKDLI IKLPKYSLEFLENG
RKRMLASAGELQKGNELALPSKYVNFLLASHYEKLGSPEDNEQKQLFVEQHKHYLDEIEEQISEFSKRVILADA
NLDKVLSAYNKHRDKPIREQAENI IHLFTLTNLGAPAAFYFDTTIDRKRYTSTKEVLDATLIHQSI TGLYETRID
LSQLGGD (SEQ ID NO:3)

[0055] 在一些实施方案中,野生型Cas9对应于,或包含SEQ ID NO:4 (核苷酸) 和/或SEQ

ID NO:5 (氨基酸)。

[0056] ATGGATAAAAAGTATTCTATTGGTTTAGACATCGGCACTAATTCCGTTGGATGGGCTGTCATAACCGA
TGAATACAAAGTACCTTCAAAGAAATTTAAGGTGTTGGGGAACACAGACCGTCATTTCGATTA AAAAGAATCTTATC
GGTGCCCTCCTATTTCGATAGTGGCGAAACGGCAGAGGCGACTCGCCTGAAACGAACCGCTCGGAGAAGGTATACAC
GTCGCAAGAACCGAATATGTTACTTACAAGAAATTTTTAGCAATGAGATGGCCAAAGTTGACGATTCTTTCTTTCA
CCGTTTGAAGAGTCCTTCCCTGTGCGAAGAGGACAAGAAACATGAACGGCACCCCATCTTTGGAAACATAGTAGAT
GAGGTGGCATATCATGAAAAGTACCCAACGATTTATCACCTCAGAAAAAGCTAGTTGACTCAACTGATAAAGCGG
ACCTGAGGTTAATCTACTTGGCTCTTGCCCATATGATAAAGTTCCGTGGGCACTTTCTCATTGAGGGTGATCTAAA
TCCGGACAACCTCGGATGTCGACAACTGTTTCATCCAGTTAGTACAAACCTATAATCAGTTGTTTGAAGAGAACCCT
ATAAATGCAAGTGGCGTGGATGCGAAGGCTATTCTTAGCGCCCGCTCTCTAAATCCCGACGGCTAGAAAACCTGA
TCGCACAATTACCCGGAGAGAAGAAAAATGGGTTGTTTCGGTAACCTTATAGCGCTCTCACTAGGCCTGACACCAAA
TTTTAAGTCGAACTTCGACTTAGCTGAAGATGCCAAATTGCAGCTTAGTAAGGACACGTACGATGACGATCTCGAC
AATCTACTGGCACA AATTTGGAGATCAGTATGCGGACTTATTTTTGGCTGCCAAAAACCTTAGCGATGCAATCCTCC
TATCTGACATACTGAGAGTTAATACTGAGATTACCAAGGCGCCGTTATCCGCTTCAATGATCAAAGGTACGATGA
ACATCACCAAGACTTGACACTTCTCAAGGCCCTAGTCCGTGAGCAACTGCCTGAGAAAATATAAGGAAATATTCTTT
GATCAGTCGAAAAACGGGTACGCAGGTTATATTGACGGCGGAGCGAGTCAAGAGGAATTCTACAAGTTTATCAAAC
CCATATTAGAGAAGATGGATGGGACGGAAGAGTTGCTTGTA AAAACTCAATCGCGAAGATCTACTGCGAAAGCAGCG
GACTTTCGACAACGGTAGCATTCCACATCAAATCCACTTAGGCGAATTGCATGCTATACTTAGAAGGCAGGAGGAT
TTTTATCCGTTCTCAAAGACAATCGTAAAAGATTGAGAAAATCCTAACCTTTCGCATACCTACTATGTGGGAC
CCCTGGCCCAGGGAACTCTCGGTTGCGATGGATGACAAGAAAGTCCGAAGAAACGATTACTCCATGGAATTTTGA
GGAAGTTGTCGATAAAGGTGCGTCAGCTCAATCGTTCATCGAGAGGATGACCAACTTTGACAAGAATTTACCGAAC
GAAAAAGTATTGCCTAAGCACAGTTTACTTTACGAGTATTTACAGTGTACAATGAACTCACGAAAGTTAAGTATG
TCACTGAGGGCATGCGTAAACCCGCCTTTCTAAGCGGAGAACAGAAGAAAGCAATAGTAGATCTGTTATTCAAGAC
CAACCGCAAAGTGACAGTTAAGCAATTGAAAGAGGACTACTTTAAGAAAATTGAATGCTTCGATTCTGTGAGATC
TCCGGGTAGAAGATCGATTTAATGCGTCACTTGGTACGTATCATGACCTCCTAAAGATAATTAAGATAAGGACT
TCCTGGATAACGAAGAGAATGAAGATATCTTAGAAGATATAGTGTGACTCTTACCCTCTTTGAAGATCGGGAAAT
GATTGAGGAAAGACTAAAAACATACGCTCACCTGTTGACGATAAGGTTATGAAACAGTTAAAGAGGCGTCGCTAT
ACGGGCTGGGGACGATTGTCGCGGAACTTATCAACGGGATAAGAGACAAGCAAAGTGGTAAAACCTATTCTCGATT
TTCTAAAGAGCGACGGCTTCGCCAATAGGAACTTTATGCAGCTGATCCATGATGACTCTTTAACCTTCAAAGAGGA
TATACAAAAGGCACAGGTTCCGGACAAGGGGACTCATTGCACGAACATATTGCGAATCTTGCTGGTTCCGACGCC
ATCAAAAAGGCATACTCCAGACAGTCAAAGTAGTGGATGAGCTAGTTAAGGTCATGGGACGTCACAAACCGGAAA
ACATTGTAATCGAGATGGCACGCGAAAATCAAACGACTCAGAAGGGGCAAAAAAACAGTCGAGAGCGGATGAAGAG
AATAGAAGAGGGTATTAAGA AACTGGGCAGCCAGATCTTAAAGGAGCATCCTGTGGA AAAATACCCAATTGCAGAAC
GAGAACTTTACCTCTATTACCTACAAAATGGAAGGGACATGTATGTTGATCAGGAACTGGACATAAACCGTTTAT
CTGATTACGACGTCGATCACATTGTACCCCAATCCTTTTTGAAGGACGATTCAATCGACAATAAAGTGCTTACACG
CTCGGATAAGAACCGAGGGAAAAGTGACAATGTTCCAAGCGAGGAAGTCGTAAAGAAAATGAAGAACTATTGGCGG
CAGCTCCTAAATGCGAACTGATAACGCAAAGAAAGTTGATAACTTAACTAAAGCTGAGAGGGGTGGCTTGTCTG
AACTTGACAAGGCCGATTTATTAACGTCAGCTCGTGAAACCCGCCAAATCACAAAGCATGTTGCACAGATACT
AGATTCCCGAATGAATACGAAATACGACGAGAACGATAAGCTGATTCGGGAAGTCAAAGTAATCACTTTAAAGTCA

AAATTGGTGTCCGACTTCAGAAAGGATTTTCAATTCTATAAAGTTAGGGAGATAAATAACTACCACCATGCGCAGC
ACGCTTATCTTAATGCCGTCGTAGGGACCGCACTCATTAAGAAATACCCGAAGCTAGAAAAGTGAGTTTGTGTATGG
TGATTACAAAGTTTATGACGTCCGTAAGATGATCGCGAAAAGCGAACAGGAGATAGGCAAGGCTACAGCCAAATAC
TTCTTTTATTCTAACATTATGAATTTCTTTAAGACGGAAATCACTCTGGCAAACGGAGAGATACGCAAACGACCTT
TAATTGAAACCAATGGGGAGACAGGTGAAATCGTATGGGATAAGGGCCGGGACTTCGCGACGGTGAGAAAAGTTTT
GTCCATGCCCCAAGTCAACATAGTAAAGAAAAGTGGAGTGCAGACCGGAGGGTTTTCAAAGGAATCGATTCTTCCA
AAAAGGAATAGTGATAAGCTCATCGCTCGTAAAAAGGACTGGGACCCGAAAAAGTACGGTGGCTTCGATAGCCCTA
CAGTTGCCTATTCTGTCTAGTAGTGGCAAAGTTGAGAAGGGAAAAATCCAAGAAACTGAAGTCAGTCAAAGAATT
ATTGGGGATAACGATTATGGAGCGCTCGTCTTTTAAAAAGAACCCCATCGACTTCCTTGAGGCGAAAGGTTACAAG
GAAGTAAAAAAGGATCTCATAATTAAGTACCAAAGTATAGTCTGTTTGTAGTTAGAAAATGGCCGAAAACGGATGT
TGGCTAGCGCCGAGAGCTTCAAAGGGGAACGAACACTCGCACTACCGTCTAAATACGTGAATTTCTGTATTTAGC
GTCCCATTACGAGAAGTTGAAAGGTTACCTGAAGATAACGAACAGAAGCAACTTTTTGTTGAGCAGCACAAACAT
TATCTCGACGAAATCATAGAGCAAATTCGGAATTCAGTAAGAGAGTCATCCTAGCTGATGCCAATCTGGACAAAG
TATTAAGCGCATAACAACAGCACAGGGATAAACCCATACGTGAGCAGGCGGAAAAATATTATCCATTTGTTTACTCT
TACCAACCTCGGCGCTCCAGCCGATTCAAGTATTTGACACAACGATAGATCGCAAACGATACACTTCTACCAAG
GAGGTGCTAGACGCGACACTGATCACCAATCCATCACGGGATTATATGAAACTCGGATAGATTTGTCACAGCTTG
GGGTGACGGATCCCCAAGAAGAAGAGGAAAGTCTCGAGCGACTACAAAGACCATGACGGTATTATAAAGATCA
TGACATCGATTACAAGGATGACGATGACAAGGCTGCAGGA (SEQ ID NO:4)

[0057] MDKKYSIGLAIGTNSVGVAVITDEYKVPKFKVLGNTDRHSIKKNIIGALLFDSGETAEATRLKRTA
RRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLEESFLVEEDKKHERHPIFGNIIVDEVAYHEKYPTIYHLRKKLVD
STDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSR
RLENLIAQLPGEKKNGLFGNLIASLGLTPNFKSNFDLAEDAQLQSKDYYDDLDNLLAQIGDQYADLFLAAKNL
SDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEF
YKFIKPILEKMDGTEELLVKLNREDLLRQRTFDNGSIPHQIHLGELHAILRRQEDFYFPLKDNREKIEKILTFRI
PYVVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPHKSLLYEYFTVYNEL
TKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLKI
IKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSG
KTILDFLKSDFANRNFMLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMG
RHKPENIVIAMARENQTTQKGQKNSRERMKRIEEDIKELGSQILKEHPVENTQLQNEKLYLYLQNGRDMYVDQEL
DINRLSDYDHDHIVPQSFLKDDSIDNKVLRSDKNRGSNDNPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAE
RGGLSELKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVI TLKSKLVSDFRKDFQFYKVI
YHHAHDAYLNAVVGTALEKYPKLESEFVYGDYKVDVRKMIKSEQEI GKATAKYFFYSNIMNFFKTEITLANGE
IRKRPLIETNGETGEIVWDKGRDFATVRKVL SMPQVNI VKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYG
GFDSPTVAYSVLVVAKEKGSKKLKSVELLGITIMERSSEFKNPIDFLEAKGYKEVKKDLI IKLPKYSLELEN
GRKRMLASAGELQKGNELALPSKYVNFYLASHYEKLGSPEDNEQKQLFVEQHKHYLDEIEQISEFSKRVI LAD
ANLDKVL SAYNKHDKPIREQAENI IHLFTLTNLGAPAAFYFDTTIDRKRYTSTKEVL DATLIHQSI TGLYETRI
DLSQLGGD (SEQ ID NO:5)

[0058] 在一些实施方案中,Cas9是指来自以下的Cas9:溃疡棒杆菌(*Corynebacterium ulcerans*) (NCBI Refs:NC_015683.1,NC_017317.1);白喉棒杆菌(*Corynebacterium*

diphtheria) (NCBI Refs:NC_016782.1,NC_016786.1);Spiroplasma syrphidicola (NCBI Ref:NC_021284.1);间型普雷沃氏菌 (Prevotella intermedia) (NCBI Ref:NC_017861.1);台湾螺原体 (Spiroplasma taiwanense) (NCBI Ref:NC_021846.1);海豚链球菌 (Streptococcus iniae) (NCBI Ref:NC_021314.1);波罗的海贝尔氏菌 (Belliella baltica) (NCBI Ref:NC_018010.1);Psychroflexus torquisI (NCBI Ref:NC_018721.1);嗜热链球菌 (Streptococcus thermophilus) (NCBI Ref:YP_820832.1)、无害李斯特氏菌 (Listeria innocua) (NCBI Ref:NP_472073.1)、空肠弯曲杆菌 (Campylobacter jejuni) (NCBI Ref:YP_002344900.1) 或脑膜炎奈瑟氏球菌 (Neisseria meningitidis) (NCBI Ref:YP_002342100.1) 或者是指来自任何其他生物体的Cas9。

[0059] Cas9识别靶DNA序列中CRISPR重复序列中的短基序 (PAM基序)。如本文所用,“PAM基序”或“前间隔区相邻基序”是指在CRISPR细菌适应性免疫系统中直接在由Cas9核酸酶靶向的DNA序列之后的DNA序列。PAM是入侵病毒或质粒的组分,但不是细菌CRISPR基因座的组分。自然地,若其后没有PAM序列,则Cas9不会成功结合或切割靶DNA序列。PAM是靶向组分 (在细菌基因组中未发现),其区分细菌自我与非自我DNA,从而防止CRISPR基因座被Cas9核酸酶活性靶向和破坏。

[0060] 野生型酿脓链球菌 (Streptococcus pyogenes) Cas9识别规范PAM序列 (例如来自嗜热链球菌 (Streptococcus thermophilus)、金黄色葡萄球菌、脑膜炎奈瑟氏球菌或齿垢密螺旋体 (Treponema denticola) 的Cas9) 及其Cas9变体已在本领域中描述为具有不同的或更放松的PAM需求。通常,Cas9蛋白,例如来自酿脓链球菌的Cas9 (spCas9),需要规范的NGG PAM序列来结合特定的核酸区域,其中“NGG”中的“N”是腺嘌呤 (A)、胸腺嘧啶 (T)、鸟嘌呤 (G) 或胞嘧啶 (C),并且G是鸟嘌呤。这可以限制在基因组内编辑期望的碱基的能力。在一些实施方案中,本文提供的碱基编辑融合蛋白需要定位于精确的位置处,例如,其中靶碱基在4碱基区域 (例如“脱氨基作用窗口”) 内,其在PAM的上游的约15个碱基。参见Komor,A.C., et al.,“Programmable editing of a target base in genomic DNA without double-stranded DNA cleavage”Nature 533,420-424 (2016),其全部内容在此通过引用并入。在一些实施方案中,脱氨基作用窗口在2、3、4、5、6、7、8、9或10碱基区域内。在一些实施方案中,脱氨基作用窗口在PAM的上游的5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、23、24或25个碱基。因此,在一些实施方案中,本文提供的任何融合蛋白可以含有能够结合不含规范的 (例如,NGG) PAM序列的核苷酸序列的Cas9域。本领域中已经描述了结合非规范PAM序列的Cas9域,并且其对于熟练技术人员而言将是显而易见的。例如,结合非规范PAM序列的Cas9域已经描述于Kleinstiver,B.P., et al.,“Engineered CRISPR-Cas9 nucleases with altered PAM specificities”Nature 523,481-485 (2015);和Kleinstiver,B.P., et al.,“Broadening the targeting range of Staphylococcus aureus CRISPR-Cas9 by modifying PAM recognition”Nature Biotechnology 33,1293-1298 (2015);每篇的全部内容在此通过引用并入。还参见:Kleinstiver et al.,Nature 529,490-495,2016;Ran et al.,Nature, Apr 9;520 (7546):186-191,2015;Hou et al., Proc Natl Acad Sci U S A,110 (39):15644-9,2014;Prykhodzhi et al., PLoS One,10 (3):e0119372,2015;Zetsche et al.,Cell 163,759-771,2015;Gao et al.,Nature Biotechnology,doi:10.1038/nbt.3547,2016;Want et al.,Nature 461,754-761,2009;

Chavez et al., doi:dx dot doi dot org/10.1101/058974; Fagerlund et al., *Genome Biol.* 2015;16:25, 2015; Zetsche et al., *Cell*, 163, 759-771, 2015; 和 Swarts et al., *Nat Struct Mol Biol*, 21 (9) :743-53, 2014, 其每一个的全部内容通过引用并入本文。

[0061] 因此,本公开的引导核苷酸序列-可编程DNA结合蛋白可以识别多种PAM序列,包括但不限于:NGG、NGAN (SEQ ID NO:741)、NGNG (SEQ ID NO:742)、NGAG (SEQ ID NO:743)、NGCG (SEQ ID NO:744)、NNGRRT (SEQ ID NO:745)、NGRRN (SEQ ID NO:746)、NNNRRT (SEQ ID NO:747)、NNGATT (SEQ ID NO:748)、NNAGAAW (SEQ ID NO:749)、NAAAC (SEQ ID NO:750)、TTN、TTTN (SEQ ID NO:751) 和YTN,其中Y是嘧啶,并且N是任何核碱基。

[0062] 具有不同的PAM特异性的RNA-可编程DNA结合蛋白的一个实例是来自普雷沃氏菌 (*Prevotella*) 和弗朗西斯菌 (*Francisella*) 1 (Cpf1) 的聚簇规则间隔短回文重复。与Cas9类似,Cpf1也是2类CRISPR效应物。已经显示,Cpf1介导了强大的DNA干扰,其具有与Cas9不同的特征。Cpf1是缺乏tracrRNA的单一RNA引导的内切核酸酶,并且它利用富含T的前间隔区相邻基序 (TTN、TTTN (SEQ ID NO:751) 或YTN)。此外,Cpf1经由交错的DNA双链断裂切割DNA。在16种Cpf1家族蛋白中,来自氨基酸球菌 (*Acidaminococcus*) 和毛螺菌 (*Lachnospiraceae*) 的两种酶显示在人细胞中具有有效的基因组编辑活性。

[0063] 本文还提供了核酸酶无活性的Cpf1 (dCpf1) 变体,其可以用作RNA-可编程DNA结合蛋白域。Cpf1蛋白具有RuvC样内切核酸酶域,其类似于Cas9的RuvC域,但不具有HNH内切核酸酶域,并且Cpf1的N端不具有Cas9的alpha螺旋识别叶 (lobe)。它在Zetsche et al., *Cell*, 163, 759-771, 2015 (其全部内容通过引用并入本文) 中显示,Cpf1的RuvC样域负责切割两条DNA链并且RuvC样域的失活使Cpf1核酸酶活性失活。例如,对应于新凶手弗朗西斯菌 (*Francisella novicida*) Cpf1 (SEQ ID NO:714) 中的D917A、E1006A或D1255A的突变使Cpf1核酸酶活性失活。在一些实施方案中,本公开的dCpf1包含对应于SEQ ID NO:714中D917A、E1006A、D1255A、D917A/E1006A、D917A/D1255A、E1006A/D1255A或D917A/E1006A/D1255A的突变。应当理解,可以根据本公开使用使Cpf1的RuvC域失活的任何突变,例如取代突变、缺失或插入。

[0064] 在一些实施方案中,本公开的引导核苷酸序列-可编程DNA结合蛋白域对PAM序列没有需求。此类引导核苷酸序列-可编程DNA结合蛋白的一个实例可以是来自格氏嗜盐碱杆菌 (*Natronobacterium gregoryi*) 的Argonaute蛋白 (NgAgo)。NgAgo是ssDNA引导的内切核酸酶。NgAgo结合约24个核苷酸的5'磷酸化ssDNA (gDNA), 将其引导至其靶位点,并将在gDNA位点处产生DNA双链断裂。与Cas9相比,NgAgo-gDNA系统不需要前间隔区相邻基序 (PAM)。使用核酸酶无活性的NgAgo (dNgAgo) 可以极大地扩展可以靶向的密码子。NgAgo的表征和使用已经描述于Gao et al., *Nat Biotechnol.* Epub 2016 May 2. PubMed PMID: 27136078; Swarts et al., *Nature*. 507 (7491) (2014) :258-61; 和 Swarts et al., *Nucleic Acids Res.* 43 (10) (2015) :5120-9, 其每个的全部内容通过引用并入本文。格氏嗜盐碱杆菌 Argonaute 的序列提供于SEQ ID NO:718中。

[0065] 本文还提供了具有放松的PAM需求的Cas9变体 (无PAM (PAMless) 的Cas9)。与如SEQ ID NO:1提供的酿脓链球菌Cas9相比,无PAM的Cas9对靶序列表现出增加的活性,所述靶序列在其3'端处不包含规范的PAM (NGG), 例如增加的活性为至少5倍、至少10倍、至少50倍、至少100倍、至少500倍、至少1,000倍、至少5,000倍、至少10,000倍、至少50,000倍、至少100,

000倍、至少500,000倍或至少1,000,000倍。因此,本公开的dCas9或Cas9切口酶可以进一步包含放松PAM需求的突变,例如,对应于SEQ ID NO:1中A262T、K294R、S409I、E480K、E543D、M694I或E1219V的突变。

[0066] 应当理解,另外的Cas9蛋白(例如,核酸酶死亡的Cas9(dCas9)、Cas9切口酶(nCas9)或核酸酶活性Cas9),包括其变体和同源物,都在本公开的范围。示例性Cas9蛋白包括但不限于下文提供的那些。在一些实施方案中,Cas9蛋白是核酸酶死亡的Cas9(dCas9)。在一些实施方案中,dCas9包含如下所示的氨基酸序列。在一些实施方案中,Cas9蛋白是Cas9切口酶(nCas9)。在一些实施方案中,nCas9包含如下所示的氨基酸序列。在一些实施方案中,Cas9蛋白是核酸酶活性Cas9。在一些实施方案中,核酸酶活性Cas9包含如下所示的氨基酸序列。

[0067] 示例性的催化无活性的Cas9(dCas9):DKKYSIGLAIGTNSVGWAVITDEYKVPKSKFKVLGN
TDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLEESFLVEEDKKH
ERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLV
QTYNQLFEENPINASGVDAKAILSARLSKSRLENLIAQLPGEKKNLFGNLIASLGLTPNFKSNFDLAEDAQLQ
LSKDTYDDDLNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQ
QLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLG
ELHAILRRQEDFYFPLKDNREKIEKILTRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIE
RMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYF
KKIECFDSVEISGVEDRFNASLGTYHDLLKIKDKDFLDNEENEDILEDIVLTLTLFEDREMIERLKYAHLFDD
KVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKITLDFLKSDFANRNFMLIHDDSLTFKEDIQKAQVSGQGDSLH
EHIANLAGSPAIKKILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGGKNSRERMKRIEEDIKELGSQILK
EHPVENTQLQNEKLYLYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLRSDKNRGKSDNVPSE
EVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKL
IREVKVITLKSCLVSDFRKDFQFYKVVREINNYHHAHDAYLNAVVGTAIIKKYPKLESEFVYGDYKVDYVRKMIAS
EQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSPQVNIIVKKTVEVQ
TGGFSKESILPKRNSDKLIARKKDWDPKKGDFDSTVAYSVLVVAKEVEKSKKLSVKELLGITIMERSSEFEKN
PIDFLEAKGYKEVKKDLIKLPHYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGSPEDNE
QKQLFVEQHKHYLDEIEQISEFSKRVLADANLKVLSAYNKHDKPIREQAENI IHLFTLTNLGAPAAFYKFD
TIDRKRYTSTKEVLDTLIHQISITGLYETRIDLSQLGGD (SEQ ID NO:752)

[0068] 示例性的Cas9切口酶(nCas9):DKKYSIGLAIGTNSVGWAVITDEYKVPKSKFKVLGNTDRHSI
KKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLEESFLVEEDKKHERHPIF
GNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQL
FEENPINASGVDAKAILSARLSKSRLENLIAQLPGEKKNLFGNLIASLGLTPNFKSNFDLAEDAQLQLSKDTY
DDDLNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAIL
RRQEDFYFPLKDNREKIEKILTRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFD
KNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECF
DSVEISGVEDRFNASLGTYHDLLKIKDKDFLDNEENEDILEDIVLTLTLFEDREMIERLKYAHLFDDKVMKQL
KRRRYTGWGRLSRKLINGIRDKQSGKITLDFLKSDFANRNFMLIHDDSLTFKEDIQKAQVSGQGDSLHEIANL

AGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEDIKELGSQILKEHPVEN
TQLQNEKLYLYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLRSDKNRSGSDNVPSEEVVKMK
KNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKV
ITLKSCLVSDFRKDFQFYKVVREINNYHHAHDAYLNAVVGITALIKKYPKLESEFVYGDYKVDVRKMIKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSPQVNIIVKKTVEVQTGGFSK
ESILPKRNSDKLIARKKDWDPKKGFFSPTVAYSVLVAVKVEKKGSKKLKSVKELLGITIMERSSEFEKNPIDFLE
AKGYKEVKKDLIIKLPKYSLELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGKSPEDNEQKQLFV
EQHKHYLDEIIEQISEFSKRIVILADANLDKVL SAYNKHRDKPIREQAENI IHLFTLTNLGAPAAFKYFDTTIDRKR
YTSTKEVL DATLIHQ SITGLYETRIDLSQLGGD (SEQ ID NO:753)

[0069] 示例性的催化活性Cas9:DKKYSIGLDIGTNSVGWAVITDEYKVPKSKFKVLGNTDRHSIKKNI
GALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLEESFLVEEDKKHERHPIFGNIVD
EVAYHEKYPTIYHLRKKLVSTDKADRLIYLALAHMIKFRGHFLIEGDLNPDNSVDKLFIQLVQTYNQLFEENP
INASGVDAKAILSARLSKSRLENLIAQLPGEKKNLFGNLIASLGLTPNFKSNFDLAEDAKLQLSKD TYDDDL
NLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFF
DQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQED
FYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAMTRKSEETITPWNFEVVDKGASAQSFIERMTNFDKNLPN
EKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEI
SGVEDRFNASLGTYHDLKI IKDKDFLDNEENEDILEDIVLTLTLFEDREMIERLKYAHLFDDKVMKQLKRRRY
TGWGRLSRKLINGIRDKQSGKITLDFLKSDFANRNFMLIHDDSLTFKEDIQKAQVSGQDLSHEHIANLAGSPA
IKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEDIKELGSQILKEHPVENTQLQN
EKLYLYLYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLRSDKNRSGSDNVPSEEVVKMKKNYWR
QLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKS
KLVSDFRKDFQFYKVVREINNYHHAHDAYLNAVVGITALIKKYPKLESEFVYGDYKVDVRKMIKSEQEIGKATAKY
FFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSPQVNIIVKKTVEVQTGGFSKESILP
KRNSDKLIARKKDWDPKKGFFSPTVAYSVLVAVKVEKKGSKKLKSVKELLGITIMERSSEFEKNPIDFLEAKGYK
EVKDLIIKLPKYSLELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGKSPEDNEQKQLFVEQHKH
YLDEIIEQISEFSKRIVILADANLDKVL SAYNKHRDKPIREQAENI IHLFTLTNLGAPAAFKYFDTTIDRKRYTSTK
EVL DATLIHQ SITGLYETRIDLSQLGGD (SEQ ID NO:754)

[0070] 在一些实施方案中,Cas9是指来自构成单细胞原核微生物的域和界的古生菌
(arehaea) (例如纳古生菌 (nanoarchaea)) 的Cas9。在一些实施方案中,Cas9是指CasX或
CasY,其已经描述于例如Burststein et al., “New CRISPR-Cas systems from
uncultivated microbes.” Cell Res.2017Feb 21.doi:10.1038/cr.2017.21,其全部内容
在此通过引用并入。使用基因组分辨的宏基因组学,鉴定了许多CRISPR-Cas系统,包括在生
命的古生菌域中首次报告的Cas9。这种趋异的Cas9蛋白在研究很少的纳古生菌中作为活性
CRISPR-Cas系统的一部分发现。在细菌中,发现了两个以前未知的系统,CRISPR-CasX和
CRISPR-CasY,它们是迄今发现的最紧凑的系统之一。在一些实施方案中,Cas9是指CasX或
CasX的变体。在一些实施方案中,Cas9是指CasY或CasY的变体。应当理解,其他RNA引导的
DNA结合蛋白可以用作引导核苷酸序列-可编程DNA结合蛋白,并且在本公开的范围內。

[0071] 在一些实施方案中,本文提供的任何融合蛋白的引导核苷酸序列-可编程DNA结合

蛋白域可以是CasX或CasY蛋白。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域是CasX蛋白。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域是CasY蛋白。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域包含与天然存在的CasX或CasY蛋白至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域是天然存在的CasX或CasY蛋白。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域包含与本文所述的示例性CasX或CasY蛋白的任一个至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域包含本文所述的示例性CasX或CasY蛋白的任一个的氨基酸序列。应当理解,根据本公开也可以使用来自其他细菌物种的CasX和CasY。

[0072] CasX(uniprot.org/uniprot/F0NN87;uniprot.org/uniprot/F0NH53)

[0073] >tr|F0NN87|F0NN87_SULIH CRISPR相关的Casx蛋白OS=冰岛硫化叶菌(*Sulfolobus islandicus*) (菌株HVE10/4) GN=SiH_0402PE=4SV=1

[0074] MEVPLYNIFGDNYIIQVATEAENSTIYNNKVEIDDEELRNVLNLAYKIAKNEDAAAERRGKAKKKKG
EEGETTTSNIILPLSGNDKNPWETLKCYNFPTTVALSEVFKNFSQVKECEEVSAPSFVKPEFYEFGRSPGMVERT
RRVKLEVEPHYLIIAAAGWVLTGLGKAKVSEG DYVGVNFTPTRGILYSLIQNVNGIVPGIKPETAFLGLWIARKVV
SSVTNPNVSVVRIYITISDAVGQNPPTINGGFSIDLTKLLEKRYLLSERLEAIARNALSISSNMRERYIVLANIYE
YLTGSKRLEDLLYFANRDLIMNLSDDGKVRDLKLISAYVNGELIRGEG (SEQ ID NO:755)

[0075] >tr|F0NH53|F0NH53_SULIR CRISPR相关的蛋白质,Casx OS=冰岛硫化叶菌(菌株REY15A) GN=SiRe_0771PE=4SV=1

[0076] MEVPLYNIFGDNYIIQVATEAENSTIYNNKVEIDDEELRNVLNLAYKIAKNEDAAAERRGKAKKKKG
EEGETTTSNIILPLSGNDKNPWETLKCYNFPTTVALSEVFKNFSQVKECEEVSAPSFVKPEFYKFGGRSPGMVERT
RRVKLEVEPHYLIIAAAGWVLTGLGKAKVSEG DYVGVNFTPTRGILYSLIQNVNGIVPGIKPETAFLGLWIARKVV
SSVTNPNVSVVSIYITISDAVGQNPPTINGGFSIDLTKLLEKRDLLSERLEAIARNALSISSNMRERYIVLANIYE
YLTGSKRLEDLLYFANRDLIMNLSDDGKVRDLKLISAYVNGELIRGEG (SEQ ID NO:756)

[0077] CasY(ncbi.nlm.nih.gov/protein/APG80656.1)

[0078] >APG80656.1CRISPR相关的蛋白质CasY[未培养的Parcubacteria组细菌]MSKRHP
RISGVKGYRLHAQRLEYTGKSGAMRTIKYPLYSSPSGGRTVPREIVSAINDDYVGLYGLSNFDDLNAEKRNEEKV
YSVLDWFYDCVQYGAVFSYTAPGLLNVAEVRGGSYELTKTLKGSPLYDELQIDKVIKFLNKKEISRANGSLDKLK
KDIIDCFKAEYRERHKDQCNKLADDIKNAKGDAGASLGERQKKLFRDFFGISEQSENDKPSFTNPLNLTCCLLPFD
TVNNNRNRGEVLFNKLKEYAQKLDKNEGSLEMWEYIGIGNSGTAFSNFLGEGFLGRLRENKITEKAMMDITDAW
RGQEQEEELKRLRILAALTIKLREP KFDNHGGYRSDINGKLSWLQNYINQTVKIKEDLKGHKDLKAKEMIN
RFGESDTKEEAVVSSLLESIEKIVPDDSDADDEKPDIPAIAIYRRFLSDGRLTLNRFVQREDVQEALIKERLEAEKK
KKPKKRKKKSDAEDEKETIDFKELFPHLAKPLKLVNPFYGD SKRELYKKYKNAAIYTDALWKAVEKIYKSAFSSSL
KNSFFDITDFDKDFFIKRLQKIFSVYRRFNTDKWKPIVKN SFAPYCDIVSLAENEVLYKPKQRSRKSAAIDKNRVR
LPSTENIAKAGIALARELSVAGFDWKDLLKKEEHEEYIDLIELHKTALALLAVTETQLDISALDFVENGTVKDFM
KTRDGNLVLEGRFLEMFSSQIVFSELRGLAGLSRKEFITRSAIQTMNGKQAEELLYIPHEFQSAKITTPKEMSRAF
LDLAPAEFATSLEPESEKSLKQMRYYPHYFGYELTRTGQIDGGVAENALRLEKSPVKKREIKCKQYKTLG

RGQNKIVLYVRSSYYQTQFLEWFLHRPKNVQTDVAVSGSFLIDEKKVKTRWNYDALTVALEPVS GSERV FVSQPFT
IFPEKSAEEEGQRYLGIDIGEYGIAYTALEITGDSAKILDQNFISDPQLKTLREEVKGLKLDQRRGTFAMPSTKIA
RIRESLVHSLRNRIHHLALKHKAKIVYELEVS RFEEGKQKIKKVYATLKKADVSEIDADKNLQTTVWGKLAVASE
ISASYTSQFCGACKKLWRAEMQVDETITTQELIGTVRVIKGGTLIDAIKDFMRPPIFDENDTPFPKYRDFCDKHHI
SKKMRGNSCLFICPFRCRANADADIQASQTIALLRYVKEEKKVEDYFERFRKLKNIKVLGQMKKI (SEQ ID NO:
757)

[0079] 术语“缀合”是指两个实体的缔合,例如两个分子,诸如两个蛋白质、两个域(例如,结合域和切割域),或蛋白质和试剂,例如蛋白质结合域和小分子的缔合。在一些方面,缔合在蛋白质(例如,RNA-可编程核酸酶)和核酸(例如,引导RNA)之间。缔合可以是例如经由直接或间接的(例如,经由接头)共价连接。在一些实施方案中,缔合是共价的。在一些实施方案中,两个分子经由连接两个分子的接头缀合。例如,在两个蛋白质彼此缀合(例如,经工程化的核酸酶的结合域和切割域)以形成蛋白质融合物的一些实施方案中,这两个蛋白质可以经由多肽接头(例如,将一个蛋白质的C端连接到另一个蛋白质的N端的氨基酸序列)缀合。

[0080] 如本文在核酸序列的背景下使用的术语“共有序列”是指表示在多个相似序列中的每个位置处发现的最频繁的核苷酸残基的经计算的序列。通常,通过序列比对确定共有序列,其中相似的序列彼此比较并计算相似的序列基序。在重组酶靶位点序列的背景下,在一些实施方案中,重组酶靶位点的共有序列可以通过给定的重组酶最频繁地结合或以最高亲和力结合的序列。

[0081] 如本文所用的术语“工程化的”是指已人为设计、生产、制备、合成和/或制造的蛋白质分子、核酸、复合物、物质或实体。因此,工程化的产品是天然不存在的产品。

[0082] 如本文所用,术语“有效量”是指足以引起期望的生物学反应的生物活性剂的量。在一些实施方案中,重组酶的有效量可以指足以诱导在由重组酶特异性结合并重组的靶位点处的重组的重组酶的量。如熟练技术人员将理解的,试剂,例如核酸酶、重组酶、杂合蛋白、融合蛋白、蛋白质二聚体、蛋白质(或蛋白质二聚体)和多核苷酸的复合物,或多核苷酸的有效量可以随各种因素而变化,诸如例如随期望的生物学反应、特定等位基因、基因组、靶位点、靶定的细胞或组织和使用的试剂而变化。

[0083] 如本文所用,“引导核苷酸序列-可编程DNA结合蛋白”是指能够结合DNA的蛋白质、多肽或域,并且与其靶DNA序列的结合由引导核苷酸序列介导。“引导核苷酸”可以是RNA或DNA分子(例如,单链DNA或ssDNA分子),其与靶序列互补并且可以将DNA结合蛋白引导至靶序列。因此,引导核苷酸序列-可编程DNA结合蛋白可以是RNA-可编程DNA结合蛋白,或ssDNA-可编程DNA结合蛋白。“可编程”意指DNA结合蛋白可以被编程以结合引导核苷酸靶标的任何DNA序列。本文提及的引导核苷酸序列-可编程DNA结合蛋白可以是本领域已知的任何引导核苷酸序列-可编程DNA结合蛋白而没有限制,其包括但不限于,二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白。术语“环状置换”是指其中蛋白质中氨基酸的顺序已经改变,导致具有改变的连接性(connectivity)但相似的(整体)三维形状的蛋白质结构的蛋白质。当原始的n和c末端氨基酸经由肽键连接时形成环状置换;然后肽序列在肽序列内的另一个位置中断裂,产生新的n和c端。环状置换可以通过许多过程发生,包括演化事件、翻译后修饰或人工工程化突变。例如,环状置换可以用于改善蛋白质的催化活性或热稳

定性。环状置换的引导核苷酸序列-可编程DNA结合蛋白可与本文所述的任何实施方案一起使用。术语“二分叉”通常是指分成两部分的单体蛋白质。通常,两个部分都是单体蛋白质的功能所需要的。二分叉的蛋白质可以或不可以自身二聚化以重建功能性蛋白质。二分叉可以通过许多过程发生,包括演化事件、翻译后修饰或人工工程化突变。当与二分叉域融合时,其他蛋白质域可以用于强制二分叉蛋白质的重新组装。在一些情况下,其相互作用依赖于小分子的蛋白质域可以与每个二分叉域融合,导致二分叉蛋白质的小分子调节的二聚化。

[0084] 如本文所用,术语“同源的”是本领域理解的术语,其是指在核苷酸和/或氨基酸序列的水平上高度相关的核酸或多肽。彼此同源的核酸或多肽称为“同源物”。两个序列之间的同源性可以通过本领域技术人员已知的序列比对方法确定。根据本发明,若两个序列对于至少20、至少30、至少40、至少50、至少60、至少70、至少80、至少90、至少100、至少120、至少150或至少200个氨基酸的至少一段(stretch),至少约50-60%相同,例如在一个或另一个序列中包含的所有残基的至少约50-60%中共享相同的残基(例如氨基酸残基)、至少约70%相同、至少约80%相同、至少约85%相同、至少约90%相同、至少约95%相同、至少约98%相同、至少约99%相同、至少约99.5%相同或至少约99.9%相同,则认为它们是同源的。

[0085] 如本文所用,术语“序列一致性”或“序列一致性百分比”可以分别指给定DNA或蛋白质中与参照序列相同的核酸或氨基酸残基的百分比。参见例如:Christopher M.Holman, Protein Similarity Score:A Simplified Version of the BLAST Score as a Superior Alternative to Percent Identity for Claiming Genuses of Related Protein Sequences,21SANTA CLARA COMPUTER&HIGH TECH.L.J.55,60(2004),以其全部内容通过引用并入本文。

[0086] 如本文所用,术语“接头”是指连接两个分子或部分,例如融合蛋白的两个域,诸如例如核酸酶无活性的Cas9域和核酸编辑域(例如腺苷脱氨酶)的键(例如共价键)、化学基团或分子。在一些实施方案中,接头连接RNA-可编程核酸酶的gRNA结合域,包括Cas9核酸酶域和核酸编辑蛋白的催化域。在一些实施方案中,接头连接dCas9和核酸编辑蛋白。通常,接头位于两个基团、分子或其他部分之间或侧翼有两个基团、分子或其他部分,并且经由共价键与每一个连接,从而连接两者。在一些实施方案中,接头是一个氨基酸或多个氨基酸(例如肽或蛋白质)。在一些实施方案中,接头是有机分子、基团、聚合物或化学部分。在一些实施方案中,接头的长度为5-100个氨基酸,例如长度为5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、23、24、25、26、27、28、29、30、30-35、35-40、40-45、45-50、50-60、60-70、70-80、80-90、90-100、100-150或150-200个氨基酸。也考虑了更长或更短的接头。在一些实施方案中,接头包含氨基酸序列SGSETPGTSESATPES(SEQ ID NO:7),其也可以称为XTEN接头。在一些实施方案中,接头包含氨基酸序列SGGS(SEQ ID NO:758)。在一些实施方案中,接头包含(SGGS)_n(SEQ ID NO:758)、(GGGS)_n(SEQ ID NO:759)、(GGGGS)_n(SEQ ID NO:722)、(G)_n、(EAAAK)_n(SEQ ID NO:723)、(GGS)_n或(XP)_n基序,或这些中任何的组合,其中n独立地是1和30之间的整数,并且其中X是任何氨基酸。在一些实施方案中,n是1、2、3、4、5、6、7、8、9、10、11、12、13、14或15。

[0087] 如本文所用,术语“突变”是指序列(例如核酸或氨基酸序列)内的残基用另一个残

基取代或序列内一个或多个残基的缺失或插入。本文通常通过鉴定初始残基,随后是序列内残基的位置和新取代的残基的身份来描述突变。用于产生本文提供的氨基酸取代(突变)的各种方法在本领域中是熟知的,并且由例如Green and Sambrook, *Molecular Cloning: A Laboratory Manual* (4th ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (2012)) 提供。

[0088] 术语“核定位序列”或“NLS”是指促进蛋白质输入细胞核(例如通过核转运)的氨基酸序列。核定位序列是本领域已知的,并且对于熟练技术人员是显而易见的。例如,NLS序列描述于2001年11月23日提交的Plank等人的国际PCT申请PCT/EP2000/011690,2001年5月31日公布为WO/2001/038547,其内容通过引用并入本文,用于其对示例性的核定位序列的公开内容。在一些实施方案中,NLS包含氨基酸序列PKKKRKV (SEQ ID NO:702) 或MDSLMLNRRKFLYQFKNVRWAKGRRETYLC (SEQ ID NO:761)。

[0089] 如本文所用,术语“核酸酶”是指能够切割连接核酸分子中两个核苷酸残基的磷酸二酯键的试剂,例如蛋白质。在一些实施方案中,“核酸酶”是指具有无活性的DNA切割域,使得核酸酶不能切割磷酸二酯键的蛋白质。在一些实施方案中,核酸酶是蛋白质,例如,可以结合核酸分子并切割连接核酸分子内的核苷酸残基的磷酸二酯键的酶。核酸酶可以是切割多核苷酸链内的磷酸二酯键的内切核酸酶,或切割在多核苷酸链的末端处的磷酸二酯键的外切核酸酶。在一些实施方案中,核酸酶是结合和/或切割特定核苷酸序列(其在本文中也称为“识别序列”、“核酸酶靶位点”或“靶位点”)内的特定磷酸二酯键的位点特异性核酸酶。在一些实施方案中,核酸酶是RNA引导的(即,RNA-可编程)核酸酶,其与具有与靶位点互补的序列的RNA(例如,引导RNA,“gRNA”)缔合(例如,结合),从而提供核酸酶的序列特异性。在一些实施方案中,核酸酶识别单链靶位点,而在其他实施方案中,核酸酶识别双链靶位点,例如双链DNA靶位点。许多天然存在的核酸酶(例如许多天然存在的DNA限制性核酸酶)的靶位点是本领域技术人员熟知的。核酸酶蛋白质通常包含介导蛋白质与核酸底物相互作用并且在一些情况下,还特异性结合靶位点的“结合域”,以及催化核酸骨架内的磷酸二酯键的切割的“切割域”。在一些实施方案中,核酸酶蛋白质可以以单体形式结合和切割核酸分子,而在其他实施方案中,核酸酶蛋白质必须二聚化或多聚化以切割靶核酸分子。天然存在的核酸酶的结合域和切割域,以及可以融合以产生结合特定靶位点的核酸酶的结合域和切割域是本领域技术人员熟知的。例如,引导核苷酸序列-可编程DNA结合蛋白(诸如RNA-可编程核酸酶(例如Cas9))或具有无活性DNA切割域的Cas9蛋白的结合域可以用作结合域(例如,结合gRNA以直接结合靶位点)以特异性结合所期望的靶位点,并与切割域融合或缀合。

[0090] 如本文所用,术语“核酸”和“核酸分子”是指包含核碱基和酸性部分(例如核苷、核苷酸或核苷酸的聚合物)的化合物。通常,聚合核酸,例如包含三个或更多个核苷酸的核酸分子是线性分子,其中相邻的核苷酸经由磷酸二酯连接彼此连接。在一些实施方案中,“核酸”是指个别的核酸残基(例如核苷酸和/或核苷)。在一些实施方案中,“核酸”是指包含三个或更多个个别核苷酸残基的寡核苷酸链。如本文所用,术语“寡核苷酸”和“多核苷酸”可以可互换地使用以指核苷酸的聚合物(例如,至少三个核苷酸的串)。在一些实施方案中,“核酸”涵盖RNA以及单链和/或双链DNA。核酸可以是天然存在的,例如在基因组、转录物、mRNA、tRNA、rRNA、siRNA、snRNA、质粒、粘粒、染色体、染色单体或其他天然存在的核酸分子的背景下。另一方面,核酸分子可以是非天然存在的分子,例如重组DNA或RNA、人工染色体、

工程化的基因组或其片段,或合成的DNA、RNA、DNA/RNA杂交体,或包括非天然存在的核苷酸或核苷。此外,术语“核酸”、“DNA”、“RNA”和/或相似术语包括核酸类似物,即具有除磷酸二酯主链之外的类似物。核酸可以从天然来源纯化,使用重组表达系统产生并任选地纯化,化学合成等。在适当的情况下,例如在化学合成分子的情况下,核酸可以包含核苷类似物,例如具有化学修饰的碱基或糖、和主链修饰的类似物。除非另有说明,核酸序列以5'至3'方向呈现。在一些实施方案中,核酸是或包含天然核苷(例如腺苷、胸苷、鸟苷、胞苷、尿苷、脱氧腺苷、脱氧胸苷、脱氧鸟苷和脱氧胞苷);核苷类似物(例如2-氨基腺苷、2-硫代胸苷、肌苷、吡咯并嘧啶、3-甲基腺苷、5-甲基胞苷、2-氨基腺苷、C5-溴尿苷、C5-氟尿苷、C5-碘尿苷、C5-丙炔基-尿苷、C5-丙炔基-胞苷、C5-甲基胞苷、2-氨基腺苷、7-脱氮腺苷、7-脱氮鸟苷、8-氧代腺苷、8-氧代鸟苷、0(6)-甲基鸟嘌呤和2-硫代胞苷);化学修饰的碱基;生物修饰的碱基(例如甲基化碱基);插入的碱基;修饰的糖(例如2'-氟核糖、核糖、2'-脱氧核糖、阿拉伯糖和己糖);和/或修饰的磷酸基团(例如硫代磷酸酯和5'-N-亚磷酰胺连接)。

[0091] 术语“正交”是指最低限度相互作用(若有的话)的生物组分。若gRNA引导的recCas9蛋白不与其他潜在的重组酶位点相互作用或最低限度地相互作用,则含有不同gRNA结合位点的重组酶靶位点是正交的。术语“正交性”指的是系统组分可以独立变化而不影响其他组分的效能的构思。复合物的gRNA引导的性质使得与recCas9蛋白复合的该组gRNA分子能够仅在gRNA引导的位点处引导重组酶活性。通过该组gRNA分子对靶定重组酶位点上的酶活性的完全或近完全的依赖性来证明该系统的正交性。

[0092] 如本文所用,术语“药物组合物”是指可以在治疗和/或预防疾病或病症的背景下施用于受试者的组合物。在一些实施方案中,药物组合物包含活性成分,例如与Cas9蛋白融合的重组酶,或其片段(或编码此类融合物的核酸)和任选的药学上可接受的赋形剂。在一些实施方案中,药物组合物包含发明的Cas9变体/融合(例如fCas9)蛋白和适于将Cas9变体/融合蛋白靶向到靶核酸的gRNA。在一些实施方案中,靶核酸是基因。在一些实施方案中,靶核酸是与疾病相关的等位基因,其中等位基因通过Cas9变体/融合蛋白的作用而被切割。在一些实施方案中,等位基因是CLTA基因、VEGF基因、PCDH15基因或FAM19A2基因的等位基因。参见例如实施例。

[0093] 如本文所用,术语“增殖性疾病”是指其中细胞或组织稳态受到干扰,使得细胞或细胞群表现出异常升高的增殖速率的任何疾病。增殖性疾病包括过度增殖性疾病,如新生前期增生性状况和新生性疾病。新生性疾病的特征是细胞的异常增殖,并包括良性和恶性新生物两者。恶性新生物也称为癌症。在一些实施方案中,本文提供的组合物和方法可用于治疗增殖性疾病。例如,在一些实施方案中,药物组合物包含Cas9(例如,fCas9)蛋白和适合于将Cas9蛋白靶向到VEGF等位基因的gRNA,其中等位基因通过Cas9蛋白的作用而失活。参见例如实施例。

[0094] 术语“蛋白质”、“肽”和“多肽”在本文中可互换使用,并且是指通过肽(酰胺)键连接在一起的氨基酸残基的聚合物。该术语是指具有任何大小、结构或功能的蛋白质、肽或多肽。通常,蛋白质、肽或多肽将是至少三个氨基酸长。蛋白质、肽或多肽可以指个别的蛋白质或蛋白质的集合。蛋白质、肽或多肽中的一个或多个氨基酸可以被修饰,例如通过添加化学实体如碳水化合物基团、羟基、磷酸基团、法呢基、异法呢基、脂肪酸基团,用于缀合、官能化或其他修饰的接头等。蛋白质、肽或多肽也可以是单个分子或者可以是多分子复合物。蛋白

质、肽或多肽可以仅仅是天然存在的蛋白质或肽的片段。蛋白质、肽或多肽可以是天然存在的、重组的或合成的,或其任何组合。如本文所用,术语“融合蛋白”是指包含来自至少两种不同蛋白质的蛋白质域的杂合多肽。一种蛋白质可以位于融合蛋白的氨基端(N端)部分或羧基端(C端)蛋白质,从而分别形成“氨基端融合蛋白”或“羧基端融合蛋白”。本文提供的任何蛋白质可以通过本领域已知的任何方法产生。例如,本文提供的蛋白质可以经由重组蛋白质表达和纯化产生,其特别适用于包含肽接头的融合蛋白。用于重组蛋白质表达和纯化的方法是熟知的,并且包括Green and Sambrook, *Molecular Cloning: A Laboratory Manual* (4th ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (2012)) 描述的那些,其全部内容通过引用并入本文。本文提及的特定融合蛋白是recCas9,一种通过将催化无活性的dCas9融合至重组酶的催化域创建的RNA编程的小丝氨酸重组酶,其能够在哺乳动物细胞中发挥功能。

[0095] 如本文所讨论的“假gix”位点或“gix假位点”是类似于Gix重组酶的天然DNA识别序列的特定的假回文核心DNA序列。参见例如N.D.F.Grindley, K.L.Whiteson, P.A.Rice, *Mechanisms of site-specific recombination. Annu Rev Biochem* 75, 567-605 (2006), 以其全部内容通过引用并入本文。类似地,“假hix”或“hix假位点”;“假six”或“six假位点”;“假resH”或“resH假位点”;“假res”或“res假位点”;“假LoxP”或“LoxP假位点”;“假att”或“att假位点”;“假FTR”或“FTR假位点”是类似于Hin重组酶、 β 重组酶、Sin重组酶、Tn3或 γ δ 重组酶、Cre重组酶、 λ 噬菌体整合酶或FLP重组酶的天然DNA识别序列的特定的假回文核心DNA序列。

[0096] 术语“RNA-可编程核酸酶”和“RNA引导的核酸酶”在本文中可互换使用,并且是指与一个或多个不是切割靶标的RNA形成复合物(例如,结合或缔合)的核酸酶。在一些实施方案中,当与RNA形成复合物时,RNA-可编程核酸酶可以称为核酸酶:RNA复合物。通常,结合的RNA称为引导RNA(gRNA)。gRNA可以作为两个或更多个RNA的复合物或者作为单个RNA分子存在。作为单个RNA分子存在的gRNA可以称为单引导RNA(sgRNA),尽管“gRNA”可互换使用以指作为单个分子或作为两个或更多个分子的复合物存在的引导RNA。通常,作为单一RNA种类存在的gRNA包含两个域:(1)与靶核酸共享同源性的域(例如,并引导Cas9复合物与靶物的结合)的域;和(2)结合Cas9蛋白的域。在一些实施方案中,域(2)对应已知为tracrRNA的序列,并且包含茎-环结构。例如,在一些实施方案中,域(2)与Jinek et al., *Science* 337:816-821 (2012)的图1E中描绘的tracrRNA同源,其全部内容通过引用并入本文。gRNA的其他实例(例如包括域2的那些)可以在2013年9月6日提交的题为“Switchable Cas9Nucleases and Uses Thereof”的美国临时专利申请U.S.S.N.61/874,682;2013年9月6号提交的题为“Delivery System For Functional Nucleases”的美国临时专利申请U.S.S.N.61/874,746;2013年3月15日提交的题为“Methods and Compositions for RNA-Directed Target DNA Modification and for RNA-Directed Modulation of Transcription”的PCT申请WO 2013/176722;和2013年3月20日提交的题为“RNA-Directed DNA Cleavage by the Cas9-crRNA Complex”的PCT申请WO 2013/142578中找到,每篇的全部内容通过引用以其整体并入本文。本文提供了gRNA的其他实例。参见例如实施例。在一些实施方案中,gRNA包含域(1)和(2)中的两个或更多个,并且可以称为“延伸的gRNA”。例如,延伸的gRNA将例如结合两个或更多个Cas9蛋白并在两个或更多个不同区域处结合靶核酸,如本文所述。gRNA包含与靶

位点互补的核苷酸序列,其介导核酸酶/RNA复合物与所述靶位点的结合,提供了核酸酶:RNA复合物的序列特异性。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是RNA-可编程核酸酶如(CRISPR相关系统)(CRISPR-associated system)Cas9内切核酸酶,例如来自酿脓链球菌的Cas9(Csn1)(参见例如“Complete genome sequence of an M1 strain of *Streptococcus pyogenes*.”Ferretti J.J.,McShan W.M.,Ajdic D.J.,Savic D.J.,Savic G.,Lyon K.,Primeaux C.,Sezate S.,Suvorov A.N.,Kenton S.,Lai H.S.,Lin S.P.,Qian Y.,Jia H.G.,Najar F.Z.,Ren Q.,Zhu H.,Song L.,White J.,Yuan X.,Clifton S.W.,Roe B.A.,McLaughlin R.E.,*Proc.Natl.Acad.Sci.U.S.A.*98:4658-4663(2001);“CRISPR RNA maturation by trans-encoded small RNA and host factor RNase III.”Deltcheva E.,Chylinski K.,Sharma C.M.,Gonzales K.,Chao Y.,Pirzada Z.A.,Eckert M.R.,Vogel J.,Charpentier E.,*Nature* 471:602-607(2011);和“A programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity.”Jinek M.,Chylinski K.,Fonfara I.,Hauer M.,Doudna J.A.,Charpentier E.*Science* 337:816-821(2012),每篇的全部内容通过引用并入本文。

[0097] 因为RNA-可编程核酸酶(例如Cas9)使用RNA:DNA杂交来确定靶DNA切割位点,所以这些蛋白质原则上能够切割由引导RNA规定的任何序列。使用RNA-可编程核酸酶例如Cas9进行位点特异性切割(例如,以修饰基因组)的方法是本领域已知的(参见例如Cong,L.et al.Multiplex genome engineering using CRISPR/Cas systems.*Science* 339,819-823(2013);Mali,P.et al.RNA-guided human genome engineering via Cas9.*Science* 339,823-826(2013);Hwang,W.Y.et al.Efficient genome editing in zebrafish using a CRISPR-Cas system.*Nature biotechnology* 31,227-229(2013);Jinek,M.et al.RNA-programmed genome editing in human cells.*eLife* 2,e00471(2013);Dicarlo,J.E.et al.Genome engineering in *Saccharomyces cerevisiae* using CRISPR-Cas systems.*Nucleic acids research*(2013);Jiang,W.et al.RNA-guided editing of bacterial genomes using CRISPR-Cas systems.*Nature biotechnology* 31,233-239(2013);每篇的全部内容通过引用并入本文)。

[0098] 如本文所用,术语“重组酶”是指介导重组酶识别序列之间DNA的重组的位点特异性酶,所述重组导致重组酶识别序列之间的DNA片段的切除、整合、倒位或交换(例如,易位)。重组酶可以分为两个不同的家族:丝氨酸重组酶(例如,解离酶和转化酶)和酪氨酸重组酶(例如整合酶)。丝氨酸重组酶的实例包括但不限于,Hin、Gin、Tn3、 β -six、CinH、ParA、 γ δ 、Bxb1、 ϕ C31、TP901、TG1、 ϕ BT1、R4、 ϕ RV1、 ϕ FC1、MR11、A118、U153和gp29。酪氨酸重组酶的实例包括但不限于,Cre、FLP、R、Lambda、HK101、HK022和pSAM2。本文提及的Gin重组酶可以是本领域已知的任何Gin重组酶,其包括但不限于T.Gaj et al.,A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells.*Nucleic Acids Research* 41,3937-3946(2013)中提出的Gin重组酶,其通过引用以其整体并入本文。在某些实施方案中,Gin重组酶催化域与SEQ ID NO:713中所示的氨基酸序列具有大于85%、90%、95%、98%或99%的序列一致性。在另一个实施方案中,Gin重组酶催化域的氨基酸序列包含对应于H106Y,和/或I127L,和/或I136R和/或G137F的突变。在另一个实施方案中,Gin重组酶催化域的氨基酸序列包含对

应于H106Y、I127L、I136R和G137F的突变。在进一步的实施方案中,Gin重组酶的氨基酸序列已得以进一步突变。在具体的实施方案中,Gin重组酶催化域的氨基酸序列包含SEQ ID NO: 713。Gin重组酶与gix靶位点(本文也称为“gix核心”、“最小gix核心”或“gix相关核心”序列)结合。最小gix核心重组酶位点是NNNNAASSWWSSTTTNNNN (SEQ ID NO:19),其中N定义为任何氨基酸,W是A或T,并且S是G或C。gix靶位点可以包括本领域已知的任何其他突变。在某些实施方案中,gix靶位点与SEQ ID NO:19中所示的氨基酸序列具有大于90%、95%或99%的序列一致性。gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离可以是1至10个碱基对、3至7个碱基对、5至7个碱基对或5至6个碱基对。gix核心或gix相关核心序列与至少一个gRNA结合位点之间的距离可以是1、2、3、4、5、6、7、8、9或10个碱基对。

[0099] 丝氨酸和酪氨酸重组酶的名称源自重组酶用来攻击DNA,并在链交换期间与DNA共价连接的保守亲核氨基酸残基。重组酶具有许多应用,包括基因敲除/敲入的创建和基因治疗应用。参见例如Brown et al.,“Serine recombinases as tools for genome engineering.”*Methods*.2011;53(4):372-9;Hirano et al.,“Site-specific recombinases as tools for heterologous gene integration.”*Appl.Microbiol.Biotechnol*.2011;92(2):227-39;Chavez and Calos,“Therapeutic applications of the Φ C31 integrase system.”*Curr.Gene Ther*.2011;11(5):375-81;Turan and Bode,“Site-specific recombinases:from tag-and-target-to tag-and-exchange-based genomic modifications.”*FASEB J*.2011;25(12):4088-107;Venken and Bellen,“Genome-wide manipulations of *Drosophila melanogaster* with transposons,Flp recombinase,and Φ C31 integrase.”*Methods Mol.Biol*.2012;859:203-28;Murphy,“Phage recombinases and their applications.”*Adv.Virus Res*.2012;83:367-414;Zhang et al.,“Conditional gene manipulation:Creating a new biological era.”*J.Zhejiang Univ.Sci.B*.2012;13(7):511-24;Karpenshif and Bernstein,“From yeast to mammals:recent advances in genetic control of homologous recombination.”*DNA Repair (Amst)*.2012;1;11(10):781-8;每篇的全部内容以其整体在此通过引用并入。本文提供的重组酶不意味着是可以用于本发明的实施方案的重组酶的排他性实例。通过挖掘新的正交重组酶的数据库或设计具有限定的DNA特异性的合成重组酶,可以扩展本发明的方法和组合物(参见例如,Groth et al.,“Phage integrases:biology and applications.”*J.Mol.Biol*.2004;335,667-678;Gordley et al.,“Synthesis of programmable integrases.”*Proc.Natl.Acad.Sci.U S A*.2009;106,5053-5058;每篇的全部内容以其整体在此通过引用并入)。

[0100] 可用于本文所述方法和组合物的重组酶的其他实例是本领域技术人员已知的,并且预期发现或产生的任何新重组酶能够用于本发明的不同实施方案中。在一些实施方案中,重组酶的催化域与核酸酶失活的RNA-可编程核酸酶(例如,dCas9或其片段)融合,使得重组酶域不包含核酸结合域或不能结合随后导致酶促催化的靶核酸(例如,重组酶域经工程化改造以使其不具有特异性DNA结合活性)。缺乏DNA结合活性的部分的重组酶和独立于辅助蛋白起作用的重组酶以及其工程化方法是已知的,并且包括Klippel et al.,“Isolation and characterisation of unusual gin mutants.”*EMBO J*.1988;7:3983-3989;Burke et al.,“Activating mutations of Tn3 resolvase marking interfaces

important in recombination catalysis and its regulation. *Mol Microbiol.* 2004; 51:937-948; Olorunniji et al., "Synapsis and catalysis by activated Tn3 resolvase mutants." *Nucleic Acids Res.* 2008; 36:7181-7191; Rowland et al., "Regulatory mutations in Sin recombinase support a structure-based model of the synaptosome." *Mol Microbiol.* 2009; 74:282-298; Akopian et al., "Chimeric recombinases with designed DNA sequence recognition." *Proc Natl Acad Sci USA.* 2003; 100:8688-8691; Gordley et al., "Evolution of programmable zinc finger-recombinases with activity in human cells." *J Mol Biol.* 2007; 367:802-813; Gordley et al., "Synthesis of programmable integrases." *Proc Natl Acad Sci USA.* 2009; 106:5053-5058; Arnold et al., "Mutants of Tn3 resolvase which do not require accessory binding sites for recombination activity." *EMBO J.* 1999; 18:1407-1414; Gaj et al., "Structure-guided reprogramming of serine recombinase DNA sequence specificity." *Proc Natl Acad Sci USA.* 2011; 108(2):498-503; 和 Proudfoot et al., "Zinc finger recombinases with adaptable DNA sequence specificity." *PLoS One.* 2011; 6(4):e19537 描述的那些; 每篇的全部内容在此通过引用并入。例如, 解离酶-转化酶组的丝氨酸重组酶 (例如 Tn3 和 γ δ 解离酶以及 Hin 和 Gin 转化酶) 具有模块化结构, 它们具有部分自主的催化和 DNA 结合域 (参见例如, Grindley et al., "Mechanism of site-specific recombination." *Ann Rev Biochem.* 2006; 75:567-605, 其全部内容通过引用并入)。因此, 这些重组酶的催化域适合于与如本文所述的核酸酶失活的 RNA-可编程核酸酶 (例如, dCas9 或其片段) 重组, 例如, 在分离不需要任何辅助因素 (例如 DNA 结合活性) 的 "激活性" 重组酶突变体之后 (参见例如, Klippel et al., "Isolation and characterisation of unusual gin mutants." *EMBO J.* 1988; 7:3983-3989; Burke et al., "Activating mutations of Tn3 resolvase marking interfaces important in recombination catalysis and its regulation. *Mol Microbiol.* 2004; 51:937-948; Olorunniji et al., "Synapsis and catalysis by activated Tn3 resolvase mutants." *Nucleic Acids Res.* 2008; 36:7181-7191; Rowland et al., "Regulatory mutations in Sin recombinase support a structure-based model of the synaptosome." *Mol Microbiol.* 2009; 74:282-298; Akopian et al., "Chimeric recombinases with designed DNA sequence recognition." *Proc Natl Acad Sci USA.* 2003; 100:8688-8691)。

[0101] 另外, 许多其他具有 N 端催化域和 C 端 DNA 结合域的天然丝氨酸重组酶是已知的 (例如, phiC31 整合酶、TnpX 转座酶、IS607 转座酶), 并且它们的催化域可以增选用于工程化如本文所述的可编程位点特异性重组酶 (参见例如, Smith et al., "Diversity in the serine recombinases." *Mol Microbiol.* 2002; 44:299-307, 其全部内容通过引用并入)。类似地, 酪氨酸重组酶 (例如 Cre、 λ 整合酶) 的核心催化域是已知的, 并且可以类似地增选以工程化如本文所述的可编程位点特异性重组酶 (参见例如 Guo et al., "Structure of Cre recombinase complexed with DNA in a site-specific recombination synapse." *Nature.* 1997; 389:40-46; Hartung et al., "Cre mutants with altered DNA binding properties." *J Biol Chem* 1998; 273:22884-22891; Shaikh et al., "Chimeras of the Flp and Cre recombinases: Tests of the mode of cleavage by Flp and Cre." *J Mol*

Biol.2000;302:27-48;Rongrong et al.,“Effect of deletion mutation on the recombination activity of Cre recombinase.”Acta Biochim Pol.2005;52:541-544; Kilbride et al.,“Determinants of product topology in a hybrid Cre-Tn3resolvase site-specific recombination system.”J Mol Biol.2006;355:185-195; Warren et al.,“A chimeric cre recombinase with regulated directionality.”Proc Natl Acad Sci USA.2008 105:18278-18283;Van Duyne,“Teaching Cre to follow directions.”Proc Natl Acad Sci USA.2009Jan 6;106(1):4-5;Numrych et al.,“A comparison of the effects of single-base and triple-base changes in the integrase arm-type binding sites on the site-specific recombination of bacteriophage λ .”Nucleic Acids Res.1990;18:3953-3959;Tirumalai et al.,“The recognition of core-type DNA sites by λ integrase.”J Mol Biol.1998;279:513-527; Aihara et al.,“A conformational switch controls the DNA cleavage activity of λ integrase.”Mol Cell.2003;12:187-198;Biswas et al.,“A structural basis for allosteric control of DNA recombination by λ integrase.”Nature.2005;435:1059-1066;和Warren et al.,“Mutations in the amino-terminal domain of λ -integrase have differential effects on integrative and excisive recombination.”Mol Microbiol.2005;55:1104-1112;每篇的全部内容通过引用并入)。

[0102] 在核酸修饰(例如,基因组修饰)的背景下,术语“重组”用于指通过重组酶蛋白质(例如,本文提供的本发明的重组酶融合蛋白)的作用修饰两个或更多个核酸分子或单个核酸分子的两个或更多个区域的过程。重组尤其可以导致核酸的插入、倒位、切除或易位,例如在一个或多个核酸分子之中或之间。

[0103] 如本文中在蛋白质或核酸的背景中使用,术语“重组体(recombinant)”是指自然界中不存在,但是作为人工程化的产物的蛋白质或核酸。例如,在一些实施方案中,重组蛋白质或核酸分子包含相比于任何天然存在的序列包含至少一个、至少两个、至少三个、至少四个、至少五个、至少六个或至少七个突变的氨基酸或核苷酸序列。

[0104] 如本文所用,术语“受试者”是指个体生物体,例如个体哺乳动物。在一些实施方案中,受试者是人。在一些实施方案中,受试者是非人哺乳动物。在一些实施方案中,受试者是非人灵长类动物。在一些实施方案中,受试者是啮齿动物。在一些实施方案中,受试者是绵羊、山羊、牛、猫或狗。在一些实施方案中,受试者是脊椎动物、两栖动物、爬行动物、鱼、昆虫、苍蝇或线虫。在一些实施方案中,受试者是研究动物。在一些实施方案中,受试者是经遗传工程化的,例如基因遗传化的非人受试者。受试者可以是任何个性别和处于任何发展阶段的。在一些实施方案中,受试者是经遗传工程化的,例如基因遗传化的非人受试者。受试者可以是任何个性别和处于任何发展阶段的。

[0105] 如本文在核酸酶的背景中使用,术语“靶核酸”和“靶基因组”分别指包含给定核酸酶的至少一个靶位点的核酸分子或基因组。在包含(核酸酶失活的)RNA-可编程核酸酶和重组酶域的融合物的背景中,“靶核酸”和“靶基因组”分别指一个或多个包含至少一个靶位点的核酸分子或基因组。在一些实施方案中,靶核酸包含至少两个、至少三个、至少四个、至少五个、至少六个、至少七个或至少八个靶位点。在一些实施方案中,靶核酸包含四个靶位点。

[0106] 术语“靶位点”是指由重组酶(例如本文提供的dCas9-重组酶融合蛋白)结合并重

组(例如在靶位点处或附近)的核酸分子内的序列。靶位点可以是单链或双链的。例如,在一些实施方案中,四个重组酶单体协调以重组靶核酸,每个单体与由gRNA引导的(核酸酶失活的)Cas9蛋白融合。在此类实例中,每个Cas9域由不同的gRNA引导以结合靶核酸,因此靶核酸包含四个靶位点,每个位点通过分开的dCas9-重组酶融合物靶向(从而协调重组靶核酸的四个重组酶单体)。对于RNA引导的核酸酶失活的Cas9(或其gRNA结合域)和本发明的Cas9的融合物,在一些实施方案中,靶位点可以是17-20个碱基对加上3个碱基对PAM(例如,NNN,其中N独立地代表任何核苷酸)。通常,PAM的第一个核苷酸可以是任何核苷酸,而两个下游核苷酸是根据特定的RNA引导的核酸酶指定的。用于RNA引导的核酸酶(例如Cas9)的示例性靶位点(例如,包含PAM)是本领域技术人员已知的并且包括但不限于NNG、NGN、NAG和NGG,其中每个N独立地是任何核苷酸。另外,来自不同物种(例如嗜热链球菌而不是酿脓链球菌)的Cas9核酸酶识别包含序列NGGNG(SEQ ID NO:763)的PAM。另外的PAM序列是已知的,包括但不限于NNAGAAW(SEQ ID NO:749)和NAAR(SEQ ID NO:771)(参见例如,Esvelt and Wang, *Molecular Systems Biology*, 9:641 (2013),其全部内容通过引用并入本文)。在一些方面, RNA引导的核酸酶(诸如例如Cas9)的靶位点可以包含结构 $[N_z]$ -[PAM],其中每个N独立地是任何核苷酸,并且z是1和50之间的整数,包括端点。在一些实施方案中,z是至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14、至少15、至少16、至少17、至少18、至少19、至少20、至少25、至少30、至少35、至少40、至少45或至少50。在一些实施方案中,z是5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、23、24、25、26、27、28、29、30、31、32、33、34、35、36、37、38、39、40、41、42、43、44、45、46、47、48、49或50。在一些实施方案中,z是20。在某些实施方案中,可以使用“无PAM的”RNA引导的核酸酶(例如,无PAM的Cas9)或如本文进一步描述的具有放松PAM需求的RNA引导的核酸酶。在一些实施方案中,“靶位点”还可以指核酸分子内被核酸酶结合但未切割的序列。例如,本文描述的某些实施方案提供包含无活性(或失活)Cas9DNA切割域的蛋白质。此类蛋白质(例如,当还包括Cas9RNA结合域时)能够结合由gRNA指定的靶位点;然而,因为DNA切割位点被灭活,所以靶位点不被特定蛋白质切割。在一些实施方案中,此类蛋白质与重组酶(或重组酶的催化域)缀合、融合或结合,所述重组酶介导靶核酸的重组。在一些实施方案中,实际上切割或重组的序列将取决于介导核酸分子的切割或重组的蛋白质(例如,重组酶)或分子,并且在一些情况下,例如可与结合失活的Cas9蛋白的接近性或距离有关。

[0107] 如本文所用,术语“转录激活物样效应物”(TALE)是指包含DNA结合域的细菌蛋白质,所述DNA结合域含有包含高度可变的双氨基酸基序(重复可变双残基,RVD)的高度保守的33-34个氨基酸的序列。RVD基序决定了对核酸序列的结合特异性,并且可以根据本领域技术人员已知的方法进行工程化以特异性结合所期望的DNA序列(参见例如,Miller, Jeffrey;et.al. (February 2011). “A TALE nuclease architecture for efficient genome editing”. *Nature Biotechnology* 29 (2):143-8; Zhang, Feng; et.al. (February 2011). “Efficient construction of sequence-specific TAL effectors for modulating mammalian transcription” *Nature Biotechnology* 29 (2):149-53; Geißler, R.; Scholze, H.; Hahn, S.; Streubel, J.; Bonas, U.; Behrens, S.E.; Boch, J. (2011), Shiu, Shin-Han. ed. “Transcriptional Activators of Human Genes with Programmable DNA-Specificity”. *PLoS ONE* 6 (5):e19509; Boch, Jens (February 2011). “TALEs of genome

targeting”.*Nature Biotechnology* 29 (2) :135-6;Boch,Jens;et.al. (December 2009). “Breaking the Code of DNA Binding Specificity of TAL-Type III Effectors”.*Science* 326 (5959) :1509-12;和Moscou,Matthew J.;Adam J.Bogdanove (December 2009). “A Simple Cipher Governs DNA Recognition by TAL Effectors”.*Science* 326 (5959) :1501;其每篇的全部内容通过引用并入本文)。氨基酸序列和DNA识别之间的简单关系允许通过选择含有适当RVD的重复区段的组合来工程化改造特定的DNA结合域。

[0108] 如本文所用,术语“转录激活物样元件核酸酶”(TALEN)是指包含转录激活物样效应物DNA结合域至DNA切割域,例如FokI域的人工核酸酶。已经报告了许多用于产生工程化TALE构建体的模块化组装方案(参见例如,Zhang,Feng;et.al. (February 2011). “Efficient construction of sequence-specific TAL effectors for modulating mammalian transcription”.*Nature Biotechnology* 29 (2) :149-53;Geißler,R.;Scholze,H.;Hahn,S.;Streubel,J.;Bonas,U.;Behrens,S.E.;Boch,J. (2011),Shiu,Shin-Han.ed. “Transcriptional Activators of Human Genes with Programmable DNA-Specificity”.*PLoS ONE* 6 (5) :e19509;Cermak,T.;Doyle,E.L.;Christian,M.;Wang,L.;Zhang,Y.;Schmidt,C.;Baller,J.A.;Somia,N.V.et al. (2011). “Efficient design and assembly of custom TALEN and other TAL effector-based constructs for DNA targeting”.*Nucleic Acids Research*;Morbiter,R.;Elsaesser,J.;Hausner,J.;Lahaye,T. (2011). “Assembly of custom TALE-type DNA binding domains by modular cloning”.*Nucleic Acids Research*;Li,T.;Huang,S.;Zhao,X.;Wright,D.A.;Carpenter,S.;Spalding,M.H.;Weeks,D.P.;Yang,B. (2011). “Modularly assembled designer TAL effector nucleases for targeted gene knockout and gene replacement in eukaryotes”.*Nucleic Acids Research*;Weber,E.;Gruetzner,R.;Werner,S.;Engler,C.;Marillonnet,S. (2011). Bendahmane,Mohammed.ed. “Assembly of Designer TAL Effectors by Golden Gate Cloning”.*PLoS ONE* 6 (5) :e19722;其每篇的全部内容通过引用并入本文)。

[0109] 术语“治疗/处理”是指如本文所述旨在逆转、缓解疾病或病症或其一种或多种症状、延迟疾病或病症或其一种或多种症状的发作或抑制疾病或病症或其一种或多种症状进展的临床干预。如本文所用,术语“治疗/处理”是指如本文所述旨在逆转、缓解疾病或病症或其一种或多种症状、延迟疾病或病症或其一种或多种症状的发作或抑制疾病或病症或其一种或多种症状进展的临床干预。在一些实施方案中,可以在一种或多种症状已经得以形成之后和/或疾病已经得到诊断之后施用治疗。在其他实施方案中,可以在没有症状的情况下施用治疗,例如用于预防或延迟症状的发作或抑制疾病的发作或进展。例如,可以在症状发作之前(例如,鉴于症状的历史和/或鉴于遗传或其他易感性因素)施用治疗于易感个体。治疗也可以在症状消退后继续进行,例如以预防或延迟其复发。

[0110] 术语“载体”是指包含一种或多种本发明的重组多核苷酸的多核苷酸,例如编码本文提供的Cas9蛋白(或其融合物)和/或gRNA的那些。载体包括但不限于质粒、病毒载体、粘粒、人工染色体和噬菌粒。载体可以能够在宿主细胞中复制,并且可以进一步通过一个或多个内切核酸酶限制性位点来表征,在该位点处可以切割载体并且可以在其中插入所期望的核酸序列。载体可以含有一种或多种适用于鉴定和/或选择已经用或尚未用载体转化或用

载体进行基因组修饰的细胞的标志物序列。标志物包括例如编码增加或降低对抗生素(例如,卡那霉素、氨基青霉素)或其他化合物的抗性或敏感性的蛋白质的基因,编码其活性可通过本领域已知的标准测定法检测的酶(例如, β -半乳糖苷酶、碱性磷酸酶或萤光素酶)的基因,以及可见地影响转化或转染细胞、宿主、菌落或噬斑的表型的基因。适用于转化本发明所囊括的宿主细胞(例如大肠杆菌、哺乳动物细胞如CHO细胞、昆虫细胞等)的任何载体,例如属于pUC系列、pGEM系列、pET系列、pBAD系列、pTET系列或pGEX系列的载体。在一些实施方案中,载体适合于转化宿主细胞用于重组蛋白质生产。用于选择和工程化用于表达蛋白质(例如,本文提供的蛋白质)的载体和宿主细胞、转化细胞和表达/纯化重组蛋白质的方法是本领域熟知的,并且由例如Green and Sambrook, *Molecular Cloning: A Laboratory Manual* (4th ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (2012)) 提供。

[0111] 如本文所用,术语“锌指”是指以折叠和一个或多个稳定化折叠的锌离子的配位为特征的小核酸结合蛋白结构基序。锌指涵盖极其多种不同的蛋白质结构(参见例如Klug A, Rhodes D (1987). “Zinc fingers: a novel protein fold for nucleic acid recognition”. *Cold Spring Harb. Symp. Quant. Biol.* 52:473-82,其全部内容通过引用并入本文)。可以设计锌指以结合核苷酸的特定序列,并且可以设计包含一系列锌指的融合物的锌指阵列以结合实际上任何所期望的靶序列。此类锌指阵列可以形成蛋白质(例如核酸酶)的结合域,例如,在与核酸切割域缀合的情况下。不同类型的锌指基序是本领域技术人员已知的,包括但不限于Cys₂His₂、Gag knuckle、Treble clef、锌带(Zinc ribbon)、Zn₂/Cys₆和TAZ2域样基序(参见例如Krishna SS, Majumdar I, Grishin NV (January 2003). “Structural classification of zinc fingers: survey and summary”. *Nucleic Acids Res.* 31 (2) :532-50)。通常,单一锌指基序结合核酸分子的3或4个核苷酸。因此,包含2个锌指基序的锌指域可以结合6-8个核苷酸,包含3个锌指基序的锌指域可以结合9-12个核苷酸,包含4个锌指基序的锌指域可以结合12-16个核苷酸,等等。可以采用任何合适的蛋白质工程化技术来改变锌指的DNA结合特异性和/或设计新的锌指融合物以结合长度为3至30个核苷酸的实际任何所期望的靶序列(参见例如Pabo CO, Peisach E, Grant RA (2001). “Design and selection of novel cys₂His₂Zinc finger proteins”. *Annual Review of Biochemistry* 70:313-340; Jamieson AC, Miller JC, Pabo CO (2003). “Drug discovery with engineered zinc-finger proteins”. *Nature Reviews Drug Discovery* 2 (5) :361-368; 和Liu Q, Segal DJ, Ghiara JB, Barbas CF (May 1997). “Design of polydactyl zinc-finger proteins for unique addressing within complex genomes”. *Proc. Natl. Acad. Sci. U.S.A.* 94 (11) ;其每篇的全部内容通过引用并入本文)。工程化锌指阵列和切割核酸的蛋白域之间的融合可以用于产生“锌指核酸酶”。锌指核酸酶通常包含结合核酸分子内特定靶位点的锌指域和切割由结合域结合的靶位点内或附近的核酸分子的核酸切割域。典型的工程化锌指核酸酶包含具有3和6个之间的个别锌指基序的结合域和长度范围为9个碱基对至18个碱基对的结合靶位点。在期望结合和切割给定基因组中独特的靶位点的情况下,较长的靶位点特别有吸引力。

[0112] 如本文所用,术语“锌指核酸酶”是指包含与包含锌指阵列的结合域缀合的核酸切割域的核酸酶。在一些实施方案中,切割域是II型限制性内切核酸酶FokI的切割域。可以设

计锌指核酸酶以靶向给定核酸分子中的实际任何所期望的序列用于切割,并且设计锌指结合域以在复杂基因组的背景下结合独特位点的可能性允许活细胞中靶向切割单个基因组位点,例如,以实现治疗价值的靶向基因组改变。由于非同源DNA修复途径的易错特性,将双链断裂靶向到期望的基因组基因座可以用于将移码突变引入基因的编码序列中。可以通过本领域技术人员熟知的方法产生锌指核酸酶以靶向感兴趣的位点。例如,可以通过组合已知特异性的个别锌指基序来设计具有所期望的特异性的锌指结合域。与DNA结合的锌指蛋白Zif268的结构已经为该领域的大量工作提供了信息,并且已经描述了为64个可能的碱基对三联体中的每一个获得锌指,然后混合和匹配这些模块化锌指以设计具有任何所期望的序列特异性的蛋白质的构思(Pavletich NP, Pabo CO (May 1991). "Zinc finger-DNA recognition: crystal structure of a Zif268-DNA complex at 2.1Å". Science 252 (5007): 809-17, 其全部内容并入本文)。在一些实施方案中,将各自识别3碱基对DNA序列的单独的锌指组合以产生3-、4-、5-或6-指阵列,其识别长度范围为9个碱基对至18个碱基对的靶位点。在一些实施方案中,考虑了更长的阵列。在其他实施方案中,组合识别6-8个核苷酸的2指模块以产生4-、6-或8-指阵列。在一些实施方案中,采用细菌或噬菌体展示来开发识别所期望的核酸序列的锌指域,例如,长度为3-30bp的所期望的核酸酶靶位点。在一些实施方案中,锌指核酸酶包含经由接头(例如多肽接头)彼此融合或以其他方式彼此缀合的锌指结合域和切割域。接头的长度决定了切口与由锌指域结合的核酸序列的距离。若使用较短的接头,则切割域将使核酸切割得更接近结合的核酸序列,而较长的接头将导致切口和结合的核酸序列之间更大的距离。在一些实施方案中,锌指核酸酶的切割域必须二聚化以切割结合的核酸。在一些此类实施方案中,二聚体是两个单体的异源二聚体,每个单体包含不同的锌指结合域。例如,在一些实施方案中,二聚体可以包含一个包含与FokI切割域缀合的锌指域A的单体,和一个包含与FokI切割域缀合的锌指域B的单体。在该非限制性实例中,锌指域A结合靶位点的一侧上的核酸序列,锌指域B结合靶位点的另一侧上的核酸序列,并且二聚化FokI域切割锌指域结合位点之间的核酸。

[0113] 发明详述

[0114] 从以下实施例将更全面地理解本发明的这些和其他实施方案的功能和优点。以下实施例旨在说明本发明的益处并描述特定实施方案,但并不旨在举例说明本发明的全部范围。因此,应当理解,实施例不意味着限制本发明的范围。

[0115] 引导核苷酸序列-可编程DNA结合蛋白

[0116] 本文描述的融合蛋白和方法可以使用任何可编程的DNA结合域。

[0117] 在一些实施方案中,可编程DNA结合蛋白域包含锌指核酸酶(ZFN)或转录激活物样效应域(TALE)的DNA结合域。在一些实施方案中,可编程DNA结合蛋白域可以由引导核苷酸序列编程,并因此称为“引导核苷酸序列-可编程DNA结合蛋白域”。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是核酸酶无活性的Cas9,或dCas9。如本文所用,dCas9涵盖Cas9,其在其核酸酶活性中完全无活性,或在其核酸酶活性中部分无活性(例如,Cas9切口酶)。因此,在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是Cas9切口酶。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是核酸酶无活性的Cpf1。在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是核酸酶无活性的Argonaute。

[0118] 在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是dCas9域。在一些实施

方案中,引导核苷酸序列-可编程DNA结合蛋白是Cas9切口酶。在一些实施方案中,dCas9域包含SEQ ID NO:2或SEQ ID NO:3的氨基酸序列。在一些实施方案中,dCas9域包含与本文提供的Cas9域的任一个至少60%、至少65%、至少70%、至少75%、至少80%、至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列,并且包含对应于SEQ ID NO:1中D10X(X是除D以外的任何氨基酸)和/或H840X(X是除H以外的任何氨基酸)的突变。在一些实施方案中,dCas9域包含与本文提供的Cas9域的任一个至少60%、至少65%、至少70%、至少75%、至少80%、至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列,并且包含对应于SEQ ID NO:1中D10A和/或H840A的突变。在一些实施方案中,Cas9切口酶包含与本文提供的Cas9域的任一个至少60%、至少65%、至少70%、至少75%、至少80%、至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列,并且包含对应于SEQ ID NO:1中D10X(X是除D以外的任何氨基酸)的突变和对应于SEQ ID NO:1中位置840的位置处的组氨酸。在一些实施方案中,Cas9切口酶包含与本文提供的Cas9域的任一个至少60%、至少65%、至少70%、至少75%、至少80%、至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同,并且包含对应于SEQ ID NO:1中D10A的突变和对应于SEQ ID NO:1中位置840的位置处的组氨酸的氨基酸序列。在一些实施方案中,提供了dCas9或Cas9切口酶的变体或同源物(例如,分别是SEQ ID NO:2或SEQ ID NO:3的变体),其分别与SEQ ID NO:2或SEQ ID NO:3至少约70%相同、至少约80%相同、至少约90%相同、至少约95%相同、至少约98%相同、至少约99%相同、至少约99.5%相同或至少约99.9%相同,并且包含对应于SEQ ID NO:1中D10A和/或H840A的突变。在一些实施方案中,提供了Cas9的变体(例如,SEQ ID NO:2的变体),其具有比SEQ ID NO:2短或长约5个氨基酸、约10个氨基酸、约15个氨基酸、约20个氨基酸、约25个氨基酸、约30个氨基酸、约40个氨基酸、约50个氨基酸、约75个氨基酸、约100个氨基酸或更多个的氨基酸序列,条件是dCas9变体包含对应于SEQ ID NO:1中的D10A和/或H840A的突变。在一些实施方案中,提供了Cas9切口酶的变体(例如,SEQ ID NO:3的变体),其具有比SEQ ID NO:3短或长约5个氨基酸、约10个氨基酸、约15个氨基酸、约20个氨基酸、约25个氨基酸、约30个氨基酸、约40个氨基酸、约50个氨基酸、约75个氨基酸、约100个氨基酸或更多个的氨基酸序列,条件是dCas9变体包含对应于D10A的突变并且包含对应于SEQ ID NO:1中位置840的位置处的组氨酸。

[0119] 基于本公开和本领域的知识,另外的合适的核酸酶无活性的dCas9域对于本领域技术人员而言将是显而易见的,并且在本公开的范围之内。此类另外的示例性合适的核酸酶无活性的Cas9域包括但不限于SEQ ID NO:1中的D10A/H840A、D10A/D839A/H840A、D10A/D839A/H840A/N863A突变体域(参见例如,Prashant et al.,*Nature Biotechnology*.2013; 31(9):833-838,其通过引用并入本文),或K603R(参见例如,Chavez et al.,*Nature Methods* 12,326-328,2015,其通过引用并入本文)。

[0120] 在一些实施方案中,本文所述的核碱基编辑器包含与野生型Cas9域相比,具有降低的Cas9域与DNA的糖-磷酸主链之间的静电相互作用的Cas9域。在一些实施方案中,Cas9域包含降低Cas9域与DNA的糖-磷酸主链之间的缔合的一个或多个突变。在一些实施方案

中,本文所述的核碱基编辑器包含dCas9(例如,具有SEQ ID NO:1中D10A和H840A突变)或Cas9切口酶(例如,具有SEQ ID NO:1中D10A突变),其中dCas9或Cas9切口酶进一步包含SEQ ID NO:10中提供的氨基酸序列的N497X、R661X、Q695X和/或Q926X突变,或SEQ ID NO:11-260中提供的任何氨基酸序列中相应的突变的一个或多个,其中X是任何氨基酸。在一些实施方案中,本文所述的核碱基编辑器包含dCas9(例如,具有SEQ ID NO:1中D10A和H840A突变)或Cas9切口酶(例如,具有SEQ ID NO:1中D10A突变),其中dCas9或Cas9切口酶进一步包含SEQ ID NO:10中提供的氨基酸序列的N497A、R661A、Q695A和/或Q926A突变,或SEQ ID NO:11-260中提供的任何氨基酸序列中相应的突变的一个或多个。在一些实施方案中,Cas9域(例如本文提供的任何核碱基编辑器的Cas9域)包含如SEQ ID NO:720中所示的氨基酸序列。在一些实施方案中,核碱基编辑器包含如SEQ ID NO:721中所示的氨基酸序列。具有高保真度的Cas9域在本领域中是已知的,并且对于本领域技术人员而言是显而易见的。例如,具有高保真度的Cas9域已经描述于Kleinstiver,B.P.,et al. “High-fidelity CRISPR-Cas9 nucleases with no detectable genome-wide off-target effects.” *Nature* 529, 490-495 (2016); 和Slaymaker,I.M.,et al. “Rationally engineered Cas9 nucleases with improved specificity.” *Science* 351,84-88 (2015) 中;其每一个的全部内容通过引用并入本文。

[0121] 具有降低的Cas9与DNA主链之间的静电相互作用的Cas9变体DKKYSIGLAIGTNSVGW AVITDEYKVPSSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVD DSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLI EGDLPDNDSDVDKLF IQLVQTYNQLFEEINP INASGVDKAIL SARLSKSRRENLI AQLPGEKKNGLFGNLI ALSL GLTPNFKSNFDLAEDAKLQLSKDTYDDDLNLLAQIGDQYADLFLAAKNLSDA ILLSDILRVNTEITKAPLSASMI KRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKNREDL LRKQRTFDNGSIPHQIHLGELHAILRRQEDFYFPLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETIT PWNFEEVVDKGASASQSFIERMTAFDKNLPNEKVLPKHSLLEYEFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVD LLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLKIKDKDFLDNEENEDILEDIVLTLTLF EDREMIERLKYAHLFDDKVMKQLKRRRYTGWALSRLKINGIRDKQSGKTILDFLKSDFANRNFMALIHDDSL TFKEDIQKAQVSGQDLSLHEHIANLAGSPAIKKGI LQTVKVVDELVKVMGRHKPENIVIAMARENQTTQKGQKNSR ERMKRIEEGKELGSQILKEHPVENTQLQNEKLYLYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDN KVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGSELDKAGFIKRQLVETRAITKH VAQILDSRMNTKYDENDKLIREVKVITLKSCLVSDFRKDFQFYKVRINNYHHAHDAYLNAVVG TALIKKYPKLES EFVYGDYKVYDVRKMIKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFAT VRKVL SMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAVSVLVAKVEKGKSKKLL SVKELLGITIMERSSEFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVN FLYLASHYEKLGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVL SAYNKHDKPIREQAENII HLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSI TGLYETRIDLSQLGGD (SEQ ID NO:720)

[0122] 高保真度核碱基编辑器

[0123] MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELKRETCCLYEINWGGRRHSIWRHTSQNTNKHVEVNFIE KFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGLRDLISSGVTIQIM TEQESGYCWRNFVNYSNEAHWPYPHLLWRLYVLELYCII LGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPP

HILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSSKFKVLGNTDRHSIKKNLIGALL
FDSGETAEATRLKRTARRRYTRRNRCYLQEIFSNEMAKVDDSFHRLVESFLVEEDKKHERHPIFGNIVDEVA
HEKYPTIYHLRKKLV DSTDKADRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQLFEENPINAS
GVDAKAILSARLSKSRRENLIAQLPGEKKNLFGNLI ALSLGLTPNFKSNFDLAEDAKLQLSKD TYDDDLNLLA
QIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPKEYKEIFFDQSK
NGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPF
LKDNREKIEKILTFRIPYYVGPLARGNSRFAMTRKSEETITPWNFEVVVKGASASQSFIERMTAFDKNLPNEKVL
PKHSLLEYEFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVE
DRFNASLGTYHDLKI IKDKDFLDNEENEDILEDIVLTLTFEDREMIEERLKYAHLFDDKVMKQLKRRRYTGWG
ALSRLKINGIRDKQSGKITLDFLKSDFANRNFMALIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKG
ILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKQKNSRERMKRIIEG IKELGSQILKEHPVENTQLQNEKLY
LYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGSNDNPSEEVVKKMKNYWRQLLN
AKLITQRKFDNLTKAERGGSELKAGFIKRQLVETRAITKHVAQILDSRMNTKYDENDKLIREVKVITLKSCLVS
DFRKDFQFYKVREINNYHHAHDAYLNAVVGTA LIKKYPKLESEFVYGDYKVYDVRKMIKSEQEIGKATAKYFFYS
NIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVL SMPQVNI VKKTEVQTGGFSKESILPKRNS
DKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSEKPNIDFLEAKGYKEVKK
DLI IKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGKSPEDNEQKQLFVEQHKHYLDE
IIEQISEFSKRVI LADANLDKVL SAYNKHDKPIREQAENI IHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLD
ATLIHQSI TGLYETRIDLSQLGGD (SEQ ID NO:721)

[0124] Cas9蛋白识别靶DNA序列内的短基序(PAM基序),其是Cas9-DNA相互作用所需要的,但不是由与引导RNA核苷酸序列的互补性确定的。如本文所用,“PAM基序”或“前间隔区相邻基序”是指直接在与引导RNA寡核苷酸序列互补的DNA序列后5’-或3’-的DNA序列。若其随后没有合适的PAM序列,Cas9将不会成功结合、切割或切口靶DNA序列。不希望受任何特定理论的束缚,Cas9酶中的特定氨基酸残基负责与PAM的碱基相互作用并确定PAM特异性。因此,这些残基或附近残基的变化导致不同或放松的PAM特异性。改变或放松PAM特异性可以转变Cas9可以结合的位置,如基于本公开对于本领域技术人员显而易见的。

[0125] 野生型酿脓链球菌Cas9识别规范PAM序列(5’-NGG-3’)。其他Cas9核酸酶(例如来自嗜热链球菌(*Streptococcus thermophiles*)、金黄色葡萄球菌、脑膜炎奈瑟氏球菌(*Neisseria meningitidis*)或齿垢密螺旋体(*Treponema denticola*)的Cas9)及其Cas9变体已在本领域中描述为具有不同的或更放松的PAM需求。例如,在Kleinstiver et al., *Nature* 523,481-485,2015;Kleinstiver et al.,*Nature* 529,490-495,2016;Ran et al.,*Nature*,Apr 9;520(7546):186-191,2015;Kleinstiver et al.,*Nat Biotechnol*,33(12):1293-1298,2015;Hou et al.,*Proc Natl Acad Sci U S A*,110(39):15644-9,2014;Prykhodzhiy et al.,*PLoS One*,10(3):e0119372,2015;Zetsche et al.,*Cell* 163,759-771,2015;Gao et al.,*Nature Biotechnology*,doi:10.1038/nbt.3547,2016;Want et al.,*Nature* 461,754-761,2009;Chavez et al.,doi:dx.doi.org/10.1101/058974;Fagerlund et al.,*Genome Biol.*2015;16:25,2015;Zetsche et al.,*Cell*,163,759-771,2015;和Swarts et al.,*Nat Struct Mol Biol*,21(9):743-53,2014中,其每一个通过引用并入本文。

[0126] 因此,本公开的引导核苷酸序列-可编程DNA结合蛋白可以识别多种PAM序列,包括但不限于在由引导RNA确定的DNA序列的3'或5'末端上的PAM序列。例如,序列可以是:NGG、NGAN (SEQ ID NO:741)、NGNG (SEQ ID NO:742)、NGAG (SEQ ID NO:743)、NGCG (SEQ ID NO:744)、NNGRRT (SEQ ID NO:745)、NGRRN (SEQ ID NO:746)、NNRRT (SEQ ID NO:747)、NNNGATT (SEQ ID NO:748)、NNAGAAW (SEQ ID NO:749)、NAAAC (SEQ ID NO:750)、TTN、TTTN (SEQ ID NO:751) 和YTN,其中Y是嘧啶,R是嘌呤,并且N是任何核碱基。

[0127] 本公开的一些方面提供了RNA-可编程DNA结合蛋白,其可以用于将蛋白质(例如碱基编辑器)引导至特定核酸(例如DNA或RNA)序列。核酸可编程DNA结合蛋白包括但不限于Cas9(例如dCas9和nCas9)、CasX、CasY、Cpf1、C2c1、C2c2、C2C3和Argonaute。具有不同的PAM特异性的RNA-可编程DNA结合蛋白的一个实例是来自普雷沃氏菌(*Prevotella*)和弗朗西斯菌(*Francisella*) 1 (Cpf1)的聚簇规则间隔短回文重复。与Cas9类似,Cpf1也是2类CRISPR效应物。已经显示,Cpf1介导了强大的DNA干扰,其具有与Cas9不同的特征。Cpf1是缺乏tracrRNA的单个RNA引导的内切核酸酶,并且它可以利用富含T的前间隔区相邻基序(例如TTN、TTTN (SEQ ID NO:751) 或YTN),其在由引导RNA确定的DNA序列的5'端上。此外,Cpf1经由交错的DNA双链断裂切割DNA。在16种Cpf1家族蛋白中,来自氨基酸球菌(*Acidaminococcus*)和毛螺菌(*Lachnospiraceae*)的两种酶显示在人细胞中具有有效的基因组编辑活性。Cpf1蛋白是本领域已知的并且先前已有描述,例如Yamano et al., "Crystal structure of Cpf1 in complex with guide RNA and target DNA." *Cell* (165) 2016, p. 949-962;其全部内容在此通过引用并入。

[0128] 在本组合物和方法中也有用的是核酸酶无活性的Cpf1 (dCpf1) 变体,其可以用作引导核苷酸序列-可编程DNA结合蛋白域。Cpf1蛋白具有RuvC样内切核酸酶域,其类似于Cas9的RuvC域,但不具有HNH内切核酸酶域,并且Cpf1的N端不具有Cas9的 α 螺旋识别叶(lobe)。它在Zetsche et al., *Cell*, 163, 759-771, 2015 (其通过引用并入本文) 中显示,Cpf1的RuvC样域负责切割两条DNA链并且RuvC样域的失活使Cpf1核酸酶活性失活。例如,对应于新凶手弗朗西斯菌(*Francisella novicida*) Cpf1 (SEQ ID NO:714) 中的D917A、E1006A或D1255A的突变使Cpf1核酸酶活性失活。在一些实施方案中,本公开的dCpf1包含对应于SEQ ID NO:714中D917A、E1006A、D1255A、D917A/E1006A、D917A/D1255A、E1006A/D1255A或D917A/E1006A/D1255A的突变。在其他实施方案中,本公开的Cpf1切口酶可以包含对应于SEQ ID NO:714中D917A、E1006A、D1255A、D917A/E1006A、D917A/D1255A、E1006A/D1255A或D917A/E1006A/D1255A的突变。有用于本公开的实施方案的Cpf1切口酶可以包含其他突变和/或本领域已知的其他突变。应当理解,可以根据本公开使用使Cpf1的RuvC域完全或部分失活的任何突变,例如取代突变、缺失或插入,以及Cpf1的这些突变可以产生例如dCpf1或Cpf1切口酶。

[0129] 因此,在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白是核酸酶无活性的Cpf1 (dCpf1)。在一些实施方案中,dCpf1包含SEQ ID NO:714-717的任一个的氨基酸序列。在一些实施方案中,dCpf1包含与SEQ ID NO:714-717的任一个至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列,并且包含对应于SEQ ID NO:714中D917A、E1006A、D1255A、D917A/E1006A、D917A/D1255A、E1006A/D1255A或D917A/E1006A/D1255A的突变。根

据本公开,来自其他细菌物种的Cpf1也可以用作dCpf1或Cpf1切口酶。

[0130] 野生型新凶手弗朗西斯菌Cpf1 (SEQ ID NO:714) (D917、E1006和D1255是粗体且加下划线的)

[0131]

MSIYQEFVNKYSLSKTLRFELIPQGKTLENIKARGLILDDEKRAKDYKKAKQIIDKYHQF
 FIEEILSSVCISEDLLQNYSDVYFKLKKSSDDDLQKDFKSAKDTIKKQISEYIKDSEKFKN
 LFNQNLIDAKKGQESDLILWLKQSKDNGIELFKANSDITDIDEALEIIKSFKGWTTYFKGF
 HENRKNVYSSNDIPTSIHYRIVDDNLPKFLNKAKYESLKDKAPEAINYEQIKKDLAEELT
 FDIDYKTSEVNQRVFSLDEVFEIANFNNYLNQSGITKFNTIIGGKVFVNGENTKRKGINEYI
 NLYSQQINDKTLKKYKMSVLFKQILSDTESKSFVIDKLEDDSDVVTTMQSFYEQIAAFK
 TVEEKSIKETLSLLFDDLKAQKLDLSKIYFKNDKSLTDLSQQVFDDYSVIGTAVLEYITQ
 QIAPKNLDNPSKKEQELIAKKTEKAKYLSLETIKLALFEFNKHRDIDKQCRFEEILANFA
 AIPMIFDEIAQNKDNLAQISIKYQNQGKDLLQASAEDDVKAIKDLLDQTNNLLHKLKIF
 HISQSEDKANILDKDEHFYLVFEECYFELANIVPLYNKIRNYITQKPYSDEKFKLNFENST
 LANGWDKNKEPDNTAILFIKDDKYLLGVMNKKNNKIFDDKAIKENKGEYKIVYKL
 LPGANKMLPKVFFSAKSIKFNPSDILRIRNHSTHTKNGSPQKGYEKFENIEDCRKFID
 FYKQSISKHPEWKDFGFRFSDTQRYNSIDEFYREVENQGYKLTFFENISESYIDSVVNQ GK
 LYLFQIYNKDFSAYSKGRPNLHTLYWKALFDERNLQDVVYKLNGEAELFYRKQSIPK KI
 THPAKEAIAKNKNDNPKKESVFEYDLIKDKRFTEDKFFFHCPITINFKSSGANKFNDEINL
 LLKEKANDVHILSIDRGERHLAYYTLVDGKGNIIKQDTFNIIGNDRMKTNYHDKLAAIE
 KDRDSARKDWKKINNIKEMKEGYLSQVVHEIAKLVIEYNAIVVFEDLNFGFKRGRFKV
 EKQVYQKLEKMLIEKLNLYLVFKDNEFDKTGGVLRAYQLTAPFETFKKMGKQTGIIYYV
 PAGFTSKICPVTGFVNQLYPKYESVSKSQEFFSKFDKICYNLDKGYFEFSFDYKNFGDKA
 AKGKWTIASFGSRLINFRNSDKNHNWDREVYPTKELEKLLKDYSIEYGHGECIKAAIC
 GESDKKFFAKLTSVLNTILQMRNSKTGTELDYLISPVADVNGNFFDSRQAPKNMPQDA
DANGAYHIGLKGLMLLGRIKNNQEGKKNLVIKNEEYFEFVQNRNN

[0132] 新凶手弗朗西斯菌Cpf1 D917A (SEQ ID NO:715)

[0133]

MSIYQEFVNKYSLSKTLRFELIPQGKTLENIKARGLILDDEKRAKDYKKAKQIIDKYHQF
 FIEEILSSVCISEDLLQNYSDVYFKLKKSSDDDLNQLKDFKSAKDTIKKQISEYIKDSEKFKN
 LFNQNLIDAKKGGQESDLILWLKQSKDNGIELFKANSDITDIDEALEIIKSFKGWTTYFKGF
 HENRKNVYSSNDIPTSIYRIVDDNLPKFLENKAKYESLKDKAPEAINYEQIKKDLAEELT
 FDIDYKTSEVNQRVFLSDEVFEIANFNNYLNQSGITKFNTIIGGKVFNGENTKRKGINEYI
 NLYSQQINDKTLKKYKMSVLFKQILSDTESKSFVIDKLEDDSDVVTMMSQSFYEQIAAFK
 TVEEKSIKETLSLLFDDLKAQKLDLSKIYFKNDKSLTDLSQQVFDDYSVIGTAVLEYITQ
 QIAPKNLDNPSKKEQELIAKKTEKAKYLSLETIKLALAEFNKHRDIDKQCRFEEILANFA
 AIPMIFDEIAQNKDNLAQISIKYQNQGKDLLQASAEDDVKAIKDLLDQTNLLHKLKIF
 HISQSEDKANILDKDEHFYLVFEECYFELANIVPLYNKIRNYITQKPYSDEKFKLNFENST
 LANGWDKNKEPDNTAILFIKDDKYLLGVMNKKNNKIFDDKAIKENKGEQYKIVYKL
 LPGANKMLPKVFFSAKSIFYNPSEDILRIRNHSTHTKNGSPQKGYEKFENIEDCRKFID
 FYKQSISKHPEWKDFGFRFSDTQRYNSIDEFYREVENQGYKLTFFENISESYIDSVVNQGK
 LYLFQIYNKDFSAYSKGRPNLHTLYWKALFDERNLQDVVYKLNGEAELFYRKQSIPKKI
 THPAKEAIAKNKNDNPKKESVFEYDLIKDKRFTEDKFFFHCPITINFKSSGANKFNDEINL
 LLKEKANDVHILSIARGERHLAYYTLVDGKGNIIKQDTFNIIGNDRMKTNYHDKLAAIE
 KDRDSARKDWKKNINNIKEMKEGYLSQVVHEIAKLVIEYNAIVVFEDLNFGFKRGRFKV
 EKQVYQKLEKMLIEKLNLYL VFKDNEFDKTGGVLRAYQLTAPFETFKKMGKQTGHIYYV
 PAGFTSKICPVTGFVNQLYPKYESVSKSQEFFSKFDKICYNLDKGYFEFSFDYKNFGDKA
 AKGKWTIASFGSRLINFRNSDKNHNWDTREVVYPTKELEKLLKDYSIEYGHGECIKAAIC
 GESDKKFFAKLTSVLNTILQMRNSKTGTELDYLISPVADVNGNFFDSRQAPKNMPQDA
DANGAYHIGLKGLMLLGRIKNNQEGKKNLVIKNEEYFEFVQNRNN

[0134] 新凶手弗朗西斯菌Cpf1 E1006A (SEQ ID NO:716)

[0135]

MSIYQEFVNKYSLSKTLRFELIPQGKTLENIKARGLILDDEKRAKDYKKAKQIIDKYHQF
 FIEEILSSVCISEDLLQNYSDVYFKLKKSSDDDLNQLKDFKSAKDTIKKQISEYIKDSEKFKN
 LFNQNLIDAKKGGQESDLILWLKQSKDNGIELFKANSDITDIDEALEIIKSFKGWTTYFKGF
 HENRKNVYSSNDIPTSIYRIVDDNLPKFLENKAKYESLKDKAPEAINYEQIKKDLAEELT
 FDIDYKTSEVNQRVFLSDEVFEIANFNNYLNQSGITKFNTIIGGKVFNGENTKRKGINEYI
 NLYSQQINDKTLKKYKMSVLFKQILSDTESKSFVIDKLEDDSDVVTMMSQSFYEQIAAFK
 TVEEKSIKETLSLLFDDLKAQKLDLSKIYFKNDKSLTDLSQQVFDDYSVIGTAVLEYITQ
 QIAPKNLDNPSKKEQELIAKKTEKAKYLSLETIKLALAEFNKHRDIDKQCRFEEILANFA
 AIPMIFDEIAQNKDNLAQISIKYQNQGKDLLQASAEDDVKAIKDLLDQTNLLHKLKIF
 HISQSEDKANILDKDEHFYLVFEECYFELANIVPLYNKIRNYITQKPYSDEKFKLNFENST
 LANGWDKNKEPDNTAILFIKDDKYLLGVMNKKNNKIFDDKAIKENKGEQYKIVYKL
 LPGANKMLPKVFFSAKSIFYNPSEDILRIRNHSTHTKNGSPQKGYEKFENIEDCRKFID
 FYKQSISKHPEWKDFGFRFSDTQRYNSIDEFYREVENQGYKLTFFENISESYIDSVVNQGK
 LYLFQIYNKDFSAYSKGRPNLHTLYWKALFDERNLQDVVYKLNGEAELFYRKQSIPKKI
 THPAKEAIAKNKNDNPKKESVFEYDLIKDKRFTEDKFFFHCPITINFKSSGANKFNDEINL
 LLKEKANDVHILSIDRGERHLAYYTLVDGKGNIIKQDTFNIIGNDRMKTNYHDKLAAIE
 KDRDSARKDWKKNINNIKEMKEGYLSQVVHEIAKLVIEYNAIVVFADLNFGFKRGRFKV
 EKQVYQKLEKMLIEKLNLYL VFKDNEFDKTGGVLRAYQLTAPFETFKKMGKQTGHIYYV
 PAGFTSKICPVTGFVNQLYPKYESVSKSQEFFSKFDKICYNLDKGYFEFSFDYKNFGDKA
 AKGKWTIASFGSRLINFRNSDKNHNWDTREVVYPTKELEKLLKDYSIEYGHGECIKAAIC
 GESDKKFFAKLTSVLNTILQMRNSKTGTELDYLISPVADVNGNFFDSRQAPKNMPQDA
DANGAYHIGLKGLMLLGRIKNNQEGKKNLVIKNEEYFEFVQNRNN

[0136] 新凶手弗朗西斯菌Cpf1 D1255A (SEQ ID NO:717)

[0137]

MSIYQEFVNKYSLSKTLRFELIPQGKTLENIKARGLILDDEKRAKDYKKAKQIIDKYHQF
 FIEEILSSVCISEDLLQNYSDVYFKLKKSSDDDLQKDFKSAKDTIKKQISEYIKDSEKFKN
 LFNQNLIDAKKGQESDLILWLKQSKDNGIELFKANSDITDIDEALEIIKSFKGWTTYFKGF
 HENRKNVYSSNDIPTSIHRYRIVDDNLPKFLNKAKYESLKDKAPEAINYEQIKKDLAEELT
 FDIDYKTSEVNQRVFLSDEVFEIANFNNYLNQSGITKFNTIIGGKRVNGENTKRKGINEYI
 NLYSQQINDKTLKKYKMSVLFKQILSDTESKSFVIDKLEDDSDVVTMMSQSFYEQIAAFK
 TVEEKSIKETLSLLFDDLKAQKLDLSKIYFKNDKSLTDLSQQVFDDYSVIGTAVLEYITQ
 QIAPKNLDNPSKKEQELIAKKTEKAKYLSLETIKLALFEFNKHRDIDKQCRFEEILANFA
 AIPMIFDEIAQNKDNLAQISIKYQNQGKDLLQASAEDDVKAIKDLLDQTNLLHKLKIF
 HISQSEDKANILDKDEHFYLVFEECYFELANIVPLYNKIRNYITQKPYSDKFKLNFENST
 LANGWDKNKEPDNTAILFIKDDKYLLGVMNKKNNKIFDDKAIKENKGEYKIVYKL
 LPGANKMLPKVFFSAKSIKFNPSDILRIRNHSTHTKNGSPQKGYEKFENIEDCRKFD
 FYKQSISKHPEWKDFGFRFSDTQRYNSIDEFYREVENQGYKLTFFENISESYIDSVVNQK
 LYLFQIYNKDFSAYSKGRPNLHTLYWKALFDERNLQDVVYKLNGEAELFYRKQSIPKKI
 THPAKEAIANKNDNPKKESVFEYDLIKDKRFTEDKFFFHCPITINFKSSGANKFNDEINL
 LLKEKANDVHILSIDRGERHLAYYTLVDGKGNIIKQDTFNIIGNDRMKTNYHDKLAAIE
 KDRDSARKDWKKINNIKEMKEGYLSQVVHEIAKLVIEYNAIVVFEDLNFGFKRGRFKV
 EKQVYQKLEKMLIEKLNLYL VKDNEFDKTGGVLRAYQLTAPFETFKKMGKQTGHIYYV
 PAGFTSKICPVTGFVNQLYPKYESVSKSQEFFSKFDKICYNLDKGYFEFSFDYKNFGDKA
 AKGKWTIASFGSRLINFRNSDKNHNWDTREVVYPTKELEKLLKDYSIEYGHGECIKAIC
 GESDKKFFAKLTSVLNTILQMRNSKTGTLDYLISPVADVNGNFFDSRQAPKNMPQDA
 AANGAYHIGLKGMLMLLGRKNNQEGKLNLVIKNEEYFEFVQNRNN

[0138] 除了Cas9和Cpf1之外,Shmakov et al.,“Discovery and Functional Characterization of Diverse Class 2 CRISPR Cas Systems”,Mol.Cell,2015 Nov 5; 60 (3) :385-397已经描述了三种不同的2类CRISPR-Cas系统 (C2c1、C2c2和C2c3),其全部内容在此通过引用并入。两个系统 (C2c1和C2c3)的效应物含有与Cpf1相关的RuvC样内切核酸酶域。第三个系统,C2c2含有具有两个预测的HEPN RNA酶域的效应物。与C2c1产生CRISPR RNA不同,成熟CRISPR RNA的产生是不依赖tracrRNA的。C2c1依赖于CRISPR RNA和tracrRNA两者用于DNA切割。已显示细菌性C2c2对于CRISPR RNA成熟具有独特的RNA酶活性,不同于其RNA激活的单链RNA降解活性。这些RNA酶功能彼此不同,并且与Cpf1的CRISPR RNA加工行为不同。参见例如,East-Seletsky,et al.,“Two distinct RNase activities of CRISPR-C2c2enable guide-RNA processing and RNA detection”,Nature,2016Oct 13; 538 (7624) :270-273,其全部内容在此通过引用并入。Leptotrichia shahii中C2c2的体外生化分析已显示,C2c2由单个CRISPR RNA引导,并且可以编程以切割携带互补前间隔区的ssRNA靶标。两个保守的HEPN域中的催化残基介导切割。催化残基中的突变产生催化无活性的RNA结合蛋白。参见例如Abudayyeh et al.,“C2c2is a single-component programmable RNA-guided RNA-targeting CRISPR effector”,Science,2016Aug5;353 (6299),其全部内容在此通过引用并入。

[0139] 已经报告了与嵌合单分子引导RNA (sgRNA)复合的酸土脂环酸芽孢杆菌 (Alicyclobaccillus acidoterrastris) C2c1 (AacC2c1)的晶体结构。参见例如,Liu et al.,“C2c1-sgRNA Complex Structure Reveals RNA-Guided DNA Cleavage Mechanism”,Mol.Cell,2017Jan 19;65 (2) :310-322,其全部内容在此通过引用并入。还已经报告了在与靶DNA结合的酸土脂环酸芽孢杆菌C2c1中作为三元复合物的晶体结构。参见例如,Yang et

al., "PAM-dependent Target DNA Recognition and Cleavage by C2C1CRISPR-Cas endonuclease", Cell, 2016 Dec 15; 167 (7): 1814-1828, 其全部内容在此通过引用并入。具有靶DNA链和非靶DNA链两者的AacC2c1的催化能力构象已被独立地捕获定位在单个RuvC催化袋内, 其中C2c1介导的切割导致靶DNA的交错的七核苷酸断裂。C2c1三元复合物与先前鉴定的Cas9和Cpf1对应物之间的结构比较证明了CRISPR-Cas9系统使用的机制的多样性。

[0140] 在一些实施方案中, 本文提供的任何融合蛋白的引导核苷酸序列-可编程DNA结合蛋白可以是C2c1、C2c2或C2c3蛋白。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白是C2c1蛋白。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白是C2c2蛋白。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白是C2c3蛋白。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白包含与天然存在的C2c1、C2c2或C2c3蛋白至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白是天然存在的C2c1、C2c2或C2c3蛋白。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白包含本文所述的任何C2c1、C2c2或C2c3蛋白至少85%、至少90%、至少91%、至少92%、至少93%、至少94%、至少95%、至少96%、至少97%、至少98%、至少99%或至少99.5%相同的氨基酸序列。在一些实施方案中, 引导核苷酸序列-可编程DNA结合蛋白包含本文所述的C2c1、C2c2或C2c3蛋白的任一个的氨基酸序列。应当理解, 根据本公开也可以使用来自其他细菌物种的C2c1、C2c2或C2c3。

[0141] C2c1 (uniprot.org/uniprot/T0D7A2#)

[0142] sp|T0D7A2|C2C1_ALIAG CRISPR相关的内切核酸酶C2c10S=酸土脂环酸芽孢杆菌 (菌株ATCC 49025/DSM 3922/CIP 106132/NCIMB 13137/GD3B) GN=c2c1PE=1SV=1

[0143] MAVKSIKVKLRLLDDMPEIRAGLWKLHKEVNAGVRYYTEWLSLLRQENLYRRSPNGDGEQECDKTAECKAELLERLRARQVENGHRGPAGSDELLQLARQYELLVPAIGAKGDAQQIARKFLSPLADKDAVGGLGIKAGNKPRWVRMREAGEPGWEEKEKAETRKSADRTADVLRALADFLKPLMRVYTDSEMSSVEWKPLRKGQAVRTWDRDMFQQAIERMMSWESWNQRVVGQYAKLVEQKNRFEQKNFVQGEHLVHLVNQLQQDMKEASPGLESKEQTAHYVTGRALRGSDKVFEEKWGLAPDAPFDLYDAEIKNVQRRNTRRFGSHDLFAKLAPEYQALWREDASFLTRYAVYNSILRKLNHAKMFATFTLPDATAHPWTRFDKLGGLNHQYTFLENEFGERRHAIRFHKLLKVENGVAREVDDVTVPISMSEQLDNLPRDPNEPIALYFRDYGAEQHFTGEFGGAKIQCRRDQLAHMHRRRGARDVYLVNSVVRVQSQSEARGERRPPYAAVFRLVGDNHRAVHFVDFKLSDYLAEHPDDGKLGSEGLLSGLRVMSVDLGLRTSASISVFRVARKDELKPNKGRVPPFFPIKGNLNLVAVHERSLLKLPGETESKDLRAIREERQRTLRLTQLAYLRLLVRCGSEDVGRRERSWAKLIEQPVDAAHMTDPDWREAFENELQKLKSLHGICSDKEWMDAVYESVRRVWRHMKGQVRDWRKDVRSGERPKIRGYAKDVGGNSIEQIEYLERQYKFLKSWFFGKVSQVIRAEKGSRAITLREHIDHAKEDRLKLDRIIMEALGYVYALDERGKGGKWKVAKYPPCQLILLEELSEYQFNDRPPSENNQLMQWSHRGVFQELINQAQVHDLLVGTMYAAFSSRFDARTGAPGIRCRRVPARCTQEHNPEFPWWLNKFVVEHTLDACPLRADDLIPTGEGEIVSPFSAEEGDFHQIHADLNAQNLQQRLWSDFDISQIRLRCDWGEVDGELVLIPLRTGKRTADSYSNKVFYTNVTGTYTYERERGGKRRKRVFAQEKLSEEEAELLVEADEAREKSVVLMRDPSTGIINRGWNTRQKEFWSMVNQRIEGYLVKQIRSRVPLQDSACENTGDI (SEQ ID NO: 762)

[0144] C2c2 (uniprot.org/uniprot/P0D0C6)

[0145] >sp|P0D0C6|C2C2_LEPSD CRISPR相关的内切核糖核酸酶C2c20S=Leptotrichia

shahii (菌株DSM 19757/CCUG 47503/CIP 107916/JCM 16776/LB37) GN=c2c2PE=1SV=1
 [0146] MGNLFGHKRWYEVDRDKKDFKIKRKVKVKNRYDGNKYILNINENNNKEKIDNNKFIKRYINYKKNNDNIL
 KEFTRKFHAGNILFKLKGKEGIIRIENNDFFLETEEVVLYIEAYGKSEKLGITKKKIIDEAIRQGITKDDKKI
 EIKRQENEEIEIDIRDEYTNKTLNDCSIIILRIIENDELETKKSIIYEIFKNINMSLYKIIIEKIIENETEKVFENRY
 YEEHLREKLLKDDKIDVILTNFMEIREKIKSNLEILGFVKFYLVGGDKKSKNKKMLVEKILNINVDLTVEDIAD
 FVIKELEFWNITKRIEKVKKVNEFLEKRRNRITYIKSYVLLDKHEKFKIERENKKDKIVKFFVENIKNNSIKEKIE
 KILAEPKIDELIKKLEKELKKGNCDEIFGIFKKHYKVNFDKSKFSKKSDEEKELYKIIYRYLKGRIEKILVNEQK
 VRLKKMEKIEIEKILNESILSEKILKRVKQYTLHEIMYLGKLRHNDIDMTTVNTDDFSRLHAKEELDLELITFFAS
 TMELNKFISRENINNDENIDFFGGDREKNYVLDKILNSKIKIIRDLDFIDNKNNITNNFIRKFTKIGTNERNRI
 LHAISKERDLQGTQDDYNKVINIIQNLKISDEEVSKALNLDVVFVKDKKNIITKINDIKISEENNDIKYLPFSKVI
 LPEILNLYRNNPKNEPFDIETEKIVLNALIYVNKELYKLLILEDDLEENESKNIFLQELKKTGLNIDEIDENIIIE
 NYKNAQISASKGNKAIKKYQKKVIECYIGYLRKNYEELDFDSDFKMNIQEIKKQIKDINDNKTYERITVKTSDK
 TIVINDDFEYIISIFALLNSNAVINKIRNRFATSVWLNTSEYQNIIDILDEIMQLNLRNECITENWNLNLEEFI
 QKMKEIEKDFDDFKIQTKKEIFNYYEDIKNNILTEFKDDINGCDVLEKKLEKIVIFDDETKFEIDKKSNIQDEQ
 RKLSNINKKDLKKKVDQYIKDKDQEKSKILCRIIFNSDFLKKYKKEIDNLIEDMESENEKFKQEIYYPKERKNEL
 YIYKKNLFLNIGNPNFDKIYGLISNDIKMADAKFLFNIDGKNIRKNKISEIDAILKNLNDKLNYSKEYKEYIKK
 LKENDDFFAKNIQKNKYKSFEDYNRVSEYKIRDLVEFNLYNKIESYLDINWKLAIQMARFERDMHYIVNGLRE
 LGI IKLSGYNTGISRAYPKRNGSDGFYTTTAYYKFFDEESYKKFEKICYGFGIDLSENSEINKPENESIRNYISHF
 YIVRNPADYSIAEQIDRVSNLLSYSTRYNNSTYASVFEVFKKDVNLDYDELKKKFKLIGNNDILERLMKPKKVS
 LELESYNSDYIKNLIIELLTKIENTNDTL (SEQ ID NO:764)

[0147] 在一些实施方案中,本公开的引导核苷酸序列-可编程DNA结合蛋白对PAM序列没有需求。此类引导核苷酸序列-可编程DNA结合蛋白的一个实例可以是来自格氏嗜盐碱杆菌 (*Natronobacterium gregoryi*) 的Argonaute蛋白 (NgAgo)。NgAgo是ssDNA引导的内切核酸酶。NgAgo结合约24个核苷酸的5'磷酸化ssDNA (gDNA), 以将其引导至其靶位点, 并将在gDNA位点处产生DNA双链断裂。与Cas9相比, NgAgo-gDNA系统不需要前间隔区相邻基序 (PAM)。使用核酸酶无活性的NgAgo (dNgAgo) 可以极大地扩展可以靶向的密码子。NgAgo的表征和使用已经描述于Gao et al., *Nat Biotechnol.*, 2016 Jul; 34 (7): 768-73. PubMed PMID: 27136078; Swarts et al., *Nature*. 507 (7491) (2014): 258-61; 和 Swarts et al., *Nucleic Acids Res.* 43 (10) (2015): 5120-9, 其每一个通过引用并入本文。格氏嗜盐碱杆菌 Argonaute 的序列提供于 SEQ ID NO: 718 中。

[0148] 野生型格氏嗜盐碱杆菌 Argonaute (SEQ ID NO: 718)

[0149] MTVIDLDSTTTADELTSGHYDYSVTLTGVDNTDEQHPRMSLAFEQDNGERRYITLWKNTTPKDVFTY
 DYATGSTYIFTNIDYEVKDGyenLTATYQTTVENATAQEVGTTDEDETFAGGEPLDHHLDDALNETPDDAETESDSG
 HVMTSFASRDQLPEWTLHTYTLTATDGAKTDEYARRTLAYTVRQELYTDHDAAPVATDGLMLLTPEPLGETPLDL
 CGVRVEADETRTLDYTTAKDRLLARELVEEGLKRSWDDYLVRGIDEVLSKEPVLTCDEFDLHERYDLSVEVGHSGR
 AYLHINFRHRFPKLTADIDDDNIYPGLRVKTTYRPRRGIHVWGLRDECATDSLNTLGNQSVVAYHRNNQTPINTD
 LLDAIEAADRRVVETRRQGHGDDAVSFPQELLAVEPNTHQIKQFASDGFHQARSKTRLSASRCSEKAQAFERLDP
 VRLNGSTVEFSSEFFTGNNEQQRLRLLYENGESVLTFRDARGAHPDETFSKGI VNPPESEFVAVVLPEQQADTCKAQ
 WDTMADLLNQAGAPPTRSETVQYDAFSSPESISLNVAGAI DPSEVDAAFVVLPPDQEGFADLASPTETYDELKALA

NMGIYSQMAYFDRFRDAKIFYTRNVALGLLAAAGGVAFTTEHAMPGDADMFIGIDVSRSPEDGASGQINIAATATA
VYKDGITILGHSSTRPQLGKELQSTDVRDIMKNAILGYQQVTGESPTHIVIHHRDGMNEDLDPATEFLNEQGVEYDIV
EIRKQPQTRLLAVSDVQYDTPVKSIAAINQNEPRATVATFGAPEYLATRDGGGLPRPIQIERVAGETDIETLTRQVY
LLSQSHIQVHNSTARLPITTAYADQASTHATKGYLVQTGAFESNVGFL

[0150] 本文还提供了具有放松的PAM需求的Cas9变体(无PAM的Cas9)。与如SEQ ID NO:1提供的酿脓链球菌Cas9相比,无PAM的Cas9对靶序列表现出增加的活性,所述靶序列在其3'端处不包括规范的PAM(例如,NGG)序列,例如活性增加至少5倍、至少10倍、至少50倍、至少100倍、至少500倍、至少1,000倍、至少5,000倍、至少10,000倍、至少50,000倍、至少100,000倍、至少500,000倍或至少1,000,000倍。此类具有放松的PAM需求的Cas9变体描述于2015年10月23日提交的美国临时申请,USSN 62/245,828;2016年1月15日提交的62/279,346;2016年3月22日提交的62/311,763;2016年4月13日提交的62/322,178;和2016年6月30日提交的62/357,332,其每一个通过引用并入本文。在一些实施方案中,可用于本公开的dCas9或Cas9切口酶可以进一步包含放松PAM需求的突变,例如,对应于SEQ ID NO:1中A262T、K294R、S409I、E480K、E543D、M694I或E1219V的突变。

[0151] 本文讨论的一般结构中使用的“-”可以表示存在任意的接头。如本文所用,术语“接头”是指连接两个分子或部分,例如融合蛋白的两个域,诸如例如引导核苷酸序列-可编程DNA结合蛋白域和重组酶催化域的化学基团或分子。通常,接头位于两个基团、分子或其他部分之间或侧翼有两个基团、分子或其他部分,并且经由共价键与每一个连接,从而连接两者。在一些实施方案中,接头是一个氨基酸或多个氨基酸(例如肽或蛋白质)。在一些实施方案中,接头是有机分子、基团、聚合物或化学部分。在一些实施方案中,接头的长度为5-100个氨基酸,例如长度为5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、23、24、25、26、27、28、29、30、30-35、35-40、40-45、45-50、50-60、60-70、70-80、80-90、90-100、100-150或150-200个氨基酸。也考虑了更长或更短的接头。接头可以是本领域已知的任何形式。例如,接头可以是来自网站诸如www[dot]ibi[dot]vu[dot]nl/programs/linkerdbwww/或来自www[dot]ibi[dot]vu[dot]nl/programs/linkerdbwww/src/database.txt的接头。接头也可以是非结构化的、结构化的、螺旋形的或延伸的。

[0152] 在一些实施方案中,引导核苷酸序列-可编程DNA结合蛋白域和重组酶催化域经由接头彼此融合。可以采用引导核苷酸序列-可编程DNA结合蛋白域和重组酶催化域之间的各种接头长度和柔性度(例如,范围为从形式(GGGS)_n(SEQ ID NO:759)、(GGGGS)_n(SEQ ID NO:722)、(GGS)_n和(G)_n的柔性接头至形式(EAAAK)_n(SEQ ID NO:723)、SGSETPGTSESATPES(SEQ ID NO:724)的更刚性接头(参见例如Guilinger et al., Nat. Biotechnol. 2014; 32(6): 577-82;其全部内容通过引用并入本文)、(XP)_n或任何这些的组合,其中X是任何氨基酸并且n独立地是1至30之间的整数,以便实现对于特定应用的活性的最佳长度。在一些实施方案中,n独立地是1、2、3、4、5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、23、24、25、26、27、28、29或30,或者若存在超过一个接头或超过一个接头基序,其任何组合。在一些实施方案中,接头包含(GGS)_n基序,其中n是1、2、3、4、5、6、7、8、9、10、11、12、13、14或15。在一些实施方案中,接头包含(GGS)_n基序,其中n是1、3或7。在一些实施方案中,接头包含XTEN接头。XTEN接头可以具有序列SGSETPGTSESATPES(SEQ ID NO:7)、SGSETPGTSESA(SEQ ID NO:8)或SGSETPGTSESATPEGGSGGS(SEQ ID NO:9)。在一些实施方案中,接头包含选自下

组的氨基酸序列,其包括但不限于AGVF (SEQ ID NO:772)、GFLG (SEQ ID NO:773)、FK、AL、ALAL (SEQ ID NO:774)和ALALA (SEQ ID NO:775)。在一些实施方案中,合适的接头基序和配置包括描述于Chen et al., Fusion protein linkers: property, design and functionality. Adv Drug Deliv Rev. 2013; 65 (10): 1357-69 (其通过引用并入本文)中的那些。在一些实施方案中,接头可以包含任何以下氨基酸序列: VPFLLEPDNINGKTC (SEQ ID NO:10)、GSAGSAAGSGEF (SEQ ID NO:11)、SIVAQLSRPDPA (SEQ ID NO:12)、MKIIEQLPSA (SEQ ID NO:13)、VRHKLKRVGS (SEQ ID NO:14)、GHGTGSTGSGSS (SEQ ID NO:15)、MSRPDPA (SEQ ID NO:16)、GSAGSAAGSGEF (SEQ ID NO:7)、SGSETPGTSESA (SEQ ID NO:8)、SGSETPGTSESATPEGGSGGS (SEQ ID NO:9)和GGSM (SEQ ID NO:17)。

[0153] 基于本公开,另外的合适的接头序列对于本领域技术人员而言将是显而易见的。在某些实施方案中,接头可以具有约33埃至约81埃的长度。在另一个实施方案中,接头可以具有约54埃至约81埃的长度。在进一步的实施方案中,接头可以具有约63至约81埃的长度。在另一个实施方案中,接头可以具有约65埃至约75埃的长度。在一些实施方案中,接头可以具有约1.20kDa至约1.85kDa的重量。在某些实施方案中,接头可以具有约1.40kDa至约1.85kDa的重量。在某些实施方案中,接头可以具有约1.60kDa至约1.7kDa的重量。在一些实施方案中,接头是一个氨基酸或多个氨基酸(例如肽或蛋白质)。在一些实施方案中,接头是有机分子、基团、聚合物或化学部分。在一些实施方案中,接头是肽接头。在一些实施方案中,肽接头是具有至少1、至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少15、至少20、至少25、至少30、至少40、至少50或更多个氨基酸的任何段的氨基酸。在某些实施方案中,肽接头为18至27个氨基酸长。在具体的实施方案中,肽接头为24个氨基酸长。在一些实施方案中,肽接头包含三肽Gly-Gly-Ser的重复,例如包含序列(GGS)_n,其中n表示至少1、2、3、4、5、6、7、8、9、10或更多个重复。在一些实施方案中,接头包含序列(GGS)₆ (SEQ ID NO:6)。在一些实施方案中,肽接头是16个残基的“XTEN”接头或其变体(参见例如,实施例;和Schellenberger et al. A recombinant polypeptide extends the in vivo half-life of peptides and proteins in a tunable manner. Nat. Biotechnol. 27, 1186-1190 (2009))。在一些实施方案中,XTEN接头包含序列SGSETPGTSESATPES (SEQ ID NO:7)、SGSETPGTSESA (SEQ ID NO:8)或SGSETPGTSESATPEGGSGGS (SEQ ID NO:9)。在一些实施方案中,肽接头选自VPFLLEPDNINGKTC (SEQ ID NO:10)、GSAGSAAGSGEF (SEQ ID NO:11)、SIVAQLSRPDPA (SEQ ID NO:12)、MKIIEQLPSA (SEQ ID NO:13)、VRHKLKRVGS (SEQ ID NO:14)、GHGTGSTGSGSS (SEQ ID NO:15)、MSRPDPA (SEQ ID NO:16);或GGSM (SEQ ID NO:17)。在一些实施方案中,接头是非肽接头。在某些实施方案中,非肽接头包含以下的一个或多个:聚乙二醇(PEG)、聚丙二醇(PPG)、共聚(乙烯/丙烯)二醇、聚氧乙烯(POE)、聚氨酯、聚磷腈、多糖、右旋糖酐、聚乙烯醇、聚乙烯吡咯烷酮、聚乙烯乙醚、聚丙烯酰胺、聚丙烯酸酯、聚氰基丙烯酸酯、脂质聚合物、甲壳质、透明质酸、肝素或烷基接头。在一个实施方案中,烷基接头具有式—NH—(CH₂)_s—C(O)—,其中s可以是任何整数。在进一步的实施方案中,s可以是1-20的任何整数。

[0154] 重组酶催化域

[0155] 用于本公开的组合物和方法的重组酶催化域可以来自任何重组酶。用于所公开的方法和组合物的合适的重组酶催化域可以获得自例如但不限于酪氨酸重组酶和丝氨酸重

组酶。本文提供的一些示例性合适的重组酶包括,但不限于,Gin重组酶(作用于gix位点)、Hin重组酶(作用于hix位点)、 β 重组酶(作用于6个位点)、Sin重组酶(作用于resh位点)、Tn3重组酶(作用于res位点)、 $\gamma\delta$ 重组酶(作用于res位点)、来自噬菌体P1的Cre重组酶(作用于LoxP位点);真菌来源的FLP重组酶(作用于FTR位点);和phiC31整合酶(作用于att位点)。示例性合适的重组酶的非限制性序列可以在下面找到。

[0156] Cre重组酶序列

[0157] MSNLLTVHQNLPALPVDATSDEVRKNLMDMFRDRQAFSEHTWKMLLSVCRSWAAWCKLNNRKWFPAEP
EDVRDYLLYLQARGLAVKTIQQHLGQLNMLHRRSGLPRPSDSNAVSLVMRRIRKENVDAGERAKQALAFERTDFDQ
VRSLMENS DRCQDIRNLAFLGIAYN TLLRIA EIARIRVKDISRTDGG RMLIHIGRTKTLVSTAGVEKALSLGVTKL
VERWISVSGVADDPNNYLFCRVRKNGVAAPSATSQ LSTRALEGIFEATHRLIYGAKDDSGQRYLAWSGHSARVGAA
RDMARAGVSIPEIMQAGGWTNVNIVMNYIRNLDSETGAMVRLLEDGD (SEQ ID NO:725)

[0158] FLP重组酶

[0159] MPQFGILCKTPPKVLVRQFVERFERPSGEKIALCAAELTYLCWMI THNGTAIKRATFMSYNTIIISNSL
SFDIVNKSLQFKYKTQKATILEASLKKLIPAWFTIIPYYGQKHQSDITDIVSSLQLQFESSEEADKGNSSHKKML
KALLSEGESIWEITEKILNSFEYTSRFTKTKTLYQFLFLATFINCGRFS DIKNVDPKSFKL VQNKYLGVI IQCLVT
ETKTSVSRHIYFFSARGRIDPLVYLDEF LRNSEPV LKRVNRTGNSSSNKQEYQLLKDNLVRSYNKALKKNAPYSIF
AIKNGPKSHIGRHLMTSFLSMKGLTELTNVVGNWSDKRASAVARTTYTHQITAIPDHYFALVSRYYAYDPISKEMI
ALKDETNPIEEWQHIEQLKGS AEGSIRYPAWNGIISQEVLDYLSSYINRRI (SEQ ID NO:726)

[0160] $\gamma\delta$ 重组酶 (Gamma Delta解离酶)

[0161] MRLFGYARVSTSQQSLDIQVRALKDAGVKANRIFTDKASGSSSDRKGLDLLRMKVEEGDVILVKKLDR
LGRDTADMIQLIKEFDAQGV SIRFIDDGISTDGEMGKMVVTILSAVAQAERQRILERTNEGRQEAMAKGVVFRGRK
(SEQ ID NO:727)

[0162] $\gamma\delta$ 重组酶 (E124Q突变)

[0163] MRLFGYARVSTSQQSLDIQVRALKDAGVKANRIFTDKASGSSSDRKGLDLLRMKVEEGDVILVKKLDR
LGRDTADMIQLIKEFDAQGV SIRFIDDGISTDGEMGKMVVTILSAVAQAERQRILQRTNEGRQEAMAKGVVFRGRK
(SEQ ID NO:728)

[0164] $\gamma\delta$ 重组酶 (E102Y/E124Q突变)

[0165] MRLFGYARVSTSQQSLDIQVRALKDAGVKANRIFTDKASGSSSDRKGLDLLRMKVEEGDVILVKKLDR
LGRDTADMIQLIKEFDAQGV SIRFIDDGISTDGYMGKMVVTILSAVAQAERQRILQRTNEGRQEAMAKGVVFRGRK
(SEQ ID NO:729)

[0166] β 重组酶

[0167] MAKIGYARVSSKEQNLDRLQALQGVSKVFSKLSGQSVERPQLQAMLNYIREGDIVVTELDRLGRN
NKELTELMNAIQK GATLEVLDLPSMNGIEDENLRRLINNLVIELYKYQAESERKR IKERQAQGIEIAKSKGKFKG
RQH (SEQ ID NO:730)

[0168] β 重组酶 (N95D突变)

[0169] MAKIGYARVSSKEQNLDRLQALQGVSKVFSKLSGQSVERPQLQAMLNYIREGDIVVTELDRLGRN
NKELTELMNAIQK GATLEVLDLPSMDGIEDENLRRLINNLVIELYKYQAESERKR IKERQAQGIEIAKSKGKFKG
RQH (SEQ ID NO:731)

[0170] Sin重组酶

[0171] MIIGYARVSSLDQNLERQLENLKTFGAEKIFTEKQSGKSIENRPILQKALNFVRMGDRFIVESIDRLG RNYNEVIHTVNYLKDKEVQLMITSLPMMNEVIGNPLLDKFMKDLIIQILAMVSEQERNESKRRQAQGIQVAKEKGV YKGRPL (SEQ ID NO:732)

[0172] Sin重组酶(Q87R/Q115R突变)

[0173] MIIGYARVSSLDQNLERQLENLKTFGAEKIFTEKQSGKSIENRPILQKALNFVRMGDRFIVESIDRLG RNYNEVIHTVNYLKDKEVRLMITSLPMMNEVIGNPLLDKFMKDLIIIRILAMVSEQERNESKRRQAQGIQVAKEKGV YKGRPL (SEQ ID NO:733)

[0174] Tn3重组酶

[0175] MRLFGYARVSTSQQSLDLQVRALKDAGVKANRIFTDKASGSSTDREGLDLLRMKVKEGDVILVKKLDR LGRDTADMLQLIKEFDAQGVAVRFIDDGISTDGMGQMVVTILSAVAQAERRRILERTNEGRQEAKLKGIFGRRR (SEQ ID NO:734)

[0176] Tn3重组酶(G70S/D102Y、E124Q突变)

[0177] MRLFGYARVSTSQQSLDLQVRALKDAGVKANRIFTDKASGSSTDREGLDLLRMKVKEGDVILVKKLDR LSRDTADMLQLIKEFDAQGVAVRFIDDGISTDGYMGQMVVTILSAVAQAERRRILQRTNEGRQEAKLKGIFGRRR (SEQ ID NO:735)

[0178] Hin重组酶

[0179] MATIGYIRVSTIDQNIDLQRNALTSANCDRIFEDRISGKIANRPGLKRALKYVNKGDTLVVWKLDRLG RSVKNLVALISELHERGAHFHSLTDSIDTSSAMGRFFFHVMSALAEMERELIVERTLAGLAAARAQRLGGRPV (SEQ ID NO:736)

[0180] Hin重组酶(H107Y突变)

[0181] MATIGYIRVSTIDQNIDLQRNALTSANCDRIFEDRISGKIANRPGLKRALKYVNKGDTLVVWKLDRLG RSVKNLVALISELHERGAHFHSLTDSIDTSSAMGRFFFYVMSALAEMERELIVERTLAGLAAARAQRLGGRPV (SEQ ID NO:737)

[0182] PhiC31重组酶

[0183] MDTYAGAYDRQSRERENSSAASPATQRSANEDKAADLQREVERDGGFRFRVGFHFSEAPGTSFAFGTAER PEFERILNECRAGRLNMIIVYDVSFRSRLKVMDAIPIVSELLALGVTIVSTQEGVFRQGNVMDLIHLIMRLDASHK ESSLKSAKILDTKNLQRELGGYVGGKAPYGFELVSETKEITRNGRMVNVVINKLAHSTTPLTGPFEFEPDVIRWWW REIKTHKHLPFKPGSQAAIHPGSITGLCKRMDADAVPTRGETIGKKTASSAWDPATVMRILRDPRIAGFAAEVIYK KKPDPGTPTTKIEGYRIQRDPITLRPVELDCGPIIEPAEWYELQAWLDGRGRGKGLSRGQAILSAMDKLYCECGAVM TSKRGEESIKDSYRCRRRVVDPSPAGQHEGTCNVSMALDKFVAERIFNKIRHAEGDEETLALLWEAARRFGKLT EAPEKSGERANLVAERADALNALEELYEDRAAGAYDGPVGRKHFRKQQAALTLRQQGAEERLAELEAAEAPKLPLD QWFPEDADADPTGPKSWWGRASVDDKRVFVGLFVDKIVVTKSTTGRGQGTPIEKRASITWAKPPTDDDEDDAQDGT EDVAATGA (SEQ ID NO:738)

[0184] 与所公开的组合物和方法一起使用的重组酶还可以包括其他突变。该公开的一些方面提供了包含与本文所讨论的重组酶序列的序列至少70%、至少80%、至少90%、至少95%或至少97%相同的氨基酸序列的重组酶,其中与本文所讨论的重组酶序列的序列相比,重组酶的氨基酸序列包含至少一个突变。在一些实施方案中,与本文所讨论的重组酶序列的序列相比,重组酶的氨基酸序列包含至少2个、至少3个、至少4个、至少5个、至少6个、至少7个、至少8个、至少9个、至少10个、至少11个、至少12个、至少13个、至少14个或至少15个

突变。

[0185] 例如, $\gamma\delta$ 重组酶可以包含来自列表的一个或多个突变:R2A、E56K、G101S、E102Y、M103I或E124Q。在一个实施方案中, $\gamma\delta$ 重组酶可以包含E102Y突变、E124Q突变或E102Y和E124Q突变两者。在另一个实施方案中, β 重组酶可以包含一个或多个突变,其包括但不限于N95D。参见例如, Sirk et al., “Expanding the zinc-finger recombinase repertoire: directed evolution and mutational analysis of serine recombinase specificity determinants” *Nucl Acids Res* (2014) 42 (7):4755-4766。在另一个实施方案中, Sin重组酶可以具有一个或多个突变,其包括但不限于:Q87R、Q115R或Q87R和Q115R。在另一个实施方案中, Tn3重组酶可以具有一个或多个突变,其包括但不限于:G70S、D102Y、E124Q及其任何组合。在另一个实施方案中, Hin重组酶可以具有一个或多个突变,其包括但不限于:H107Y。在另一个实施方案中, Sin重组酶可以具有一个或多个突变,其包括但不限于:H107Y。与所公开的组合物和方法一起使用的任何重组酶催化域可以与天然(或野生型)氨基酸序列具有大于85%、90%、95%、98%或99%的序列一致性。例如,在某些实施方案中, Gin重组酶催化域与SEQ ID NO:713中所示的氨基酸序列具有大于85%、90%、95%、98%或99%的序列一致性。在另一个实施方案中, Gin重组酶催化域的氨基酸序列包含对应于H106Y, 和/或I127L, 和/或I136R和/或G137F的突变。在另一个实施方案中, Gin重组酶催化域的氨基酸序列包含对应于H106Y、I127L、I136R和G137F的突变。在进一步的实施方案中, Gin重组酶的氨基酸序列已得以进一步突变。在具体的实施方案中, Gin重组酶催化域的氨基酸序列包含SEQ ID NO:713。

[0186] 用于本公开的组合物和方法的重组酶催化域可以来自演化的重组酶。如本文所用, 术语“演化的重组酶”是指已经改变(例如, 通过突变)以识别非天然DNA靶序列的重组酶。

[0187] 可以演化的合适的重组酶包括例如但不限于酪氨酸重组酶和丝氨酸重组酶(例如, 本文讨论的任何重组酶)。可以通过本文提供的方法和策略演化的一些示例性合适的重组酶包括例如但不限于, Gin重组酶(作用于gix位点)、Hin重组酶(作用于hix位点)、 β 重组酶(作用于6个位点)、Sin重组酶(作用于resH位点)、Tn3重组酶(作用于res位点)、 $\gamma\delta$ 重组酶(作用于res位点)、来自噬菌体P1的Cre重组酶(作用于LoxP位点); λ 噬菌体整合酶(作用于att位点); 真菌来源的FLP重组酶(作用于FTR位点); phiC31整合酶; Dre重组酶, BxB1; 和原核 β -重组酶。

[0188] 例如, 可以改变与本公开的组合物和方法一起使用的演化的重组酶以与非规范重组酶靶序列相互作用(例如, 结合和重组)。作为非限制性实例, 非规范重组酶靶序列可以是天然存在的, 诸如例如哺乳动物基因组中“安全港(safe harbor)”基因组基因座内的序列, 例如已知对基因修饰耐受而没有任何不期望的影响的基因组基因座。靶向此类序列的重组酶允许例如在特定基因组位置处靶向插入核酸构建体, 而不需要常规的时间和劳动密集型基因靶向规程, 例如经由同源重组技术。此外, 本文提供的定向演化策略可以用于演化具有改变的活性谱的重组酶, 例如, 有利于核酸序列的整合而不是该序列的切除的重组酶, 或反之亦然。

[0189] 与其野生型对应物相比, 演化的重组酶表现出改变的靶序列偏好, 可以用于靶向用于重组酶活性的实际任何靶序列。因此, 演化的重组酶可以用于修饰例如细胞或受试者

的基因组内的任何序列。因为重组酶可以影响异源核酸分子插入到靶核酸分子中、从核酸分子中切除核酸序列、倒位或替换核酸序列,所以本文提供的技术能够以多种方式(例如,整合、缺失、倒位、交换核酸序列)有效修饰基因组靶标。

[0190] 来自与本公开的方法和组合物一起使用的演化的重组酶的催化域包含与野生型重组酶的序列至少70%、至少80%、至少90%、至少95%或至少97%相同的氨基酸序列,其中与野生型重组酶的序列相比,演化的重组酶的氨基酸序列包含至少一个突变,并且其中演化的重组酶识别DNA重组酶靶序列,其与规范重组酶靶序列相差至少一个核苷酸。在一些实施方案中,演化的重组酶识别DNA重组酶靶序列,其与规范重组酶靶序列(例如,res、gix、hix、six、resH、LoxP、FTR或att核心或相关核心序列)相差至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14、至少15、至少16、至少17、至少18、至少19、至少20至少25或至少30个核苷酸。在一些实施方案中,演化的重组酶识别DNA重组酶靶序列,其与规范重组酶靶序列相差1、2、3、4、5、6、7、8、9、10、11、12、13、14、15、16、17、18、19、20、21、22、23、24、25、26、27、28、29或30个核苷酸。

[0191] 在一些实施方案中,仅重组酶的一部分用于本文所述的融合蛋白和方法中。作为非限制性实施方案,仅重组酶的C端部分可以用于本文所述的融合蛋白和方法中。在具体的实施方案中,Cre重组酶的25kDa羧基末端域可以用于组合物和方法中。参见例如,Hoess et al,“DNA Specificity of the Cre Recombinase Resides in the 25kDa Carboxyl Domain of the Protein,”J.Mol.Bio.1990Dec 20,216(4):873-82,其出于所有目的通过引用并入本文。Cre重组酶的25kDa羧基末端域是从蛋白质的R118延伸至羧基末端的部分。在一些实施方案中,用于本发明融合蛋白和方法的Cre重组酶的25kDa羧基末端域可以与Cre重组酶规范25kDa羧基末端域相差至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14、至少15、至少16、至少17、至少18、至少19或至少20个氨基酸。在一些实施方案中,用于本发明融合蛋白和方法的Cre重组酶的25kDa羧基末端域可以与Cre重组酶规范25kDa羧基末端域相差1、2、3、4、5、6、7、8、9、10、11、12、13、14、15、16、17、18、19或20个氨基酸。在某些实施方案中,仅Cre重组酶的25kDa羧基末端域的一部分可以用于本文所述的融合蛋白和方法中。例如,使用的Cre重组酶的部分可以是R130至蛋白质的羧基末端、T140至蛋白质的羧基末端、E150至蛋白质的羧基末端、N160至蛋白质的羧基末端、T170至蛋白质的羧基末端、I180至蛋白质的羧基末端、G190至蛋白质的羧基末端、T200至蛋白质的羧基末端、E210至蛋白质的羧基末端、L220至蛋白质的羧基末端、V230至蛋白质的羧基末端、C240至蛋白质的羧基末端、P250至蛋白质的羧基末端、A260至蛋白质的羧基末端、R270至蛋白质的羧基末端、G280至蛋白质的羧基末端、S290至蛋白质的羧基末端、A300至蛋白质的羧基末端或M310至蛋白质的羧基末端。作为另一组非限制性实例,使用的Cre重组酶的部分可以是R118-E340、R118-S330、R118-I320、R118-M310、R118-A300、R118-S290、R118-G280、R118-R270、R118-A260、R118-P250、R118-C240、R118-V230、R118-L220或R118-E210。作为进一步的一组非限制性实例,使用的Cre重组酶的部分可以是R118-E210、G190-R270、E210-S290、P250-M310或R270至蛋白质的羧基末端。

[0192] 在一些实施方案中,本文所述的融合蛋白和方法中使用的Cre重组酶可以在任何位置截短。在具体的实施方案中,本文所述的融合蛋白和方法中使用的Cre重组酶可以截短,使得它以氨基酸R118、A127、E138或R154开始(每种情况下其之前都是甲硫氨酸)。在另

一组非限制性实施方案中,本文所述的融合蛋白和方法中使用的Cre重组酶可以在R118、A127、E138或R154的10个氨基酸、9个氨基酸、8个氨基酸、7个氨基酸、6个氨基酸、5个氨基酸、4个氨基酸、3个氨基酸、2个氨基酸或1个氨基酸内截短。

[0193] 在一些实施方案中,重组酶靶序列为10-50个核苷酸之间长。在一些实施方案中,重组酶是Cre重组酶、Hin重组酶或FLP重组酶。在一些实施方案中,规范重组酶靶序列是LoxP位点(5'-ATAACTTCGTATA GCATACAT TATACGAAGTTAT-3'(SEQ ID NO:739)。在一些实施方案中,规范重组酶靶序列是FRT位点(5'-GAAGTTCCTATTCTCTAGAAA GTATAGGAAGTTC-3')(SEQ ID NO:740)。在一些实施方案中,与野生型重组酶的序列相比,演化的重组酶的氨基酸序列包含至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14或至少15个突变。在一些实施方案中,演化的重组酶识别包含左半位点、间隔区序列和右半位点的DNA重组酶靶序列,并且其中左半位点不是右半位点的回文序列。

[0194] 在一些实施方案中,演化的重组酶识别包含天然存在的序列的DNA重组酶靶序列。在一些实施方案中,演化的重组酶识别包含在哺乳动物的基因组中的DNA重组酶靶序列。在一些实施方案中,演化的重组酶识别包含在人的基因组中的DNA重组酶靶序列。在一些实施方案中,演化的重组酶识别在哺乳动物的基因组中仅发生一次的DNA重组酶靶序列。在一些实施方案中,演化的重组酶识别哺乳动物的基因组中的DNA重组酶靶序列,其与基因组中的任何其他位点相差至少1、至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14或至少15个核苷酸。在一些实施方案中,演化的重组酶识别位于安全港基因组基因座中的DNA重组酶靶序列。在一些实施方案中,安全港基因组基因座是Rosa26基因座。在一些实施方案中,演化的重组酶识别位于与疾病或病症相关的基因组基因座中的DNA重组酶靶序列。

[0195] 在某些实施方案中,演化的重组酶可以靶向人基因组的Rosa基因座中的位点(例如36C6)。此类重组酶的非限制性组可以在例如,2017年1月26日公开的国际PCT公开WO 2017/015545A1,题为“Evolution of Site Specific Recombinases,”中找到,其为此目的通过引用并入本文。在一些实施方案中,与野生型重组酶的序列相比,演化的重组酶的氨基酸序列包含至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14或至少15个突变。编码36C6的核苷酸序列以粗体显示如下;编码GGG接头的那些以斜体显示;编码dCas9接头的那些为黑色;编码FLAG

[0196] 标签和NLS的那些分别用下划线和小写字母表示。

[0197] dCas9-36C6(核苷酸)(SEQ ID NO:765)

[0198]

ATGTCCAACCTCCTTACCGTCCACCAGAATCTCCCTGCCCTTCCGGTGGATGC
CACCTCTGATGAAGTGCGAAAAACCTGATGGATATGTTTCGCGATAGGCAAG
CTTTTTCTGAACACACGTGGAAGATGCTCCTGTGAGTGTGTAGAAGCTGGGCA
GCTTGGTGCAAGTTGAACAACCGAAAATGGTTTCCTGCCGAACCCGAAGATGT
GAGAGACTACCTCCTTACCTGCAGGCTCGAGGGCTCGCCGTGAAAAAATCC
AACAACTTGGGTCAGCTCAACATGCTGCACAGGAGATCTGGGCTGCCCGG
CCGAGTGACTCTAATGCCGTTAGTCTCGTAATGCGGCGCATTGCAAAGAGAA
TGTGGATGCTGGAGAACGGGCGAAACAGGCACTGGCTTTTGAACGGACCGAC
TTCGATCAGGTGCGGAGTCTTATGGAGAATAGTGACAGATGCCAGGACATTCG
GAACCTTGCATTCCTGGGTATCGCGTATAAATACCCTGCTGAGAATCGCTGAGA
TCGCCAGAATCAGGGTAAAGGATATTTCTCGAACGGACGGGGGACGGATGTTG
ATTCATATCGGTGCGACTAAAACACTTGTGAGTACCGCCGGGGTAGAGAAAGC
CCTGAGCCTTGGAGTTACTAAACTGGTGGAGCGGTGGATTAGCGTGTCCGGCG
TGGCGGATGACCCAAACAATTACTTGTTTTGTAGGGTGC GGAAAAATGGTGT
GCCGCTCCATCCGCTACCTCACAGTTGAGTACACGCGCGTTGGAGGGGATTTT
CGAAGCCACACATCGCTTGATCTACGGCGCCAAGGACGATTCAGGCCAGCGAT
ATCTTGCCCTGGAGCGGGCATAAGTGC CCGGGTGGGTGCCGCCGAGACATGGC
AAGGGCTGGCGTGTCAATTCCTGAAATCATGCAGGCCGGCGGGTGGACCAAC
GTGAACATTGTGATGAACTATATCCGGAACCTGGATAGCGAGACCGGAGCAAT
GGTCAGACTGCTTGAGGATGGCGACGGTGGATCCGGAGGGTCCGGAGGTAGTGGC
GGCAGCGGTGGTTCAGGTGGCAGCGGAGGGTCCAGGAGGCTCTGATAAAAAGTATTCT
ATTGGTTTAGCTATCGGCACTAATTCCGTTGGATGGGCTGTCATAACCGATGAATAC
AAAGTACCTTCAAAGAAATTTAAGGTGTTGGGGAACACAGACCGTCATTCGATTAA
AAAGAATCTTATCGGTGCCCTCCTATTCGATAGTGGCGAAACGGCAGAGGCGACTC
GCCTGAAACGAACCGCTCGGAGAAGGTATACACGTCGCAAGAACCGAATATGTTAC
TTACAAGAAATTTTTAGCAATGAGATGGCCAAAGTTGACGATTCTTTCTTCCACCGT
TTGGAAGAGTCCCTTCCTTGTGCAAGAGGACAAGAAACATGAACGGCACCCCATCTT
TGGAACATAGTAGATGAGGTGGCATATCATGAAAAGTACCCAACGATTTATCACC
TCAGAAAAAAGCTAGTTGACTCAACTGATAAAGCGGACCTGAGGTTAATCTACTTG
GCTCTTGCCCATATGATAAAGTTCCGTGGGCACTTTCTCATTGAGGGTGTCTAAAT
CCGGACAACCTCGGATGTGACAAACTGTTTCATCCAGTTAGTACAAACCTATAATCA
GTTGTTTGAAGAGAACCCTATAAATGCAAGTGGCGTGGATGCGAAGGCTATTCTTA
GCGCCCGCCTCTCTAAATCCCGACGGCTAGAAAACCTGATCGCACAATTACCCGGA
GAGAAGAAAAATGGGTTGTTTCGGTAACCTTATAGCGCTCTCACTAGGCCTGACACC
AAATTTTAAGTCGAACTTCGACTTAGCTGAAGATGCCAAATTGCAGCTTAGTAAGG
ACACGTACGATGACGATCTCGACAATCTACTGGCACAATTGGAGATCAGTATGCG
GACTTATTTTTGGCTGCCAAAACCTTAGCGATGCAATCCTCCTATCTGACATACTG
AGAGTTAATACTGAGATTACCAAGGCGCCGTTATCCGCTTCAATGATCAAAAAGGTA
CGATGAACATACCAAGACTTGACACTTCTCAAGGCCCTAGTCCGTCAGCAACTGC
CTGAGAAATATAAGGAAATATTCTTTGATCAGTCGAAAAACGGGTACGCAGGTTAT
ATTGACGGCGGAGCGAGTCAAGAGGAATTCTACAAGTTTATCAAACCCATATTAGA
GAAGATGGATGGGACGGAAGAGTTGCTTGTA AACTCAATCGCGAAGATCTACTGC
GAAAGCAGCGGACTTTCGACAACGGTAGCATTCCACATCAAATCCACTTAGGCGAA
TTGCATGCTATACTTAGAAGGCAGGAGGATTTTTATCCGTTCTCAAAGACAATCGT
GAAAAGATTGAGAAAATCCTAACCTTTCGCATACCTTACTATGTGGGACCCCTGGC
CCGAGGGAACCTCTCGGTTTCGCATGGATGACAAGAAAGTCCGAAGAAACGATTACTC
CATGGAATTTTGAGGAAGTTGTGATAAAGGTGCGTCAGCTCAATCGTTCATCGAG

[0199]

AGGATGACCAACTTTGACAAGAATTTACCGAACGAAAAAGTATTGCCTAAGCACAG
TTTACTTTACGAGTATTTACAGTGTACAATGAACTCACGAAAGTTAAGTATGTCAC
TGAGGGCATGCGTAAACCCGCCTTTCTAAGCGGAGAACAGAAGAAAGCAATAGTA
GATCTGTTATTCAAGACCAACCGCAAAGTGACAGTTAAGCAATTGAAAGAGGACTA
CTTTAAGAAAATTGAATGCTTCGATTCTGTCGAGATCTCCGGGGTAGAAGATCGATT
TAATGCGTCACTTGGTACGTATCATGACCTCCTAAAGATAATTAAGATAAAGGACTT
CCTGGATAACGAAGAGAATGAAGATATCTTAGAAGATATAGTGTTGACTCTTACCC
TCTTTGAAGATCGGGAAATGATTGAGGAAAGACTAAAAACATACGCTCACCTGTTC
GACGATAAGGTTATGAAACAGTTAAAGAGGGCGTCGCTATACGGGCTGGGGACGATT
GTCGCGGAAACTTATCAACGGGATAAGAGACAAGCAAAGTGGTAAAACCTATTCTCG
ATTTTCTAAAGAGCGACGGCTTCGCCAATAGGAACTTTATGCAGCTGATCCATGATG
ACTCTTTAACCTTCAAAGAGGATATACAAAAGGCACAGGTTTCCGGACAAGGGGAC
TCATTGCACGAACATATTGCGAATCTTGCTGGTTCGCCAGCCATCAAAAAGGGCAT
ACTCCAGACAGTCAAAGTAGTGGATGAGCTAGTTAAGGTCATGGGACGTCACAAAC
CGGAAAACATTGTAATCGAGATGGCACGCGAAAATCAAACGACTCAGAAGGGGCA
AAAAACAGTCGAGAGCGGATGAAGAGAATAGAAGAGGGTATTAAGAAGTGGGC
AGCCAGATCTTAAAGGAGCATCCTGTGGAAAATACCCAATTGCAGAACGAGAACT
TTACCTCTATTACCTACAAAATGGAAGGGACATGTATGTTGATCAGGAACTGGACA
TAAACCGTTTATCTGATTACGACGTCGATGCCATTGTACCCCAATCCTTTTTGAAGG
ACGATTCAATCGACAATAAAGTGCTTACACGCTCGGATAAGAACCGAGGGAAAAGT
GACAATGTTCCAAGCGAGGAAGTCGTAAGAAAATGAAGAAGTATTGGCGGCAGC
TCCTAAATGCGAAACTGATAACGCAAAGAAAGTTCGATAACTTAACTAAAGCTGAG
AGGGGTGGCTTGTCTGAAGTTGACAAGGCCGGATTTATTAACGTCAGCTCGTGGA
AACCCGCCAAATCACAAGCATGTTGCACAGATACTAGATTCCCGAATGAATACGA
AATACGACGAGAACGATAAGCTGATTTCGGGAAGTCAAAGTAATCACTTTAAAGTCA
AAATTGGTGTTCGGACTTCAGAAAGGATTTTCAATTCTATAAAGTTAGGGAGATAAA
TAACTACCACCATGCGCACGACGCTTATCTTAATGCCGTCGTAGGGACCGCACTCAT
TAAGAAATACCCGAAGCTAGAAAGTGAGTTTGTGTATGGTGATTACAAAGTTTATG
ACGTCCGTAAGATGATCGCGAAAAGCGAACAGGAGATAGGCAAGGCTACAGCCAA
ATACTTCTTTTATTCTAACATTATGAATTTCTTTAAGACGGAAATCACTCTGGCAAA
CGGAGAGATACGCAAACGACCTTTAATTGAAACCAATGGGGAGACAGGTGAAATC
GTATGGGATAAGGGCCGGGACTTCGCGACGGTGAGAAAAGTTTTGTCCATGCCCA
AGTCAACATAGTAAAGAAAAGTGAAGTGCAGACCGGAGGGTTTTCAAAGGAATCG
ATTCTTCCAAAAGGAATAGTGATAAGCTCATCGCTCGTAAAAAGGACTGGGACCC
GAAAAGTACGGTGGCTTCGATAGCCCTACAGTTGCCTATTCTGTCTAGTAGTGGC
AAAAGTTGAGAAGGGAAAATCCAAGAACTGAAGTCAGTCAAAGAATTATTGGGG
ATAACGATTATGGAGCGCTCGTCTTTTGAAAAGAACCCCATCGACTTCCTTGAGGCG
AAAGGTTACAAGGAAGTAAAAAAGGATCTCATAATTAAGTACCAAAGTATAGTCT
GTTTGAGTTAGAAAATGGCCGAAAACGGATGTTGGCTAGCGCCGGAGAGCTTCAA
AGGGGAACGAACTCGCACTACCGTCTAAATACGTGAATTTCTGTATTTAGCGTCCC
ATTACGAGAAGTTGAAAGGTTACCTGAAGATAACGAACAGAAGCAACTTTTTGTT
GAGCAGCACAAACATTATCTCGACGAAATCATAGAGCAAATTCGGAATTCAGTAA
GAGAGTCATCCTAGCTGATGCCAATCTGGACAAAGTATTAAGCGCATACAACAAGC
ACAGGGATAAACCATACGTGAGCAGGCGGAAAATATTATCCATTTGTTTACTCTT
ACCAACCTCGGCGCTCCAGCCGATTCAAGTATTTGACACAACGATAGATCGCAA
ACGATACACTTCTACCAAGGAGGTGCTAGACGCGACACTGATTCACCAATCCATCA
CGGGATTATATGAAACTCGGATAGATTTGTCACAGCTTGGGGGTGACGGTGGCTCC
GATTATAAGGATGATGACGACAAGGGAGGTTCCccaaagaagaaaaggaaggtcTGA

[0200] dCas9-36C6 (氨基酸) (SEQ ID NO: 766)

[0201]

MSNLLTVHQNLPALPVDATSDEVRKNLMDMFRDRQAFSEHTWKMLLSVCRSWA
 AWCKLNNRKWFPAEPEDVRDYLLYLQARGLAVKTIQQHLGQLNMLHRRSGLPRP
 SDSNAVSLVMRRIRKENVDAGERAKQALAFERTDFDQVRSLMENS DRCQDIRNLA
 FLGIAYNTLLRIAIEIARIRVKDISRTDGG RMLIHIGRTKTLVSTAGVEKALS LGVTK
 LVERWISVSGVADDPNNYLF CRVRKNGVAAPSATSQ LSTRALEGIFEATHRLIYGA
 KDDSGQRYLAWSGHSARVGAARDMARAGVSIPEIMQAGGWTNVNIVMNYIRNLD
 SETGAMVRLLEDG DGGSGGSGGSGGSGGSGGSGGSGGSDKKYSIGLAIGTNSVGWAVI
 TDEYKVP SKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRIC
 YLQEIFSNEMAKVDDSFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRK
 KLV DSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQLFEEN
 PINASGVDAKAILSARLSKSRLENLIAQLPGEKKNGLFGNLIASLGLTPNFKSNFDLAE
 DAKLQLSKD TYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSA
 SMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPI
 LEKMDGTEELLVKNLREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFY PFLKDNRE
 KIEKILTRIPYYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNF
 DKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTN
 RKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGT YHDLKIIKDKDFLDNEENEDIL
 EDIVLTLTLFEDREMIEERLKYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQS
 GKTILDFLKS DGFANRNFMLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAICK
 GILQTVKVVDELVKVMGRHKPENIVIAMARENQTTQKGQKNSRERMKRIEEGIKELGS
 QILKEHPVENTQLQNEKLYLYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSI
 DNKVLTRSDKNRGKSDNVPSEE VVKMKKNYWRQLLNAKLITQRKFDNLTKAERGGLS
 ELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKS KLVSDFRK
 DFQFYKVREINNYHHAHDAYLNAVVG TALIKKYPKLESEFVYGDYKVYDVRKMIAS
 EQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRK
 VLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPK KYGGFDSPTVAYSVL
 VVAKVEK GKSKKLKSVKELLGITIMERS SFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLF
 ELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGSPEDNEQKQLFVEQH
 KHYLDEIIEQISEFSKR VILADANLDKVL SAYNKHRDKPIREQAENIIHLFTLNLGAPAA
 FKYFDTTIDRKRYTSTKEVLDATLIHQ SITGLYETRIDLSQLGGDGGSDYKDDDDKGGSp
 kkkrv 终止

[0202] 本公开的一些方面提供了演化的重组酶(例如Cre重组酶),其包含与本文讨论的重组酶序列(例如Cre重组酶)的序列至少70%、至少80%、至少90%、至少95%或至少97%相同的氨基酸序列,其中与本文讨论的重组酶(例如Cre重组酶)的序列相比,重组酶(例如Cre重组酶)的氨基酸序列包含至少一个突变,并且其中重组酶(例如Cre重组酶)识别与规范LoxP位点5'-ATAACTTCGTATA GCATACAT TATACGAAGTTAT-3'(SEQ ID NO:739)相差至少一个核苷酸的DNA重组酶靶序列。

[0203] 在一些实施方案中,与本文讨论的重组酶(例如Cre重组酶)序列的序列相比,演化的重组酶(例如Cre重组酶)的氨基酸序列包含至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14或至少15个突变,并且识别与规范靶位点(例如LoxP位点)相差至少1、至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14或至少15个核苷酸的DNA重组酶靶序列。

[0204] 在一些实施方案中,演化的Cre重组酶识别包含左半位点、间隔区序列和右半位点的DNA重组酶靶序列,并且其中左半位点不是右半位点的回文序列。在一些实施方案中,演化的Cre重组酶识别包含天然存在的序列的DNA重组酶靶序列。在一些实施方案中,演化的

Cre重组酶识别包含在哺乳动物的基因组中的DNA重组酶靶序列。

[0205] 在一些实施方案中,演化的Cre重组酶识别包含在人的基因组中的DNA重组酶靶序列。在一些实施方案中,演化的Cre重组酶识别在哺乳动物的基因组中仅包含一次的DNA重组酶靶序列。在一些实施方案中,演化的Cre重组酶识别哺乳动物的基因组中的DNA重组酶靶序列,其与基因组中的任何其他位点相差至少1、至少2、至少3、至少4、至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14或至少15个核苷酸。在一些实施方案中,演化的Cre重组酶识别位于安全港基因组基因座中的DNA重组酶靶序列。在一些实施方案中,安全港基因组基因座是Rosa26基因座。在一些实施方案中,演化的Cre重组酶识别位于与疾病或病症相关的基因组基因座中的DNA重组酶靶序列。

[0206] 与本发明方法和组合物一起使用的另外的演化的重组酶(及其制备方法)可以在例如美国专利申请号15/216,844中找到,其通过引用并入本文。

[0207] 对于提供重组酶催化域或演化的重组酶催化域两者,另外的合适的重组酶对于本领域技术人员而言将是显而易见的,并且此类合适的重组酶包括但不限于描述于Hirano et al., Site-specific recombinases as tools for heterologous gene integration. *Appl Microbiol Biotechnol.* 2011 Oct; 92(2):227-39; Fogg et al., New applications for phage integrases. *J Mol Biol.* 2014 Jul 29; 426(15):2703; Brown et al., Serine recombinases as tools for genome engineering. *Methods.* 2011 Apr; 53(4):372-9; Smith et al., Site-specific recombination by phiC31 integrase and other large serine recombinases. *Biochem Soc Trans.* 2010 Apr; 38(2):388-94; Grindley et al., Mechanisms of site-specific recombination. *Annu Rev Biochem.* 2006; 75:567-605; Smith et al., Diversity in the serine recombinases. *Mol Microbiol.* 2002 Apr; 44(2):299-307; Grainge et al., The integrase family of recombinase: organization and function of the active site. *Mol Microbiol.* 1999 Aug; 33(3):449-56; Gopaul et al., Structure and mechanism in site-specific recombination. *Curr Opin Struct Biol.* 1999 Feb; 9(1):14-20; Cox et al., Conditional gene expression in the mouse inner ear using Cre-loxP. *J Assoc Res Otolaryngol.* 2012 Jun; 13(3):295-322; Birling et al., Site-specific recombinases for manipulation of the mouse genome. *Methods Mol Biol.* 2009; 561:245-63; 和 Mishina M, Sakimura K. Conditional gene targeting on the pure C57BL/6 genetic background. *Neurosci Res.* 2007 Jun; 58(2):105-12中的那些;其每一个的全部内容通过引用并入本文。

[0208] 融合蛋白的结构

[0209] 本公开的融合蛋白可以是本文所述的元件的任何组合和顺序。示例性融合蛋白包括但不限于任何以下结构: NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[任选的亲亲和标签]-COOH。在另一个实施方案中,融合蛋白具有结构NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲亲和标签]-COOH。在另一个实施方案中,融合蛋白具有结构NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-[任

选的接头序列)-[亲和标签]-COOH。在另一个实施方案中,融合蛋白具有结构NH₂-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[NLS域]-[接头序列]-[亲和标签]-COOH。

[0210] 在另一个实施方案中,融合蛋白具有结构NH₂-[重组酶催化域]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[亲和标签]-COOH、NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH、NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH、NH₂-[亲和标签]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[任选的接头序列]-[NLS域]-COOH、NH₂-[亲和标签]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-COOH或NH₂-[亲和标签]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-COOH。

[0211] 在另一个实施方案中,融合蛋白具有结构:NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[亲和标签]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-[接头序列]-[NLS域]-[接头序列]-[亲和标签]-COOH。

[0212] 在另一个实施方案中,融合蛋白具有结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接

头序列)-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[接头序列]-[NLS域]-[接头序列]-[亲和标签]-COOH。

[0213] 在另一个实施方案中,融合蛋白具有结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[接头序列]-[NLS域]-[接头序列]-[亲和标签]-COOH。

[0214] 在另一个实施方案中,融合蛋白具有结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[接头序列]-[NLS域]-[接头序列]-[亲和标签]-COOH。

[0215] 在另一个实施方案中,融合蛋白具有结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[任选的亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[任选的

亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[亲和标签]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-[接头序列]-[NLS域]-[接头序列]-[亲和标签]-COOH。

[0216] 在一个实施方案中,融合蛋白具有结构NH₂-[任选的亲和标签]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[任选的亲和标签]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[亲和标签]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[亲和标签]-[接头序列]-[NLS域]-[接头序列]-[重组酶催化域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-COOH。

[0217] 在一个实施方案中,融合蛋白具有结构NH₂-[任选的亲和标签]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[任选的亲和标签]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[亲和标签]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-COOH。在一个实施方案中,融合蛋白包含结构NH₂-[亲和标签]-[接头序列]-[NLS域]-[接头序列]-[引导核苷酸序列-可编程DNA结合蛋白域]-[接头序列]-[重组酶催化域]-COOH。

[0218] 在另一个实施方案中,融合蛋白具有结构NH₂-[任选的亲和标签]-[任选的接头序列]-[任选的NLS域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[任选的亲和标签]-[任选的接头序列]-[NLS域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[亲和标签]-[接头序列]-[NLS域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[任选的接头序列]-[重组酶催化域]-[任选的接头序列]-[二分叉或环状置换的引导

头序列]-[NLS域]-[任选的接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-COOH。在另一个实施方案中,融合蛋白包含结构NH₂-[亲和标签]-[接头序列]-[NLS域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的N端部分]-[接头序列]-[重组酶催化域]-[接头序列]-[二分叉或环状置换的引导核苷酸序列-可编程DNA结合蛋白域的C端部分]-COOH。

[0222] 融合蛋白可以进一步包含一个或多个亲和标签。本文提供的合适的亲和标签包括但不限于生物素羧化酶载体蛋白(BCCP)标签、myc标签、钙调蛋白标签、FLAG标签、血凝素(HA)标签、多组氨酸标签(也称为组氨酸标签或His标签)、多精氨酸(多Arg)标签、麦芽糖结合蛋白(MBP)-标签、nus标签、谷胱甘肽-S-转移酶(GST)标签、绿色荧光蛋白(GFP)标签、硫氧还蛋白标签、S标签、Softag(例如,Softag 1、Softag 3)、strep标签、生物素连接酶标签、FlAsH标签、V5标签和SBP标签。另外的合适的序列对于本领域技术人员而言将是显而易见的。FLAG标签可以具有序列PKKKRKV(SEQ ID NO:702)。一个或多个亲和标签经由一个或多个第三接头与引导核苷酸序列-可编程DNA结合蛋白域、重组酶催化域或NLS域结合。第三接头可以是本文所述的任何肽接头。例如,第三接头可以是肽接头。

[0223] 作为非限制性组的实例,第三接头可以包含XTEN接头SGSETPGTSESATPES(SEQ ID NO:7)、SGSETPGTSESA(SEQ ID NO:8)或SGSETPGTSESATPEGGSGGS(SEQ ID NO:9),包含三肽GGG的一个或多个重复的氨基酸序列,或任何以下氨基酸序列:VPFLLEPDNINGKTC(SEQ ID NO:10)、GSAGSAAGSGEF(SEQ ID NO:11)、SIVAQLSRPDPA(SEQ ID NO:12)、MKIIEQLPSA(SEQ ID NO:13)、VRHKLKRVGS(SEQ ID NO:14)、GHGTGSTGSGSS(SEQ ID NO:15)、MSRPDPA(SEQ ID NO:16)或GGSM(SEQ ID NO:17)。在某些实施方案中,第三接头包含三肽GGG的一个或多个重复。在实施方案中,第三接头包含三肽GGG的一个至五个重复。在实施方案中,第三接头包含三肽GGG的一个重复。在具体的实施方案中,第三接头具有序列GGG。

[0224] 第三接头也可以是非肽接头。在某些实施方案中,非肽接头包含聚乙二醇(PEG)、聚丙二醇(PPG)、共聚(乙烯/丙烯)二醇、聚氧乙烯(POE)、聚氨酯、聚磷腈、多糖、右旋糖酐、聚乙烯醇、聚乙烯吡咯烷酮、聚乙烯乙醚、聚丙烯酰胺、聚丙烯酸酯、聚氰基丙烯酸酯、脂质聚合物、甲壳质、透明质酸、肝素或烷基接头。在其他实施方案中,烷基接头具有式—NH—(CH₂)_s—C(O)—,其中s可以是1和100之间的任何整数,包括端点。在具体的实施方案中,s是1和20之间的任何整数,包括端点。

[0225] 本公开的融合蛋白与SEQ ID NO:185的氨基酸1-1544中所示的氨基酸序列(其与SEQ ID NO:719中所示的序列相同)具有大于90%、95%或99%的序列一致性。

[0226] MLIGYVRVSTNDQNTDLQRNALVCAGCEQIFEDKLSGTRTDRPGLKRALKRLQKGDTLVWKLDRDLGRSMKHLISLVGELRERGINFRSLTDSIDTSSPMGRFFFYVMGALAEMERELI IERTMAGLAAARNKRRFRPPKGGSGGSGGSGGSGGSGGSGGSDKKYSIGLAIGTNSVGVAVITDEYKVPSSKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEKAVDSSFFHRLEESFLVEEDKKHERHP IFGNIVDEVAYHEK YPTIYHLRKKLVSDTKADLR LIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQLFEENPINASGVD AKAILSARLSKSRRENLI AQLPGEKKNLFGNLI ALSLGLTPNFKSNFDLAEDAKLQLSKD TYDDDLDNLLAQIG DQYADLFLAAKNLSDA ILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGY AGYIDGGASQEEFYKFIKPILEKMDGTEELLVKNLREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYFPFLKD

NREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEVVDKGASAQSFIERMTNFDKNLPNEKVLPHK
SLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRF
NASLGTYHDLKIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKYAHLFDDKVMKQLKRRRYTGWRSL
RKLINGIRDKQSGKTILDFLKSDFANRNFMLIHDDSLTFKEDIQKAQVSGQDLSLHEHIANLAGSPAIIKKGILQ
TVKVVDLVKVMGRHKPENIVIAMARENQTTQKGQKNSRERMKRIEEG IKELGSQILKEHPVENTQLQNEKLYLY
LQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKL
ITQRKFDNLTKAERGGSELKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSCLVSDFR
KDFQFYKREINNYHHAHDAYLNAVVGTAIIKKYPKLESEFVYGDYKVVYDVRKMIKSEQEIGKATAKYFFYSNIM
NFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVL SMPQVNIKKTEVQTGGFSKESILPKRNSDKL
IARKKDWDPKKYGGFDSPTVAYSVLVAKVEKGKSKKLKSVKELGITIMERSSEFKNPIDFLEAKGYKEVKKDLI
IKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGKSPEDNEQKQLFVEQHKHYLDEIIE
QISEFSKRVILADANLDKVL SAYNKHRDKPIREQAENI IHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVL DATL
IHQSITGLYETRIDLSQLGGDGGSDYKDDDDK终止 (SEQ ID NO: 719)

[0227] 在二聚化(或多聚化)的蛋白质,诸如例如核酸酶失活的Cas9(或Cas9gRNA结合域)与重组酶(或重组酶的催化域)之间的融合的背景下,靶位点通常包含左半位点(由一种蛋白质结合)、右半位点(由第二种蛋白质结合),以及在其中进行重组的半位点之间的间隔区序列。在一些实施方案中,重组左半位点或右半位点(而非间隔区序列)。在其他实施方案中,重组间隔区序列。该结构([左半位点]-[间隔区序列]-[右半位点])在本文中称为LSR结构。在一些实施方案中,左半位点和/或右半位点对应于RNA引导的靶位点(例如,Cas9靶位点)。在一些实施方案中,半位点中的任一个或两个比例如Cas9靶向的典型区域更短或更长,例如短于或长于20个核苷酸。在一些实施方案中,左半位点和右半位点包含不同的核酸序列。在一些实施方案中,间隔区序列长度为至少5、至少6、至少7、至少8、至少9、至少10、至少11、至少12、至少13、至少14、至少15、至少16、至少17、至少18、至少19、至少20、至少25、至少30、至少35、至少40、至少45、至少50、至少60、至少70、至少80、至少90、至少100、至少125、至少150、至少175、至少200或至少250bp。在一些实施方案中,间隔区序列为约15bp和约25bp之间长。在一些实施方案中,间隔区序列为约15bp长。在一些实施方案中,间隔区序列为约25bp长。

实施例

[0228] 实施例1:可编程Cas9-丝氨酸重组酶融合蛋白,其在哺乳动物细胞中的DNA序列上起作用

[0229] 材料和方法

[0230] 寡核苷酸和PCR

[0231] 所有寡核苷酸购自Integrated DNA Technologies (IDT,Coralville,CA)并列于表1-5中。除非另有说明,否则酶购自New England Biolabs (Ipswich,MA)。Plasmid Safe ATP依赖性DNA酶购自Epicenter (Madison,WI)。将所有组装的载体转化到One Shot Mach1-T1噬菌体抗性化学感受态细胞 (Fisher Scientific,Waltham,MA)。除非另有说明,否则所有PCR反应均使用Q5热启动高保真 (Hot Start High-Fidelity) 2X Master Mix进行。Phusion聚合酶用于环状聚合酶延伸克隆 (CPEC) 组装。

[0232] 表1:用于gRNA构建的寡核苷酸

[0233]

寡核苷酸名称	序列	SEQ ID NO:
R.pHU6.TSS(-1).univ	GGTGTTCGTCCTTTCCACAAG	20
F.非靶	GCACACTAGTTAGGGATAACAGTTTTAG AGCTAGAAATAGC	21
F.Chr10-1	GCCCATGACCCTTCTCCTCTGTTTTAGAG CTAGAAATAGC	22
F.Chr10-1-rev	GCTCAGGGCCTGTGATGGGAGGTTTTAG AGCTAGAAATAGC	23
F.Chr10-2	GGCCCATGACCCTTCTCCTCGTTTTAGAG CTAGAAATAGC	24
F.Chr10-2rev	GCCTCAGGGCCTGTGATGGGAGTTTTAG AGCTAGAAATAGC	25
F.Centromere_Chr_1_5_19-g RNA-for	GACTTGAAACACTCTTTTTCGTTTTAGAG CTAGAAATAGC	26
F.Centromere_Chr_1_5_19-g RNA-rev	GAGTTGAAGACACACAACACAGTTTTAG AGCTAGAAATAGC	27
F.Ch5_155183064-gRNA-for	GGAACTCATGTGATTAAGTGGTTTTAGA GCTAGAAATAGC	28
F.Ch5_155183064-gRNA-rev- 1	GTCTACCTCTCATGAGCCGGTGGTTTTAGA GCTAGAAATAGC	29
F.Ch5_169395198-gRNA-for	GTTTCCCGCAGGATGTGGGATGTTTTAG AGCTAGAAATAGC	30
F.Ch5_169395198-gRNA-rev	GCCTGGGGATTTATGTTCTTAGTTTTAGA GCTAGAAATAGC	31
F.Ch12_62418577-gRNA-for	GAAATAGCACAATGAATGGAAGTTTTAG AGCTAGAAATAGC	32
F.Ch12_62418577-gRNA-rev	GACTTTTTGGGGGAGAGGGAGGTTTTAG AGCTAGAAATAGC	33
F.Ch13_102010574-gRNA-for	GGAGACTTAAGTCCAAAACCGTTTTAGA GCTAGAAATAGC	34
F.Ch13_102010574-gRNA-re v	GTCAGCTATGATCACTTCCCTGTTTTAGA GCTAGAAATAGC	35

[0234] 表2:用于报告物构建的寡核苷酸和gBlock

[0235]

构建体名称	序列	SEQ ID NO:
1-0bp-for	TCGTCTCGGCGTCCCAATTTCCCAAACAGAG GTCTGTAAACCGAGGTGAGACGG	36

[0236]

1-0bp-rev	CCGTCTCACCTCGGTTTACAGACCTCTGTTTGG GAAAATTGGGGACGCCGAGACGA	37
1-1bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTtCTGTAAACCGAGGTGAGACGG	38
1-1bp-rev	CCGTCTCACCTCGGTTTACAGaACCTCTGTTTGG GGAAAATTGGGGACGCCGAGACGA	39
1-2bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTatCTGTAAACCGAGGTGAGACGG	40
1-2bp-rev	CCGTCTCACCTCGGTTTACAGatACCTCTGTTTGG GGAAAATTGGGGACGCCGAGACGA	41
1-3bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTaatCTGTAAACCGAGGTGAGACGG	42
1-3bp-rev	CCGTCTCACCTCGGTTTACAGattACCTCTGTTTGG GGAAAATTGGGGACGCCGAGACGA	43
1-4bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTaaatCTGTAAACCGAGGTGAGACGG	44
1-4bp-rev	CCGTCTCACCTCGGTTTACAGatttACCTCTGTTT GGGAAAATTGGGGACGCCGAGACGA	45
1-5bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTgaaatCTGTAAACCGAGGTGAGACGG	46
1-5bp-rev	CCGTCTCACCTCGGTTTACAGatttcACCTCTGTTT GGGAAAATTGGGGACGCCGAGACGA	47
1-6bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTcgaatCTGTAAACCGAGGTGAGACGG	48
1-6bp-rev	CCGTCTCACCTCGGTTTACAGatttcgACCTCTGTT TGGGAAAATTGGGGACGCCGAGACGA	49
1-7bp-for	TCGTCTCGGCGTCCCAATTTTCCCAAACAGAG GTtcgaaatCTGTAAACCGAGGTGAGACGG	50
1-7bp-rev	CCGTCTCACCTCGGTTTACAGatttgaACCTCTGT TTGGGAAAATTGGGGACGCCGAGACGA	51
2-0bp-for	TCGTCTCGGAGGTTTTGGAACCTCTGTTTGGGA AAATTGGGGAGTCTGAGACGG	52
2-0bp-rev	CCGTCTCAGACTCCCAATTTTCCCAAACAGAG GTTCCAAAACCTCCGAGACGA	53
2-1bp-for	TCGTCTCGGAGGTTTTGGACACCTCTGTTTGGG AAAATTGGGGAGTCTGAGACGG	54
2-1bp-rev	CCGTCTCAGACTCCCAATTTTCCCAAACAGAG GTGTCCAAAACCTCCGAGACGA	55
2-2bp-for	TCGTCTCGGAGGTTTTGGACTACCTCTGTTTGG GAAAATTGGGGAGTCTGAGACGG	56
2-2bp-rev	CCGTCTCAGACTCCCAATTTTCCCAAACAGAG GTAGTCCAAAACCTCCGAGACGA	57
2-3bp-for	TCGTCTCGGAGGTTTTGGACTTACCTCTGTTTGG GGAAAATTGGGGAGTCTGAGACGG	58
2-3bp-rev	CCGTCTCAGACTCCCAATTTTCCCAAACAGAG GTAAGTCCAAAACCTCCGAGACGA	59
2-4bp-for	TCGTCTCGGAGGTTTTGGACTTAACTCTGTTT GGGAAAATTGGGGAGTCTGAGACGG	60
2-4bp-rev	CCGTCTCAGACTCCCAATTTTCCCAAACAGAG GTTAAGTCCAAAACCTCCGAGACGA	61

[0237]

2-5bp-for	TCGTCTCGGAGGTTTTGGACTTAGACCTCTGTT TGGGAAAATTGGGGAGTCTGAGACGG	62
2-5bp-rev	CCGTCTCAGACTCCCCAATTTCCCAAACAGAG GTCTAAGTCCAAAACCTCCGAGACGA	63
2-6bp-for	TCGTCTCGGAGGTTTTGGACTTAGCACCTCTGT TTGGGAAAATTGGGGAGTCTGAGACGG	64
2-6bp-rev	CCGTCTCAGACTCCCCAATTTCCCAAACAGAG GTGCTAAGTCCAAAACCTCCGAGACGA	65
2-7bp-for	TCGTCTCGGAGGTTTTGGACTTAGCTACCTCTG TTTGGGAAAATTGGGGAGTCTGAGACGG	66
2-7bp-rev	CCGTCTCAGACTCCCCAATTTCCCAAACAGAG GTAGCTAAGTCCAAAACCTCCGAGACGA	67
4-0bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTCTGTAAACCGATGAGACGG	68
4-0bp-rev	CCGTCTCATCGGTTTACAGACCTCTGTTTGGGA AAATTGGGGGTGCAGAGACGA	69
4-1bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTtCTGTAAACCGATGAGACGG	70
4-1bp-rev	CCGTCTCATCGGTTTACAGaACCTCTGTTTGGG AAAATTGGGGGTGCAGAGACGA	71
4-2bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTatCTGTAAACCGATGAGACGG	72
4-2bp-rev	CCGTCTCATCGGTTTACAGatACCTCTGTTTGGG AAAATTGGGGGTGCAGAGACGA	73
4-3bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTaatCTGTAAACCGATGAGACGG	74
4-3bp-rev	CCGTCTCATCGGTTTACAGattACCTCTGTTTGGG AAAATTGGGGGTGCAGAGACGA	75
4-4bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTaaatCTGTAAACCGATGAGACGG	76
4-4bp-rev	CCGTCTCATCGGTTTACAGatttACCTCTGTTTGG GAAAATTGGGGGTGCAGAGACGA	77
4-5bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTgaaatCTGTAAACCGATGAGACGG	78
4-5bp-rev	CCGTCTCATCGGTTTACAGatttcACCTCTGTTTGG GAAAATTGGGGGTGCAGAGACGA	79
4-6bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTcgaaatCTGTAAACCGATGAGACGG	80
4-6bp-rev	CCGTCTCATCGGTTTACAGatttcgACCTCTGTTTGG GGAAAATTGGGGGTGCAGAGACGA	81
4-7bp-for	TCGTCTCTGCACCCCCAATTTCCCAAACAGAG GTtcgaaatCTGTAAACCGATGAGACGG	82
4-7bp-rev	CCGTCTCATCGGTTTACAGatttcgaACCTCTGTTT GGGAAAATTGGGGGTGCAGAGACGA	83
5-0bp-for	TCGTCTCGCCGAGGTTTTGGAACTCTGTTTGG GAAAATTGGGGCTCGTGAGACGG	84
5-0bp-rev	CCGTCTCACGAGCCCCAATTTCCCAAACAGA GGTTCCAAAACCTCGGCGAGACGA	85
5-1bp-for	TCGTCTCGCCGAGGTTTTGGACACCTCTGTTTGG GGAAAATTGGGGCTCGTGAGACGG	86

[0238]

5-1bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTGTCCAAAACCTCGGCGAGACGA	87
5-2bp-for	TCGTCTCGCCGAGGTTTTGGACTACCTCTGTT GGGAAAATTGGGGCTCGTGAGACGG	88
5-2bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTAGTCCAAAACCTCGGCGAGACGA	89
5-3bp-for	TCGTCTCGCCGAGGTTTTGGACTTACCTCTGTT TGGGAAAATTGGGGCTCGTGAGACGG	90
5-3bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTAAGTCCAAAACCTCGGCGAGACGA	91
5-4bp-for	TCGTCTCGCCGAGGTTTTGGACTTAACCTCTGT TTGGGAAAATTGGGGCTCGTGAGACGG	92
5-4bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTTAAGTCCAAAACCTCGGCGAGACGA	93
5-5bp-for	TCGTCTCGCCGAGGTTTTGGACTTAGACCTCTG TTTGGGAAAATTGGGGCTCGTGAGACGG	94
5-5bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTCTAAGTCCAAAACCTCGGCGAGACGA	95
5-6bp-for	TCGTCTCGCCGAGGTTTTGGACTTAGCACCTCT GTTTGGGAAAATTGGGGCTCGTGAGACGG	96
5-6bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTGCTAAGTCCAAAACCTCGGCGAGACGA	97
5-7bp-for	TCGTCTCGCCGAGGTTTTGGACTTAGCTACCTC TGTTTGGGAAAATTGGGGCTCGTGAGACGG	98
5-7bp-rev	CCGTCTCACGAGCCCCAATTTTCCCAAACAGA GGTAGCTAAGTCCAAAACCTCGGCGAGACGA	99
1-Chr10--54913298-5 4913376-for	TCGTCTCGGCGTCCCCTCCCATCACAGGCCCTG AGGTTTAAGAGAAAACCTGAGACGG	100
1-Chr10-54913298-54 913376-rev	CCGTCTCAGGTTTTCTCTTAAACCTCAGGGCCT GTGATGGGAGGGGACGCCGAGACGA	101
2-Chr10--54913298-5 4913376-for	TCGTCTCGAACCATGGTTTTGTGGGCCAGGCC ATGACCCTTCTCCTCTGGGAGTCTGAGACGG	102
2-Chr10--54913298-5 4913376-rev	CCGTCTCAGACTCCCAGAGGAGAAGGGTCATG GGCCTGGCCACAAAACCATGGTTCGAGACGA	103
4-Chr10-54913298-54 913376-for	TCGTCTCTGCACCCCCTCCCATCACAGGCCCTG AGGTTTAAGAGAAAACCATTGAGACGG	104
4-Chr10-54913298-54 913376-rev	CCGTCTCAATGGTTTTCTCTTAAACCTCAGGGC CTGTGATGGGAGGGGTGCAGAGACGA	105
5-Chr10-54913298-54 913376-for	TCGTCTCGCCATGGTTTTGTGGGCCAGGCCCAT GACCCTTCTCCTCTGGGCTCGTGAGACGG	106
5-Chr10-54913298-54 913376-rev	CCGTCTCACGAGCCAGAGGAGAAGGGTCATG GGCCTGGCCACAAAACCATGGCGAGACGA	107
3-for	ATCCGTCTCCAGTCGAGTCGGATTTGATCTGAT CAAGAGACAG	108
3-rev	AACCGTCTCGGTGCGTTCGGATTTGATCCAGAC ATGATAAGATAC	109
Esp3I-插入-for	/Phos/CGCGTTGAGACGCTGCCATCCGTCTCGC	110
Esp3I-插入-rev	/Phos/TCGAGCGAGACGGATGGCAGCGTCTCAA	111
着 丝 粒 _Chr_1_5_19-1_2*	GTTGTTCTGCTCGGCGTCCTTGTGTTGTGTGTCT TCAACTCACAGAGTTAAACGATGCTTTACACA GAGTAGACTTGAAACACTCTTTTTCTGGAGTCT	112

[0239]

	GAGACGGTCTGTTTTGGTGTGATTAGTTAT	
着 丝 粒 _Chr_1_5_19-4_5*	GTTGGTCGTCTCTGCACCCTTGTGTTGTGTGTC TTCAACTCACAGAGTTAAACGATGCTTTACACA GAGTAGACTTGAAACACTCTTTTTCTGGCTCGT GAGACGGTCTGTTTTGGTGTGATTAGTTAT	113
Ch5_155183064-1551 83141-1_2*	GTTGTTCGTCTCGGCGTCCCACCGGCTCATGAG AGGTAGAGCTAAGGTCCAAACCTAGGTTTATC TGAGACCGGAACTCATGTGATTAAGTGTGGAG TCTGAGACGGTCTGTTTTGGTGTGATTAGTTA T	114
Ch5_155183064-1551 83141-4_5*	GTTGGTCGTCTCTGCACCCACCGGCTCATGAG AGGTAGAGCTAAGGTCCAAACCTAGGTTTATC TGAGACCGGAACTCATGTGATTAAGTGTGGCT CGTGAGACGGTCTGTTTTGGTGTGATTAGTTA T	115
Ch5_169395198-1693 95274-1_2*	GTTGTTCGTCTCGGCGTCCTTAAGAACATAAAT CCCCAGGAATTCACAGAAACCTTGGTTTGAGC TTTGGATTTCCCGCAGGATGTGGGATAGGAGT CTGAGACGGTCTGTTTTGGTGTGATTAGTTAT	116
Ch5_169395198-1693 95274-4_5*	GTTGGTCGTCTCTGCACCCTTAAGAACATAAAT CCCCAGGAATTCACAGAAACCTTGGTTTGAGC TTTGGATTTCCCGCAGGATGTGGGATAGGCTCG TGAGACGGTCTGTTTTGGTGTGATTAGTTAT	117
Ch12_62418577-6241 8652-1_2*	GTTGTTCGTCTCGGCGTCCACTCCCTCTCCCC AAAAAGTAAAGGTAGAAAACCAAGGTTTACAG GCAACAAATAGCACAATGAATGGAATGGAGTC TGAGACGGTCTGTTTTGGTGTGATTAGTTAT	118
Ch12_62418577-6241 8652-4_5*	GTTGGTCGTCTCTGCACCCACTCCCTCTCCCC AAAAAGTAAAGGTAGAAAACCAAGGTTTACAG GCAACAAATAGCACAATGAATGGAATGGCTCG TGAGACGGTCTGTTTTGGTGTGATTAGTTAT	119
chr13_102010574-102 010650-1_2*	GTTGTTCGTCTCGGCGTCCTAGGGAAGTGATCA TAGCTGAGTTTCTGGAAAAACCTAGGTTTAAA GTTGAGGAGACTTAAGTCCAAAACCTGGAGTC TGAGACGGTCTGTTTTGGTGTGATTAGTTAT	120
chr13_102010574-102 010650-4_5*	GTTGGTCGTCTCTGCACCCTAGGGAAGTGATCA TAGCTGAGTTTCTGGAAAAACCTAGGTTTAAA GTTGAGGAGACTTAAGTCCAAAACCTGGCTCG TGAGACGGTCTGTTTTGGTGTGATTAGTTAT	121
<p>将寡核苷酸序列退火以产生图1中所示的片段。名称对应于片段编号(1、2、4或5)，并然 后对应于将Cas9结合位点与gix核心位点分开的碱基对间隔区核苷酸的数量。</p> <p>*如支持材料文件中的方法中所述的双链gBlock。</p>		

[0240] 表3:用于recCas9构建的寡核苷酸

[0241]

寡核苷酸名称	序列	SEQ ID NO:
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[0242]

1GGS-连接-for_BamHI	TTCATCGGATCCGATAAAAAGTATTCTATTG GTTTAGCTATCGGCAC	122
5GGS-连接-for_BamHI	TTCATCGGATCCGGTGGTTCAGGTGGCAGC GGAG	123
8GGS-连接-for_BamHI	TTCATCGGATCCGGAGGGTCCGGAGGTAGT GGCGGCAGCGGTGGTTCAGGTGGCAGCGGA G	124
Cas9-rev-FLAG-NLS-AgeI	AATAACCGGTTTCAGACCTTCCTTTTCTTCTT TGGGGAACCTCCCTTGTCTCATCATCCTTA TAATCGGAGCCACCGTCACCCCAAGCTGT GACAAATC	125
1GGS-rev-BamHI	TGATAAGGATCCACCCTTTGGTGGTCTTCCA AACCGCC	126
2GGS-rev-BamH	TGATAAGGATCCACCGCTACCACCCTTTGG TGGTCTTC	127
Gin-for_NotI	AGATCCGCGGCCGCTAATAC	128
Esp3I-for-质粒	TTGAGTcgtctcTATACTCTTCCTTTTCAATAT TATTGAAGCATTATCAGGG	129
Esp3I-rev-质粒	CTGGAcgtctcACTGTCAGACCAAGTTTACTC ATATATACTTTAGATTG	130
spec-Esp3I-for	GGTGTGcgtctcTACAGTTATTTGCCGACTACC TTGGTGATCTCGC	131
spec-Esp3I-rev	ACACCAcgtctcTGTATGAGGGAAGCGGTGAT CGCC	132
cpec组装-for-质粒	CATACTCTTCCTTTTCAATATTATTGAAGC ATTTATCAGGG	133
cpec 组装-rev-质粒	CTGTCAGACCAAGTTTACTCATATATACTTT AGATTG	134
cpec 组装-for-spec	CAATCTAAAGTATATATGAGTAAACTTGGT CTGACAGTTTGCCGACTACCTTGGTGATCTC G	135
cpec 组装-for-spec2	CAATCTAAAGTATATATGAGTAAACTTGGT CTGACAGTTATTTGCCGACTACCTTGGTGAT CTCG	136
cpec 组装-rev-spec	CCCTGATAAATGCTTCAATAATATTGAAAA AGGAAGAGTATG	137

[0243] 表4:定制测序寡核苷酸

[0244]

寡核苷酸名称	序列	SEQ ID NO:
Fwd CMV	CGCAAATGGGCGGTAGGCGTG	138
Cas9coRevE1	CCGTGATGGATTGGTGAATC	139
Cas9coRevE2	CCCATACGATTTACCTGTC	140
Cas9coRevE3	GGGTATTTTCCACAGGATGC	141
Cas9coRevE4	CTTAGAAAGGCGGGTTTACG	142
Cas9coRevE5	CTTACTAAGCTGCAATTTGG	143
Cas9coRevE6	TGTATTCATCGGTTATGACAG	144
bGH_PArev seq1	CAGGGTCAAGGAAGGCACG	145

[0245]

pHU6-gRNA_for	GTTCCGCGCACATTTCC	146
pHU6-gRNA_rev	GCGGAGCCTATGGAAAAC	147
pCALNL-for1	GCCTTCTTCTTTTCCTACAGC	148
pCALNL-for2	CGCATCGAGCGAGCAC	149

[0246] 表5:基因组PCR引物

[0247]

寡核苷酸名称	序列	SEQ ID NO:
FAM19A2-F1	TCAAGTAGCAAAAAGAAGTAGGAGTCAG	150
FAM19A2-F2	TTAGATGCATTCGTGCTTGAAG	151
FAM19A2-C1	TTAATTTCTGCTGCTAGAACTAAATCTGG	152
FAM19A2-R1	GGGAAGAAAAGTGGATGGAGAATG	153
FAM19A2-R2	CATAAATGACCTAGTGGAGCTG	154
FAM19A2-C2	TGGTTATTTGCCATTAGTTGATGC	155

[0248] 报告物构建

[0249] 使用五件式Golden Gate组装构建下述的报告物。片段1-5侧翼为Esp3I位点；Esp3I消化产生互补的5'突出端，指定片段组装的顺序(图6)。通过退火表5中列出的正向和反向互补寡核苷酸产生片段1、2、4和5。通过将20 μ l的分子级水中的10 μ l的每种寡核苷酸(100 μ M)混合，在95 $^{\circ}$ C温育3分钟，并以-0.1 $^{\circ}$ C/秒的速率将温度降至16 $^{\circ}$ C来使片段退火。用引物3-for和3-rev通过PCR扩增含有kanR和PolyA终止密码子的区域产生片段3。这些引物还在该序列的5'和3'末端附加了Esp3I。

[0250] 将退火的片段1、2、4和5稀释12,000倍并将0.625 μ l的每个片段添加至含有以下的混合物：

[0251] 1) 40-50ng片段3

[0252] 2) 100ng pCALNL EGFP-Esp3I

[0253] 3) 1 μ L Tango缓冲液(10X)[0254] 4) 1 μ L DTT(10mM)[0255] 5) 1 μ L ATP(10mM)[0256] 6) 0.25 μ L T7连接酶(3,000U/ μ L)

[0257] 7) 0.75uL Esp3I (10U/ μ L)

[0258] 8) H₂O至10 μ L

[0259] 将反应在编程进行20个循环(37°C5分钟,20°C)的热循环仪中温育。

[0260] 在完成GoldenGate反应后,将7 μ L的每个反应物与1 μ L的ATP (10mM)、1 μ L的10X Plasmid Safe ATP依赖性DNA酶缓冲液(10X)和1 μ L的Plasmid Safe ATP依赖性DNA酶(10U/ μ L) (Epicentre, Madison, WI)混合以去除线性DNA并减少背景。将DNA酶消化物在37°C温育30分钟,并在70°C加热杀死30分钟。将每个反应物的一半(5 μ L)转化到Mach1-T1细胞中。通过菌落PCR分析菌落并测序。

[0261] 针对图4中使用的报告物修改了方案。使用编码PolyA终止子的5'或3'的靶位点的两个gBlock代替片段1、2、4和5。将这些gBlock(10ng)加入到MMX中,其循环10次(37°C5分钟,20°C)并如上所述继续进行。

[0262] 质粒

[0263] 除非另有说明,否则使用QIAquick凝胶提取试剂盒(Qiagen, Valencia, CA)从琼脂糖凝胶分离DNA片段并使用DNA Clean&Concentrator-5(Zymo Research, Irvine, CA)或Qiaquick PCR纯化试剂盒(Qiagen, Valencia, CA)进一步纯化。使用上面列出的试剂盒之一分离不需要凝胶纯化的PCR片段。

[0264] 使用pCALNL-GFP亚克隆载体pCALNL-EGFP-Esp3I克隆所有recCas9报告物质粒,并基于先前描述的pCALNL-GFP载体(Matsuda and Cepko, Controlled expression of transgenes introduced by in vivo electroporation. Proceedings of the National Academy of Sciences of the United States of America 104, 1027-1032 (2007), 其通过引用并入本文)。为了产生pCALNL-EGFP-Esp3I,用XhoI和MluI消化pCALNL-GFP载体并凝胶纯化以除去loxP位点、卡那霉素抗性标志物和poly-A终止子。退火的寡核苷酸形成EspI插入物,其含有反向的Esp3I位点以及XhoI和MluI相容的突出端;将该插入物连接到XhoI和MluI消化的质粒中并转化。

[0265] 用退火的寡聚物和含有相容的Esp3I突出端的PCR产物通过Golden Gate组装产生pCALNL-GFP recCas9报告物质粒。如前所述用Esp3I(ThermoFisher Scientific, Waltham, MA)建立并进行Golden Gate反应(Sanjana et al., A transcription activator-like effector toolbox for genome engineering. Nature protocols 7, 171-192 (2012), 其全部内容在此通过引用并入)。图6概述了用于报告物组装的一般组装方案和相关引物以及所有recCas9靶位点的序列分别列于表2和6中。侧翼为两个recCas9靶位点的含有KanR(粗体和下划线)和PolyA终止子(斜体和下划线)的代表性DNA序列如下所示。显示的靶位点均为PAM_NT1-0bp-gix_核心-0bp-NT1_PAM(见表6)。原相邻间隔区基序(protoadjacent spacer motif) (PAM)以粗体显示。碱基对间隔区是小写的。gix位点或gix相关位点以斜体显示,并且dCas9结合位点以下划线标出。对于图4的测定中使用的基因组报告物质粒,在卡那霉素抗性标志物中观察到G至T颠换,在下面的序列中由G/T表示。这存在于该图中使用的所有报告物中,并且预计不会影响结果,因为它远离PolyA终止子和recCas9靶位点。

[0266]

ACGCGTCCCAATTTTCCCAAACAGAGGTCTGTAAACCGAGGTTTTGGAACCTCTGTT
TGGGAAAATTGGGGAGTCGAGTCGGATTTGATCTGATCAAGAGACAGGATGAGGA
TCGTTTCGCATGATTGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTGG
GTGGAGAGGCTATTCGGCTATGACTGGGCACAACAGACAATCGGCTGCTCTGA
TGCCGCCGTGTTCCGGCTGTCAG/TCGCAGGGGCGCCCGGTTCTTTTTGTCAAG
ACCGACCTGTCCGGTGCCCTGAATGAACTGCAGGACGAGGCAGCGCGGCTAT
CGTGGCTGGCCACGACGGGCGTTCCTTGCGCAGCTGTGCTCGACGTTGTCACT
GAAGCGGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCAGGATCTCC
TGTCATCTCACCTTGCTCCTGCCGAGAAAGTATCCATCATGGCTGATGCAATG
CGGCGGCTGCATACGCTTGATCCGGCTACCTGCCATTCGACCACCAAGCGAA
ACATCGCATCGAGCGAGCACGTACTCGGATGGAAGCCGGTCTTGTCGATCAGG
ATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGAACTGTTCCGCCAG
GCTCAAGGCGCGCATGCCCGACGGCGAGGATCTCGTCGTGACCCATGGCGAT
GCCTGCTTGCCGAATATCATGGTGGAAAATGGCCGCTTTTCTGGATTCATCGA
CTGTGGCCGGCTGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACC
CGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCGTGCT
TTACGGTATCGCCGCTCCCGATTCGCAGCGCATCGCCTTCTATCGCCTTCTTGA
CGAGTTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCCA
ACCTGCCATCACGAGATTTGATTCCACCGCCGCTTCTATGAAAGGTTGGGCTTCG
GAATCGTTTTCCGGGACGCCGGCTGGATGATCCTCCAGCGCGGGGATCTCATGCTG
GAGTTCTTCGCCACCCCATCGATAACTTGTTATTGCAGCTTATAATGGTTACAAATAA
AGCAATAGCATCAAAATTTCAAAATAAAGCATTTTTTTCACTGCATTCTAGTTGTGGTTT
GTCCAAACTCATCAATGTATCTTATCATGTCTGGATCAAATCCGAACGCACCCCAATT
TTCCCAAACAGAGGTCTGTAAACCGAGGTTTTGGAACCTCTGTTTGGGAAAATTGGG
GCTCGAG (SEQ ID NO: 156)

[0267] 表6:用于报告物测定法的靶位点序列的列表

[0268]

靶位点名称	序列	SEQ ID NO:
PAM_NT1-0bp-gix_核心-0bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTtCTGTAAACCGAGGTTTTGGAACCTCTGTTTGGGAAAATTGGGG	157
PAM_NT1-1bp-gix_核心-1bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTtCTGTAAACCGAGGTTTTGGcAACCTCTGTTTGGGAAAATTGGGG	158
PAM_NT1-2bp-gix_核心-2bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTatCTGTAAACCGAGTTTTGGctAACCTCTGTTTGGGAAAATTGGGG	159
PAM_NT1-3bp-gix_核心-3bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTaatCTGTAAACCGAGTTTTGGcttAACCTCTGTTTGGGAAAATTGGGG	160
PAM_NT1-4bp-gix_核心-4bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTaaatCTGTAAACCGAGTTTTGGcttAACCTCTGTTTGGGAAAATTGGGG	161

[0269]

核心-4bp-NT1_PAM	<i>AGGTTTTGG</i> cctaAACCTCTGTTTGGGAAAATTGGGG	
PAM_NT1-5bp-gix_核心-5bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTgaaatCTGTAAACC <i>GAGGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	162
PAM_NT1-6bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTcgaaatCTGTAAAC <i>CGAGGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	163
PAM_NT1-7bp-gix_核心-7bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTtcgaaatCTGTAAAC <i>CGAGGTTTTGG</i> ccttagctAACCTCTGTTTGGGAAAATTGGGG	164
PAM_NT1-6bp-gix_核心-0bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTtcgaaatCTGTAAAC <i>CGAGGTTTTGGA</i> ACCTCTGTTTGGGAAAATTGGGG	165
PAM_NT1-6bp-gix_核心-1bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTtcgaaatCTGTAAAC <i>CGAGGTTTTGGc</i> AACCTCTGTTTGGGAAAATTGGGG	166
PAM_NT1-6bp-gix_核心-2bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTcgaaatCTGTAAAC <i>CGAGGTTTTGGct</i> AACCTCTGTTTGGGAAAATTGGGG	167
PAM_NT1-6bp-gix_核心-4bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTcgaaatCTGTAAAC <i>CGAGGTTTTGG</i> cctaAACCTCTGTTTGGGAAAATTGGGG	168
PAM_NT1-6bp-gix_核心-5bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTcgaaatCTGTAAAC <i>CGAGGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	169
PAM_NT1-0bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTCTGTAAACCGAG <i>GTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	170
PAM_NT1-1bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTtCTGTAAACCGAG <i>GTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	171
PAM_NT1-2bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTatCTGTAAACCGAG <i>GGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	172
PAM_NT1-3bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTaatCTGTAAACCG <i>AGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	173
PAM_NT1-4bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTaaatCTGTAAACCG <i>AGGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	174
PAM_NT1-5bp-gix_核心-6bp-NT1_PAM	CCCCAATTTTCCCAAACAGAGGTgaaatCTGTAAACC <i>GAGGTTTTGG</i> ccttagAACCTCTGTTTGGGAAAATTGGGG	175
染色体_10-54913298-54913376*	CCCTCCCATCACAGGCCCTGAGggttaaGAGAAAAC <i>CATGTTTTGTG</i> ggccagGCCCATGACCCTTCTCCTCTGGG	176
着丝粒_染色体s_1_5_19	CCTTGTGTTGTGTGTCTTCAACTcacagAGTTAAACGATGCTTTACACagagtaGACTTGAAACACTCTTTTCTGG	177
染色体_5_155183064-155183141(位点1)	CCACCGGCTCATGAGAGGTAGAGctaagGTCCAACCTAGGTTTATCTgagaccGGAACCTCATGTGATTAACCTGG	178
染色体	CCTTAAGAACATAAATCCCCAGGaattcACAGAAACC	179

[0270]

<u>_5_169395198-169395274 (位点2)</u>	<i>TTGGTTTGAGC</i> <i>tttga</i> TTTCCCGCAGGATGTGGGATA GG	
染色 体 <u>_12_62418577-62418652</u>	CCACTCCCTCTCCCCAAAAAGTaaag <i>TAGAAAACC</i> <i>AAGGTTTACAG</i> <i>gcaac</i> AAATAGCACAATGAATGGAA TGG	180
染色 体 <u>_13_102010574-102010650 (FGF14)</u>	CCTAGGGAAGTGATCATAGCTGA <i>gtttct</i> GGAAAAAC <i>CTAGGTTTTAAA</i> <i>gttga</i> GGAGACTTAAGTCCAAAACCT GG	181
原相邻间隔区基序(PAM)以粗体显示。碱基对间隔区是小写的。gix位点或gix相关位点用斜体显示，dCas9结合位点用下划线标出。		
*染色体_10报告物在gix位点的5'和3'末端含有两个重叠的PAM位点和dCas9结合位点。		

[0271] 通过用寡核苷酸1GGs-rev-BamHI或2GGs-rev-BamHI (使用接头SEQ ID NO:182) 和Gin-for-NotI对编码演化的、过度激活的Gin变体(Ginβ) (Gaj et al., A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells. *Nucleic acids research* 41,3937-3946 (2013), 其全部内容在此通过引用并入) 的gBlock进行PCR扩增来构建含有recCas9基因的质粒。用BamHI和NotI消化PCR片段, 纯化并连接到先前描述的表达载体(Addgene质粒43861) (参见例如, Fu et al., High-frequency off-target mutagenesis induced by CRISPR-Cas nucleases in human cells. *Nature biotechnology* 31,822-826 (2013), 其全部内容在此通过引用并入) 中以产生亚克隆载体pGin-1GGs和pGin-2GGs (使用接头SEQ ID NO:182)。与Cas9-rev-FLAG-NLS-AgeI一起使用寡核苷酸1GGs-连接-for-BamHI、5GGs-连接-for-BamHI (使用接头SEQ ID NO:701) 或8GGs-连接-for-BamHI (使用接头SEQ ID NO:183) 构建编码具有1、5或8个GGs接头的Cas9-FLAG-NLS的PCR片段(见表3)。对于编码GGs氨基酸接头的DNA序列, 见表7。用BamHI和AgeI消化PCR片段和亚克隆质粒并连接以产生质粒pGinβ-2xGGs-dCas9-FLAG-NLS (使用接头SEQ ID NO:182)、pGinβ-5xGGs-dCas9-FLAG-NLS (使用接头SEQ ID NO:701) 和pGinβ-8xGGs-dCas9-FLAG-NLS (使用接头SEQ ID NO:183)。对于pGinβ-8xGGs-dCas9-FLAG-NLS (即recCas9) 的DNA和氨基酸序列, 参见下文。编码Ginβ的序列以粗体显示; 编码GGs接头的序列以斜体显示; 编码dCas9接头的序列为黑色; 编码FLAG标记和NLS的序列分别用下划线和小写字母表示。

[0272]

ATGCTCATTGGCTACGTGCGCGTCTCAACTAACGACCAGAATACCGATCTTCA
GAGGAACGCACTGGTTTGTGCAGGCTGCGAACAGATTTTCGAGGACAACTCA
GCGGGACACGGACGGACAGACCTGGCCTCAAGCGAGCACTCAAGAGGCTGCA
GAAAGGAGACACTCTGGTGGTCTGGAAATTGGACCGCCTGGGTCGAAGCATG
AAGCATCTCATTCTCTGGTTGGCGAACTGCGAGAAAGGGGGATCAACTTTCG
AAGTCTGACGGATTCCATAGATAACAAGCAGCCCCATGGGCCGGTCTTCTTCT
ACGTGATGGGTGCACTGGCTGAAATGGAAAGAGAACTCATTATAGAGCGAACC
ATGGCAGGGCTTGC GGCTGCCAGGAATAAAGGCAGGCGGTTTGGAAAGACCAC
CAAAGGGTGGATCCGGAGGGTCCGGAGGTAGTGGCGGCAGCGGTGGTTCAGGTGGCA
GCGGAGGGTTCAGGAGGCTCTGATAAAAAGTATTCTATTGGTTTAGCTATCGGCACTA
ATTCCGTTGGATGGGCTGTCATAACCGATGAATACAAAGTACCTTCAAAGAAATTT
AAGGTGTTGGGGAACACAGACCGTCATTTCGATTA AAAAGAATCTTATCGGTGCCCT
CCTATTTCGATAGTGGCGAAACGGCAGAGGCGACTCGCCTGAAACGAACCGCTCGGA
GAAGGTATACACGTCGCAAGAACCGAATATGTTACTTACAAGAAATTTTTAGCAAT
GAGATGGCCAAAGTTGACGATTCTTCTTTCACCGTTTGGAAAGAGTCCTTCCTTGTC
GAAGAGGACAAGAAACATGAACGGCACCCCATCTTTGGAAACATAGTAGATGAGG
TGGCATATCATGAAAAGTACCCAACGATTTATCACCTCAGAAAAAAGCTAGTTGAC
TCAACTGATAAAGCGGACCTGAGGTTAATCTACTTGGCTCTTGCCCATATGATAAAG
TTCCGTGGGCACTTTCTCATTGAGGGTGTCTAAATCCGGACAACCTCGGATGTCGAC
AAACTGTTTCATCCAGTTAGTACAAACCTATAATCAGTTGTTTGAAGAGAACCCTATA
AATGCAAGTGGCGTGGATGCGAAGGCTATTCTTAGCGCCCGCCTCTCTAAATCCCG
ACGGCTAGAAAACCTGATCGCACAATTACCCGGAGAGAAGAAAAATGGGTTGTTTCG
GTAACCTTATAGCGCTCTCACTAGGCCTGACACCAAATTTTAAGTCGAACTTCGACT
TAGCTGAAGATGCCAAATTGCAGCTTAGTAAGGACACGTACGATGACGATCTCGAC
AATCTACTGGCACAAATTGGAGATCAGTATGCGGACTTATTTTTGGCTGCCAAAA
CCTTAGCGATGCAATCCTCTATCTGACATACTGAGAGTTAATACTGAGATTACCAA
GGCGCCGTTATCCGCTTCAATGATCAAAAGGTACGATGAACATACCAAGACTTGA
CACTTCTCAAGGCCCTAGTCCGTCAGCAACTGCCTGAGAAATATAAGGAAATATTC
TTTGATCAGTCGAAAAACGGGTACGCAGGTTATATTGACGGCGGAGCGAGTCAAGA
GGAATTCTACAAGTTTATCAAACCCATATTAGAGAAGATGGATGGGACGGAAGAGT
TGCTTGTA AAACTCAATCGCGAAGATCTACTGCGAAAGCAGCGGACTTTCGACAAC
GGTAGCATTCCACATCAAATCCACTTAGGCGAATTGCATGCTATACTTAGAAGGCA
GGAGGATTTTTATCCGTTCTCAAAGACAATCGTGAAAAGATTGAGAAAAATCCTAA
CCTTTCGCATACCTTACTATGTGGGACCCCTGGCCCGAGGGAACCTCTCGGTTTCGCAT
GGATGACAAGAAAGTCCGAAGAAACGATTACTCCATGGAATTTTGAGGAAGTTGTC
GATAAAGGTGCGTCAGCTCAATCGTTCATCGAGAGGATGACCAACTTTGACAAGAA
TTTACCGAACGAAAAAGTATTGCCTAAGCACAGTTTACTTTACGAGTATTTACAGT
GTACAATGAACTCACGAAAGTTAAGTATGTCACTGAGGGCATGCGTAAACCCGCCT
TTCTAAGCGGAGAACAGAAGAAAGCAATAGTAGATCTGTTATTCAAGACCAACCGC
AAAGTGACAGTTAAGCAATTGAAAGAGGACTACTTTAAGAAAATTGAATGCTTCGA
TTCTGTGAGATCTCCGGGGTAGAAGATCGATTTAATGCGTCACTTGGTACGTATCA
TGACCTCCTAAAGATAATTAAGATAAGGACTTCTTGATAACGAAGAGAATGAAG
ATATCTTAGAAGATATAGTGTGACTCTTACCCTCTTTGAAGATCGGGAAATGATTG
AGGAAAGACTAAAAACATACGCTCACCTGTTTCGACGATAAGGTTATGAAACAGTTA
AAGAGGCGTCGCTATACGGGCTGGGGACGATTGTGCGCGAAACTTATCAACGGGAT
AAGAGACAAGCAAAGTGGTAAACTATTCTCGATTTTCTAAAGAGCGACGGCTTCG
CCAATAGGAACTTTATGCAGCTGATCCATGATGACTCTTTAACCTTCAAAGAGGATA
TACAAAAGGCACAGGTTTCCGGACAAGGGGACTCATTGCACGAACATATTGCGAAT
CTTGCTGGTTCGCCAGCCATCAAAAAGGGCATACTCCAGACAGTCAAAGTAGTGGA
TGAGCTAGTTAAGGTCATGGGACGTCACAAACCGGAAAACATTGTAATCGAGATGG

[0273]

CACGCGAAAATCAAACGACTCAGAAGGGGCAAAAAACAGTCGAGAGCGGATGAA
 GAGAATAGAAGAGGGTATTAAGAAGCTGGGCAGCCAGATCTTAAAGGAGCATCCT
 GTGGAAAATACCCAATTGCAGAACGAGAACTTTACCTCTATTACCTACAAAATGG
 AAGGGACATGTATGTTGATCAGGAACTGGACATAAACCGTTTATCTGATTACGACG
 TCGATGCCATTGTACCCCAATCCTTTTTGAAGGACGATTCAATCGACAATAAAGTGC
 TTACACGCTCGGATAAGAACCGAGGGAAAAGTGACAATGTTCCAAGCGAGGAAGT
 CGTAAAGAAAATGAAGAACTATTGGCGGCAGCTCCTAAATGCGAACTGATAACGC
 AAAGAAAAGTTCGATAACTTAACTAAAGCTGAGAGGGGTGGCTTGTCTGAACTTGAC
 AAGGCCGGATTTATTAACGTCAGCTCGTGGAACCCGCCAAATCACAAAGCATGT
 TGCACAGATACTAGATTCCCGAATGAATACGAAATACGACGAGAACGATAAGCTGA
 TTCGGGAAGTCAAAGTAATCACTTTAAAGTCAAATTGGTGTCTGGACTTCAGAAAG
 GATTTTCAATTCTATAAAGTTAGGGAGATAAATAACTACCACCATGCGCACGACGC
 TTATCTTAATGCCGTCGTAGGGACCGCACTCATTAAAGAAATACCCGAAGCTAGAAA
 GTGAGTTTGTGTATGGTGATTACAAAGTTTATGACGTCGGTAAGATGATCGCGAAA
 AGCGAACAGGAGATAGGCAAGGCTACAGCCAAATACTTCTTTTATTCTAACATTAT
 GAATTTCTTTAAGACGGAATCACTCTGGCAAACGGAGAGATACGCAAACGACCTT
 TAATTGAAACCAATGGGGAGACAGGTGAAATCGTATGGGATAAAGGCCGGGACTT
 CGCGACGGTGAGAAAAGTTTTGTCCATGCCCAAGTCAACATAGTAAAGAAAAGT
 AGGTGCAGACCGGAGGGTTTTCAAAGGAATCGATTCTTCCAAAAGGAATAGTGAT
 AAGTCATCGCTCGTAAAAAGGACTGGGACCCGAAAAAGTACGGTGGCTTCGATAG
 CCCTACAGTTGCCTATTCTGTCTAGTAGTGCCAAAAGTTGAGAAGGGAAAAATCCA
 AGAACTGAAAGTCAGTCAAAGAATTATTGGGGATAACGATTATGGAGCGCTCGTCT
 TTTGAAAAGAACCCCATCGACTTCCTTGAGGCGAAAAGGTTACAAGGAAGTAAAAAA
 GGATCTCATAATTAAGTACCAAAGTATAGTCTGTTGAGTTAGAAAATGGCCGAA
 AACGGATGTTGGCTAGCGCCGGAGAGCTTCAAAGGGGAACGAACTCGCACTACC
 GTCTAAATACGTGAATTTCTGTATTTAGCGTCCCATTACGAGAAGTTGAAAGGTTT
 ACCTGAAGATAACGAACAGAAGCAACTTTTTGTTGAGCAGCACAAACATTATCTCG
 ACGAAATCATAGAGCAAATTTCCGAATTCAGTAAGAGAGTCATCCTAGCTGATGCC
 AATCTGGACAAAGTATTAAGCGCATACAACAAGCACAGGGATAAACCCATACGTG
 AGCAGGCGGAAAATATTATCCATTTGTTTACTCTTACCAACCTCGGCGCTCCAGCCG
 CATTCAAGTATTTTGACACAACGATAGATCGCAAACGATACACTTCTACCAAGGAG
 GTGCTAGACGCGCACTGATTCACCAATCCATCACGGGATTATATGAACTCGGAT
 AGATTTGTCACAGCTTGGGGGTGACGGTGGCTCCGATTATAAGGATGATGACGACA
AGGGAGGTTCCccaaagaagaaaaggaaggtcTGA (SEQ ID NO: 184)

MLIGYVRVSTNDQNTDLQRNALVCAGCEQIFEDKLSGTRTRDRPGLKRALKRLQKG
DTLVVWKLDRDLGRSMKHLISLVGELRERGINFRSLTDSIDTSSPMGRFFFFYVMGAL
AEMERELIERTMAGLAAARNKGRFRFRPPKGGSGGSGGSGGSGGSGGSGGSDK
 KYSIGLAIGTNSVGWAVITDEYKVPSKFKVLGNTDRHSIKKNLIGALLFDSGETAEATR
 LKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFHRLVESFLVEEDKKHERHPIFGNIV
 DEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVD
 KLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRLENLIAQLPGEKKNLFGNLI
 ALSLGLTPNFKSNFDLAEDAQLQSKDYYDDLDNLLAQIGDQYADLFLAAKNLSDAIL
 LSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYA
 GYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELH
 AILRRQEDFYFPLKDNREKIEKILFRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEE
 VVDKGASAQSFIERMTNFDKNLPNEKVLPHKSLLYEYFTVYNELTKVKYVTEGMRKPA
 FLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDL
 LKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYT
 GWGRLSRKLINGIRDKQSGKTILDFLKSDFANRNFQMQLIHDDSLTFKEDIQKAQVSGQ
 GDSLHEHIANLAGSPAIKKILQTVKVVDELVKVMGRHKPENIVIAMARENQTTQKGQ
 KNSRERMKRIEIGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRL

[0274]

SDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAK
LITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKL
IREVKVITLKSCLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVG TALIKKYPKLESEF
VYGDYKVYDVRKMIKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGE
TGEIVWDKGRDFATVRKVL SMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDW
DPKKYGGFDSPTVAYSVLVVAKVEKGKSKLKS VKELLGITIMERSSEKNPIDFLEAK
GYKEVKKDLIHKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYE
KLGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVLADANLDKVL SAYNKHRDKPIR
EQAENIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQ SITGLYETRIDSQ
LGGDGGSDYKDDDDKGGSpkkrkv 终止 (SEQ ID NO: 185)

[0275] Gin重组酶催化域(其是SEQ ID NO:185的氨基酸1-142)与SEQ ID NO:713的序列相同。dCas9域(其是SEQ ID NO:185的氨基酸167-1533)与SEQ ID NO:712的序列相同。

[0276] MLIGYVRVSTNDQNTDLQRNALVCAGCEQIFEDKLSGTRTDRPGLKRALKRLQKGD TLVWKLDR LGR
SMKHLISLVGELRERGINFRSLTDSIDTSSPMGRFFFYVMGALAE MERELI IERTMAGLAAARNKGRRFGRPPK
(SEQ ID NO:713)

[0277] DKKYSIGLAIGTNSVGWAVITDEYKVP SKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTAR
RRYTRRKNRICYLQE IFSNEMAKVDDSFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDS
TDKADRLIYLALAHMIKFRGHFLIEGDLNPDNSDVKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSR
LENLIAQLPGEKKNLFGNLIALSLGLTPNFKSNFDLAEDAQLLSKDTYDDDLNLLAQIGDQYADFLAAKNLS
DAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLP EKYKEIFFDQSKNGYAGYIDGGASQEEFY
KFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFY PFLKDNREKIEKILTFRIP
YYVGPLARGNSRFAMTRKSEETITPWNFEVV DKGASQSFIERMTNFDKNLPNEKVL PKHSLLYEYFTVYNELT
KVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLKII
KDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKYAHLFDDKVMKQLKRRRYTGWRLSRKLINGIRDKQSGK
TILDFLKSDGFANRFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPA I KKGILQTVKVVDELVKVMGR
HKPENIVIEMARENQTTQKGQKNSRERMKRIE EG IKELGSQILKEHPVENTQLQNEKLYLYLQNGRDMYVDQELD
INRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAER
GGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSCLVSDFRKDFQFYKVREINNY
HHAHDAYLNAVVG TALIKKYPKLESEFVYGDYKVYDVRKMIKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEI
RKRPLIETNGETGEIVWDKGRDFATVRKVL SMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGG
FDSPTVAYSVLVVAKVEKGKSKLKS VKELLGITIMERSSEKNPIDFLEAKGYKEVKKDLI IKLPKYSLFELENG
RKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVLADA
NLDKVL SAYNKHRDKPIREQAENI IHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQ SITGLYETRID
LSQLGGD (SEQ ID NO:712)

[0278] 表7: 编码GGG接头的DNA序列

[0279]

GGG 接 头	SEQ ID NO:	用于GGG接头的DNA序列	SEQ ID NO:
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[0280]

2XGGS	182	GGTGGTAGCGGTGGATCC	186
5XGGS	701	GGTGGATCCGGTGGTTCAGGTGGCAGCGGAGGGTCAG GAGGCTCT	187
8XGGS	183	GGTGGATCCGGAGGGTCCGGAGGTAGTGGCGGCAGC GGTGGTTCAGGTGGCAGCGGAGGGTCAGGAGGCTCT	188

[0281] 对于质粒测序实验,通过用PCR片段的golden gate克隆,用SpecR替换pGinβ-8xGGS-dCas9-FLAG-NLS(使用接头SEQ ID NO:183)中的AmpR基因。通过用Esp3I-for-质粒和Esp3I-rev-质粒进行PCR,在位于AmpR基因侧翼的位点处将Esp3I位点引入pGinβ-8xGGS-dCas9-FLAG-NLS(使用接头SEQ ID NO:183)质粒中。引物spec-Esp3I-for和spec-Esp3I-rev用于扩增SpecR标志物以及引入Esp3I位点和Esp3I产生的突出端,所述突出端与由Esp3I切割的质粒PCR产物产生的突出端相容。按照用于产生如本文所述的报告物质粒的方案对两个片段进行golden gate组装。

[0282] pHU6-NT1 引导RNA表达载体基于先前描述的pFYF1328(Fu et al., High-frequency off-target mutagenesis induced by CRISPR-Cas nucleases in human cells. *Nature biotechnology* 31,822-826 (2013),其全部内容在此通过引用并入),其被改变以靶向细菌萤光素酶基因LuxAB内的区域。通过用通用引物R.pHU6.TSS(-1).univ和编码独特引导RNA序列的引物(表1)进行整个载体的PCR扩增来产生引导RNA表达载体。表8中给出了引导RNA序列的列表。这些引物用T4多核苷酸激酶磷酸化。将PCR反应产物和线性引导RNA表达载体平端连接并转化。用于初始优化的引导RNA表达载体、脱靶对照引导RNA序列和靶向染色体10基因座的那些包含AmpR。本研究中所述的所有其他质粒含有specR以促进测序实验。最初经由CPEC将壮观霉素抗性引入到引导RNA表达载体中,基本上如所述(Quan et al., Circular polymerase extension cloning of complex gene libraries and pathways. *PloS one* 4,e6441 (2009);和Hillson(2010),vol.2015,pp.CPEC protocol;其每篇通过引用并入本文),并且然后如上所述通过载体的PCR扩增构建引导RNA质粒。将反应物在37℃与40U的DpnI温育过夜,纯化并转化。通过用寡核苷酸cpec-组装-for-spec2和cpec-组装-rev进行引导RNA表达载体的PCR扩增产生CPEC的片段。通过经由寡核苷酸cpec-组装-for-spec和cpec-组装-rev-spec对SpecR基因进行PCR扩增产生specR片段。pUC19(ThermoFisher Scientific,Waltham,MA)进行了类似的修饰。

[0283] 表8:gRNA序列的列表

[0284]

gRNA名称	gRNA序列	SEQ ID NO:
中靶_gRNA	ACCTCTGTTTGGGAAAATTG	189
非靶_gRNA	gCACACTAGTTAGGGATAACA	190
染色 体 _10-54913298-54913376_gRNA-rev-5	gCCTCAGGGCCTGTGATGGGA	191
染色 体 _10-54913298-54913376_gRNA-rev-6	gCTCAGGGCCTGTGATGGGAG	192
染色 体 _10-54913298-54913376_gRNA-for-5	GGCCCATGACCCTTCTCCTC	193
染色 体 _10-54913298-54913376_gRNA-for-6	GCCCATGACCCTTCTCCTCT	194
着丝粒_染色体s_1_5_19-gRNA-for	GACTTGAAACACTCTTTTTC	195
着丝粒_染色体s_1_5_19-gRNA-rev	gAGTTGAAGACACACAACACA	196
染色体_5_155183064-155183141_(位 点1)_gRNA-for	GGAACTCATGTGATTAAGT	197
染色体_5_155183064-155183141_(位 点1)_gRNA-rev	gTCTACCTCTCATGAGCCGGT	198
染色体_5_169395198-169395274_(位 点2)_gRNA-for	gTTTCCCGCAGGATGTGGGAT	199
染色体_5_169395198-169395274_(位 点2)_gRNA-rev	gCCTGGGGATTTATGTTCTTA	200
染色 体 _12_62418577-62418652_gRNA-for	gAAATAGCACAAATGAATGGAA	201
染色 体 _12_62418577-62418652_gRNA-rev	gACTTTTTGGGGGAGAGGGAG	202
染色 体 _13_102010574-102010650_(FGF14) _gRNA-for	GGAGACTTAAGTCCAAAACC	203
染色 体 _13_102010574-102010650_(FGF14) _gRNA-rev	gTCAGCTATGATCACTTCCCT	204
脱靶-for (CLTA)	GCAGATGTAGTGTTCACACA	205
脱靶-rev(VEGF)	GGGTGGGGGGAGTTTGCTCC	206
染色 体 _12_62098359-62098434_(FAM19A2) _gRNA-rev	gATATCCGTTTATCAGTGTC	207
染色 体 _12_62098359-62098434_(FAM19A2) _gRNA-for	gTTCCTAAGCTTGGGCTGCAG	208
染色 体 _12_62112591-62112668_(FAM19A2) _gRNA-rev	gCCTAAAAGTGACTGGGAGAA	209

[0285]

染色 _12_62112591-62112668_(FAM19A2) _gRNA-for	gCACAGTCCCATATTTCTTGG	210
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[0286] 细胞培养和转染

[0287] HEK293T细胞购自美国典型培养物保藏中心(American Type Culture Collection)(ATCC,Manassas,VA)。将细胞在补充有10%胎牛血清(FBS,Life Technologies,Carlsbad,CA)的Dulbecco改良的Eagle培养基(DMEM)+GlutaMAX-1(4.5g/L D葡萄糖+110mg/mL丙酮酸钠)中培养。将细胞在37°C,5%CO₂在潮湿的培养箱中培养。

[0288] 用于转染的质粒从PureYield Plasmid Miniprep System(Promega,Madison,WI)中分离。转染前一晚,在48孔胶原处理的平板(Corning,Corning,NY)中以每孔 3×10^5 个细胞的密度接种HEK293T细胞。在25 μ L的Opti-MEM(ThermoFisher Scientific,Waltham,MA)中制备转染反应。对于每次转染,将45ng的每种引导RNA表达载体、9ng的报告物质粒、9ng的pIRFP670-N1(Addgene Plasmid 45457)和160ng的recCas9表达载体混合,与Opti-MEM(ThermoFisher Scientific,Waltham,MA)中的0.8 μ L lipofectamine 2000组合并加入各孔中。

[0289] 流式细胞术

[0290] 转染后60-72小时后,用磷酸盐缓冲盐水洗涤细胞,并用50 μ L的0.05%胰蛋白酶-EDTA(Life Technologies,Carlsbad,CA)在37°C收获5-10分钟。将细胞在250 μ L培养基中稀释,并在BD Fortessa分析仪上运行。使用635nm激光激发iRFP荧光,并使用670/30带通滤光器收集发射。使用488nm激光激发EGFP并用505长通和530/30带通滤光器获得发射荧光。在FlowJo软件上分析数据,门控活的且经转染的事件(表达iRFP)。以从至少6,000个活事件门控的转染细胞的百分比测量阳性GFP表达细胞。对于优化实验,通过测量在用报告物质粒和pUC(没有recCas9或引导RNA表达载体)共转染时产生eGFP的转染细胞的百分比来确定测定背景。然后,当与recCas9和中靶或非靶引导RNA表达载体共转染报告物质粒时从观察到的eGFP阳性细胞的百分比中减去该背景。

[0291] 基因组靶位点的鉴定

[0292] 使用Bioconductor(使用R统计程序的开源生物信息学包)进行搜索适当的靶位点(Fu et al.,High-frequency off-target mutagenesis induced by CRISPR-Cas nucleases in human cells.Nature biotechnology 31,822-826(2013),其全部内容在此通过引用并入)。由Genome Reference Consortium发布的人参考基因组的最新版本(GRCh38)用于搜索与Cas9的PAM需求和如文本中描述的演化的gix序列两者相匹配的位点。在将基因组加载到R中的情况下,每个搜索模式都表示为Biostring(允许字符串匹配和操作的R中的容器)。当使用GRCh38参考组装进行搜索时,使用所述参数扫描整个基因组的两条DNA链揭示了人基因组中大约450个潜在靶标(表9)。

[0293] 表9:计算机中鉴定的recCas9基因组靶标

[0294]

染色体	开始	结束	序列	模式 ID	SEQ ID NO:
chr1	3416902 7	3416910 3	CCTTTAGTGAAAAGTAGACAGCT CTGAATATGAAAGGTAGGTTTTC ATTTCTGGGAAAGAGACGCCAAG TGATGTGG	2	211
chr1	5100670 3	5100678 0	CCTCCAATAAATATGGGACTATG TGGAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACGG GAAGAATGG	1	212
chr1	8922937 3	8922945 0	CCATTCTGCCCGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTAG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	213
chr1	1156380 77	1156381 54	CCATTCTCCCGTCACTTTCAGGT ACAACAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	214
chr1	1225524 02	1225524 78	CCTTGTAGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT GTTGTGG	2	215
chr1	1226098 74	1226099 50	CCTTGTAGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCATACTTGAAACACTCTT TTTGTGG	2	216
chr1	1226686 77	1226687 53	CCTTGTGTTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	217
chr1	1234224 19	1234224 95	CCTTGTGTTGTGTTTATTCAACTC ACAGAGTTAAACGATCCTTTACA	2	218

[0295]

			CAGAGCAGACTTGAAATACTCTT TTTGTGG		
chr1	1236486 14	1236486 90	CCTTGTAGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCATACTTGAAACACTCTT TTTGTGG	2	219
chr1	1238063 35	1238064 11	CCTTGTATTGTGAGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	220
chr1	1240782 28	1240783 04	CCTTGTGTTGTGTGTCTTCAACTC ACAGAGTTAAACGATGCTTTACA CAGAGTAGACTTGAAACACTCTT TTTCTGG	2	221
chr1	1242310 74	1242311 50	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGTAACACTCTT TTTGTGG	2	222
chr1	1242324 35	1242325 11	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACGTGAAACACTCTT TTTGTGG	2	223
chr1	1243447 81	1243448 57	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	224
chr1	1244357 16	1244357 92	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGGAGACTTGTAACACTCTT TTTGTGG	2	225
chr1	1586771 86	1586772 62	CCTGAGGTTTTCCAGGTTTTAAA AGGAAACCTAAAGGTAGGTTTAG CATTAAAGTGTCTTGAAGTTTATTT TAAAAGG	2	226
chr1	1676294 79	1676295 54	CCAAAATTCACAAAACCGAAT GCATCAGTCAAAGCAAGGTTTGA AGAAAAGATTTACCACTTCAGGG AGCTTGG	4	227
chr1	1677834 28	1677835 04	CCTTTTCTGGATATCGTTGATGCT CTGTATGCAAAGGTAGGTTTTT GGGTTATGTTGTTAAACAGTGAT TGAATGG	3	228
chr1	1694093 67	1694094 44	CCTCCAAGAAATATGGAACATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG AGAGAATGG	1	229
chr1	1741453 46	1741454 23	CCTCCAAGAAATATGGGACTATG TGAGAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	230
chr1	1837501	1837502	CCATTCTCCCATCGCTTTCAGGT	1	231

[0296]

	68	45	ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTTCCATATTCT TTGGAGG		
chr1	2008015 40	2008016 17	CCATTCTCCCCATCACTTTCAGGT GTACCGATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTC TTGGAGG	1	232
chr1	2075899 36	2075900 13	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACGG GGAGAATGG	1	233
chr1	2097683 70	2097684 45	CCTTCAGGGCAGAAACAGCTCTA CTAGCAGAGAAAGCAAGCTTTC ATATTGTGCAATACAAAACGAG AGCAGGG	4	234
chr1	2186523 78	2186524 55	CCATTCTCCTCATCTCCTTCTGGT ACTCCAATCAAACGTAGGTTTGG TCTTTTCTCATAGTCTCATATTTC TTGGAGG	1	235
chr1	2221472 50	2221473 27	CCTCCAAGACATATAGGACTATG TGAAAATACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGTATGG	1	236
chr1	2458707 10	2458707 85	CCTGCCAGATACCAGTAGTCACT GTGAATTACAAAGCTACGTTTCT TCCATAGGGAAAGTTTGGAGTCC AGCCAGG	4	237
chr2	2376037	2376114	CCATTCTCCCTGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTC TTGGAGG	1	238
chr2	4119629	4119706	CCATTCTCCCCACCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTC TTGTAGG	1	239
chr2	4909047	4909124	CCTAACCAGAACTAACTAATAG ATATGGGCAGAAAGCATCCTTTC ACTTTTGTCTGGGAGAGGGAAG AAGCAAAGG	1	240
chr2	2898487 7	2898495 3	CCATTTTGGGGAGGCCTTGATGG GAAGCTGGAAAAGGAAGCTTTC TCCCAGTCTGCTGAAGGCCTTG CCAGCTGG	2	241
chr2	3175583 3	3175591 0	CCTCCAAGAAACACAGGACTATG TGAAAAGATCAAACCTACGTTTG ATTGGTGTTCCTGAAAGTGATGG GGAGAATGG	1	242
chr2	3982958 3	3982966 0	CCATTCTCTTCATGACTTTCAGGT ACACCATTGAAACGTAGGTTTGG TCTTTTCACATTGTCCCATATTTC TTGGAGG	1	243

[0297]

chr2	6020594 7	6020602 4	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCGTATTTTC TTGGTGG	1	244
chr2	7908236 2	7908243 9	CCATTCTCCCTGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGGGG	1	245
chr2	7908236 2	7908243 8	CCATTCTCCCTGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGGG	3	246
chr2	1084309 15	1084309 92	CCTCCAAGAAATATGAGATTATA TGAAAAGACCAAACCTACGTTTG ATTGGTGTACTTTAAAGTGACGG GGAGAATGG	1	247
chr2	1158936 85	1158937 62	CCATTCTCCCCGTCATTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCAAATTTTC TTGGAGG	1	248
chr2	1196200 68	1196201 45	CCCCAAGAAATGTGGGACTATA TGAAAAGACCAAACCTACGTTTG ACTGGTGTACCTAAAAGTGATGG GGAGAATGG	1	249
chr2	1196200 69	1196201 45	CCCCAAGAAATGTGGGACTATAT GAAAAGACCAAACCTACGTTTGA CTGGTGTACCTAAAAGTGATGGG GAGAATGG	2	250
chr2	1284950 68	1284951 44	CCCATTGGTGCTGACCAGATGGT GAAGGAGGCAAAGGTTGCTTTGA ATGACTGTGCTCTGGGGTGAGCC AGGCCTGG	2	251
chr2	1331335 59	1331336 34	CCCTTACAGAGGTGAGCTTTGT TATTAGTAAAAAGGTAGGTTTCC CTGTTTTTCTGAAGAAAAGCTGT GAGTGGG	4	252
chr2	1341749 83	1341750 60	CCACTGCCATTGACAGAGTGGC GAGGTGGGTGAAACCTTGCTTTC CTCCTGGCCCATGGGCAGGGTGG GGCTGTGGG	1	253
chr2	1341749 83	1341750 59	CCACTGCCATTGACAGAGTGGC GAGGTGGGTGAAACCTTGCTTTC CTCCTGGCCCATGGGCAGGGTGG GGCTGTGG	3	254
chr2	1380699 45	1380700 22	CCATTCTCCCTGTCACTTTTAGAT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATGTTTC TTGGAGG	1	255
chr2	1387974 20	1387974 96	CCTCCAAGAAATATCAACTGTGT GAAAAGACGAAACCTACGTTTGA TTAATGTACCTGAAAGTGACAGG	2	256

[0298]

			GAGAATGG		
chr2	1452124 34	1452125 11	CCATTCTCCCATTAACCTTTCAAGT ACACCAATCAAAGGTAGGTTTGG TGTTTTCCCATAGTCCCGTATTTT TTGGAGG	1	257
chr2	1478378 42	1478379 19	CCTTTTCATCATGCCCCTTTCACT TTAAGGTGAAAACCTTGCTTTAC ATGTCAGAGAAAAGAAGAGCCC TCAGCTGGG	1	258
chr2	1478378 42	1478379 18	CCTTTTCATCATGCCCCTTTCACT TTAAGGTGAAAACCTTGCTTTAC ATGTCAGAGAAAAGAAGAGCCC TCAGCTGG	3	259
chr2	1541525 40	1541526 17	CCATTCACCCCGTCACTTTCAGG TACACCAATCAAACGTAGGTTT GTCTTTTCACATAGTCCCATATTT CTTGGAGG	1	260
chr2	1577059 43	1577060 19	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTT ATGGTGTACCCGAAAGTGACAGG GAGAATGG	3	261
chr2	1583611 52	1583612 29	CCACCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTT ATAGGTATACCTGAAAGTGACAG GGAGAATGG	1	262
chr2	1614610 06	1614610 83	CCATTCTCCCATCACTTTCAGGT GCACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTT TTGGAGG	1	263
chr2	1790773 76	1790774 53	CCCTCAAGAAATATGAGACTATG TGAAAAGACCAAACCTACGTTT ACTGGTATACCTGAAAGTGACAG GGAGAATGG	1	264
chr2	1790773 77	1790774 53	CCTCAAGAAATATGAGACTATGT GAAAAGACCAAACCTACGTTTGA CTGGTATACCTGAAAGTGACAGG GAGAATGG	2	265
chr2	1810906 99	1810907 76	CCTCCAACAAATATGGGACTATG TGAAAAGACCAAACCTACGTTT ATTGGTGTACCTGAAAGTGACGG GGATAATGG	1	266
chr2	1823319 57	1823320 34	CCATTCTCTCCCTCACTTTCAAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCTTATATTT TTGGCGG	1	267
chr2	1836205 62	1836206 38	CCATTCTCCCTGTCACTGTCAGTA CACCAATCAAACGTAGGTTTGGT CTCTTCACATAGTCCCATATTTCT TGGAGG	2	268
chr2	2073459 27	2073460 03	CCTCCAAGAAATATGGGACTATG TGAACAGACCAAACCTACGTTT	3	269

[0299]

			ATTGGTGTACCTGAAAGTGATGG CAGAATGG		
chr2	2166520 47	2166521 23	CCACCATGCCTGGCCACCACACA TTTTTTTCTAAAGCTTGGTTTTGG CCACAGTGAGAGTTTCTTGGGCT GTCAGGG	2	270
chr2	2166520 47	2166521 22	CCACCATGCCTGGCCACCACACA TTTTTTTCTAAAGCTTGGTTTTGG CCACAGTGAGAGTTTCTTGGGCT GTCAGG	4	271
chr2	2237800 40	2237801 16	CCCCTAGGTGGCGATATCTGAG GGTCCAATGAAACCATGCTTTTT ACTCAGATCTTCCACTAACCACC TCCCCCGG	2	272
chr2	2244865 95	2244866 72	CCTCTAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ACTGGTGTACCTGAAAGTGACGG GGAGAATGG	1	273
chr2	2305269 02	2305269 79	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTAGTGTACCTGAAAGTGACGG GGAGAATGG	1	274
chr2	2320361 27	2320362 04	CCATTCTCCCTGTCACTTTCAGGT ACATCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	275
chr3	4072812	4072889	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ACTGGTGTACCTGAAAGGGATGG GGAGAATGG	1	276
chr3	9261677	9261754	CCCCAAGAAATATGAGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	277
chr3	9261678	9261754	CCCCAAGAAATATGAGACTATGT GAAAAGACCAAACCTACGTTTGA TTGGTGTACCTGAAAGTGACAGG GAGAATGG	2	278
chr3	1673214 6	1673222 3	CCTCTAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTAACTGAAAGTGACAG GGAGAATGG	1	279
chr3	1745071 2	1745078 9	CCTCCAAGAAATATGCGCCTATG TGAAAAGACCAAACCTACGTTTG ATTGGTATACCTGAAAGTGATGG AGAGAATGG	1	280
chr3	2155976 9	2155984 6	CCATTCTCCCTGTCACTTTGAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATATTCGCATATTTT TTGGAGG	1	281
chr3	2341665	2341673	CCATTCTCCCCGTCACTTTCAGGT	1	282

[0300]

	8	5	ACACCAACCAAACGTTGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGAGG		
chr3	2998401 9	2998409 6	CCATTCTCCCTGTCACCTTCCAGT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACATACTCCCATATTTTC TTGGAGG	1	283
chr3	3826955 1	3826962 7	CCTGGCCTAATTTTAAATCCTTAG TTTGACTTAAACCTTGCTTTTAGT GTGATGGCGACAAAAGCTGAGCT GAAAGG	2	284
chr3	4051521 3	4051528 8	CCAGTGCTTTTTGGTTTTAAAGG CAAGCCTCCAAACCTTCCTTTCTC CTGGATGCTGTGGTGGTTGCCAT GCATGG	4	285
chr3	4923361 2	4923368 7	CCCAACTCCTGCGAGAAGTAGCT CACCATGACAAAGCTACCTTTGC TTTTATCGTTTTGCAAAACAAAA AAGGGGG	4	286
chr3	6629289 4	6629297 1	CCATTCTCCCCGTCACCTTTCAGGT GTGCCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCTATATTTTC TTGGAGG	1	287
chr3	6754149 3	6754157 0	CCTCCAATAAATATGGGACTACG TAAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAACTGACAG GGAGAATGG	1	288
chr3	8227301 1	8227308 8	CCATTCTCCCCGTCACCTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGAGG	1	289
chr3	9868334 9	9868342 6	CCTACAAGATATATGGGACTATG TGAAAAGACCAAACCTACGTTTT ACTGGTGTGCCTGAAACTGACGG GGAGAATGG	1	290
chr3	1019236 53	1019237 30	CCATTCTCTCTGTCACCTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGAGG	1	291
chr3	1145334 67	1145335 44	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTC ATTGGTGTACCTGAAAGTGATAG GGAGAATGG	1	292
chr3	1326076 02	1326076 79	CCTCCAATAAATATGGGATGATG TGAAAAGACCAAACCTAGGTTTG ACTGGTGTACCTGAAAATGATGG GGAGAATGG	1	293
chr3	1375451 76	1375452 53	CCTCCAAGAAATATGAGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	294

[0301]

chr3	1376556 79	1376557 56	CCTCCAAGAAATATGGGACTACG TGAAAAGATCAAACCTACGTTTG ATTGTTGTACCTGAAAGTGATGG GGAGAATGG	1	295
chr3	1376620 40	1376621 17	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGTTGTACCTGAAAGTGATGG GGAGAATGG	1	296
chr3	1421337 96	1421338 73	CCTCAAAGTGTCTGGTTTTGTT TTGTTTTTTAAACCATGGTTTTAC CTCTGGCTTAGTGGGACTAAAA TAGGAGG	1	297
chr3	1467269 49	1467270 26	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ACTGGTGTACCTGAAAGTGATGG GGAAAATGG	1	298
chr3	1524210 96	1524211 73	CCTCCAAGAAATATGGGACTGTG TGTAAGACCAAACCTACGTTTG ATTGGTGTACCTCAAAGTGATGG GGAGAATGG	1	299
chr3	1706202 47	1706203 24	CCATTCTCCCATCACATTCAGG TACACCAATCAAACGTAGGTTTG GTCTTTTCACATAGTCCCATATTT CTTGGAGG	1	300
chr3	1811668 73	1811669 49	CCCCTGGAAAAGTTGGAGCATCA CAGGAAAAGCAAACCAACCTTTT TTCTCCCCTAGGTAAACTGGGGA GCCAGGGG	3	301
chr3	1811668 74	1811669 49	CCCTGGAAAAGTTGGAGCATCAC AGGAAAAGCAAACCAACCTTTT TCTCCCCTAGGTAAACTGGGGAG CCAGGGG	4	302
chr4	6604233	6604309	CCTTCCCCAGTTGCAGCAGACAA GAGTCTCGAAAAGCTTGCTTTGG TTGCTGCAGTGGATGGGTTGGTA GGCACAGG	2	303
chr4	6626269	6626344	CCCCACCTCCCAAGCTGCTGGC TTCTCGAATAAAGCTACCTTTCCT TTTACCAAACCTTGTCTCTCGAA TGTCGG	4	304
chr4	8155396	8155472	CCTTGGCCCTGGACAGCTGCTTT TCCTTCCCTAAACCTTGTTTCCC CCTTTGTGCAGGTGGGTGGGTTT GGGCTGG	2	305
chr4	1038680 3	1038688 0	CCTCTTCTAGTGAACCCATGGGG TTACCAAGGGAAAGCAACCTTTT GATAAATATCCCATCTTTTTATG TTGTCTGG	1	306
chr4	2070157 9	2070165 6	CCACTTGAAAGGGTTACCAAGGA TAAGATTTTTAAAGCTTGCTTCA CAAACAACCTCATGCTCCAGGCTT	1	307

[0302]

			GTCAGTGG		
chr4	2959428 6	2959436 3	CCTTTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGA TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	308
chr4	5366842 2	5366849 9	CCATTCTCCCCATCAATTTTCAGTT ACACCAATGAAACGTAGGTTTGG CCTTTTCACATAGTCCCATATTTT TTAGAGG	1	309
chr4	7491480 2	7491487 9	CCATTCTCCCTGTCACTCTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCATATAGTCCCATATTTT TTGGAGG	1	310
chr4	7533278 3	7533285 9	CCTCCAAGAAAATTGGGACTATG TGAAAAACCAAACCTACGTTT ATTGATGTACCTGAAAGTGACAG GAGAATGG	3	311
chr4	8812364 3	8812372 0	CCTTCAAGAAATATGGGACTATG TGAAAGGACAAAACCTACGTTTT ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	312
chr4	8956719 2	8956726 9	CCATTCTCCCCATCACTTTCAGGT ACGCTAATCAAACGTAGGTTTGA TCTTTTCACATAGTCTTATATTTT TTGGAGG	1	313
chr4	9355657 7	9355665 4	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTT ACTGGTGTACCTCAATGTGACAG GGAGAATGG	1	314
chr4	1002663 79	1002664 56	CCATTCTCCCTGTCACTTTTAGGT ACACCAATCAAACGTACGTTTGG TCTTTTCACATAGACCCATATTTT TTGGAGG	1	315
chr4	1034862 34	1034863 11	CCTTCAAGAAATATGGGACTGTG TGAAAAGACCAAAGCTAGGTTT ATTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	316
chr4	1059231 29	1059232 04	CCTACTATTCACAGAGTAATGCA GTTTGCTGAAAAGTTGGTTTTT GCTGACCTCTGAGAGCTCACATT ACAGTGG	4	317
chr4	1068747 11	1068747 88	CCATTCTCTCTGTCACTTTCTGGT ACACCAATCAAACGTAGGTTTGC TCTTTTCACATAATCCCATATTTA TTGAAGG	1	318
chr4	1158057 91	1158058 67	CCATAACATGTATTTGCTGGTGC TAGACTCTCAAAGCTAGGTTT TTTCTACAACAATGGCTGGAAGT CTTCTTGG	3	319
chr4	1220332 77	1220333 54	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG	1	320

[0303]

			TCTTCTCACACAGTCCCATATTTCTTGGAGG		
chr4	1291251 32	1291252 09	CCATTCTTCCCATTACTTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCACATTTCTTGGAGG	1	321
chr4	1354725 62	1354726 39	CCATTCTCCCCCTCACTTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATTGTCCCATATTTCTTGGAGG	1	322
chr4	1385070 99	1385071 76	CCATTCTCCCCAGCACTTACAGG TACACCAATCAAACGTAGGTTTGG GTCATTTACATAGTCCCATATTTCTTGGAGG	1	323
chr4	1442490 93	1442491 70	CCATTCTCCCTGTCACTTTTCAGGT ACAGCAATCAAACGTAGGTTTGG TCTTTTCACATGGTCCCATATTTCTTGGAGG	1	324
chr4	1444364 06	1444364 83	CCTCCAAGAAATATGAGACTATG TGAAAAGACCAAACCTACGTTTGG ATTGGTGTACCTGAAAGTGACGG GGAAGATGG	1	325
chr4	1541102 59	1541103 36	CCTCCAAGAAATATGAGACTATG TGAAAAGACCAAACCTACGTTTGG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	326
chr4	1548934 38	1548935 15	CCTCCAAGAGATATGAGACTATG TAAATAGACCAAACCTACCTTTGG ATTGGTGTACGTGAAAGTGACAG GAAGAATGG	1	327
chr4	1611168 54	1611169 31	CCATTCTCCCCATCACTTTTCAGGT ACACCAACCAAACGTAGGTTTGG TCTTTTCACATAGTCTCATATTTCTTGGAGG	1	328
chr4	1651407 48	1651408 23	CCTCCATTGACTACTCCTTATCAT TGGCTAGAAAACCTACCTTTCAA CCAGTTTCTAAGGCCAAGAAACT TGGAGG	4	329
chr4	1819285 08	1819285 85	CCACCAAGAAATATGGGACTACG TGAAAAGACCAAACCTACGTTTGG ATGGGTGTGCCTGAAAGTGACGG GAAGAATGG	1	330
chr4	1875219 58	1875220 35	CCTCCAAGAAATAAGGGACTATG TGAAAAGACCAAACCTACGTTTGG ATTGGTGTACCTGAAGGTGACAG GGAGAATGG	1	331
chr5	1267563 9	1267571 5	CCAAAGGGCCTTTGTGATTCTAC TTTGTAATATAAAGGATGGTTTC TTACTACGGTTGGTGTCTTGCA GGAGTGGG	3	332
chr5	2927180	2927188	CCTCCAAGAAATATGGGACTATG	1	333

[0304]

	4	1	TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGATGG GGAGAATGG		
chr5	3535266 0	3535273 7	CCATTCTCCCCGTTACTTTCAGGT ACACCAATAAAACCTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	334
chr5	3872323 5	3872331 0	CCCATATCTCTGGCAAGGGCAGC TCTCTGGCTAAACCAAGCTTTCC TGTAGAGCTTGAGTTCCAAGGCA GCGTTGG	4	335
chr5	4735833 9	4735841 5	CCTTGTAGTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT GTTGTGG	2	336
chr5	4741581 1	4741588 7	CCTTGTAGTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCATACTTGAAACACTCTT TTTGTGG	2	337
chr5	4747461 4	4747469 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	338
chr5	4822835 6	4822843 2	CCTTGTGTTGTGTTTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAATACTCTT TTTGTGG	2	339
chr5	4845455 1	4845462 7	CCTTGTAGTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCATACTTGAAACACTCTT TTTGTGG	2	340
chr5	4861227 2	4861234 8	CCTTGTATTGTGAGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	341
chr5	4888416 5	4888424 1	CCTTGTGTTGTGTGTCTTCAACTC ACAGAGTTAAACGATGCTTTACA CAGAGTAGACTTGAAACACTCTT TTTCTGG	2	342
chr5	4903701 1	4903708 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGTAACACTCTT TTTGTGG	2	343
chr5	4903837 2	4903844 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACGTGAAACACTCTT TTTGTGG	2	344
chr5	4915071 8	4915079 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	345

[0305]

chr5	4924165 3	4924172 9	CCTTGTGTTGTGTGATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGGAGACTTGTAACACTCTT TTTGTGG	2	346
chr5	8858271 4	8858279 0	CCTTTTCATAAGAAGAAAATCGA CTCATCATTGAAACCAAGCTTTG GTACAATTTTCATTGATGTTTCCA GAAGCAGG	3	347
chr5	9349715 6	9349723 1	CCCATAGACTATGATAGAAACAA AATAACCCAAAAGCTAGCTTTCT GATTGAGTTTCCATAAATGCAAT GTGAAGG	4	348
chr5	9429502 9	9429510 5	CCATTCACTTGTCACTTTCTGGTA CACCAATCAAACGTAGGTTTGGT CTTTTCACATAGTCTCATATTTCT TGGAGG	2	349
chr5	9495674 6	9495682 3	CCTCCAAGAAATATGGGACTCTG TAAAGAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGAAGG GGAGAATGG	1	350
chr5	1060034 88	1060035 65	CCATTCTCCCCGTCATTTTCAGGT ACACCAATCAAACCTAGGTTTGG TCTTTTACATAGTCCCATATTTT TTGGAGG	1	351
chr5	1187279 05	1187279 82	CCTCCACGAAACATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	352
chr5	1321560 32	1321561 09	CCAATTTCCCCCTCACTTTCAGAT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTT CTGGAGG	1	353
chr5	1520379 51	1520380 28	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATATTCCCATATGTC TTGGAGG	1	354
chr5	1551830 64	1551831 41	CCCACCGGCTCATGAGAGGTAGA GCTAAGGTCCAAACCTAGGTTTA TCTGAGACCGGAACTCATGTGAT TAACTGTGG	1	355
chr5	1551830 65	1551831 41	CCACCGGCTCATGAGAGGTAGAG CTAAGGTCCAAACCTAGGTTTAT CTGAGACCGGAACTCATGTGATT AACTGTGG	2	356
chr5	1631482 11	1631482 88	CCTTCAAGAAATATGGGACTATG TGAAGAGACCAAACCTACGTTTG ATTGGTGTAGCCAAAAGTGATGG GGAAAATGG	1	357
chr5	1658895 37	1658896 14	CCTCAGATTAGATTTACTTGCAA AGAGACATTTAAAGGATCGTTTT GATACTATTTTGAAAGTACTATA	1	358

[0306]

			CAAAGATGG		
chr5	1693951 98	1693952 74	CCTTAAGAACATAAATCCCCAGG AATTCACAGAAACCTTGGTTTGA GCTTTGGATTTCCTCGCAGGATGT GGGATAGG	2	359
chr5	1710213 80	1710214 57	CCATTCTCTCTGTCACCTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCTCATAGTCCCATATTTT TTGGAGG	1	360
chr5	1730598 98	1730599 73	CCATTTACCATCATTCTCTGTCAT GGCAGGTGAAAGCAAGCTTTTAT ATAGACAATGTTCTACTTAGTTT ACAGGG	4	361
chr5	1741023 59	1741024 35	CCCAAAGTTAATTTTACTCTTTTT CTGAATCAAAGGAACCTTTCCT CCATGAGAAGAATCCTGCCATAT TTCTAGG	2	362
chr5	1809278 11	1809278 88	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGCTATACATGAAAGTGACGG GGAGAATGG	1	363
chr6	1752363	1752440	CCTTCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACCTTTG ATTGGTGTACCTGAAAGTGATGG GAAGAATGG	1	364
chr6	2059527 9	2059535 6	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTACATAGTCCCATAGTTC TTGGAGG	1	365
chr6	2343137 0	2343144 7	CCATTCTCCCCGTCACCTTTCAGG GACAACAATCAAACGTAGGTTTGG GCCTTTGCACATAGTCTTATATTT CTTGGAGG	1	366
chr6	2919062 4	2919070 1	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTACATAGTCCCATATTTT TTGGAGG	1	367
chr6	6153326 6	6153334 3	CCTCCAAAAAATATGGGACTATG TGAGAAGACCAAACCTACGTTTT ATTAGTGTACCTCAAAGTGACAG GGAGGATGG	1	368
chr6	1010527 64	1010528 41	CCATTCTCCCCATCACTTTCAGGT ACACCAATGAAACGTAGGTTTGG CCTTTTACATAGTTTCATATTTT TTGGAGG	1	369
chr6	1171763 55	1171764 32	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	370
chr6	1177470	1177471	CCTACAAGAAATATGGAACCTTGT	2	371

[0307]

	73	49	AAAAAGACCAAACCTACGTTTGA TTGGTGTACCTGAAAGTGACGGG GAGAATGG		
chr6	1184225 08	1184225 85	CCTCCAAGAAATATGGGACAATG TGAAAAGGCCAAAGCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	372
chr6	1220350 19	1220350 96	CCTTTCAACTTAGAGGTAAACA AAAGTCCTGAAAACCTAGGTTTG ACCATAAGTTGGGACCATACGAG CATAGAAGG	1	373
chr6	1344452 10	1344452 87	CCAAAAATAAAAAAAAAATTGAC TTATAAGTAAGAAAGGTTTCGTTT TCTCACATTCAGAAAGAGAACCC ACATGTTGGG	1	374
chr6	1344452 10	1344452 86	CCAAAAATAAAAAAAAAATTGAC TTATAAGTAAGAAAGGTTTCGTTT TCTCACATTCAGAAAGAGAACCC ACATGTTGG	3	375
chr6	1351549 44	1351550 21	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	376
chr6	1378899 95	1378900 72	CCATTCTCCCCGTCCTTTCAGGT ACACCAATCAAACGTGGTTTAG TCTATTACATAGTCCCATATTTT TTGGAGG	1	377
chr6	1439939 04	1439939 81	CCGAAAAGAATAAGACTATCAG CTGAAGTCTTAAAACGATCCTTT GGCCCCAGTACTCTATATGCAG GATAGAAAGG	1	378
chr6	1526104 73	1526105 49	CCTACAAAATAGGGGACTATGT GATAAGACCAAACCTACGTTTGA TTGGTGTACCTGAAAGTGATGGG GAGAATGG	2	379
chr6	1603726 04	1603726 81	CCATTCTACCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG CCTTTTCATATAGTCTCATATTTT TTGGAGG	1	380
chr6	1693524 78	1693525 55	CCATTCTCCCCATCACTTTCAGGT ATACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTAGAGG	1	381
chr6_GL 000251v 2_alt	677196	677273	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	382
chr6_GL 000252v 2_alt	456242	456319	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	383

[0308]

chr6_GL 000253v 2_alt	456202	456279	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	384
chr6_GL 000254v 2_alt	456371	456448	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	385
chr6_GL 000255v 2_alt	456225	456302	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	386
chr6_GL 000256v 2_alt	500011	500088	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	387
chr7	5256551	5256627	CCACCACACCCAGCCTTATGGGA TGGTTTTCAAAGCATCCTTTTTT AGAAGTGGATTCTGATATATAAT CGGATGG	2	388
chr7	7392583	7392660	CCATTCTCAATGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	389
chr7	8737741	8737818	CCATTCTCTCTGTCACTTTCAGGT ACACCAGTCAAAGGTAGGTTTGT TTTATTACACGTTACATATTTT TTGGAGG	1	390
chr7	1135222 6	1135230 3	CCATTTCGCCCCATCACTTTCAGG TACACTAGTAAAACGTAGGTTTGG GTCTTTTCACATAGTTCCATATTT CTTGGAGG	1	391
chr7	1551914 5	1551922 2	CCTCCAAGAAATATGGGACTATG TGAAGAGATCAAACCTAGGTTTGG ATTGTTGTACCTGAAAGTGATAA GAAGAATGG	1	392
chr7	1922834 1	1922841 8	CCTCCAATAAATATGGGGCTATG TGAAAAGACCAAACCTACGTTTGG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	393
chr7	2377844 5	2377852 2	CCTTTTCCCTGTCACTTTCAGGT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACATAGTCGAATATTTT TCAAGG	1	394
chr7	2377844 6	2377852 2	CCTTTTCCCTGTCACTTTCAGGTA CACCAGTCAAACGTAGGTTTGGT CTTTTCACATAGTCGAATATTTCT TCAAGG	2	395
chr7	2676906 5	2676914 2	CCATTCTCCCTGTCACTTTCAGGT ACACTAATCAAACGTAGGTTTGG TGTATTACACAGTCCCATATTTT	1	396

[0309]

			TTGGAGG		
chr7	4286403 5	4286411 2	CCATTCTTCCTGTCACCTTCAGGT ATACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATGTTTC TTGGAGG	1	397
chr7	4649892 3	4649900 0	CCTCCAAGAAATATGAGACTATA TGAAAATACCAAACCTACGTTTG ATTGGTGTACCTGAAAGAGACAG GGAGAATGG	1	398
chr7	5153536 0	5153543 7	CCATTCTCCCTATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCATGTAGTCCCATATTC TTGGAGG	1	399
chr7	5192710 6	5192718 3	CCATTCTGCCCCTGTCACCTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	400
chr7	5697694 2	5697701 8	CCGTCCGATTATATATCAGAATC TACTTCTAAAAAAGGATGCTTTT GAAAACCATCCCATAAGGCTGGG TGTGGTGG	3	401
chr7	8002159 8	8002167 5	CCTACAAGGAATATAGGACTATG TGAAAATACCAAACCTACGTTTC ACTGCTGTACCTGAAGGTGACAG GGAGAATGG	1	402
chr7	8967385 3	8967393 0	CCATTCTCCCCATCATTTCAGGT AAACCAATCAAAGGTAGGTTTGG TCATTTTCACATAGTCCCATATTC TTGGAGG	1	403
chr7	1034047 90	1034048 67	CCATTCTCCCCCTGTCACCTTCAGGT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACACAGTCCCATATTC CTGGAGG	1	404
chr7	1130536 51	1130537 28	CCATTCTCCCCATCATTTCAGGT ACAGCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	405
chr7	1257652 04	1257652 79	CCACTACAGATTCTTGGGTCAAG ATGTGTGCAAAAGGATGCTTTAG GGTGATGGATATGAGTGGGATGA AATGAGG	4	406
chr7	1280421 58	1280422 34	CCTGAAAAAAAACCTGCCAGCC AGCAACTCTGAAAGGATGCTTTG TGTGAGTGAGCAGTGTCTGAGAT GGACAGGG	3	407
chr7	1306373 32	1306374 09	CCATTCTCCCCATCATTTCAGGT ACGCCAATCAAACGTAGGTTTGG TCTTTTGACATAGTCCCATATTC TTGGAGG	1	408
chr7	1369830 50	1369831 27	CCGTTCTCCCCATCATTTTAGGT ACACCAATCAAACGTAGGTTTGG	1	409

[0310]

			TCTTTTCACATAGTCTCATATTTCTTGGAGG		
chr7	1435795 07	1435795 84	CCATTCTCCTGGTCACTTTCAGGT ATACCAATCAAACGTAGGTTTGG TCTTTTCATGTAGTCCCATATTTCTTGGAGG	1	410
chr7	1437498 81	1437499 58	CCTCCAAGAAATATGGGACTACATGAAAAGACCAAACCTACGTTTATTGGTATACCTGAAAGTGACCA GGAGAATGG	1	411
chr8	2338364	2338441	CCTCCAAGAACTATGGGACTATGTGAAAAGACCAAACCTACGTTTATTGGTGTACCTGAAAGTGACGG GGAGAATGG	1	412
chr8	2383289	2383366	CCATTCTCCCCGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATAGTTCTTGGAGG	1	413
chr8	8414568	8414645	CCATTCTCCCCGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACAGAGTCCCATATTTCTTGGAGG	1	414
chr8	2416314 2	2416321 9	CCATTCTCCCCGTCACTTTCATGT ACACCAAGCAAACGTAGGTTTGA TCTTTCCACATAGTCCCGTGTTCTTGGAGG	1	415
chr8	3429905 1	3429912 8	CCTCCAAGAAATATGGGACTATGTGAAAAGACCAAACCTACGTTTATTGGTGTACTTGAAAGTGACAG GGAGAATGG	1	416
chr8	4096548 5	4096556 2	CCTCCAAGAAATATGGGACTATGTGAAAAGACAAAACCTACGTTTACTGGTGTACCTGAAAGTGACAG GGAGGATGG	1	417
chr8	4837165 9	4837173 5	CCCCACCTTTTAAAAACATGCATACATACGGAAACGTTGCTTTCTGCACGATTTCAATTTAATGGAAC AGAACAGG	2	418
chr8	8253496 0	8253503 7	CCATTTCCCCTGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTATCATATTTCTTGGAGG	1	419
chr8	1092176 24	1092177 00	CCATTCTCCCCGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTCTGGAGG	3	420
chr8	1347902 85	1347903 61	CCTTTTGTAAAGTAATAGAATTCTGCTTCTTAAAGGAACCTTTCA GGCAAGATGGTGGTTAGAGCACC TAAATGGG	2	421
chr8	1347902	1347903	CCTTTTGTAAAGTAATAGAATT	4	422

[0311]

	85	60	CTGCTTCTTAAAGGAACCTTTCA GGCAAGATGGTGGTTAGAGCACC TAAATGG		
chr8_KI2 70821v1 _alt	519635	519712	CCTCCAAGAAGAACTATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACGG GGAGAATGG	1	423
chr8_KI2 70821v1 _alt	564557	564634	CCATTCTCCCCGTCACCTTCAGGT ACACCAATCAAACGTAGGTTTGG CCTTTTCACATAGTCCCATAGTTC TTGGAGG	1	424
chr9	1495120 7	1495128 3	CCTCCAAGAAATATGGGACTGGT GAAAAGACCAAACCTACGTTTGA CTGGTGTACCTGAAAGTGACGGG GAGACTGG	2	425
chr9	2324921 8	2324929 5	CCTCCAAGAAACATGGGAATGTG TGAAAAGACCAAACCTACGTTTG ATTGGCGTACCTGAAAGTGACGG GGAGTATGG	1	426
chr9	2627889 6	2627897 3	CCTCCAAGAAATATGGGACTGTG TGAAAAGACCAAACCTACGTTTG ATTGGTATACCTGAAAGTGACAG AGAGAATGG	1	427
chr9	2732323 7	2732331 4	CCATTCTCCCCTTCACTATCAGGT ACACCAATCAAACGTAGGTTTAG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	428
chr9	3151799 3	3151807 0	CCATTCTCCCCGTCACCTTCAGAT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	429
chr9	3969486 0	3969493 7	CCATCTTACTTTGTACTIONACTGT TCTTTAGAGAAAGCTTCCTTTTG GAGACCAACCAGGACTCCTTAGA AGCAGAGG	1	430
chr9	4245113 2	4245120 9	CCATCTTACTTTGTACTIONACTGT TCTTTAGAGAAAGCTTCCTTTTG GAGACCAACCAGGACTCCTTAGA AGCAGAGG	1	431
chr9	6077657 3	6077665 0	CCTCTGCTTCTAAGGAGTCCTGG TTGGTCTCCAAAAGGAAGCTTTC TCTAAAGAACAGTGTAGTACAAA GTAAGATGG	1	432
chr9	6264748 2	6264755 9	CCTCTGCTTCTAAGGAGTCCTGG TTGGTCTCCAAAAGGAAGCTTTC TCTAAAGAACAGTGTAGTACAAA GTAAGATGG	1	433
chr9	6668203 0	6668210 7	CCTCTGCTTCTAAGGAGTCCTGG TTGGTCTCCAAAAGGAAGCTTTC TCTAAAGAACAGTGTAGTACAAA GTAAGATGG	1	434

[0312]

chr9	8226442 7	8226450 3	CCACCCTGTGCCTGGCCATTTT CACTATTCTTAAAGGAAGCTTTG GTTTACAAAGGTTTGCTACTGTA CTTCCAGG	3	435
chr9	8404268 4	8404276 1	CCATTCTCCCTGTCACTTTCAGGT ACACCATTCAAACGTAGGTTTGG TCTTTTCTCATAGTCCCATATTTT TTGGAGG	1	436
chr9	9525601 2	9525608 9	CCTCCAAGAAATTCGGGACTATG TGAAAAGACAAAACCTACGTTTA ATTGGTGTGTGGTGTACCTGAAA GTGACAAGG	1	437
chr9	1018169 88	1018170 65	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACCA GAAGAATGG	1	438
chr9	1358423 27	1358424 03	CCTCCAAGAAATATGGGACTATG TGAAAAGCCCAAACCTACGTTTG ACTGATGTACCTAAAGTGACGGG GAGAATGG	3	439
chr9	1369108 65	1369109 40	CCCGCACTGTGAGCTTGGCCGAG TGCTGTCTGAAAGCATCCTTTCC CTTACCTGGAGACTGGAGCGCC ATAGAGG	4	440
chr10	1371031 2	1371038 9	CCTGTCTCCCCCATTCCATGCAA AATAAAACACAAACCAAGCTTTG CTTTAAGTGCTCCCTGATGCAGT TCAGCGTGG	1	441
chr10	1893812 9	1893820 6	CCATTCTTCCCGTCACATTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCCCATAGTCCCATATTTT TTAGAGG	1	442
chr10	2271283 8	2271291 4	CCCCCTGCTCAGCTTGGGGAAGA AAAATACAAAACGATGCTTTTA GGCATTTTAAACAACCTTACTAC ATTGAGGG	2	443
chr10	2271283 8	2271291 3	CCCCCTGCTCAGCTTGGGGAAGA AAAATACAAAACGATGCTTTTA GGCATTTTAAACAACCTTACTAC ATTGAGG	4	444
chr10	4016093 2	4016100 9	CCTTTGTGTTGTGTGTATTCAACT CACAGAGTGAAACCTTCCTTTAT TCAGAGCAGTTTTGAAACACTCT TTTTGTGG	1	445
chr10	4039013 6	4039021 3	CCTTTGTGTTGTGTGTATTCAACT CACAGAGTGAAACCTTCCTTTAT TCAGAGCAGTTTTGAAAAACACT TTTTGTGG	1	446
chr10	4040915 2	4040922 9	CCTTTGTGTTGTGTGTATTCAACT CACAGAGTGAAACCTTCCTTTAT TCAGAGCAGTTTTGAAAAACTCT	1	447

[0313]

			TTTTGTGG		
chr10	4043394 0	4043401 7	CCTTTGTGTTGTGTGTATTCAACT CACAGAGTGAAACCTTCCTTTAT TCAGAGCAGTTTTGAAACACTCT TTTTGTGG	1	448
chr10	4058815 5	4058823 2	CCTTTGTGTTGTGTGTATTCAACT CACAGAGTGAAACCTTCCTTTAT TCAGAGCAGTTTTGAAATACTCT TTTTGTGG	1	449
chr10	4114620 7	4114628 4	CCTTTGTGTTGTGTGTATTCAACT CACAGAGTGAAACCTTCCTTTAT TCAGAGCAGTTTTGAAACACTCT TTTTGTGG	1	450
chr10	4383518 3	4383526 0	CCATTCTCCCTGTCACCTTCAAGT ACACCAATCAAACCTAGGTTTGG TCTTTTCACATAGTCCATATTTT TTGGAGG	1	451
chr10	5491322 2	5491329 9	CCCCTCCCATCACAGGCCCTGAG GTTTAAGAGAAAACCATGGTTTT GTGGGCCAGGCCCATGACCCTTC TCCTCTGGG	1	452
chr10	5491322 2	5491329 8	CCCCTCCCATCACAGGCCCTGAG GTTTAAGAGAAAACCATGGTTTT GTGGGCCAGGCCCATGACCCTTC TCCTCTGG	3	453
chr10	5491322 3	5491329 9	CCCTCCCATCACAGGCCCTGAGG TTTAAGAGAAAACCATGGTTTTG TGGGCCAGGCCCATGACCCTTCT CCTCTGGG	2	454
chr10	5491322 3	5491329 8	CCCTCCCATCACAGGCCCTGAGG TTTAAGAGAAAACCATGGTTTTG TGGGCCAGGCCCATGACCCTTCT CCTCTGG	4	455
chr10	5803595 1	5803602 8	CCATTCTCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTCA TCTTTTCACATAGTCCCACGTTT TTGGAGG	1	456
chr10	5867752 5	5867760 2	CCTCCAAGATATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAATTGATGG GGAGAATGG	1	457
chr10	8402139 0	8402146 7	CCTCCAAGAAATATGGGACTGTG TGAAAAGAACAAACCTACGTTTG ATTGGTGTACGTGAAAGTGATGG GGAGAATGG	1	458
chr10	9144269 2	9144276 9	CCATTCTCCCGTCACTTTCAGAT ACACCAAAAAACGTAGGTTTGG TCTTTCACATAGTCCCACATTTT TTGGAGG	1	459
chr10	9144684 8	9144692 5	CCTCCAAGAAATGTGGGACTATG TGAAGAGACCAAACCTACGTTTT	1	460

[0314]

			TTTGGTGTATCTGAAAGTGACGG GAGGAATGG		
chr10	1169287 84	1169288 60	CCTCCAAGGGGAATCTGAGTTCT CTGAAGACAAAAAGCATGGTTTC TTTTCTTCTGTATTTCTTATTGTTT CCTAGG	3	461
chr10	1169377 71	1169378 48	CCATTCTCCCTATCACTTTCCAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGAGG	1	462
chr11	3118207 0	3118214 7	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTATACTTGAAATTGACAA GGAGAATGG	1	463
chr11	3473927 3	3473935 0	CCTCCAAGAAATATGGGACTATG TGGAAAGACCAAACCTACGTTTG ACTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	464
chr11	8664652 9	8664660 6	CCTCTAAGAAATATGGGACTATG TGAAGAGATGAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACGA GGAGAATGG	1	465
chr11	9046979 1	9046986 7	CCCTCGTATACTACATGCTATAG TCAAAGCAGTAAACCTTCCTTTTC CTTAAGCAGACCACACTCTTTCA TGCCTGGG	3	466
chr11	9046979 2	9046986 7	CCTCGTATACTACATGCTATAGT CAAAGCAGTAAACCTTCCTTTCC TTAAGCAGACCACACTCTTTTCAT GCCTGGG	4	467
chr11	9242998 5	9243006 2	CCATTCTCCCCATCACTTTCAGGT ATACTAATCAAAGGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC ATGGAGG	1	468
chr11	1028184 98	1028185 74	CCATTCCCCCGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTTC TTGGAGG	2	469
chr11	1207650 65	1207651 42	CCATTCTCCCCGTCCTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCTTATAGTCCCATATTTTC TTGGAGG	1	470
chr11	1231319 01	1231319 78	CCACTGCACCTGACCAAGATCCT TAATTTTTCTAAACCTACGTTTAT CATCTATAAAATGAGCCATCTTT TCACATGG	1	471
chr11	1294685 20	1294685 97	CCTCCGAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGTTGTACCTGAAAGTGACAG GGAGAATGG	1	472
chr11	1312723	1312724	CCATTCTCCCCATCACTTTTAGGT	1	473

[0315]

	61	38	ACACCAATCAAACGTAGGTTTGG TCCTTTTGCATAGACCCATATTTT TTGGAGG		
chr11	1327614 15	1327614 92	CCATTTTCCCCGTCAGTTTCATAT ACACCTATCAAACGTAGGTTTAC TGTTTTACATAGTCCCTTATTTT TTGGAGG	1	474
chr12	2236741 6	2236749 3	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACCTTTG ATTGGTGTACCTGAAAGTGACGG GCAGGATGG	1	475
chr12	3314638 4	3314646 1	CCATTCTTCTCGTCATTTTCAAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCGCATAGTCCCATATTTT TTGGAGG	1	476
chr12	3319847 6	3319855 3	CCATTCTTCTCGTCACTTTCAAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	477
chr12	4603833 2	4603840 9	CCTCCAAGAAATATAGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACTTGAAAGTGACAG GGAGAATGG	1	478
chr12	6023612 6	6023620 3	CCTCCAAGAAATGTGGAACATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	479
chr12	6209835 9	6209843 4	CCCTGACACTGATAAACGGATAT GAAGAGAAAAAAGCTAGGTTTTT GCTGGAATTCCTAAGCTTGGGCT GCAGTGG	4	480
chr12	6211259 1	6211266 8	CCCTTCTCCAGTCACTTTTAGGT ACACCAATGAAACGTAGGTTTGG TCTTTTCACACAGTCCCATATTTT TTGGAGG	1	481
chr12	6211259 2	6211266 8	CCTTCTCCCAGTCACTTTTAGGTA CACCAATGAAACGTAGGTTTGGT CTTTTCACACAGTCCCATATTTCT TGGAGG	2	482
chr12	6241857 7	6241865 2	CCACTCCCTCTCCCCAAAAAGT AAAGGTAGAAAACCAAGGTTTA CAGGCAACAAATAGCACAATGA ATGGAATGG	4	483
chr12	7173231 1	7173238 8	CAAACCCGCATCGCACACCCTG TGAGGGGGACAAAGGAACCTTTC CGTTCCAACATCAAGGTTGTTTT GACCCAAGG	1	484
chr12	7804781 6	7804789 3	CCATTCTTTCTGTCACTTTCAGGT ATACCAGTCAAACCTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	485

[0316]

chr12	8148001 6	8148009 3	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	486
chr12	9684023 1	9684030 7	CCACACGGTAGAGGATAAACTA GGTGGATTCTCAAAGCAACCTTT GAAATAATCTATGCAGTTTTTCT GGTACTGG	3	487
chr12	9918716 5	9918724 2	CCACCAAGAAACATGGGACTATG TGAAAAGACCAAACCTACGTTTG GTTGGTGTACCTGGAAGTGACGG GGAGAGTGG	1	488
chr12	1078608 41	1078609 18	CCTCCAAGAAATATGGGACCATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	489
chr12	1108828 09	1108828 85	CCTGTAAAAAGGTCACATGGTCA GGTGTGCCTAAACGATCCTTTTA TTTATTTATTTATTTATTTTAAAG AAACAGG	2	490
chr12	1190633 21	1190633 97	CCAGCCCCAAAATGTCAGGGGCT TAGAACAAACAAAGGTTCTTTTC ATGTTTATACTACATGTTTGTCAT GGGCTGG	2	491
chr13	3532070 4	3532078 1	CCGTTTTCCCATCACTTTCAGGT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACATGGTCCCACATTTT TTGGAGG	1	492
chr13	5313347 7	5313355 4	CCTGGAATAGCTTTCCTGACTGT CTGACTTCAAAAACCTTGGTTTG ACCACTTCGTCTATATCATGAGG AAGGACTGG	1	493
chr13	5318488 0	5318495 6	CCCTACTCTGAACCTACCTTGAT AAAGCCTAGAAAACCAAGCTTTG ACAAGATTTGACAAGAGATGGA ATTTGGAGG	3	494
chr13	5318488 1	5318495 6	CCTACTCTGAACCTACCTTGATA AAGCCTAGAAAACCAAGCTTTGA CAAGATTTGACAAGAGATGGAAT TTGGAGG	4	495
chr13	5789696 2	5789703 8	CCCTTATAAAACTGAAAACCTTA ACCTTTTTTAAAGCATGCTTTTGA ATAAATTCTTTTATTACAAAAAA GACCAGG	2	496
chr13	6261010 0	6261017 7	CCATTCTCCCTGTCACCTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACGTAGTCCCATATTTT TTGGAGG	1	497
chr13	7700438 2	7700445 8	CCCTTTATTATCCAAGTGGTTTCC TGCTCTTCAAACCTTCTTTCAA ATTTTGTCTCCTACTTAAAACAA	2	498

[0317]

			GTTAGG		
chr13	8164607 5	8164615 1	CCTTCTGTTGAGACCTACTGCTA AGAAAACAAAAAAGGTTTCCTTTC AAATATTATTGTGAATCAATAAT GTACCTGG	3	499
chr13	8375585 4	8375593 1	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTC ATTGATGGACCTGAAAGTGATGG GGAGAATGG	1	500
chr13	8971919 9	8971927 5	CCATTCTCCCTTCACTTTCAGTTA CACCAATCAAACGTAGGTTTGGT CTTTTCACATAGTCCCATATTTCT TGGAGG	2	501
chr13	1020105 74	1020106 50	CCTAGGGAAGTGATCATAGCTGA GTTTCTGGAAAACCTAGGTTTT AAAGTTGAGGAGACTTAAGTCCA AACCTGG	3	502
chr13_KI 270841v 1_alt	124240	124316	CCATTCTCCCTTCACTTTCAGTTA CACCAATCAAACGTAGGTTTGGT CTTTTCACATAGTCCCATATTTCT TGGAGG	2	503
chr14	2598064 6	2598072 3	CCTCCAAGAAATATGGGACTATG TGAAAAGACTAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	504
chr14	3584278 6	3584286 3	CCATTCTCCCTGTCACTTTCAGGT ATGCCAGTCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	505
chr14	4264640 0	4264647 7	CCTCCAAGAAATATGGGACTATG TAAAAGACGAAACCTACGTTTG ATTGGTGTACTTAAAAGTGACGA GGAGAATGG	1	506
chr14	4906324 2	4906331 9	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTTGTGTACCTGAAAGTGATGG GGAGAATGG	1	507
chr14	4913037 9	4913045 6	CCATTCTCCCCGTCACTTTCAGGC ACACCAATCAAACGTAGGTTTAG TCTTTTCACATAGTCCCATATTTT TTAGAGG	1	508
chr14	5135234 2	5135241 8	CCTTAATGCATTCATATTTTCATAT TTAAATAAAACCATGGTTTCCC ACAGAGTGACTTCTACTCTAAGA AATGGGG	2	509
chr14	5135234 2	5135241 7	CCTTAATGCATTCATATTTTCATAT TTAAATAAAACCATGGTTTCCC ACAGAGTGACTTCTACTCTAAGA AATGGG	4	510
chr14	6083584 2	6083591 9	CCGTTCTTTCCGTCACTTTCAGGT ACACCAGTCAAACGTAGGTTTGG	1	511

[0318]

			TCTTTTCACATAGTCCCATATTTCTTGGAGG		
chr14	6652907 2	6652914 8	CCATTCTCCCCATCACTTTCATGTACACCAATCAAACGTAGGTTTGGTCTTTGTAAACATAGTCCCATATTCTTTGG	3	512
chr14	7921087 3	7921094 9	CCCTATAAAGCTTAGAGAAACACAGGGCTCTTTAAACGATCCTTTTTCTCTTTTCTGTTTTAAATTCATCACTTGG	3	513
chr14	7921087 4	7921094 9	CCTATAAAGCTTAGAGAAACACAGGGCTCTTTAAACGATCCTTTTTCTCTTTTCTGTTTTAAATTCATCACTTGG	4	514
chr14	8537154 1	8537161 8	CCATTCTCCCCATCACTTTCAGGTACACTAATCAAAGGTAGGTTTGGTCTTTTCACATGGTCCTATATTTCTTGGAGG	1	515
chr14	9291871 3	9291879 0	CCCCATAGCACGATCACATGGGACATTCAGGGGAAAGCAACCTTTTCCAGGAAGGAAAACCCAATGCTGGGACCCAGG	1	516
chr14	9291871 4	9291879 0	CCCATAGCACGATCACATGGGACATTCAGGGGAAAGCAACCTTTTCAGGAAGGAAAACCCAATGCTGGGACCCAGG	2	517
chr14	1033868 21	1033868 97	CCTTTTAGCGCTCACAGGCTATGGTTTTATAAAAGGAACCTTTGATTTTGTTCATGTGAAACTACAAAATGCCAGG	2	518
chr14_KI 270847v 1_alt	33275	33352	CCCCATAGCACGATCACATGGGACATTCAGGGGAAAGCAACCTTTTCCAGGAAGGAAAACCCAATGCTGGGACCCAGG	1	519
chr14_KI 270847v 1_alt	33276	33352	CCCATAGCACGATCACATGGGACATTCAGGGGAAAGCAACCTTTTCAGGAAGGAAAACCCAATGCTGGGACCCAGG	2	520
chr15	2063056 6	2063064 3	CCTCCAAGAAATATTGGAGTATGTGATAAGACCAAACCTTCGTTTACTGGTGTACCTGAAAGTGATGGGAGAATGG	1	521
chr15	2167510 3	2167518 0	CCATTCTCCCCGTCACTTTCAGGTACACCAATCAAACGTAGGTTTGGTCTTTTCACATAGTCCCATATTTCTTGGAGG	1	522
chr15	2211757 1	2211764 8	CCATTCTCCCCGTCACTTTCAGGTACACCAATCAAACGTAGGTTTGGTCTTTTCACATAGTCCCATATTTCTTGGAGG	1	523
chr15	2236974	2236982	CCATTCTCCCCATCACTTTCAGGT	1	524

[0319]

	4	1	ACACCAGTCAAACGAAGGTTTGG TCTTATCACATACTCCAATATTTT TTGGAGG		
chr15	4230283 2	4230290 9	CCTCCAAGATATATGGGACTATG TGAAAAGGCCAAACCTACCTTTG ATTGATACACCTGAAAATGACAG GGAGAATGG	1	525
chr15	4996760 1	4996767 8	CCTCCAAGAAATATGCGACTATG TGAAAAGACCAAACCTACGTTTC ATTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	526
chr15	8396450 1	8396457 7	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG TTTGGTGTACCTGAAAGTGAGGG GAGAATGG	3	527
chr15	8726138 8	8726146 5	CCATTCTCCTCATCACTTTCAAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCTTATATTTT TTGGAGG	1	528
chr15_KI 270727v 1_rando m	409348	409425	CCATTCTCCCGTCACTTTCAAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	529
chr15_KI 270851v 1_alt	14235	14312	CCATTCTCCCGTCACTTTCAAGT ACACCAGTCAAACGAAGGTTTGG TCTTATCACATACTCCAATATTTT TTGGAGG	1	530
chr15_KI 270852v 1_alt	440099	440176	CCATTCTCCCGTCACTTTCAAGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	531
chr16	2212367 1	2212374 8	CCAGCAGAAGAATCTGGGGCAC AGTCTGTGAAAAAAGGTACCTTT CTTAAGCAGGGTTCTTATCCTTC ATGGGTCTGG	1	532
chr16	2555762 3	2555770 0	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGTTGTACCTGAAAGTGAGGG GGAGAATGG	1	533
chr16	3642717 9	3642725 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	534
chr16	3647645 0	3647652 6	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	535
chr16	3651246 9	3651254 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	536

[0320]

chr16	3652096 4	3652104 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CACAGCAGATTTGAAACACTGTT TTTCTGG	2	537
chr16	3652470 4	3652478 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	538
chr16	3656681 2	3656688 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	539
chr16	3657360 3	3657367 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	540
chr16	3666769 4	3666777 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	541
chr16	3667732 0	3667739 6	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	542
chr16	3668309 6	3668317 2	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	543
chr16	3669125 1	3669132 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	544
chr16	3671095 1	3671102 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	545
chr16	3675036 4	3675044 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	546
chr16	3679145 5	3679153 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CACAGCAGATTTGAAACACTGTT TTTCTGG	2	547
chr16	3685668 3	3685675 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	548
chr16	3692665 5	3692673 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT	2	549

[0321]

			TTTCTGG		
chr16	3693175 2	3693182 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	550
chr16	3694805 8	3694813 4	CCTTGTGTTGTGTGTATTCAACTC ACCGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	551
chr16	3697454 1	3697461 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	552
chr16	3698133 1	3698140 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	553
chr16	3699083 9	3699091 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	554
chr16	3702107 5	3702115 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	555
chr16	3704281 2	3704288 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	556
chr16	3708597 1	3708604 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	557
chr16	3712946 2	3712953 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	558
chr16	3714611 0	3714618 6	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CACAGCAGATTTGAAACACTGTT TTTCTGG	2	559
chr16	3715730 9	3715738 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	560
chr16	3718311 8	3718319 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	561
chr16	3719092 4	3719100 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA	2	562

[0322]

			CAGAGCAGATTTGAAACACTGTT TTTCTGG		
chr16	3722180 8	3722188 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	563
chr16	3725950 1	3725957 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	564
chr16	3727240 9	3727248 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	565
chr16	3728192 3	3728199 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGTAACACTGTT TTTCTGG	2	566
chr16	3734647 2	3734654 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	567
chr16	3735700 0	3735707 6	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	568
chr16	3737330 1	3737337 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	569
chr16	3741949 8	3741957 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	570
chr16	3743071 4	3743079 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	571
chr16	3745584 5	3745592 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	572
chr16	3745855 8	3745863 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	573
chr16	3748612 7	3748620 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	574
chr16	3752518	3752525	CCTTGTGTTGTGTGTATTCAACTC	2	575

[0323]

	3	9	ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTGTGG		
chr16	3753673 5	3753681 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	576
chr16	3755473 0	3755480 6	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	577
chr16	3757578 4	3757586 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	578
chr16	3757748 3	3757755 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	579
chr16	3758359 8	3758367 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	580
chr16	3769636 8	3769644 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTCCA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	581
chr16	3770452 4	3770460 0	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	582
chr16	3770622 3	3770629 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	583
chr16	3770894 1	3770901 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	584
chr16	3776362 2	3776369 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	585
chr16	3777211 5	3777219 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	586
chr16	3779181 5	3779189 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	587

[0324]

chr16	3779622 9	3779630 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	588
chr16	3779792 8	3779800 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	589
chr16	3784345 3	3784352 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	590
chr16	3784854 8	3784862 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	591
chr16	3786484 6	3786492 2	CCTTGTGTTGTGTGTATTCAACTC ACCGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	592
chr16	3790255 0	3790262 6	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	593
chr16	3790730 7	3790738 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	594
chr16	3792803 3	3792810 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	595
chr16	3795926 2	3795933 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	596
chr16	3796435 5	3796443 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	597
chr16	3797488 1	3797495 7	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAAACTGTT TTTCTGG	2	598
chr16	3798778 9	3798786 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTAAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	599
chr16	3799458 6	3799466 2	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT	2	600

[0325]

			TTTGTGG		
chr16	3800647 9	3800655 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	601
chr16	3801156 7	3801164 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	602
chr16	3804009 6	3804017 2	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	603
chr16	3804145 6	3804153 2	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	604
chr16	3806217 9	3806225 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	605
chr16	3810293 7	3810301 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	606
chr16	3812841 2	3812848 8	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	607
chr16	3813180 9	3813188 5	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	608
chr16	3814472 3	3814479 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	609
chr16	3816884 5	3816892 1	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	610
chr16	3820928 7	3820936 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	611
chr16	3821098 6	3821106 2	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGATTTGAAACACTGTT TTTCTGG	2	612
chr16	3822966 7	3822974 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA	2	613

[0326]

			CAGAGCAGATTTGAAACACTGTT TTTCTGG		
chr16	4742403 7	4742411 4	CCATTCTCCCTATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	614
chr16	6073054 9	6073062 5	CCTCGTCACTGCCAGATTTTGTG GCTACCAGCAAAGGATCGTTTTA AGCTGCAACTCAGGAAATTGAGA AAATATGG	2	615
chr16	7254501 4	7254509 1	CCTCCAAGAAATATGGGACTATG TGAAAAACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	616
chr16	8194550 3	8194557 9	CCCTGTGTTCTTTTATACTAAAAC AAGCCAGCAAACCAACCTTTGAG ATGTGTTGCCTTAAACATTACTG AATGGGG	2	617
chr16	8194550 3	8194557 8	CCCTGTGTTCTTTTATACTAAAAC AAGCCAGCAAACCAACCTTTGAG ATGTGTTGCCTTAAACATTACTG AATGGG	4	618
chr17	1647402 4	1647410 0	CCGAGAAACGGCTTTAGCAACAA ATAAATATCAAAGGATGCTTTC TCTTCAGAATAATCTAAAGTAAG TTGGGAGG	3	619
chr17	3443851 2	3443858 9	CCATGTTACTCCGGATAAGGACA GCAAAGGAGGAAAGGAACCTTT TCTGGGCCACCAGAAGGATGAGC TTGGGCTTGG	1	620
chr17	4369078 2	4369085 9	CCCAGGGATATGCTGGCCACGGG GAGGAGCCGAAACCAACCTTTG TGTCCTGTGTAGTGACAAGTGC CTTTGGAGG	1	621
chr17	4369078 3	4369085 9	CCAGGGATATGCTGGCCACGGGG AGGAGCCGAAACCAACCTTTGT GTCCTGTGTAGTGACAAGTGCC TTTGGAGG	2	622
chr17	6915629 8	6915637 5	CCTTAGGGACCCATAATGGCCAC AACCAGGAGAAAAGCAAGCTTT GATGCTTAAACACTACTTACAGA CATGTACAGG	1	623
chr17	7459522 8	7459530 5	CCTGCCTCTGTTCCCTCCTCCTGA TGGTGGCGGAAAGGATGCTTTTG CCAGATCAACAGTCACACACAAC ACACCAGG	1	624
chr17	8319164 4	8319172 1	CCTGACTCCAGCCCTCCTTGACA AGGTCTCCGTAAGCATGCTTTC TCTTAGGGACCCTCAGAGGGAGG CTTGGTGGG	1	625
chr17	8319164	8319172	CCTGACTCCAGCCCTCCTTGACA	3	626

[0327]

	4	0	AGGTCTCCGTAAAGCATGCTTTC TCTTAGGGACCCTCAGAGGGAGG CTTGGTGG		
chr18	3513522 4	3513530 0	CCTTATTTGGAATGTGACAAGAC CCATTTGTTTAAACCTTGGTTTTT ATGCAGAAAGAAAAGGAAGGCT GCAGTGGG	3	627
chr18	3891886 1	3891893 8	CCATTCTCCCTGTCACCTTCAGGT ACACTAATCAAACGTAGGTTTGC TGTTTTTACATAGGCTCATATTC TTGGAGG	1	628
chr18	4547658 9	4547666 6	CCATTCTCCCATCACTTTCAGGT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	629
chr18	4864082 1	4864089 6	CCTGTTTGTATTTTAGCTAATGT CAAAAAGAAAACCTTGCTTTTTT TGAACCCTTTCAGAGGCAGAAAG TGGGGG	4	630
chr18	7109673 2	7109680 8	CCATTTTCCCCACCACTTTCACGT ACAGCAATCAAACGTAGGTTTGG TCTTTTACTAGTCCCATATTTCT TGGAGG	3	631
chr19	2495784 4	2495792 0	CCTTGTAGTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT GTTGTGG	2	632
chr19	2501531 6	2501539 2	CCTTGTAGTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCATACTTGAAACACTCTT TTTGTGG	2	633
chr19	2507411 9	2507419 5	CCTTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	634
chr19	2582786 1	2582793 7	CCTTGTTGTGTGTATTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAATACTCTT TTTGTGG	2	635
chr19	2605405 6	2605413 2	CCTTGTAGTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCATACTTGAAACACTCTT TTTGTGG	2	636
chr19	2621177 7	2621185 3	CCTTGATTGTGAGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	637
chr19	2648367 0	2648374 6	CCTTGTTGTGTGTCTTCAACTC ACAGAGTTAAACGATGCTTTACA CAGAGTAGACTTGAAACACTCTT TTTCTGG	2	638

[0328]

chr19	2663651 6	2663659 2	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGTAACACTCTT TTTGTGG	2	639
chr19	2663787 7	2663795 3	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACGTGAAACACTCTT TTTGTGG	2	640
chr19	2675022 3	2675029 9	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGCAGACTTGAAACACTCTT TTTGTGG	2	641
chr19	2684115 8	2684123 4	CCTTGTGTTGTGTGTATTCAACTC ACAGAGTTAAACGATCCTTTACA CAGAGGAGACTTGTAACACTCTT TTTGTGG	2	642
chr19	2851722 0	2851729 7	CCAGGAAAAAATTTAAACTTTCT TAACTTGATAAAAGGTAGCTTTC AAAACCTACAATAAATAACATAC TTAGAGTGG	1	643
chr19	3456682 1	3456689 8	CCATTCTCCTCGTCACTTTCAGGT ACACCAAACAAACGTAGGTTTGG TCTTTTACGTAGTCCCATATTTT TTGGAGG	1	644
chr19	5226177 0	5226184 7	CCCTCTTGAAGTTAGGGAAGTAG CATTTAAGGGAAACGTAGCTTTA CTATTAAGAATTTCAAACAGCAC TTGTCAGGG	1	645
chr19	5226177 0	5226184 6	CCCTCTTGAAGTTAGGGAAGTAG CATTTAAGGGAAACGTAGCTTTA CTATTAAGAATTTCAAACAGCAC TTGTCAGG	3	646
chr19	5226177 1	5226184 7	CCTCTTGAAGTTAGGGAAGTAGC ATTTAAGGGAAACGTAGCTTTAC TATTAAGAATTTCAAACAGCACT TGTCAGGG	2	647
chr19	5226177 1	5226184 6	CCTCTTGAAGTTAGGGAAGTAGC ATTTAAGGGAAACGTAGCTTTAC TATTAAGAATTTCAAACAGCACT TGTCAGG	4	648
chr20	1115139 2	1115146 9	CCATTCTCCCGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTACATATTCCCATATTTT TTGGAGG	1	649
chr20	1402706 7	1402714 3	CCATTCTCCCTTCACTTTCAGGTA CACCAATCAAACGTAGGTTTGGT CTTTTACATAGTCCCATATTTT TGGAGG	2	650
chr20	5061539 9	5061547 6	CCTATAGTCTCAGTTACTTGGGA GGCTGAGGTAAGGATCGTTTG AGCCAGGAGGTGGAGGTTGCA	1	651

[0329]

			GTGAGCCGGG		
chr20	5061539 9	5061547 5	CCTATAGTCTCAGTACTTGGGA GGCTGAGGTAAAAGGATCGTTTG AGCCAGGAGGTGGAGGTTGCA GTGAGCCGG	3	652
chr20	6090941 4	6090949 0	CCTTTCCCAACTCTGCTATTGCC CCACATCCTAAAGGAACCTTTCT TTTTTATATATTTTATTTTAAGT TCCAGG	3	653
chr21	1622608 6	1622616 3	CCTCCAAGAAATATGGAACATG TGAAAAGACCAAACCTACGTTTG ATTGACGTACCTGAAAGTGACAG GGAGAATGG	1	654
chr21	1783523 4	1783530 9	CCTCTTCTGAAAGCATTGATAAT CAACATTTTAAACGTAGCTTTTC CCCATATTGCTAGGAAGGCTCAT TCCCGG	4	655
chr21	1942563 6	1942571 3	CCTCCAAGAAATATGGGACTATG TGAAAAGGCCAAACCTACGTTTG ATTGCTGTACCCGAGAGTGACGG GGAGAATGG	1	656
chr21	3222095 8	3222103 5	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	657
chr21	3433587 7	3433595 3	CCCGGGCCTGGGTGCCCAGTGC CAGTGGTCAGAAAGGTTGCTTTG GTGTTTTTCATTGTTAGTGAGAC AGAGATGG	3	658
chr21	3433587 8	3433595 3	CCGGGGCCTGGGTGCCCAGTGCC AGTGGTCAGAAAGGTTGCTTTGG TGTTTTTCATTGTTAGTGAGACA GAGATGG	4	659
chr21	3631527 6	3631535 3	CCATTCTCCCATCATTTTCAGGT ACACCAATCAAACGTAGGTTTGA TCTTTTCACATAGCCCCATATTC TTGGAGG	1	660
chr21	4154795 2	4154802 8	CCACCAGCACTTCTGTTAGAAGT TGCAGCAGAGAAAGGATCCTTTA GGCACATCTCCAGATCCTTGCG AAGAGGGG	3	661
chr22	1897319 4	1897327 1	CCTGTGCCAGGGTCCTTCCACTG GGACTGGCAGAAACGTAGGTTTG CATGGAGTGAGAAGCAGGGGAG AGGTTGAGGG	1	662
chr22	1897319 4	1897327 0	CCTGTGCCAGGGTCCTTCCACTG GGACTGGCAGAAACGTAGGTTTG CATGGAGTGAGAAGCAGGGGAG AGGTTGAGG	3	663
chr22	2026546 2	2026553 9	CCCTCAGCCTCTCCCCTGCTTCTC ACTCCATGCAAACCTACGTTTCT	1	664

[0330]

			GCCAGTCCCAGCAGAAGGACCCT GGCACGGG		
chr22	2026546 2	2026553 8	CCCTCAGCCTCTCCCCTGCTTCTC ACTCCATGCAAACCTACGTTTCT GCCAGTCCCAGCAGAAGGACCCT GGCACGGG	3	665
chr22	2026546 3	2026553 9	CCTCAGCCTCTCCCCTGCTTCTCA CTCCATGCAAACCTACGTTTCTG CCAGTCCCAGCAGAAGGACCCTG GCACGGG	2	666
chr22	2026546 3	2026553 8	CCTCAGCCTCTCCCCTGCTTCTCA CTCCATGCAAACCTACGTTTCTG CCAGTCCCAGCAGAAGGACCCTG GCACGGG	4	667
chrX	2730099 8	2730107 5	CCTCCAAGAAATATGGGGCTATG TGAAAAGACCAAACCTACCTTTG ATTGGTGTATCTGAAAGTGACGG GGAGAATGG	1	668
chrX	2845666 6	2845674 3	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTTGTGTACCTGAAAGTGATGG GGAGAATGG	1	669
chrX	3563498 5	3563506 2	CCATTCTCCCCGTCACCTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCTCATTGTCCCATATTC TTGGAGG	1	670
chrX	3946014 8	3946022 3	CCCATCAAGAGCGGTTGTGCATG GCAACAGTAAAAGGATGGTTTGT TACACTAGTACAAAAAGAGGTG GCCAGAGG	4	671
chrX	4392640 3	4392648 0	CCATTCTCTCTGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTC TTGGAGG	1	672
chrX	4425460 0	4425467 7	CCTCCAAGAAATACGGGACTATG TGAAAAGACCAAACGTACGTTTG ATTGGTGTACCTGAAAGTGATAG GGAGAATGG	1	673
chrX	4608860 2	4608867 9	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACTG GGAGAATGG	1	674
chrX	5022287 4	5022295 1	CCATTCTCCCTGTCACTTTCAGGT ACACGAATCAAACGTAGGTTTCA TCTTTTCACATAGTCCCATATTC TTAGAGG	1	675
chrX	5741683 5	5741691 1	CCATTCTCTCTGTCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTTTCACATATT TCTTGG	3	676
chrX	5785646	5785654	CCTCCAAGAAATATGGGACTATG	1	677

[0331]

	6	3	TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAA GGAAAATGG		
chrX	6270247 9	6270255 6	CCTGAAAAACATTGTTTCCAACC TGGTAAATCAAAGGAAGGTTTA ACTTTGTTAGATAAGTCCACATA TCACCAAGG	1	678
chrX	6306712 9	6306720 6	CCTCCAAGAAATGTGGGACTATG GGAAAAGACCAAACCTACCTTTG TTTGGTGTACCTGAAAGTGACGG GGAGAAAGG	1	679
chrX	6493625 0	6493632 7	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTC ATTGGTGTACCTGAAAGTGATGG GTAGAATGG	1	680
chrX	6672009 9	6672017 6	CCTACAAGAAATATGGGACTATG GGAAAAGACCAAACCTACGTTTG ATTGGTACACTGGAAAGTGACAG GGATAATGG	1	681
chrX	6852908 6	6852916 3	CCATTCTCCCTGTCACCTTCTGGT ACACCAATCAAAGGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	682
chrX	7389399 4	7389407 1	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGATGG GGAGAATGG	1	683
chrX	7572320 1	7572327 8	CCATTCTCTTTGTCACCTTTCAGGT ATACCAATCAAACGTTGGTTTGG TCTTTTGCATAGTCCCATATTTT GTGGAGG	1	684
chrX	7581565 9	7581573 6	CCTCCAAGAAATATGAGACTATG TGAAAAGACCAAACCTACGTTTG ATTAGTGTACCTGAAAATGATGG GGAGAATGG	1	685
chrX	8096710 3	8096718 0	CCATTCTTTCTGTCACCTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	686
chrX	8993642 5	8993650 2	CCATTCTCCCTGTCACCTTTCAGGT ACACCAATCAAACGTAGGTTTGT TCTTTTCACATAGTCCCATATTTT TTGGAGG	1	687
chrX	9103876 8	9103884 5	CCATTATCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TTTTTTCACATAGTTCAATATTTT TTTGAGG	1	688
chrX	9147127 1	9147134 8	CCTCCAAGAAATATGGGACTATC TGAAAAGATCAAACCTACGTTTG ATTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	689

[0332]

chrX	9642818 0	9642825 7	CCTTTCTCCCCATCACTTTCAGGT ACACCAATCAAACGTAGGTTTGG TCTTTTCATATAGTCCCATATTTT TTGGAGG	1	690
chrX	1002682 91	1002683 68	CCTCCAAGAAATATGGGACTATG TGCAAAGATCAAACCTACGTTTG ATTGCTGTACCTGAAAGTGATGG GGAGAATGG	1	691
chrX	1058110 46	1058111 23	CCATTCTCCCCATCACTTTCAGGT ACACCAGTCAAACGTAGGTTTGG TCTTTTCACATAATCCCATATTTT TTGGAGG	1	692
chrX	1156730 65	1156731 41	CCTCCAAGAAGTATGGGACCATG GAAAAGATCAAACCTACGTTTGA CTGGTGTACCTGAAAGTGACTGG GAGAATGG	2	693
chrX	1172698 46	1172699 23	CCTCCAAGAAATATGGGACTATG TGAAAAGACCAAACCTACGTTTG ATTGGAGTACTTGAAAATGACAG GGATAATGG	1	694
chrX	1391913 69	1391914 45	CCTTTAAAGACATGCTCTTTGTG CCAGAAATTCAAAGGTTGCTTTT ATGTCCAGTGGGGTGGAGGGAG GAAGCTCGG	3	695
chrX	1479886 14	1479886 91	CCATTCTCCCCGTCCTTTCAGG GACCTCAATCAAACGTAGGTTTT GTCTTTTCACATAGTCCCATATTT CTTGGAGG	1	696
chrX	1553210 41	1553211 18	CCTCCAAGAAATATAGGACTATG TGAAAAGACCAAACCTACGTTTG ACTGGTGTACCTGAAAGTGACAG GGAGAATGG	1	697
chrY	1510939 1	1510946 8	CCATTCTCCCCATCACTTTCAGGT ACACCAATCAAAGGTAGGTTTGG TCTTTTCACATAGTCCGATATTTT CTGCAGG	1	698
通过搜索CCX ₍₃₀₋₃₁₎ -AAASSWWSSTTT-X ₍₃₀₋₃₁₎ -GG (SEQ ID NO: 699)鉴定染色体位点, 其中W是T或A并且S是G或C。模式1是CCX ₍₃₁₎ -AAASSWWSSTTT-X ₍₃₁₎ -GG (SEQ ID NO: 699), 2是CCX ₍₃₀₎ -AAASSWWSSTTT-X ₍₃₁₎ -GG (SEQ ID NO: 699), 3是CCX ₍₃₁₎ -AAASSWWSSTTT-X ₍₃₀₎ -GG (SEQ ID NO: 699), 并且4是CCX ₍₃₀₎ -AAASSWWSSTTT-X ₍₃₀₎ -GG (SEQ ID NO: 699)。仅显示+链并且开始和结束对应于染色体(GRCh38)中的第一个和最后一个碱基对或适用时的备用组装。					

[0333] DNA测序

[0334] 如上述一式六份进行293T细胞的转染并温育72小时。收获细胞并合并重复。使用涉及碱性裂解和旋转柱纯化的改良HIRT提取来提取附加型DNA,基本上如所述(Quan et al., Circular polymerase extension cloning of complex gene libraries and pathways. PloS one 4, e6441 (2009); 和Hillson (2010), vol. 2015, pp. CPEC protocol; 其每篇的全部内容在此通过引用并入)。简而言之,收获后,将HEK293T细胞在500 μ L的冰冷的PBS中洗涤,重悬于250 μ L GTE缓冲液(50mM葡萄糖,25mM Tris-HCl,10mM EDTA并且pH8.0)

中,在室温温育5分钟,并用200 μ L裂解缓冲液(200mM NaOH,1%十二烷基硫酸钠)在冰上裂解5分钟。用150 μ L的乙酸钾溶液(5M乙酸盐,3M钾,pH 6.7)中和裂解。通过在21,130g离心15分钟沉淀细胞碎片,并将裂解物应用于Econospin旋转柱(Epoch Life Science, Missouri City, TX)。用750 μ L洗涤缓冲液(Omega Bio-tek, Norcross, GA)洗涤柱两次,并在45 μ L TE缓冲液, pH8.0中洗脱。

[0335] 用RecBCD(10U)按照制造商的说明将分离的附加型DNA在37 $^{\circ}$ C消化2小时,并用MinElute Reaction Cleanup试剂盒(Qiagen, Valencia, CA)纯化到10 μ LEB中。用5 μ L的附加型提取物转化Mach1-T1化学感受态细胞,并在选择羧苄青霉素抗性的琼脂糖平板(含有50 μ g/mL羧苄青霉素)上铺板。用引物pCALNL-for-1对个别菌落进行测序以确定重组率。测序读段揭示了“左”完整的非重组recCas9位点、预期的重组产物、具有小插入/缺失的“左”非重组位点的罕见实例、或大缺失产物的一个情况。

[0336] 对recCas9催化基因组缺失的分析

[0337] 在24孔胶原处理的平板中以每孔 6×10^5 个细胞的密度接种HEK293T细胞并生长过夜(Corning, Corning, NY)。在Opti-MEM(ThermoFisher Scientific, Waltham, MA)中使转染反应物达到100 μ L的终体积。对于每次转染,将90ng的每种引导RNA表达载体、20ng的pmaxGFP(Lonza, Allendale, NJ)和320ng的recCas9表达载体与Opti-MEM(ThermoFisher Scientific, Waltham, MA)中2 μ L Lipofectamine2000组合并加入各孔中。48小时后,收获细胞并在BD FACS AriaIIIu细胞分选仪上分选GFP转染对照。使用100 μ m喷嘴在纯度模式下分选细胞,并通过与未转染的细胞比较确定背景荧光。在PBS中在冰上收集分选的细胞,沉淀并用冷PBS洗涤两次。使用E.Z.N.A.组织DNA试剂盒(Omega Bio-Tek, Norcross, GA)收获基因组DNA并在100 μ L Eb中洗脱。使用在Tecan Infinite M1000Pro荧光酶标仪(plate reader)上测量的Quant-iT PicoGreen dsDNA试剂盒(ThermoFisher Scientific, Waltham, MA)定量基因组DNA。

[0338] 使用补充有3%DMSO的Q5热启动聚合酶2xMaster Mix进行巢式PCR,并用HyClone水,分子生物学级(GE Life Sciences, Logan, UT)稀释。使用引物对FAM19A2-F1和FAM19A2-R1(表5)用20ng的基因组DNA作为模板以25 μ L规模进行初级PCR。初级PCR条件如下:98 $^{\circ}$ C1分钟,35个循环(98 $^{\circ}$ C10秒、59 $^{\circ}$ C30秒、72 $^{\circ}$ C30秒),72 $^{\circ}$ C1分钟。使用引物FAM19A2-F2和FAM19A2-R2,将1:50稀释的初级PCR用作次级PCR的模板。次级PCR条件如下:98 $^{\circ}$ C1分钟,30个循环(98 $^{\circ}$ C10秒、59 $^{\circ}$ C20秒、72 $^{\circ}$ C20秒),72 $^{\circ}$ C1分钟。通过在TAE中的1%琼脂糖凝胶上与1Kb Plus DNA梯(ladder)(ThermoFisher Scientific, Waltham, MA)一起电泳来分析DNA。在Qiagen Minelute柱(Valencia, CA)上使用制造商的方案纯化待Sanger测序的材料。来自3个生物学重复的模板DNA用于三个独立的基因组巢式PCR。

[0339] 考虑到一组完整的人染色体重约3.6pg(3.3×10^9 bp $\times 1 \times 10^{-1}$,计算检出限。因此,用20ng的基因组DNA模板接种的PCR反应物含有约5500组染色体。

[0340] 为了定量基因组缺失,对于3个生物学重复中的每一个,一式三份进使用上述条件行巢式PCR。使用两倍稀释系列的基因组DNA作为模板,从未稀释的储液开始(对于样品1,47.17ng/ μ L;对于样品2,75.96ng/ μ L;并且对于样品3,22.83ng/ μ L)以减少潜在的移液误差来源。假定可以观察到缺失PCR产物的最低DNA浓度含有每个总基因组DNA的单一缺失产物。

[0341] 可以推断给定量的模板DNA中存在的基因组的数量,并因此可以确定FAM19A2基因

座处recCas9的最小缺失效率的估计值。例如,以20ng基因组DNA模板开始,以两倍稀释系列为例。巢式PCR后,只有接种20ng的孔产生正确的PCR产物。每个基因组3.6pg,该PCR含有大约5500个基因组,并且由于必须存在至少一个重组基因组,因此最小缺失效率为5500分之一或0.018%。

[0342] 使用基因组模板的有限稀释来定量基因组DNA的水平,因为使用定量PCR(qPCR)来确定基因组编辑的绝对水平将需要一组PCR条件,其仅明确地且仅特异性地扩增自重组后基因组DNA。如图5B中所示,使用基因组DNA作为模板的初级PCR导致大约2.5kb脱靶带作为优势种类;需要使用巢式引物进行第二轮的PCR以揭示引导RNA-和recCas9依赖性基因组编辑。

[0343] 结果

[0344] 将Gin重组酶融合至dCas9

[0345] 最近已经证明dCas9的N端可以融合至FokI核酸酶催化域,产生二聚体dCas9-FokI融合物,其切割侧翼为两个引导RNA指定序列的DNA位点(参见例如,Guilinger et al., Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. Nature biotechnology, (2014); Tsai et al., Dimeric CRISPR RNA-guided FokI nucleases for highly specific genome editing. Nature biotechnology, (2014); 其每篇的全部内容在此通过引用并入)。使用相同的融合方向将dCas9与Gin β (Barbas及其同事先前演化的二聚体Gin转化酶的高活性催化域)连接(Gaj et al., A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells. Nucleic acids research 41, 3937-3946 (2013), 其全部内容在此通过引用并入)。Gin β 混杂地重组与天然核心序列CTGTAAACCGAGGTTTTGGA (SEQ ID NO:700) 相关的几个20-bp核心“gix”序列(Gaj et al., A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells. Nucleic acids research 41, 3937-3946 (2013); Klippel et al., The DNA Invertase Gin of Phage Mu-Formation of a Covalent Complex with DNA Via a Phosphoserine at Amino-Acid Position-9. Embo Journal 7, 1229-1237 (1988); Mertens et al., Site-specific recombination in bacteriophage Mu: characterization of binding sites for the DNA invertase Gin. The EMBO journal 7, 1219-1227 (1988); Plasterk et al., DNA inversions in the chromosome of Escherichia coli and in bacteriophage Mu: relationship to other site-specific recombination systems. Proceedings of the National Academy of Sciences of the United States of America 80, 5355-5358 (1983); 其每篇的全部内容在此通过引用并入)。引导RNA将recCas9二聚体定位于侧翼为两个引导RNA指定的序列的gix位点,使得Gin β 域能够以引导RNA编程的方式催化DNA重组(图1D)。

[0346] 为了测定所得的dCas9-Gin β (recCas9) 融合物,构建了含有两个recCas9靶位点的报告物质粒,两个recCas9靶位点位于阻断EGFP转录的poly-A终止子的侧翼(图1A-1C)。每个recCas9靶位点由侧翼为与引导RNA前间隔区序列匹配的位点的gix核心假位点组成。重组酶介导的缺失去除了终止子,恢复了EGFP的转录。用该报告物质粒、转录引导RNA的质粒和产生候选dCas9-Gin β 融合蛋白的质粒共转染HEK293T细胞,并使用表现出EGFP荧光的细

胞级分评估每种融合构建体的相对活性。

[0347] 影响recCas9组分的结构的参数,包括核心gix位点和引导RNA结合位点之间的间距(0至7bp),以及dCas9和Gin β 部分之间的接头长度((GGS)₂(SEQ ID NO:182)、(GGS)₅(SEQ ID NO:701)或(GGS)₈(SEQ ID NO:183))是变化的(图2A-2F)。大多数融合结构导致无观察到的引导RNA依赖性EGFP表达(图1C-1D)。然而,当存在匹配但不错配的引导RNA时,一个含有8个GGS重复的接头和3-6个碱基对间隔区的融合构建体导致约1%的重组(图2E-2F)。当5-6个碱基对将dCas9结合位点与核心分开时,重组活性始终较高(图2F)。这些结果共同揭示了dCas9和Gin β 之间的特异性融合结构可以在人细胞中与间隔区侧翼的gix相关的核心位点处产生引导RNA依赖性重组活性。8xGGS接头融合构建体称为“recCas9”。

[0348] 用recCas9靶向人基因组中发现的DNA序列

[0349] 观察到的活性的低水平可以由次优的引导RNA序列或核心gix序列引起,与表明引导RNA:Cas9结合的效率呈序列依赖性的之前的报告(参见例如,Xu et al., Sequence determinants of improved CRISPR sgRNA design. *Genome research* 25,1147-1157 (2015),其全部内容在此通过引用并入)一致。此外,尽管目前的优化是用天然gix核心序列进行的(参见例如,Klippel et al., The DNA Invertase Gin of Phage Mu-Formation of a Covalent Complex with DNA Via a Phosphoserine at Amino-Acid Position-9. *Embo Journal* 7,1229-1237 (1988); Mertens et al., Site-specific recombination in bacteriophage Mu: characterization of binding sites for the DNA invertase Gin. *The EMBO journal* 7,1219-1227 (1988); Plasterk et al., DNA inversions in the chromosome of *Escherichia coli* and in bacteriophage Mu: relationship to other site-specific recombination systems. *Proceedings of the National Academy of Sciences of the United States of America* 80,5355-5358 (1983); 其每篇的全部内容在此通过引用并入),但是一些研究已表明锌指-Gin或TALE-Gin融合对略微改变的核心位点是有活性的,并且在某些情况下更具活性。参见例如,Gordley et al., 3rd, Synthesis of programmable integrases. *Proceedings of the National Academy of Sciences of the United States of America* 106,5053-5058 (2009); Gersbach et al., Targeted plasmid integration into the human genome by an engineered zinc-finger recombinase. *Nucleic acids research* 39,7868-7878 (2011); Mercer et al., Chimeric TALE recombinases with programmable DNA sequence specificity. *Nucleic acids research* 40,11163-11172 (2012); Gaj et al., A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells. *Nucleic acids research* 41,3937-3946 (2013); Gordley et al., 3rd, Evolution of programmable zinc finger-recombinases with activity in human cells. *J Mol Biol* 367,802-813 (2007); Gersbach et al., 3rd, Directed evolution of recombinase specificity by split gene reassembly. *Nucleic acids research* 38,4198-4206 (2010); 和Gaj et al., Structure-guided reprogramming of serine recombinase DNA sequence specificity. *Proceedings of the National Academy of Sciences of the United States of America* 108,498-503 (2011); 其每篇的全部内容在此通过引用并入)。因此,靶向在人基因组内发现的序列以测试未修饰的人基因组序列是否能够被recCas9靶

向并测试是否改变引导RNA和核心序列将增加recCas9活性。

[0350] 为了鉴定潜在的靶位点,使用了表征演化的Gin变体的先前发现(参见例如,Gaj et al.,A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells.Nucleic acids research 41,3937-3946 (2013),其全部内容在此通过引用并入)以及上述观察结果。使用该信息,在人基因组中搜索含有CCN₍₃₀₋₃₁₎-AAASSWWSSTTT-N₍₃₀₋₃₁₎-GG (SEQ ID NO:699)的位点,其中W是A或T,S是G或C,N是任何核苷酸。N₍₃₀₋₃₁₎包括以下各项的N:NGG前间隔区相邻基序(PAM)、20碱基对Cas9结合位点、Cas9和gix位点之间5-6碱基对间隔以及gix核心位点的四个最外碱基对。预先确定gix核心位点的内部12个碱基对(AAASSWWSSTTT,SEQ ID NO:699)对Ginβ活性是重要的(参见例如,Gaj et al.,Nucleic acids research 41,3937-3946 (2013))。

[0351] 该搜索揭示了人基因组中约450个此类基因座(表9)。产生报告物构建体,其含有与PCDH15中发现的这些基因组基因座之一相同的序列,并且然后构建引导RNA表达载体以将recCas9导向该序列(图3A)。这些载体编码两对引导RNA,其每对含有与PCDH15假gix位点侧翼的5'和3'区域匹配的间隔区序列。报告物质粒、这些侧翼引导RNA表达载体的组合和recCas9表达载体的共转染导致在11%-13%的转染细胞中的EGFP表达(图3B),表明与图2中所示结果相比,活性提高了10倍以上。这些发现表明,recCas9靶序列的更明智的选择可以导致与人基因组中发现的DNA序列匹配的DNA序列处的重组效率显著提高。

[0352] 接下来,确定是否需要这两个引导RNA序列引起recCas9介导的缺失。用靶向PCDH15假gix核心位点的5'或3'侧翼序列的引导RNA载体的仅一个、PCDH15报告物质粒和recCas9表达载体共转染HEK293T细胞。这些共转染导致2.5-3%EGFP表达(图3B)。靶向性引导RNA的仅一个和recCas9的表达时观察到的低活性水平可能是由过度激活的gix单体形成二聚体的倾向引起的(参见例如,Gaj et al.,Enhancing the Specificity of Recombinase-Mediated Genome Engineering through Dimer Interface Redesign.J Am Chem Soc 136,5047-5056 (2014),其全部内容在此通过引用并入);瞬时二聚化可以偶尔允许单一前间隔区序列将二聚体定位于靶位点。当使用脱靶引导RNA载体时或当用pUC替换recCas9载体时,未检测到高于背景的活性(图3B)。

[0353] 这些发现表明,通过选择不同的靶位点和匹配引导RNA序列,recCas9活性可以比初始实验中观察到的适度活性显著增加。与原始靶序列相比,观察到PCDH15位点上的活性增加大于10倍(比较图3B和图2F)。此外,最大重组活性取决于两个引导RNA和recCas9的存在。

[0354] recCas9的正交性

[0355] 接下来,测试recCas9是否可以以正交方式靶向在人基因组中发现的序列匹配的多个分开的基因座。基于其作为基因组整合的安全港基因座的潜在用途,或者在一种情况下基于其在遗传疾病中涉及的基因内的位置,选择人基因组中recCas9靶位点的子集。

[0356] 为了鉴定这些位点,搜索ENSEMBL(版本81)以鉴定哪些预测的recCas9靶位点落入注释的基因内(参见例如,Cunningham et al.,Ensembl 2015.Nucleic acids research 43,D662-669 (2015),其全部内容在此通过引用并入)。一个此类位点落入FGF14的内含子区域。据信FGF14内的突变会引起脊髓小脑性共济失调(spinocerebellar ataxia) 27(SCA 27)(参见例如,van Swieten et al.,A mutation in the fibroblast growth factor

14gene is associated with autosomal dominant cerebellar ataxia[corrected]. *Am J Hum Genet* 72,191-199 (2003);Brusse et al.,Spinocerebellar ataxia associated with a mutation in the fibroblast growth factor 14gene (SCA27):A new phenotype. *Mov Disord* 21,396-401 (2006);Choquet et al.,A novel frameshift mutation in FGF14causes an autosomal dominant episodic ataxia. *Neurogenetics* 16,233-236 (2015);Coebergh et al.,A new variable phenotype in spinocerebellar ataxia 27 (SCA 27) caused by a deletion in the FGF14gene. *Eur J Paediatr Neurol* 18,413-415 (2014);Shimajima et al.,Spinocerebellar ataxias type 27derived from a disruption of the fibroblast growth factor 14gene with mimicking phenotype of paroxysmal non-kinesigenic dyskinesia. *Brain Dev* 34,230-233 (2012);其每篇的全部内容通过引用并入本文)。最后,手动询问未落入基因内的预测的recCas9靶位点的级分,以确定一些序列是否落入安全港基因座内。使用ENSEMBL中的注释鉴定了基因组靶标,该靶标与Bushman及其同事描述的安全港基因座的五个标准的大多数相匹配 (Cunningham et al.,Ensembl 2015. *Nucleic acids research* 43,D662-669 (2015);和Sadelain et al.,Safe harbours for the integration of new DNA in the human genome. *Nat Rev Cancer* 12,51-58 (2012),其每篇的全部内容通过引用并入本文)。构建了五种报告物和含有与基因组中序列相同的序列的相应的引导RNA载体对。为了评估recCas9在用不同的引导RNA编程时的正交性,测试了五种引导RNA对与五种报告物的所有组合。

[0357] 报告物、引导RNA质粒和recCas9表达载体的共转染显示,所测试的五种报告物中的三种导致与recCas9介导的重组一致的显著水平的EGFP阳性细胞。该EGFP表达严格依赖于recCas9表达载体和与报告物构建体上的靶位点序列匹配的引导RNA质粒的共转染(图4A)。当与非关联报告物质粒共转染时,与关联报告物质粒和recCas9载体共转染时引起重组的相同的引导RNA对不能介导重组(图4A)。这些结果证明recCas9活性是正交的,并且当用与侧翼序列匹配的一对引导RNA编程时,仅催化gix相关核心位点处的重组。当未用表达recCas9和引导RNA的载体转染报告物质粒时,没有观察到高于测定的背景水平的重组酶活性。

[0358] recCas9产物的特性

[0359] 表征recCas9介导的报告物质粒重组的产物以证实EGFP表达是recCas9介导的poly-A终止子序列除去的结果。在与recCas9表达载体和与产生关联或非关联引导RNA对的质粒共转染后,对报告物质粒进行染色体5位点1、染色体12和染色体13 (FGF14基因座)的测序。温育72小时后,提取附加型DNA(如上所述)并转化到大肠杆菌中以分离报告物质粒。对含有报告物质粒的单菌落进行测序(图4B)。

[0360] 预期个别菌落含有未修饰的或重组的报告物质粒(图4C)。对于每个生物学重复,对用从每个转染条件分离的报告物质粒转化的平均97个菌落进行测序。若报告物质粒先前与关联引导RNA质粒和recCas9表达载体共转染,则仅观察到重组质粒(图4D)。在两个单独的实验中,重组质粒的百分比范围从染色体5中的位点1的12%到染色体13中的FGF14基因座的平均32%。因此,测序数据与图4A中更早的流式细胞术分析一致。重组质粒的绝对水平略高于EGFP阳性细胞的百分比(图4)。这种差异可能是由于流式细胞术测定没有报告当在单一细胞中存在多个拷贝的报告物质粒时可发生的多个重组事件而出现;甚至单一重组事

件可以导致EGFP荧光。结果,EGFP阳性细胞的百分比可以对应于重组报告物质粒的实际百分比的下限。或者,差异可以反映质粒大小和转化效率之间的负相关性(参见例如,Hanahan,Studies on transformation of Escherichia coli with plasmids.J Mol Biol 166,557-580(1983),其全部内容在此通过引用并入);重组质粒约为5,700个碱基对,并且可以转化得略好于完整质粒(其约为6,900个碱基对)。

[0361] 由于已经报告锌指重组酶在重组酶核心位点连接处引起突变(参见例如,e.g.,Gaj et al.,A comprehensive approach to zinc-finger recombinase customization enables genomic targeting in human cells.Nucleic acids research 41,3937-3946(2013),其全部内容在此通过引用并入),因此测试了此类诱变是否自recCas9处理发生。在报告物构建体中,recCas9应通过首先切割这两个gix核心位点的中心二核苷酸并然后将两个核心彼此再连接来删除kanR和poly-A终止子(图4C)。因此,重组产物应当是由“左”靶位点的第一半和“右”靶位点的第二半组成的单一重组位点。错误或不完整的反应可以导致其他产物。引人注目的是,所检查的所有134个重组序列都含有预期的重组产物。此外,来自两组分开的转染实验的总共2,317个测序读段显示仅三个含有在其他情况下非重组质粒上潜在的缺失产物的测序读段。

[0362] 在用pUC对照转染并且缺少两个recCas9靶位点以及polyA终止子的染色体12报告物质粒中观察到这些含有缺失的读段之一。该产物归因于在转染、分离或随后操作期间发生的DNA损伤。因为当与报告物和关联引导RNA表达载体共转染时recCas9仅可以定位于序列,所以更相关的度量可以是测量当与关联引导RNA载体和recCas9表达载体共转染报告物质粒时观察到的缺失产物的总数。从与染色体5-位点1报告物和关联引导RNA的共转染测序的总共185个质粒中观察到单一插入/缺失。类似地,在用关联引导RNA和recCas9表达载体转染后,从来自染色体12报告物的204个质粒中观察到一个插入/缺失。值得注意的是,在202个测序读段中,在关联引导RNA和recCas9共转染后,从染色体13报告物中未观察到插入/缺失,尽管导致观察到的最高重组水平。这些观察结果共同表明recCas9主要介导无错重组。

[0363] 总之,这些结果建立了recCas9可以在最小的交叉反应性或副产物形成的情况下靶向人基因组内发现的多个位点。仅在存在关联引导RNA序列和recCas9的情况下,底物经历有效重组,在人细胞中产生干净的重组产物,并且通常不导致核心位点连接处的突变或产物诸如由细胞DNA修复产生的插入/缺失。

[0364] RecCas9介导的基因组缺失

[0365] 最后,研究了recCas9是否能够直接在培养的人细胞的基因组DNA上操作。使用人基因组中潜在的recCas9识别位点的列表(表9),搜索在由recCas9靶向的情况下将产生通过PCR可检测的染色体缺失事件的位点对。设计引导RNA表达载体以将recCas9引导至最接近染色体5-位点1或染色体13(FGF14基因座)的那些recCas9位点,即均显示在瞬时转染测定中重组的位点(图4)。新的靶位点范围为染色体5-位点1的上游约3至23Mbp和下游7至10Mbp,以及染色体13-FGF14位点的上游的12至44Mbp。与这些新的引导RNA对之每种和用于染色体5-位点1或染色体13-FGF14的经验证的引导RNA对共转染recCas9表达载体,但通过基因组PCR未观察到染色体缺失的证据。

[0366] 若recCas9靶位点在基因组上彼此更接近,则认为基因组缺失可能更有效。鉴定了

在FAM19A2的内含子区域内以14.2kb分开的两个recCas9位点;这些位点还含有相同的二核苷酸核心,这应当有助于缺失。FAM19A2是五个密切相关的TAF1A家族基因之一,其编码认为在免疫和神经细胞中具有调节作用的小的分泌蛋白(参见例如Parker et al., Admixture mapping identifies a quantitative trait locus associated with FEV1/FVC in the COPD Gene Study. *Genet Epidemiol* 38,652-659 (2014),其全部内容在此通过引用并入)。位于FAM19A2的内含子序列中的小核苷酸多态性在全基因组关联研究与系统性红斑狼疮(SLE)和慢性阻塞性肺病(COPD)的风险升高有关(参见例如,Parker et al., Admixture mapping identifies a quantitative trait locus associated with FEV1/FVC in the COPD Gene Study. *Genet Epidemiol* 38,652-659 (2014),其全部内容在此通过引用并入);因此,该基因的内含子区域的缺失可以了解这些疾病的原因。将四个引导RNA序列克隆到表达载体中,所述表达载体设计以介导这两个FAM19A2位点之间的recCas9缺失。与recCas9表达载体共转染表达这些引导RNA的载体(图5A)。RecCas9介导的两个位点之间的重组应当导致14.2kb居间区域的缺失。实际上,使用位于两个FAM19A2recCas9靶侧翼的基因特异性引物,通过巢式PCR检测到该缺失事件。仅在从用recCas9和所有四种引导RNA表达载体共转染的细胞分离的基因组DNA中观察到与recCas9介导的缺失一致的预期PCR产物(图5B)。在没有单独的引导RNA表达载体的上游或下游对的情况下,在没有recCas9表达质粒的情况下转染的细胞的基因组DNA中,或对于未转染的对照细胞的基因组DNA,未检测到缺失PCR产物(图5B)。这些巢式PCR产物的估计检测限为每5,500个染色体拷贝约1个缺失事件。分离对应于预测的基因组缺失的415-bp PCR产物并测序。测序证实PCR产物与自重组酶介导的基因组缺失预期的预测连接相匹配,并且不含任何提示NHEJ的插入或缺失(图5C)。

[0367] 在基因组模板的连续稀释液上使用巢式PCR估计最小基因组缺失效率的下限(参见上文或例如Sykes et al., Quantitation of targets for PCR by use of limiting dilution. *Biotechniques* 13,444-449 (1992),其全部内容在此通过引用并入,以获得更多细节)。产生recCas9特异性巢式PCR产物的给定量的基因组DNA必须含有至少一个经编辑的染色体。为了建立该recCas9介导的基因组缺失事件的下限,在基因组DNA(从用recCas9和四种FAM19A2引导RNA表达载体转染的细胞中分离)的连续稀释液上进行巢式PCR,以确定导致可检测的缺失产物的基因组模板DNA的最低浓度。这些实验揭示了 $0.023 \pm 0.017\%$ (三个生物学重复的平均值)的缺失效率的下限(图5D),表明recCas9介导的基因组缺失以至少该效率进行。未转染细胞的基因组DNA的巢式PCR不产生产物,估计检测限为 $<0.0072\%$ 重组。

[0368] 其他替代重组酶的使用

[0369] 称为“36C6”的演化为靶向人基因组的Rosa基因座中位点的Cre重组酶与dCas9融合。然后,将该融合物用于以引导RNA依赖性方式重组含有Rosa靶位点的基于质粒的报告物。图7A证明了使用野生型Cre和36C6的接头优化的结果。显示的1x、2x、5x和8x接头是接头中GGG重复的数量。回复分析证明了对与dCas9融合的36C6进行突变可以影响嵌合融合物的相对引导依赖性(图7B)。回复用其未突变的氨基酸标记。例如,已经突变为M的位置306在进行测定之前回复为I。靶向其关联报告物的GinB构建体用作图7A和7B中所示实验数据的对照。中靶引导是chr13-102010574引导(质粒BC165和166)。显示的缩写是GGG-36C6:dCas9-GGG-36C6;2GGG-36C6(使用接头SEQ ID NO:182):sdCas9-GGGGGG-36C6(使用接头SEQ ID NO:182)。

[0370] 用于36C6和所有变体转染的靶序列如下所示:(引导物-斜体;Rosa位点-粗体):

[0371] CCTAGGGAAGTGATCATAGCTGAGTTTCTATCTCATGGTTTATGCTAAACTATATGTTGACATGTTGAGGAGACTTAAGTCCAAAACCTGG (SEQ ID NO:760)

[0372] 在图7A、7B、8、9A和9B中,GinB的中靶引导是chr13-102010574引导(质粒BC165和166)。图7A、7B、8、9A和9B中的所有脱靶引导由chr12-62418577引导(BC163和BC164)构成。

[0373] 鉴定了人基因组中Rosa26位点侧翼的PAM,其可以支持dCas9结合(图8,上图)。然后设计引导RNA和质粒报告物以测试内源性前间隔区是否可以支持dCas9-36C6活性。靶向其关联报告物的GinB构建体用作对照。见图8。混合:Cas9和36C6之间所有5种接头变体的等份混合物。对于hRosa,靶序列(包括引导RNA靶标)如下:(引导-斜体;Rosa位点-粗体)

[0374] CCTGAAATAATGCAAGTGTAATAACTTTTTAAATCTCATGGTTTATGCTAAACTATATGTTGACATAAGAGTGTTGATAAGGCAACAGTAGG (SEQ ID NO:767)

[0375] 用于hRosa的中靶引导质粒与其他gRNA表达质粒相同,只是前间隔区被上文所示的那些替换(图8)。

[0376] dCas9-Cre重组酶融合物的几个测试的Cre截短显示在图9A中。与dCas9融合的Cre重组酶的截短的变体显示出可察觉的重组酶活性以及对Lox质粒报告物系统中引导RNA的存在严格依赖(图9B)。截短的变体用截短的Cre开始的残基标记。图9A和9B中所示的所有融合蛋白的接头是8xGGS。与dCas9融合的野生型Cre用作阳性对照。用于36C6和所有变体转染的靶序列如下所示:(引导-斜体;Rosa位点-粗体):

[0377] CCTAGGGAAGTGATCATAGCTGAGTTTCTATCTCATGGTTTATGCTAAACTATATGTTGACATGTTGAGGAGACTTAAGTCCAAAACCTGG (SEQ ID NO:768)

[0378] 使用的中靶引导是chr13-102010574引导(质粒BC165和166),并且脱靶引导是chr12-62418577引导(BC163和BC164)。

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[0450] 等同实施方案和范围

[0451] 本领域技术人员将认识到或能够仅仅使用常规的实验确定本文所述的本发明具体实施方案的许多等同实施方案。本发明的范围不意图限于以上说明书,而是如所附权利要求中所述。

[0452] 在权利要求中,诸如“一种”、“一个”和“该”的冠词可以表示一个或超出一个,除非相反地指出或者从上下文中显而易见。若一个、超出一个或所有组成员在给定产物或过程中存在、使用或以其他方式相关,则认为在组中的一个或多个成员之间包括“或”的权利要求或说明书是满足的,除非另有说明或从上下文中显而易见。本发明包括实施方案,其中组的恰好一个成员在给定产物或过程中存在、使用或以其他方式相关。本发明还包括实施方案,其中超出一个或所有组成员在给定产物或过程中存在、使用或以其他方式相关。

[0453] 此外,应理解,本发明涵盖所有变型、组合和置换,其中来自一个或多个权利要求或来自说明书的相关部分的一个或多个限制、元素、条款、描述性术语等被引入另一个权利要求中。例如,可以修改依赖于另一个权利要求的任何权利要求以包括在依赖于相同基本权利要求的任何其他权利要求中找到的一个或多个限制。此外,在权利要求叙述组合物的情况下,应当理解包括将组合物用于本文公开的任何目的的方法,并且包括根据本文公开的任何制备方法或本领域中已知的其他方法制备组合物的方法,除非另有说明或者除非本领域普通技术人员明白会出现矛盾或不一致。

[0454] 在将元素呈现为列表(例如,以马库什群组格式)的情况下,应当理解,还公开了元素的每个子群,并且可以从群组中移除任何元素。还应注意,术语“包含”旨在是开放的并且允许包含另外的元素或步骤。应当理解,通常,在本发明或本发明的方面称为包含特定元素、特征、步骤等的情况下,本发明或本发明的方面的某些实施方案由此类元素、特征、步骤等组成,或基本上由之组成。出于简化的目的,那些实施方案未在本文中用同样的词语具体阐述。因此,对于包含一个或多个元素、特征、步骤等的本发明的每个实施方案,本发明还提供了由这些元素、特征、步骤等组成或基本上由之组成的实施方案。

[0455] 在给出范围的情况下,端点包括在内。此外,应当理解,除非另有说明或从上下文和/或本领域普通技术人员的理解中明显看出,否则表示为范围的值可以假定在本发明的不同实施方案中的所述范围内的任何特定值,至该范围下限的单位的十分之一,除非上下文另有明确规定。还应当理解,除非另有说明或从上下文和/或本领域普通技术人员的理解中明显看出,否则表示为范围的值可以假定给定范围内的任何子范围,其中子范围的端点表示为与范围的下限的单位的十分之一相同的精度。

[0456] 此外,应当理解,本发明的任何具体实施方案可以明确地从任何一个或多个权利

要求中排除。在给出范围的情况下,该范围内的任何值可以明确地从任何一个或多个权利要求中排除。本发明的组合物和/或方法的任何实施方案、元素、特征、应用或方面可以从任何一个或多个权利要求中排除。出于简洁的目的,本文未明确阐述其中排除一个或多个元素、特征、目的或方面的所有实施方案。

[0457] 本文提及的所有出版物、专利和序列数据库条目,包括上面列出的那些项,通过引用整体并入本文,如同每个单独的出版物或专利被具体和单独地指出通过引用并入。在冲突的情况下,以本申请(包括本文中的任何定义)为准。

序列表

- <110> 哈佛大学的校长及成员们
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           35           40           45
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           50           55           60
Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys
65           70           75           80
Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser
           85           90           95
Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys
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His Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr
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His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp
           130          135          140
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	210	215	220
Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu Phe Gly Asn			
225	230	235	240
Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe			
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Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp			
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Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp			
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Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp			
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Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe			
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Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp			
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Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg			
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Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe			
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Leu Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile			
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 Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr
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 Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp
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 Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly
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 Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr
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 His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr
 645 650 655
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 Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe
 690 695 700
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 705 710 715 720
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Gly Lys Ser Asp Asn	Val Pro Ser Glu Glu	Val Val Lys Lys Met Lys
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Tyr Ser Asn Ile Met	Asn Phe Phe Lys Thr	Glu Ile Thr Leu Ala
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Asn Gly Glu Ile Arg	Lys Arg Pro Leu Ile	Glu Thr Asn Gly Glu
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Thr Gly Glu Ile Val	Trp Asp Lys Gly Arg	Asp Phe Ala Thr Val
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Arg Lys Val Leu Ser Met Pro Gln Val Asn Ile Val Lys Lys Thr 1085	1090	1095
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Glu Leu Gln Lys Gly Asn Glu Leu Ala Leu Pro Ser Lys Tyr Val 1220	1225	1230
Asn Phe Leu Tyr Leu Ala Ser His Tyr Glu Lys Leu Lys Gly Ser 1235	1240	1245
Pro Glu Asp Asn Glu Gln Lys Gln Leu Phe Val Glu Gln His Lys 1250	1255	1260
His Tyr Leu Asp Glu Ile Ile Glu Gln Ile Ser Glu Phe Ser Lys 1265	1270	1275
Arg Val Ile Leu Ala Asp Ala Asn Leu Asp Lys Val Leu Ser Ala 1280	1285	1290
Tyr Asn Lys His Arg Asp Lys Pro Ile Arg Glu Gln Ala Glu Asn 1295	1300	1305
Ile Ile His Leu Phe Thr Leu Thr Asn Leu Gly Ala Pro Ala Ala 1310	1315	1320
Phe Lys Tyr Phe Asp Thr Thr Ile Asp Arg Lys Arg Tyr Thr Ser 1325	1330	1335
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<211> 4104

<212> DNA

<213> 酿脓链球菌

<400> 2

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<211> 1367

<212> PRT

<213> 酿脓链球菌

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Lys	Val	Leu	Gly	Asn	Thr	Asp	Arg	His	Ser	Ile	Lys	Lys	Asn	Leu	Ile
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Lys	Arg	Thr	Ala	Arg	Arg	Arg	Tyr	Thr	Arg	Arg	Lys	Asn	Arg	Ile	Cys
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Tyr	Leu	Gln	Glu	Ile	Phe	Ser	Asn	Glu	Met	Ala	Lys	Val	Asp	Asp	Ser
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Phe	Phe	His	Arg	Leu	Glu	Glu	Ser	Phe	Leu	Val	Glu	Glu	Asp	Lys	Lys
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His	Glu	Arg	His	Pro	Ile	Phe	Gly	Asn	Ile	Val	Asp	Glu	Val	Ala	Tyr
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His	Glu	Lys	Tyr	Pro	Thr	Ile	Tyr	His	Leu	Arg	Lys	Lys	Leu	Ala	Asp
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Ser	Thr	Asp	Lys	Ala	Asp	Leu	Arg	Leu	Ile	Tyr	Leu	Ala	Leu	Ala	His
145					150						155				160
Met	Ile	Lys	Phe	Arg	Gly	His	Phe	Leu	Ile	Glu	Gly	Asp	Leu	Asn	Pro
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Asp	Asn	Ser	Asp	Val	Asp	Lys	Leu	Phe	Ile	Gln	Leu	Val	Gln	Ile	Tyr
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Leu	Ile	Ala	Leu	Ser	Leu	Gly	Leu	Thr	Pro	Asn	Phe	Lys	Ser	Asn	Phe
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Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser			
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Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp			
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Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg			
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Lys Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu			
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Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe			
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Leu Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile			
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Pro Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp			
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Met Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu			
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Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser			
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Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln			
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Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp			
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Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly			
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Ala Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp			
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Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr			
610	615	620	

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His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr																		
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Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp																		
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Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe																		
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Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe																		
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Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly His Ser Leu																		
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His Glu Gln Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly																		
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Ile Leu Gln Thr Val Lys Ile Val Asp Glu Leu Val Lys Val Met Gly																		
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His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln Thr																		
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Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met Lys Arg Ile Glu																		
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Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys Glu His Pro Val																		
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Ser Asp Tyr Asp Val Asp His Ile Val Pro Gln Ser Phe Ile Lys Asp																		
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Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly																		
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Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn																		
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Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe																		
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Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys																		
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Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys																		
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His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu																		

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Gly Thr Ala Leu Ile Lys	Lys Tyr Pro Lys Leu Glu	Ser Glu Phe Val
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<223> 合成多肽

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			20					25					30		
Lys	Val	Leu	Gly	Asn	Thr	Asp	Arg	His	Ser	Ile	Lys	Lys	Asn	Leu	Ile
			35					40					45		
Gly	Ala	Leu	Leu	Phe	Asp	Ser	Gly	Glu	Thr	Ala	Glu	Ala	Thr	Arg	Leu
			50				55					60			
Lys	Arg	Thr	Ala	Arg	Arg	Arg	Tyr	Thr	Arg	Arg	Lys	Asn	Arg	Ile	Cys
65				70						75				80	
Tyr	Leu	Gln	Glu	Ile	Phe	Ser	Asn	Glu	Met	Ala	Lys	Val	Asp	Asp	Ser
				85						90				95	
Phe	Phe	His	Arg	Leu	Glu	Glu	Ser	Phe	Leu	Val	Glu	Glu	Asp	Lys	Lys
			100						105					110	

His Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr
 115 120 125
 His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp
 130 135 140
 Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His
 145 150 155 160
 Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro
 165 170 175
 Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val Gln Thr Tyr
 180 185 190
 Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala
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 Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg Leu Glu Asn
 210 215 220
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 Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe
 245 250 255
 Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp
 260 265 270
 Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp
 275 280 285
 Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp
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 Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser
 305 310 315 320
 Met Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys
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 Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe
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 Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser
 355 360 365
 Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp
 370 375 380
 Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg
 385 390 395 400
 Lys Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu
 405 410 415
 Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe

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Pro Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp		
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Met Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu		
465	470	475
Val Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr		
485	490	495
Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser		
500	505	510
Leu Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys		
515	520	525
Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln		
530	535	540
Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr		
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Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp		
565	570	575
Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly		
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Thr Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp		
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Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr		
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625	630	635
His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr		
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Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp		
660	665	670
Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe		
675	680	685
Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe		
690	695	700
Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly Asp Ser Leu		
705	710	715
His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly		
725	730	735

Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys Val Met Gly
 740 745 750
 Arg His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln
 755 760 765
 Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met Lys Arg Ile
 770 775 780
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 Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu Tyr Tyr Leu
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 820 825 830
 Leu Ser Asp Tyr Asp Val Asp His Ile Val Pro Gln Ser Phe Leu Lys
 835 840 845
 Asp Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg
 850 855 860
 Gly Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys
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 Asn Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys
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 Phe Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp
 900 905 910
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 915 920 925
 Lys His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp
 930 935 940
 Glu Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser
 945 950 955 960
 Lys Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg
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 Glu Ile Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val
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 Val Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe
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Thr Gly Glu Ile Val Trp	Asp Lys Gly Arg Asp	Phe Ala Thr Val
1070	1075	1080
Arg Lys Val Leu Ser Met	Pro Gln Val Asn Ile	Val Lys Lys Thr
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Glu Val Gln Thr Gly Gly	Phe Ser Lys Glu Ser	Ile Leu Pro Lys
1100	1105	1110
Arg Asn Ser Asp Lys Leu	Ile Ala Arg Lys Lys	Asp Trp Asp Pro
1115	1120	1125
Lys Lys Tyr Gly Gly Phe	Asp Ser Pro Thr Val	Ala Tyr Ser Val
1130	1135	1140
Leu Val Val Ala Lys Val	Glu Lys Gly Lys Ser	Lys Lys Leu Lys
1145	1150	1155
Ser Val Lys Glu Leu Leu	Gly Ile Thr Ile Met	Glu Arg Ser Ser
1160	1165	1170
Phe Glu Lys Asn Pro Ile	Asp Phe Leu Glu Ala	Lys Gly Tyr Lys
1175	1180	1185
Glu Val Lys Lys Asp Leu	Ile Ile Lys Leu Pro	Lys Tyr Ser Leu
1190	1195	1200
Phe Glu Leu Glu Asn Gly	Arg Lys Arg Met Leu	Ala Ser Ala Gly
1205	1210	1215
Glu Leu Gln Lys Gly Asn	Glu Leu Ala Leu Pro	Ser Lys Tyr Val
1220	1225	1230
Asn Phe Leu Tyr Leu Ala	Ser His Tyr Glu Lys	Leu Lys Gly Ser
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Pro Glu Asp Asn Glu Gln	Lys Gln Leu Phe Val	Glu Gln His Lys
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His Tyr Leu Asp Glu Ile	Ile Glu Gln Ile Ser	Glu Phe Ser Lys
1265	1270	1275
Arg Val Ile Leu Ala Asp	Ala Asn Leu Asp Lys	Val Leu Ser Ala
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Tyr Asn Lys His Arg Asp	Lys Pro Ile Arg Glu	Gln Ala Glu Asn
1295	1300	1305
Ile Ile His Leu Phe Thr	Leu Thr Asn Leu Gly	Ala Pro Ala Ala
1310	1315	1320
Phe Lys Tyr Phe Asp Thr	Thr Ile Asp Arg Lys	Arg Tyr Thr Ser
1325	1330	1335

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Gly Ser

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Ser Gly Ser Glu Thr Pro Gly Thr Ser Glu Ser Ala Thr Pro Glu Ser
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ga 62

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- <210> 135
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 35 40 45
 Leu Lys Arg Leu Gln Lys Gly Asp Thr Leu Val Val Trp Lys Leu Asp
 50 55 60
 Arg Leu Gly Arg Ser Met Lys His Leu Ile Ser Leu Val Gly Glu Leu
 65 70 75 80
 Arg Glu Arg Gly Ile Asn Phe Arg Ser Leu Thr Asp Ser Ile Asp Thr
 85 90 95
 Ser Ser Pro Met Gly Arg Phe Phe Phe Tyr Val Met Gly Ala Leu Ala
 100 105 110
 Glu Met Glu Arg Glu Leu Ile Ile Glu Arg Thr Met Ala Gly Leu Ala
 115 120 125
 Ala Ala Arg Asn Lys Gly Arg Arg Phe Gly Arg Pro Pro Lys Gly Gly
 130 135 140
 Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser
 145 150 155 160
 Gly Gly Ser Gly Gly Ser Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile
 165 170 175
 Gly Thr Asn Ser Val Gly Trp Ala Val Ile Thr Asp Glu Tyr Lys Val
 180 185 190
 Pro Ser Lys Lys Phe Lys Val Leu Gly Asn Thr Asp Arg His Ser Ile
 195 200 205
 Lys Lys Asn Leu Ile Gly Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala
 210 215 220
 Glu Ala Thr Arg Leu Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg
 225 230 235 240
 Lys Asn Arg Ile Cys Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala
 245 250 255
 Lys Val Asp Asp Ser Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val

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Asp Glu Val Ala Tyr His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg		
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Lys Lys Leu Val Asp Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr		
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Leu Ala Leu Ala His Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu		
325	330	335
Gly Asp Leu Asn Pro Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln		
340	345	350
Leu Val Gln Thr Tyr Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala		
355	360	365
Ser Gly Val Asp Ala Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser		
370	375	380
Arg Arg Leu Glu Asn Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn		
385	390	395
Gly Leu Phe Gly Asn Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn		
405	410	415
Phe Lys Ser Asn Phe Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser		
420	425	430
Lys Asp Thr Tyr Asp Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly		
435	440	445
Asp Gln Tyr Ala Asp Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala		
450	455	460
Ile Leu Leu Ser Asp Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala		
465	470	475
Pro Leu Ser Ala Ser Met Ile Lys Arg Tyr Asp Glu His His Gln Asp		
485	490	495
Leu Thr Leu Leu Lys Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr		
500	505	510
Lys Glu Ile Phe Phe Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile		
515	520	525
Asp Gly Gly Ala Ser Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile		
530	535	540
Leu Glu Lys Met Asp Gly Thr Glu Glu Leu Leu Val Lys Leu Asn Arg		
545	550	555
Glu Asp Leu Leu Arg Lys Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro		
565	570	575

His Gln Ile His Leu Gly Glu Leu His Ala Ile Leu Arg Arg Gln Glu
 580 585 590
 Asp Phe Tyr Pro Phe Leu Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile
 595 600 605
 Leu Thr Phe Arg Ile Pro Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn
 610 615 620
 Ser Arg Phe Ala Trp Met Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro
 625 630 635 640
 Trp Asn Phe Glu Glu Val Val Asp Lys Gly Ala Ser Ala Gln Ser Phe
 645 650 655
 Ile Glu Arg Met Thr Asn Phe Asp Lys Asn Leu Pro Asn Glu Lys Val
 660 665 670
 Leu Pro Lys His Ser Leu Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu
 675 680 685
 Leu Thr Lys Val Lys Tyr Val Thr Glu Gly Met Arg Lys Pro Ala Phe
 690 695 700
 Leu Ser Gly Glu Gln Lys Lys Ala Ile Val Asp Leu Leu Phe Lys Thr
 705 710 715 720
 Asn Arg Lys Val Thr Val Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys
 725 730 735
 Ile Glu Cys Phe Asp Ser Val Glu Ile Ser Gly Val Glu Asp Arg Phe
 740 745 750
 Asn Ala Ser Leu Gly Thr Tyr His Asp Leu Leu Lys Ile Ile Lys Asp
 755 760 765
 Lys Asp Phe Leu Asp Asn Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile
 770 775 780
 Val Leu Thr Leu Thr Leu Phe Glu Asp Arg Glu Met Ile Glu Glu Arg
 785 790 795 800
 Leu Lys Thr Tyr Ala His Leu Phe Asp Asp Lys Val Met Lys Gln Leu
 805 810 815
 Lys Arg Arg Arg Tyr Thr Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile
 820 825 830
 Asn Gly Ile Arg Asp Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu
 835 840 845
 Lys Ser Asp Gly Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp
 850 855 860
 Asp Ser Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly
 865 870 875 880
 Gln Gly Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro

	885	890	895
Ala Ile Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu			
	900	905	910
Val Lys Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met			
	915	920	925
Ala Arg Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu			
	930	935	940
Arg Met Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile			
945	950	955	960
Leu Lys Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu			
	965	970	975
Tyr Leu Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val Asp Gln Glu			
	980	985	990
Leu Asp Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp Ala Ile Val Pro			
	995	1000	1005
Gln Ser Phe Leu Lys Asp Asp Ser Ile Asp Asn Lys Val Leu Thr			
	1010	1015	1020
Arg Ser Asp Lys Asn Arg Gly Lys Ser Asp Asn Val Pro Ser Glu			
	1025	1030	1035
Glu Val Val Lys Lys Met Lys Asn Tyr Trp Arg Gln Leu Leu Asn			
	1040	1045	1050
Ala Lys Leu Ile Thr Gln Arg Lys Phe Asp Asn Leu Thr Lys Ala			
	1055	1060	1065
Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys Ala Gly Phe Ile Lys			
	1070	1075	1080
Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys His Val Ala Gln			
	1085	1090	1095
Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu Asn Asp Lys			
	1100	1105	1110
Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys Leu Val			
	1115	1120	1125
Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu Ile			
	1130	1135	1140
Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val			
	1145	1150	1155
Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe			
	1160	1165	1170
Val Tyr Gly Asp Tyr Lys Val Tyr Asp Val Arg Lys Met Ile Ala			
	1175	1180	1185

Lys Ser Glu Gln Glu Ile Gly Lys Ala Thr Ala Lys Tyr Phe Phe	1190	1195	1200
Tyr Ser Asn Ile Met Asn Phe Phe Lys Thr Glu Ile Thr Leu Ala	1205	1210	1215
Asn Gly Glu Ile Arg Lys Arg Pro Leu Ile Glu Thr Asn Gly Glu	1220	1225	1230
Thr Gly Glu Ile Val Trp Asp Lys Gly Arg Asp Phe Ala Thr Val	1235	1240	1245
Arg Lys Val Leu Ser Met Pro Gln Val Asn Ile Val Lys Lys Thr	1250	1255	1260
Glu Val Gln Thr Gly Gly Phe Ser Lys Glu Ser Ile Leu Pro Lys	1265	1270	1275
Arg Asn Ser Asp Lys Leu Ile Ala Arg Lys Lys Asp Trp Asp Pro	1280	1285	1290
Lys Lys Tyr Gly Gly Phe Asp Ser Pro Thr Val Ala Tyr Ser Val	1295	1300	1305
Leu Val Val Ala Lys Val Glu Lys Gly Lys Ser Lys Lys Leu Lys	1310	1315	1320
Ser Val Lys Glu Leu Leu Gly Ile Thr Ile Met Glu Arg Ser Ser	1325	1330	1335
Phe Glu Lys Asn Pro Ile Asp Phe Leu Glu Ala Lys Gly Tyr Lys	1340	1345	1350
Glu Val Lys Lys Asp Leu Ile Ile Lys Leu Pro Lys Tyr Ser Leu	1355	1360	1365
Phe Glu Leu Glu Asn Gly Arg Lys Arg Met Leu Ala Ser Ala Gly	1370	1375	1380
Glu Leu Gln Lys Gly Asn Glu Leu Ala Leu Pro Ser Lys Tyr Val	1385	1390	1395
Asn Phe Leu Tyr Leu Ala Ser His Tyr Glu Lys Leu Lys Gly Ser	1400	1405	1410
Pro Glu Asp Asn Glu Gln Lys Gln Leu Phe Val Glu Gln His Lys	1415	1420	1425
His Tyr Leu Asp Glu Ile Ile Glu Gln Ile Ser Glu Phe Ser Lys	1430	1435	1440
Arg Val Ile Leu Ala Asp Ala Asn Leu Asp Lys Val Leu Ser Ala	1445	1450	1455
Tyr Asn Lys His Arg Asp Lys Pro Ile Arg Glu Gln Ala Glu Asn	1460	1465	1470
Ile Ile His Leu Phe Thr Leu Thr Asn Leu Gly Ala Pro Ala Ala			

1475 1480 1485
 Phe Lys Tyr Phe Asp Thr Thr Ile Asp Arg Lys Arg Tyr Thr Ser
 1490 1495 1500
 Thr Lys Glu Val Leu Asp Ala Thr Leu Ile His Gln Ser Ile Thr
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 Gly Leu Tyr Glu Thr Arg Ile Asp Leu Ser Gln Leu Gly Gly Asp
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<211> 77

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<210> 212

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<212> DNA

<213> 人工序列

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<400> 212

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aagtgacggg aagaatgg 78

<210> 213

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<400> 213

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<210> 215

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<400> 215

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aaacactctt gttgtgg 77

<210> 216

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<210> 217

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<400> 217

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aaacactcct tttgtgg 77

<210> 221

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- <210> 224
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- <210> 225
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- <210> 226
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<400> 229

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<400> 230

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<210> 231

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ttccatattc tttggagg 78

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<400> 232

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<210> 233

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<400> 233

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<210> 234

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- <210> 236
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aagtgacagg gagtatgg 78
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- <210> 238
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tcccatatatt cttggagg 78
- <210> 239
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agaggggaaga agcaaagg 78

<210> 241

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<210> 242

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aagtgatggg gagaatgg 78

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<210> 244

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<400> 244

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<210> 246

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tcccatattt cttggggg 77

<210> 247

<211> 78

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<400> 247

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aagtgacggg gagaatgg 78

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- <211> 78
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tcccaaattt cttggagg 78
- <210> 249
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- <210> 407
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<211> 78

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<211> 78

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<210> 624

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<211> 78
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<210> 630

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<210> 632

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<210> 633

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<210> 636

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<400> 636

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aaacactcct tttgtgg 77

<210> 638

- <211> 77
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aaacactcct tttctgg 77
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aacagcact gtcagg 77

<210> 647

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<212> DNA

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<400> 647

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aacagcactt gtcagg 77

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<213> 人工序列

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aacagcactt gtcagg 76

<210> 649

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tcccatatatt cttggagg 78

<210> 650

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- <211> 78
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aggttgacagt gagccggg 78
- <210> 652
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<210> 658

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<210> 664

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gaaggaccct ggcacggg 78
- <210> 665
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gaaggaccct ggcacgg 77
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aaggaccctg gcacggg 77
- <210> 667
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aaggaccctg gcacgg 76
- <210> 668
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gtccacatat caccaagg 78
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aagtgacggg gagaaagg 78
- <210> 680
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aagtgatggg tagaatgg 78
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aagtgatggg gagaatgg 78
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 35 40 45
 Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala Thr Arg Leu Lys
 50 55 60
 Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys Tyr
 65 70 75 80
 Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser Phe
 85 90 95
 Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys His
 100 105 110
 Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr His
 115 120 125
 Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp Ser
 130 135 140
 Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His Met
 145 150 155 160
 Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro Asp

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Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala Lys			
	195	200	205
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	210	215	220
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225	230	235	240
Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe Asp			
	245	250	255
Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp Asp			
	260	265	270
Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp Leu			
	275	280	285
Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp Ile			
	290	295	300
Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser Met			
305	310	315	320
Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys Ala			
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Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe Asp			
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Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser Gln			
	355	360	365
Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp Gly			
	370	375	380
Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg Lys			
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Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu Gly			
	405	410	415
Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe Leu			
	420	425	430
Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile Pro			
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Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp Met			
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Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu Val			
465	470	475	480

Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr Asn
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 Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser Leu
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 Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys Tyr
 515 520 525
 Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln Lys
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 580 585 590
 Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp Asn
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 625 630 635 640
 Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr Thr
 645 650 655
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 660 665 670
 Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe Ala
 675 680 685
 Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe Lys
 690 695 700
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 705 710 715 720
 Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly Ile
 725 730 735
 Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys Val Met Gly Arg
 740 745 750
 His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln Thr
 755 760 765
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 Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys Glu His Pro Val

785	790	795	800
Glu Asn Thr Gln Leu	Gln Asn Glu Lys Leu	Tyr Leu Tyr Tyr Leu	Gln
	805	810	815
Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp Ile Asn Arg Leu			
	820	825	830
Ser Asp Tyr Asp Val Asp Ala Ile Val Pro Gln Ser Phe Leu Lys Asp			
	835	840	845
Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly			
	850	855	860
Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn			
865	870	875	880
Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe			
	885	890	895
Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys			
	900	905	910
Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys			
	915	920	925
His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu			
	930	935	940
Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys			
945	950	955	960
Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu			
	965	970	975
Ile Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val			
	980	985	990
Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe Val			
	995	1000	1005
Tyr Gly Asp Tyr Lys Val Tyr Asp Val Arg Lys Met Ile Ala Lys			
	1010	1015	1020
Ser Glu Gln Glu Ile Gly Lys Ala Thr Ala Lys Tyr Phe Phe Tyr			
	1025	1030	1035
Ser Asn Ile Met Asn Phe Phe Lys Thr Glu Ile Thr Leu Ala Asn			
	1040	1045	1050
Gly Glu Ile Arg Lys Arg Pro Leu Ile Glu Thr Asn Gly Glu Thr			
	1055	1060	1065
Gly Glu Ile Val Trp Asp Lys Gly Arg Asp Phe Ala Thr Val Arg			
	1070	1075	1080
Lys Val Leu Ser Met Pro Gln Val Asn Ile Val Lys Lys Thr Glu			
	1085	1090	1095

Val Gln Thr Gly Gly Phe Ser Lys Glu Ser Ile Leu Pro Lys Arg	1100	1105	1110
Asn Ser Asp Lys Leu Ile Ala Arg Lys Lys Asp Trp Asp Pro Lys	1115	1120	1125
Lys Tyr Gly Gly Phe Asp Ser Pro Thr Val Ala Tyr Ser Val Leu	1130	1135	1140
Val Val Ala Lys Val Glu Lys Gly Lys Ser Lys Lys Leu Lys Ser	1145	1150	1155
Val Lys Glu Leu Leu Gly Ile Thr Ile Met Glu Arg Ser Ser Phe	1160	1165	1170
Glu Lys Asn Pro Ile Asp Phe Leu Glu Ala Lys Gly Tyr Lys Glu	1175	1180	1185
Val Lys Lys Asp Leu Ile Ile Lys Leu Pro Lys Tyr Ser Leu Phe	1190	1195	1200
Glu Leu Glu Asn Gly Arg Lys Arg Met Leu Ala Ser Ala Gly Glu	1205	1210	1215
Leu Gln Lys Gly Asn Glu Leu Ala Leu Pro Ser Lys Tyr Val Asn	1220	1225	1230
Phe Leu Tyr Leu Ala Ser His Tyr Glu Lys Leu Lys Gly Ser Pro	1235	1240	1245
Glu Asp Asn Glu Gln Lys Gln Leu Phe Val Glu Gln His Lys His	1250	1255	1260
Tyr Leu Asp Glu Ile Ile Glu Gln Ile Ser Glu Phe Ser Lys Arg	1265	1270	1275
Val Ile Leu Ala Asp Ala Asn Leu Asp Lys Val Leu Ser Ala Tyr	1280	1285	1290
Asn Lys His Arg Asp Lys Pro Ile Arg Glu Gln Ala Glu Asn Ile	1295	1300	1305
Ile His Leu Phe Thr Leu Thr Asn Leu Gly Ala Pro Ala Ala Phe	1310	1315	1320
Lys Tyr Phe Asp Thr Thr Ile Asp Arg Lys Arg Tyr Thr Ser Thr	1325	1330	1335
Lys Glu Val Leu Asp Ala Thr Leu Ile His Gln Ser Ile Thr Gly	1340	1345	1350
Leu Tyr Glu Thr Arg Ile Asp Leu Ser Gln Leu Gly Gly Asp	1355	1360	1365
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<211> 142			
<212> PRT			

<213> 人工序列

<220>

<223> 合成多肽

<400> 713

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Met Leu Ile Gly Tyr Val Arg Val Ser Thr Asn Asp Gln Asn Thr Asp
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Leu Gln Arg Asn Ala Leu Val Cys Ala Gly Cys Glu Gln Ile Phe Glu
           20           25           30
Asp Lys Leu Ser Gly Thr Arg Thr Asp Arg Pro Gly Leu Lys Arg Ala
           35           40           45
Leu Lys Arg Leu Gln Lys Gly Asp Thr Leu Val Val Trp Lys Leu Asp
           50           55           60
Arg Leu Gly Arg Ser Met Lys His Leu Ile Ser Leu Val Gly Glu Leu
65           70           75           80
Arg Glu Arg Gly Ile Asn Phe Arg Ser Leu Thr Asp Ser Ile Asp Thr
           85           90           95
Ser Ser Pro Met Gly Arg Phe Phe Phe Tyr Val Met Gly Ala Leu Ala
           100          105          110
Glu Met Glu Arg Glu Leu Ile Ile Glu Arg Thr Met Ala Gly Leu Ala
           115          120          125
Ala Ala Arg Asn Lys Gly Arg Arg Phe Gly Arg Pro Pro Lys
           130          135          140

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<210> 714

<211> 1300

<212> PRT

<213> 新凶手弗朗西斯菌

<400> 714

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Met Ser Ile Tyr Gln Glu Phe Val Asn Lys Tyr Ser Leu Ser Lys Thr
1           5           10           15
Leu Arg Phe Glu Leu Ile Pro Gln Gly Lys Thr Leu Glu Asn Ile Lys
           20           25           30
Ala Arg Gly Leu Ile Leu Asp Asp Glu Lys Arg Ala Lys Asp Tyr Lys
           35           40           45
Lys Ala Lys Gln Ile Ile Asp Lys Tyr His Gln Phe Phe Ile Glu Glu
           50           55           60
Ile Leu Ser Ser Val Cys Ile Ser Glu Asp Leu Leu Gln Asn Tyr Ser
65           70           75           80
Asp Val Tyr Phe Lys Leu Lys Lys Ser Asp Asp Asp Asn Leu Gln Lys
           85           90           95

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Asp Phe Lys Ser Ala Lys Asp Thr Ile Lys Lys Gln Ile Ser Glu Tyr			
100	105	110	
Ile Lys Asp Ser Glu Lys Phe Lys Asn Leu Phe Asn Gln Asn Leu Ile			
115	120	125	
Asp Ala Lys Lys Gly Gln Glu Ser Asp Leu Ile Leu Trp Leu Lys Gln			
130	135	140	
Ser Lys Asp Asn Gly Ile Glu Leu Phe Lys Ala Asn Ser Asp Ile Thr			
145	150	155	160
Asp Ile Asp Glu Ala Leu Glu Ile Ile Lys Ser Phe Lys Gly Trp Thr			
	165	170	175
Thr Tyr Phe Lys Gly Phe His Glu Asn Arg Lys Asn Val Tyr Ser Ser			
	180	185	190
Asn Asp Ile Pro Thr Ser Ile Ile Tyr Arg Ile Val Asp Asp Asn Leu			
	195	200	205
Pro Lys Phe Leu Glu Asn Lys Ala Lys Tyr Glu Ser Leu Lys Asp Lys			
	210	215	220
Ala Pro Glu Ala Ile Asn Tyr Glu Gln Ile Lys Lys Asp Leu Ala Glu			
225	230	235	240
Glu Leu Thr Phe Asp Ile Asp Tyr Lys Thr Ser Glu Val Asn Gln Arg			
	245	250	255
Val Phe Ser Leu Asp Glu Val Phe Glu Ile Ala Asn Phe Asn Asn Tyr			
	260	265	270
Leu Asn Gln Ser Gly Ile Thr Lys Phe Asn Thr Ile Ile Gly Gly Lys			
	275	280	285
Phe Val Asn Gly Glu Asn Thr Lys Arg Lys Gly Ile Asn Glu Tyr Ile			
	290	295	300
Asn Leu Tyr Ser Gln Gln Ile Asn Asp Lys Thr Leu Lys Lys Tyr Lys			
305	310	315	320
Met Ser Val Leu Phe Lys Gln Ile Leu Ser Asp Thr Glu Ser Lys Ser			
	325	330	335
Phe Val Ile Asp Lys Leu Glu Asp Asp Ser Asp Val Val Thr Thr Met			
	340	345	350
Gln Ser Phe Tyr Glu Gln Ile Ala Ala Phe Lys Thr Val Glu Glu Lys			
	355	360	365
Ser Ile Lys Glu Thr Leu Ser Leu Leu Phe Asp Asp Leu Lys Ala Gln			
	370	375	380
Lys Leu Asp Leu Ser Lys Ile Tyr Phe Lys Asn Asp Lys Ser Leu Thr			
385	390	395	400
Asp Leu Ser Gln Gln Val Phe Asp Asp Tyr Ser Val Ile Gly Thr Ala			

	405	410	415
Val Leu Glu Tyr Ile Thr Gln Gln Ile Ala Pro Lys Asn Leu Asp Asn			
	420	425	430
Pro Ser Lys Lys Glu Gln Glu Leu Ile Ala Lys Lys Thr Glu Lys Ala			
	435	440	445
Lys Tyr Leu Ser Leu Glu Thr Ile Lys Leu Ala Leu Glu Glu Phe Asn			
	450	455	460
Lys His Arg Asp Ile Asp Lys Gln Cys Arg Phe Glu Glu Ile Leu Ala			
465	470	475	480
Asn Phe Ala Ala Ile Pro Met Ile Phe Asp Glu Ile Ala Gln Asn Lys			
	485	490	495
Asp Asn Leu Ala Gln Ile Ser Ile Lys Tyr Gln Asn Gln Gly Lys Lys			
	500	505	510
Asp Leu Leu Gln Ala Ser Ala Glu Asp Asp Val Lys Ala Ile Lys Asp			
	515	520	525
Leu Leu Asp Gln Thr Asn Asn Leu Leu His Lys Leu Lys Ile Phe His			
	530	535	540
Ile Ser Gln Ser Glu Asp Lys Ala Asn Ile Leu Asp Lys Asp Glu His			
545	550	555	560
Phe Tyr Leu Val Phe Glu Glu Cys Tyr Phe Glu Leu Ala Asn Ile Val			
	565	570	575
Pro Leu Tyr Asn Lys Ile Arg Asn Tyr Ile Thr Gln Lys Pro Tyr Ser			
	580	585	590
Asp Glu Lys Phe Lys Leu Asn Phe Glu Asn Ser Thr Leu Ala Asn Gly			
	595	600	605
Trp Asp Lys Asn Lys Glu Pro Asp Asn Thr Ala Ile Leu Phe Ile Lys			
	610	615	620
Asp Asp Lys Tyr Tyr Leu Gly Val Met Asn Lys Lys Asn Asn Lys Ile			
625	630	635	640
Phe Asp Asp Lys Ala Ile Lys Glu Asn Lys Gly Glu Gly Tyr Lys Lys			
	645	650	655
Ile Val Tyr Lys Leu Leu Pro Gly Ala Asn Lys Met Leu Pro Lys Val			
	660	665	670
Phe Phe Ser Ala Lys Ser Ile Lys Phe Tyr Asn Pro Ser Glu Asp Ile			
	675	680	685
Leu Arg Ile Arg Asn His Ser Thr His Thr Lys Asn Gly Ser Pro Gln			
	690	695	700
Lys Gly Tyr Glu Lys Phe Glu Phe Asn Ile Glu Asp Cys Arg Lys Phe			
705	710	715	720

Ile Asp Phe Tyr Lys Gln Ser Ile Ser Lys His Pro Glu Trp Lys Asp			
	725	730	735
Phe Gly Phe Arg Phe Ser Asp Thr Gln Arg Tyr Asn Ser Ile Asp Glu			
	740	745	750
Phe Tyr Arg Glu Val Glu Asn Gln Gly Tyr Lys Leu Thr Phe Glu Asn			
	755	760	765
Ile Ser Glu Ser Tyr Ile Asp Ser Val Val Asn Gln Gly Lys Leu Tyr			
	770	775	780
Leu Phe Gln Ile Tyr Asn Lys Asp Phe Ser Ala Tyr Ser Lys Gly Arg			
785	790	795	800
Pro Asn Leu His Thr Leu Tyr Trp Lys Ala Leu Phe Asp Glu Arg Asn			
	805	810	815
Leu Gln Asp Val Val Tyr Lys Leu Asn Gly Glu Ala Glu Leu Phe Tyr			
	820	825	830
Arg Lys Gln Ser Ile Pro Lys Lys Ile Thr His Pro Ala Lys Glu Ala			
	835	840	845
Ile Ala Asn Lys Asn Lys Asp Asn Pro Lys Lys Glu Ser Val Phe Glu			
	850	855	860
Tyr Asp Leu Ile Lys Asp Lys Arg Phe Thr Glu Asp Lys Phe Phe Phe			
865	870	875	880
His Cys Pro Ile Thr Ile Asn Phe Lys Ser Ser Gly Ala Asn Lys Phe			
	885	890	895
Asn Asp Glu Ile Asn Leu Leu Leu Lys Glu Lys Ala Asn Asp Val His			
	900	905	910
Ile Leu Ser Ile Asp Arg Gly Glu Arg His Leu Ala Tyr Tyr Thr Leu			
	915	920	925
Val Asp Gly Lys Gly Asn Ile Ile Lys Gln Asp Thr Phe Asn Ile Ile			
	930	935	940
Gly Asn Asp Arg Met Lys Thr Asn Tyr His Asp Lys Leu Ala Ala Ile			
945	950	955	960
Glu Lys Asp Arg Asp Ser Ala Arg Lys Asp Trp Lys Lys Ile Asn Asn			
	965	970	975
Ile Lys Glu Met Lys Glu Gly Tyr Leu Ser Gln Val Val His Glu Ile			
	980	985	990
Ala Lys Leu Val Ile Glu Tyr Asn Ala Ile Val Val Phe Glu Asp Leu			
	995	1000	1005
Asn Phe Gly Phe Lys Arg Gly Arg Phe Lys Val Glu Lys Gln Val			
	1010	1015	1020
Tyr Gln Lys Leu Glu Lys Met Leu Ile Glu Lys Leu Asn Tyr Leu			

1025	1030	1035
Val Phe Lys Asp Asn Glu Phe Asp Lys Thr Gly Gly Val Leu Arg		
1040	1045	1050
Ala Tyr Gln Leu Thr Ala Pro Phe Glu Thr Phe Lys Lys Met Gly		
1055	1060	1065
Lys Gln Thr Gly Ile Ile Tyr Tyr Val Pro Ala Gly Phe Thr Ser		
1070	1075	1080
Lys Ile Cys Pro Val Thr Gly Phe Val Asn Gln Leu Tyr Pro Lys		
1085	1090	1095
Tyr Glu Ser Val Ser Lys Ser Gln Glu Phe Phe Ser Lys Phe Asp		
1100	1105	1110
Lys Ile Cys Tyr Asn Leu Asp Lys Gly Tyr Phe Glu Phe Ser Phe		
1115	1120	1125
Asp Tyr Lys Asn Phe Gly Asp Lys Ala Ala Lys Gly Lys Trp Thr		
1130	1135	1140
Ile Ala Ser Phe Gly Ser Arg Leu Ile Asn Phe Arg Asn Ser Asp		
1145	1150	1155
Lys Asn His Asn Trp Asp Thr Arg Glu Val Tyr Pro Thr Lys Glu		
1160	1165	1170
Leu Glu Lys Leu Leu Lys Asp Tyr Ser Ile Glu Tyr Gly His Gly		
1175	1180	1185
Glu Cys Ile Lys Ala Ala Ile Cys Gly Glu Ser Asp Lys Lys Phe		
1190	1195	1200
Phe Ala Lys Leu Thr Ser Val Leu Asn Thr Ile Leu Gln Met Arg		
1205	1210	1215
Asn Ser Lys Thr Gly Thr Glu Leu Asp Tyr Leu Ile Ser Pro Val		
1220	1225	1230
Ala Asp Val Asn Gly Asn Phe Phe Asp Ser Arg Gln Ala Pro Lys		
1235	1240	1245
Asn Met Pro Gln Asp Ala Asp Ala Asn Gly Ala Tyr His Ile Gly		
1250	1255	1260
Leu Lys Gly Leu Met Leu Leu Gly Arg Ile Lys Asn Asn Gln Glu		
1265	1270	1275
Gly Lys Lys Leu Asn Leu Val Ile Lys Asn Glu Glu Tyr Phe Glu		
1280	1285	1290
Phe Val Gln Asn Arg Asn Asn		
1295	1300	
<210> 715		
<211> 1300		

<212> PRT

<213> 新凶手弗朗西斯菌

<400> 715

Met	Ser	Ile	Tyr	Gln	Glu	Phe	Val	Asn	Lys	Tyr	Ser	Leu	Ser	Lys	Thr
1				5					10					15	
Leu	Arg	Phe	Glu	Leu	Ile	Pro	Gln	Gly	Lys	Thr	Leu	Glu	Asn	Ile	Lys
			20					25					30		
Ala	Arg	Gly	Leu	Ile	Leu	Asp	Asp	Glu	Lys	Arg	Ala	Lys	Asp	Tyr	Lys
		35					40					45			
Lys	Ala	Lys	Gln	Ile	Ile	Asp	Lys	Tyr	His	Gln	Phe	Phe	Ile	Glu	Glu
	50					55				60					
Ile	Leu	Ser	Ser	Val	Cys	Ile	Ser	Glu	Asp	Leu	Leu	Gln	Asn	Tyr	Ser
65				70					75					80	
Asp	Val	Tyr	Phe	Lys	Leu	Lys	Lys	Ser	Asp	Asp	Asp	Asn	Leu	Gln	Lys
				85					90					95	
Asp	Phe	Lys	Ser	Ala	Lys	Asp	Thr	Ile	Lys	Lys	Gln	Ile	Ser	Glu	Tyr
			100					105					110		
Ile	Lys	Asp	Ser	Glu	Lys	Phe	Lys	Asn	Leu	Phe	Asn	Gln	Asn	Leu	Ile
	115						120					125			
Asp	Ala	Lys	Lys	Gly	Gln	Glu	Ser	Asp	Leu	Ile	Leu	Trp	Leu	Lys	Gln
	130					135						140			
Ser	Lys	Asp	Asn	Gly	Ile	Glu	Leu	Phe	Lys	Ala	Asn	Ser	Asp	Ile	Thr
145				150						155				160	
Asp	Ile	Asp	Glu	Ala	Leu	Glu	Ile	Ile	Lys	Ser	Phe	Lys	Gly	Trp	Thr
				165					170					175	
Thr	Tyr	Phe	Lys	Gly	Phe	His	Glu	Asn	Arg	Lys	Asn	Val	Tyr	Ser	Ser
			180					185						190	
Asn	Asp	Ile	Pro	Thr	Ser	Ile	Ile	Tyr	Arg	Ile	Val	Asp	Asp	Asn	Leu
	195					200							205		
Pro	Lys	Phe	Leu	Glu	Asn	Lys	Ala	Lys	Tyr	Glu	Ser	Leu	Lys	Asp	Lys
	210					215							220		
Ala	Pro	Glu	Ala	Ile	Asn	Tyr	Glu	Gln	Ile	Lys	Lys	Asp	Leu	Ala	Glu
225					230					235				240	
Glu	Leu	Thr	Phe	Asp	Ile	Asp	Tyr	Lys	Thr	Ser	Glu	Val	Asn	Gln	Arg
				245					250					255	
Val	Phe	Ser	Leu	Asp	Glu	Val	Phe	Glu	Ile	Ala	Asn	Phe	Asn	Asn	Tyr
			260					265					270		
Leu	Asn	Gln	Ser	Gly	Ile	Thr	Lys	Phe	Asn	Thr	Ile	Ile	Gly	Gly	Lys
	275						280						285		

Phe Val Asn Gly Glu Asn Thr Lys Arg Lys Gly Ile Asn Glu Tyr Ile
 290 295 300
 Asn Leu Tyr Ser Gln Gln Ile Asn Asp Lys Thr Leu Lys Lys Tyr Lys
 305 310 315 320
 Met Ser Val Leu Phe Lys Gln Ile Leu Ser Asp Thr Glu Ser Lys Ser
 325 330 335
 Phe Val Ile Asp Lys Leu Glu Asp Asp Ser Asp Val Val Thr Thr Met
 340 345 350
 Gln Ser Phe Tyr Glu Gln Ile Ala Ala Phe Lys Thr Val Glu Glu Lys
 355 360 365
 Ser Ile Lys Glu Thr Leu Ser Leu Leu Phe Asp Asp Leu Lys Ala Gln
 370 375 380
 Lys Leu Asp Leu Ser Lys Ile Tyr Phe Lys Asn Asp Lys Ser Leu Thr
 385 390 395 400
 Asp Leu Ser Gln Gln Val Phe Asp Asp Tyr Ser Val Ile Gly Thr Ala
 405 410 415
 Val Leu Glu Tyr Ile Thr Gln Gln Ile Ala Pro Lys Asn Leu Asp Asn
 420 425 430
 Pro Ser Lys Lys Glu Gln Glu Leu Ile Ala Lys Lys Thr Glu Lys Ala
 435 440 445
 Lys Tyr Leu Ser Leu Glu Thr Ile Lys Leu Ala Leu Glu Glu Phe Asn
 450 455 460
 Lys His Arg Asp Ile Asp Lys Gln Cys Arg Phe Glu Glu Ile Leu Ala
 465 470 475 480
 Asn Phe Ala Ala Ile Pro Met Ile Phe Asp Glu Ile Ala Gln Asn Lys
 485 490 495
 Asp Asn Leu Ala Gln Ile Ser Ile Lys Tyr Gln Asn Gln Gly Lys Lys
 500 505 510
 Asp Leu Leu Gln Ala Ser Ala Glu Asp Asp Val Lys Ala Ile Lys Asp
 515 520 525
 Leu Leu Asp Gln Thr Asn Asn Leu Leu His Lys Leu Lys Ile Phe His
 530 535 540
 Ile Ser Gln Ser Glu Asp Lys Ala Asn Ile Leu Asp Lys Asp Glu His
 545 550 555 560
 Phe Tyr Leu Val Phe Glu Glu Cys Tyr Phe Glu Leu Ala Asn Ile Val
 565 570 575
 Pro Leu Tyr Asn Lys Ile Arg Asn Tyr Ile Thr Gln Lys Pro Tyr Ser
 580 585 590
 Asp Glu Lys Phe Lys Leu Asn Phe Glu Asn Ser Thr Leu Ala Asn Gly

595	600	605
Trp Asp Lys Asn Lys Glu Pro Asp Asn Thr Ala Ile Leu Phe Ile Lys		
610	615	620
Asp Asp Lys Tyr Tyr Leu Gly Val Met Asn Lys Lys Asn Asn Lys Ile		
625	630	635
640		
Phe Asp Asp Lys Ala Ile Lys Glu Asn Lys Gly Glu Gly Tyr Lys Lys		
645	650	655
Ile Val Tyr Lys Leu Leu Pro Gly Ala Asn Lys Met Leu Pro Lys Val		
660	665	670
Phe Phe Ser Ala Lys Ser Ile Lys Phe Tyr Asn Pro Ser Glu Asp Ile		
675	680	685
Leu Arg Ile Arg Asn His Ser Thr His Thr Lys Asn Gly Ser Pro Gln		
690	695	700
Lys Gly Tyr Glu Lys Phe Glu Phe Asn Ile Glu Asp Cys Arg Lys Phe		
705	710	715
720		
Ile Asp Phe Tyr Lys Gln Ser Ile Ser Lys His Pro Glu Trp Lys Asp		
725	730	735
Phe Gly Phe Arg Phe Ser Asp Thr Gln Arg Tyr Asn Ser Ile Asp Glu		
740	745	750
Phe Tyr Arg Glu Val Glu Asn Gln Gly Tyr Lys Leu Thr Phe Glu Asn		
755	760	765
Ile Ser Glu Ser Tyr Ile Asp Ser Val Val Asn Gln Gly Lys Leu Tyr		
770	775	780
Leu Phe Gln Ile Tyr Asn Lys Asp Phe Ser Ala Tyr Ser Lys Gly Arg		
785	790	795
800		
Pro Asn Leu His Thr Leu Tyr Trp Lys Ala Leu Phe Asp Glu Arg Asn		
805	810	815
Leu Gln Asp Val Val Tyr Lys Leu Asn Gly Glu Ala Glu Leu Phe Tyr		
820	825	830
Arg Lys Gln Ser Ile Pro Lys Lys Ile Thr His Pro Ala Lys Glu Ala		
835	840	845
Ile Ala Asn Lys Asn Lys Asp Asn Pro Lys Lys Glu Ser Val Phe Glu		
850	855	860
Tyr Asp Leu Ile Lys Asp Lys Arg Phe Thr Glu Asp Lys Phe Phe Phe		
865	870	875
880		
His Cys Pro Ile Thr Ile Asn Phe Lys Ser Ser Gly Ala Asn Lys Phe		
885	890	895
Asn Asp Glu Ile Asn Leu Leu Leu Lys Glu Lys Ala Asn Asp Val His		
900	905	910

Ile Leu Ser Ile Ala Arg Gly Glu Arg His Leu Ala Tyr Tyr Thr Leu
 915 920 925
 Val Asp Gly Lys Gly Asn Ile Ile Lys Gln Asp Thr Phe Asn Ile Ile
 930 935 940
 Gly Asn Asp Arg Met Lys Thr Asn Tyr His Asp Lys Leu Ala Ala Ile
 945 950 955 960
 Glu Lys Asp Arg Asp Ser Ala Arg Lys Asp Trp Lys Lys Ile Asn Asn
 965 970 975
 Ile Lys Glu Met Lys Glu Gly Tyr Leu Ser Gln Val Val His Glu Ile
 980 985 990
 Ala Lys Leu Val Ile Glu Tyr Asn Ala Ile Val Val Phe Glu Asp Leu
 995 1000 1005
 Asn Phe Gly Phe Lys Arg Gly Arg Phe Lys Val Glu Lys Gln Val
 1010 1015 1020
 Tyr Gln Lys Leu Glu Lys Met Leu Ile Glu Lys Leu Asn Tyr Leu
 1025 1030 1035
 Val Phe Lys Asp Asn Glu Phe Asp Lys Thr Gly Gly Val Leu Arg
 1040 1045 1050
 Ala Tyr Gln Leu Thr Ala Pro Phe Glu Thr Phe Lys Lys Met Gly
 1055 1060 1065
 Lys Gln Thr Gly Ile Ile Tyr Tyr Val Pro Ala Gly Phe Thr Ser
 1070 1075 1080
 Lys Ile Cys Pro Val Thr Gly Phe Val Asn Gln Leu Tyr Pro Lys
 1085 1090 1095
 Tyr Glu Ser Val Ser Lys Ser Gln Glu Phe Phe Ser Lys Phe Asp
 1100 1105 1110
 Lys Ile Cys Tyr Asn Leu Asp Lys Gly Tyr Phe Glu Phe Ser Phe
 1115 1120 1125
 Asp Tyr Lys Asn Phe Gly Asp Lys Ala Ala Lys Gly Lys Trp Thr
 1130 1135 1140
 Ile Ala Ser Phe Gly Ser Arg Leu Ile Asn Phe Arg Asn Ser Asp
 1145 1150 1155
 Lys Asn His Asn Trp Asp Thr Arg Glu Val Tyr Pro Thr Lys Glu
 1160 1165 1170
 Leu Glu Lys Leu Leu Lys Asp Tyr Ser Ile Glu Tyr Gly His Gly
 1175 1180 1185
 Glu Cys Ile Lys Ala Ala Ile Cys Gly Glu Ser Asp Lys Lys Phe
 1190 1195 1200
 Phe Ala Lys Leu Thr Ser Val Leu Asn Thr Ile Leu Gln Met Arg

1205	1210	1215
Asn Ser Lys Thr Gly Thr	Glu Leu Asp Tyr Leu	Ile Ser Pro Val
1220	1225	1230
Ala Asp Val Asn Gly Asn	Phe Phe Asp Ser Arg	Gln Ala Pro Lys
1235	1240	1245
Asn Met Pro Gln Asp Ala	Asp Ala Asn Gly Ala	Tyr His Ile Gly
1250	1255	1260
Leu Lys Gly Leu Met Leu	Leu Gly Arg Ile Lys	Asn Asn Gln Glu
1265	1270	1275
Gly Lys Lys Leu Asn Leu	Val Ile Lys Asn Glu	Glu Tyr Phe Glu
1280	1285	1290
Phe Val Gln Asn Arg Asn	Asn	
1295	1300	
<210> 716		
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<213> 新凶手弗朗西斯菌		
<400> 716		
Met Ser Ile Tyr Gln Glu	Phe Val Asn Lys Tyr Ser	Leu Ser Lys Thr
1	5	10
15		
Leu Arg Phe Glu Leu Ile	Pro Gln Gly Lys Thr	Leu Glu Asn Ile Lys
	20	25
		30
Ala Arg Gly Leu Ile Leu	Asp Asp Glu Lys Arg	Ala Lys Asp Tyr Lys
	35	40
		45
Lys Ala Lys Gln Ile Ile	Asp Lys Tyr His Gln	Phe Phe Ile Glu Glu
	50	55
		60
Ile Leu Ser Ser Val Cys	Ile Ser Glu Asp Leu	Leu Gln Asn Tyr Ser
65	70	75
		80
Asp Val Tyr Phe Lys Leu	Lys Lys Ser Asp Asp	Asp Asn Leu Gln Lys
	85	90
		95
Asp Phe Lys Ser Ala Lys	Asp Thr Ile Lys Lys	Gln Ile Ser Glu Tyr
	100	105
		110
Ile Lys Asp Ser Glu Lys	Phe Lys Asn Leu Phe	Asn Gln Asn Leu Ile
	115	120
		125
Asp Ala Lys Lys Gly Gln	Glu Ser Asp Leu Ile	Leu Trp Leu Lys Gln
	130	135
		140
Ser Lys Asp Asn Gly Ile	Glu Leu Phe Lys Ala	Asn Ser Asp Ile Thr
145	150	155
		160
Asp Ile Asp Glu Ala Leu	Glu Ile Ile Lys Ser	Phe Lys Gly Trp Thr

	165		170		175
Thr Tyr Phe Lys Gly Phe His Glu Asn Arg Lys Asn Val Tyr Ser Ser					
	180		185		190
Asn Asp Ile Pro Thr Ser Ile Ile Tyr Arg Ile Val Asp Asp Asn Leu					
	195		200		205
Pro Lys Phe Leu Glu Asn Lys Ala Lys Tyr Glu Ser Leu Lys Asp Lys					
	210		215		220
Ala Pro Glu Ala Ile Asn Tyr Glu Gln Ile Lys Lys Asp Leu Ala Glu					
225		230		235	240
Glu Leu Thr Phe Asp Ile Asp Tyr Lys Thr Ser Glu Val Asn Gln Arg					
	245		250		255
Val Phe Ser Leu Asp Glu Val Phe Glu Ile Ala Asn Phe Asn Asn Tyr					
	260		265		270
Leu Asn Gln Ser Gly Ile Thr Lys Phe Asn Thr Ile Ile Gly Gly Lys					
	275		280		285
Phe Val Asn Gly Glu Asn Thr Lys Arg Lys Gly Ile Asn Glu Tyr Ile					
	290		295		300
Asn Leu Tyr Ser Gln Gln Ile Asn Asp Lys Thr Leu Lys Lys Tyr Lys					
305		310		315	320
Met Ser Val Leu Phe Lys Gln Ile Leu Ser Asp Thr Glu Ser Lys Ser					
	325		330		335
Phe Val Ile Asp Lys Leu Glu Asp Asp Ser Asp Val Val Thr Thr Met					
	340		345		350
Gln Ser Phe Tyr Glu Gln Ile Ala Ala Phe Lys Thr Val Glu Glu Lys					
	355		360		365
Ser Ile Lys Glu Thr Leu Ser Leu Leu Phe Asp Asp Leu Lys Ala Gln					
	370		375		380
Lys Leu Asp Leu Ser Lys Ile Tyr Phe Lys Asn Asp Lys Ser Leu Thr					
385		390		395	400
Asp Leu Ser Gln Gln Val Phe Asp Asp Tyr Ser Val Ile Gly Thr Ala					
	405		410		415
Val Leu Glu Tyr Ile Thr Gln Gln Ile Ala Pro Lys Asn Leu Asp Asn					
	420		425		430
Pro Ser Lys Lys Glu Gln Glu Leu Ile Ala Lys Lys Thr Glu Lys Ala					
	435		440		445
Lys Tyr Leu Ser Leu Glu Thr Ile Lys Leu Ala Leu Glu Glu Phe Asn					
	450		455		460
Lys His Arg Asp Ile Asp Lys Gln Cys Arg Phe Glu Glu Ile Leu Ala					
465		470		475	480

Asn Phe Ala Ala Ile Pro Met Ile Phe Asp Glu Ile Ala Gln Asn Lys
 485 490 495
 Asp Asn Leu Ala Gln Ile Ser Ile Lys Tyr Gln Asn Gln Gly Lys Lys
 500 505 510
 Asp Leu Leu Gln Ala Ser Ala Glu Asp Asp Val Lys Ala Ile Lys Asp
 515 520 525
 Leu Leu Asp Gln Thr Asn Asn Leu Leu His Lys Leu Lys Ile Phe His
 530 535 540
 Ile Ser Gln Ser Glu Asp Lys Ala Asn Ile Leu Asp Lys Asp Glu His
 545 550 555 560
 Phe Tyr Leu Val Phe Glu Glu Cys Tyr Phe Glu Leu Ala Asn Ile Val
 565 570 575
 Pro Leu Tyr Asn Lys Ile Arg Asn Tyr Ile Thr Gln Lys Pro Tyr Ser
 580 585 590
 Asp Glu Lys Phe Lys Leu Asn Phe Glu Asn Ser Thr Leu Ala Asn Gly
 595 600 605
 Trp Asp Lys Asn Lys Glu Pro Asp Asn Thr Ala Ile Leu Phe Ile Lys
 610 615 620
 Asp Asp Lys Tyr Tyr Leu Gly Val Met Asn Lys Lys Asn Asn Lys Ile
 625 630 635 640
 Phe Asp Asp Lys Ala Ile Lys Glu Asn Lys Gly Glu Gly Tyr Lys Lys
 645 650 655
 Ile Val Tyr Lys Leu Leu Pro Gly Ala Asn Lys Met Leu Pro Lys Val
 660 665 670
 Phe Phe Ser Ala Lys Ser Ile Lys Phe Tyr Asn Pro Ser Glu Asp Ile
 675 680 685
 Leu Arg Ile Arg Asn His Ser Thr His Thr Lys Asn Gly Ser Pro Gln
 690 695 700
 Lys Gly Tyr Glu Lys Phe Glu Phe Asn Ile Glu Asp Cys Arg Lys Phe
 705 710 715 720
 Ile Asp Phe Tyr Lys Gln Ser Ile Ser Lys His Pro Glu Trp Lys Asp
 725 730 735
 Phe Gly Phe Arg Phe Ser Asp Thr Gln Arg Tyr Asn Ser Ile Asp Glu
 740 745 750
 Phe Tyr Arg Glu Val Glu Asn Gln Gly Tyr Lys Leu Thr Phe Glu Asn
 755 760 765
 Ile Ser Glu Ser Tyr Ile Asp Ser Val Val Asn Gln Gly Lys Leu Tyr
 770 775 780
 Leu Phe Gln Ile Tyr Asn Lys Asp Phe Ser Ala Tyr Ser Lys Gly Arg

785	790	795	800
Pro Asn Leu His Thr Leu Tyr Trp Lys Ala Leu Phe Asp Glu Arg Asn			
	805	810	815
Leu Gln Asp Val Val Tyr Lys Leu Asn Gly Glu Ala Glu Leu Phe Tyr			
	820	825	830
Arg Lys Gln Ser Ile Pro Lys Lys Ile Thr His Pro Ala Lys Glu Ala			
	835	840	845
Ile Ala Asn Lys Asn Lys Asp Asn Pro Lys Lys Glu Ser Val Phe Glu			
	850	855	860
Tyr Asp Leu Ile Lys Asp Lys Arg Phe Thr Glu Asp Lys Phe Phe Phe			
865	870	875	880
His Cys Pro Ile Thr Ile Asn Phe Lys Ser Ser Gly Ala Asn Lys Phe			
	885	890	895
Asn Asp Glu Ile Asn Leu Leu Leu Lys Glu Lys Ala Asn Asp Val His			
	900	905	910
Ile Leu Ser Ile Asp Arg Gly Glu Arg His Leu Ala Tyr Tyr Thr Leu			
	915	920	925
Val Asp Gly Lys Gly Asn Ile Ile Lys Gln Asp Thr Phe Asn Ile Ile			
	930	935	940
Gly Asn Asp Arg Met Lys Thr Asn Tyr His Asp Lys Leu Ala Ala Ile			
945	950	955	960
Glu Lys Asp Arg Asp Ser Ala Arg Lys Asp Trp Lys Lys Ile Asn Asn			
	965	970	975
Ile Lys Glu Met Lys Glu Gly Tyr Leu Ser Gln Val Val His Glu Ile			
	980	985	990
Ala Lys Leu Val Ile Glu Tyr Asn Ala Ile Val Val Phe Ala Asp Leu			
	995	1000	1005
Asn Phe Gly Phe Lys Arg Gly Arg Phe Lys Val Glu Lys Gln Val			
	1010	1015	1020
Tyr Gln Lys Leu Glu Lys Met Leu Ile Glu Lys Leu Asn Tyr Leu			
	1025	1030	1035
Val Phe Lys Asp Asn Glu Phe Asp Lys Thr Gly Gly Val Leu Arg			
	1040	1045	1050
Ala Tyr Gln Leu Thr Ala Pro Phe Glu Thr Phe Lys Lys Met Gly			
	1055	1060	1065
Lys Gln Thr Gly Ile Ile Tyr Tyr Val Pro Ala Gly Phe Thr Ser			
	1070	1075	1080
Lys Ile Cys Pro Val Thr Gly Phe Val Asn Gln Leu Tyr Pro Lys			
	1085	1090	1095

Tyr Glu Ser Val Ser Lys Ser Gln Glu Phe Phe Ser Lys Phe Asp
 1100 1105 1110
 Lys Ile Cys Tyr Asn Leu Asp Lys Gly Tyr Phe Glu Phe Ser Phe
 1115 1120 1125
 Asp Tyr Lys Asn Phe Gly Asp Lys Ala Ala Lys Gly Lys Trp Thr
 1130 1135 1140
 Ile Ala Ser Phe Gly Ser Arg Leu Ile Asn Phe Arg Asn Ser Asp
 1145 1150 1155
 Lys Asn His Asn Trp Asp Thr Arg Glu Val Tyr Pro Thr Lys Glu
 1160 1165 1170
 Leu Glu Lys Leu Leu Lys Asp Tyr Ser Ile Glu Tyr Gly His Gly
 1175 1180 1185
 Glu Cys Ile Lys Ala Ala Ile Cys Gly Glu Ser Asp Lys Lys Phe
 1190 1195 1200
 Phe Ala Lys Leu Thr Ser Val Leu Asn Thr Ile Leu Gln Met Arg
 1205 1210 1215
 Asn Ser Lys Thr Gly Thr Glu Leu Asp Tyr Leu Ile Ser Pro Val
 1220 1225 1230
 Ala Asp Val Asn Gly Asn Phe Phe Asp Ser Arg Gln Ala Pro Lys
 1235 1240 1245
 Asn Met Pro Gln Asp Ala Asp Ala Asn Gly Ala Tyr His Ile Gly
 1250 1255 1260
 Leu Lys Gly Leu Met Leu Leu Gly Arg Ile Lys Asn Asn Gln Glu
 1265 1270 1275
 Gly Lys Lys Leu Asn Leu Val Ile Lys Asn Glu Glu Tyr Phe Glu
 1280 1285 1290
 Phe Val Gln Asn Arg Asn Asn
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<210> 717

<211> 1300

<212> PRT

<213> 新凶手弗朗西斯菌

<400> 717

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 Leu Arg Phe Glu Leu Ile Pro Gln Gly Lys Thr Leu Glu Asn Ile Lys
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 Ala Arg Gly Leu Ile Leu Asp Asp Glu Lys Arg Ala Lys Asp Tyr Lys
 35 40 45

Lys Ala Lys Gln Ile Ile Asp Lys Tyr His Gln Phe Phe Ile Glu Glu
 50 55 60
 Ile Leu Ser Ser Val Cys Ile Ser Glu Asp Leu Leu Gln Asn Tyr Ser
 65 70 75 80
 Asp Val Tyr Phe Lys Leu Lys Lys Ser Asp Asp Asp Asn Leu Gln Lys
 85 90 95
 Asp Phe Lys Ser Ala Lys Asp Thr Ile Lys Lys Gln Ile Ser Glu Tyr
 100 105 110
 Ile Lys Asp Ser Glu Lys Phe Lys Asn Leu Phe Asn Gln Asn Leu Ile
 115 120 125
 Asp Ala Lys Lys Gly Gln Glu Ser Asp Leu Ile Leu Trp Leu Lys Gln
 130 135 140
 Ser Lys Asp Asn Gly Ile Glu Leu Phe Lys Ala Asn Ser Asp Ile Thr
 145 150 155 160
 Asp Ile Asp Glu Ala Leu Glu Ile Ile Lys Ser Phe Lys Gly Trp Thr
 165 170 175
 Thr Tyr Phe Lys Gly Phe His Glu Asn Arg Lys Asn Val Tyr Ser Ser
 180 185 190
 Asn Asp Ile Pro Thr Ser Ile Ile Tyr Arg Ile Val Asp Asp Asn Leu
 195 200 205
 Pro Lys Phe Leu Glu Asn Lys Ala Lys Tyr Glu Ser Leu Lys Asp Lys
 210 215 220
 Ala Pro Glu Ala Ile Asn Tyr Glu Gln Ile Lys Lys Asp Leu Ala Glu
 225 230 235 240
 Glu Leu Thr Phe Asp Ile Asp Tyr Lys Thr Ser Glu Val Asn Gln Arg
 245 250 255
 Val Phe Ser Leu Asp Glu Val Phe Glu Ile Ala Asn Phe Asn Asn Tyr
 260 265 270
 Leu Asn Gln Ser Gly Ile Thr Lys Phe Asn Thr Ile Ile Gly Gly Lys
 275 280 285
 Phe Val Asn Gly Glu Asn Thr Lys Arg Lys Gly Ile Asn Glu Tyr Ile
 290 295 300
 Asn Leu Tyr Ser Gln Gln Ile Asn Asp Lys Thr Leu Lys Lys Tyr Lys
 305 310 315 320
 Met Ser Val Leu Phe Lys Gln Ile Leu Ser Asp Thr Glu Ser Lys Ser
 325 330 335
 Phe Val Ile Asp Lys Leu Glu Asp Asp Ser Asp Val Val Thr Thr Met
 340 345 350
 Gln Ser Phe Tyr Glu Gln Ile Ala Ala Phe Lys Thr Val Glu Glu Lys

355	360	365
Ser Ile Lys Glu Thr Leu	Ser Leu Leu Phe Asp	Asp Leu Lys Ala Gln
370	375	380
Lys Leu Asp Leu Ser Lys	Ile Tyr Phe Lys Asn	Asp Lys Ser Leu Thr
385	390	395
Asp Leu Ser Gln Gln Val	Phe Asp Asp Tyr Ser	Val Ile Gly Thr Ala
405	410	415
Val Leu Glu Tyr Ile Thr	Gln Gln Ile Ala Pro	Lys Asn Leu Asp Asn
420	425	430
Pro Ser Lys Lys Glu Gln	Glu Leu Ile Ala Lys	Lys Thr Glu Lys Ala
435	440	445
Lys Tyr Leu Ser Leu Glu	Thr Ile Lys Leu Ala	Leu Glu Glu Phe Asn
450	455	460
Lys His Arg Asp Ile Asp	Lys Gln Cys Arg Phe	Glu Glu Ile Leu Ala
465	470	475
Asn Phe Ala Ala Ile Pro	Met Ile Phe Asp Glu	Ile Ala Gln Asn Lys
485	490	495
Asp Asn Leu Ala Gln Ile	Ser Ile Lys Tyr Gln	Asn Gln Gly Lys Lys
500	505	510
Asp Leu Leu Gln Ala Ser	Ala Glu Asp Asp Val	Lys Ala Ile Lys Asp
515	520	525
Leu Leu Asp Gln Thr Asn	Asn Leu Leu His Lys	Leu Lys Ile Phe His
530	535	540
Ile Ser Gln Ser Glu Asp	Lys Ala Asn Ile Leu	Asp Lys Asp Glu His
545	550	555
Phe Tyr Leu Val Phe Glu	Glu Cys Tyr Phe Glu	Leu Ala Asn Ile Val
565	570	575
Pro Leu Tyr Asn Lys Ile	Arg Asn Tyr Ile Thr	Gln Lys Pro Tyr Ser
580	585	590
Asp Glu Lys Phe Lys Leu	Asn Phe Glu Asn Ser	Thr Leu Ala Asn Gly
595	600	605
Trp Asp Lys Asn Lys Glu	Pro Asp Asn Thr Ala	Ile Leu Phe Ile Lys
610	615	620
Asp Asp Lys Tyr Tyr Leu	Gly Val Met Asn Lys	Lys Lys Asn Asn Lys Ile
625	630	635
Phe Asp Asp Lys Ala Ile	Lys Glu Asn Lys Gly	Glu Gly Tyr Lys Lys
645	650	655
Ile Val Tyr Lys Leu Leu	Pro Gly Ala Asn Lys	Met Leu Pro Lys Val
660	665	670

Phe Phe Ser Ala Lys Ser Ile Lys Phe Tyr Asn Pro Ser Glu Asp Ile
 675 680 685
 Leu Arg Ile Arg Asn His Ser Thr His Thr Lys Asn Gly Ser Pro Gln
 690 695 700
 Lys Gly Tyr Glu Lys Phe Glu Phe Asn Ile Glu Asp Cys Arg Lys Phe
 705 710 715 720
 Ile Asp Phe Tyr Lys Gln Ser Ile Ser Lys His Pro Glu Trp Lys Asp
 725 730 735
 Phe Gly Phe Arg Phe Ser Asp Thr Gln Arg Tyr Asn Ser Ile Asp Glu
 740 745 750
 Phe Tyr Arg Glu Val Glu Asn Gln Gly Tyr Lys Leu Thr Phe Glu Asn
 755 760 765
 Ile Ser Glu Ser Tyr Ile Asp Ser Val Val Asn Gln Gly Lys Leu Tyr
 770 775 780
 Leu Phe Gln Ile Tyr Asn Lys Asp Phe Ser Ala Tyr Ser Lys Gly Arg
 785 790 795 800
 Pro Asn Leu His Thr Leu Tyr Trp Lys Ala Leu Phe Asp Glu Arg Asn
 805 810 815
 Leu Gln Asp Val Val Tyr Lys Leu Asn Gly Glu Ala Glu Leu Phe Tyr
 820 825 830
 Arg Lys Gln Ser Ile Pro Lys Lys Ile Thr His Pro Ala Lys Glu Ala
 835 840 845
 Ile Ala Asn Lys Asn Lys Asp Asn Pro Lys Lys Glu Ser Val Phe Glu
 850 855 860
 Tyr Asp Leu Ile Lys Asp Lys Arg Phe Thr Glu Asp Lys Phe Phe Phe
 865 870 875 880
 His Cys Pro Ile Thr Ile Asn Phe Lys Ser Ser Gly Ala Asn Lys Phe
 885 890 895
 Asn Asp Glu Ile Asn Leu Leu Leu Lys Glu Lys Ala Asn Asp Val His
 900 905 910
 Ile Leu Ser Ile Asp Arg Gly Glu Arg His Leu Ala Tyr Tyr Thr Leu
 915 920 925
 Val Asp Gly Lys Gly Asn Ile Ile Lys Gln Asp Thr Phe Asn Ile Ile
 930 935 940
 Gly Asn Asp Arg Met Lys Thr Asn Tyr His Asp Lys Leu Ala Ala Ile
 945 950 955 960
 Glu Lys Asp Arg Asp Ser Ala Arg Lys Asp Trp Lys Lys Ile Asn Asn
 965 970 975
 Ile Lys Glu Met Lys Glu Gly Tyr Leu Ser Gln Val Val His Glu Ile

980	985	990
Ala Lys Leu Val Ile Glu Tyr Asn	Ala Ile Val Val Phe Glu Asp Leu	
995	1000	1005
Asn Phe Gly Phe Lys Arg Gly Arg Phe Lys Val Glu Lys Gln Val		
1010	1015	1020
Tyr Gln Lys Leu Glu Lys Met Leu Ile Glu Lys Leu Asn Tyr Leu		
1025	1030	1035
Val Phe Lys Asp Asn Glu Phe Asp Lys Thr Gly Gly Val Leu Arg		
1040	1045	1050
Ala Tyr Gln Leu Thr Ala Pro Phe Glu Thr Phe Lys Lys Met Gly		
1055	1060	1065
Lys Gln Thr Gly Ile Ile Tyr Tyr Val Pro Ala Gly Phe Thr Ser		
1070	1075	1080
Lys Ile Cys Pro Val Thr Gly Phe Val Asn Gln Leu Tyr Pro Lys		
1085	1090	1095
Tyr Glu Ser Val Ser Lys Ser Gln Glu Phe Phe Ser Lys Phe Asp		
1100	1105	1110
Lys Ile Cys Tyr Asn Leu Asp Lys Gly Tyr Phe Glu Phe Ser Phe		
1115	1120	1125
Asp Tyr Lys Asn Phe Gly Asp Lys Ala Ala Lys Gly Lys Trp Thr		
1130	1135	1140
Ile Ala Ser Phe Gly Ser Arg Leu Ile Asn Phe Arg Asn Ser Asp		
1145	1150	1155
Lys Asn His Asn Trp Asp Thr Arg Glu Val Tyr Pro Thr Lys Glu		
1160	1165	1170
Leu Glu Lys Leu Leu Lys Asp Tyr Ser Ile Glu Tyr Gly His Gly		
1175	1180	1185
Glu Cys Ile Lys Ala Ala Ile Cys Gly Glu Ser Asp Lys Lys Phe		
1190	1195	1200
Phe Ala Lys Leu Thr Ser Val Leu Asn Thr Ile Leu Gln Met Arg		
1205	1210	1215
Asn Ser Lys Thr Gly Thr Glu Leu Asp Tyr Leu Ile Ser Pro Val		
1220	1225	1230
Ala Asp Val Asn Gly Asn Phe Phe Asp Ser Arg Gln Ala Pro Lys		
1235	1240	1245
Asn Met Pro Gln Asp Ala Ala Ala Asn Gly Ala Tyr His Ile Gly		
1250	1255	1260
Leu Lys Gly Leu Met Leu Leu Gly Arg Ile Lys Asn Asn Gln Glu		
1265	1270	1275

Gly Lys Lys Leu Asn Leu Val Ile Lys Asn Glu Glu Tyr Phe Glu
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 Phe Val Gln Asn Arg Asn Asn
 1295 1300
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 Ser Gly His Thr Tyr Asp Ile Ser Val Thr Leu Thr Gly Val Tyr Asp
 20 25 30
 Asn Thr Asp Glu Gln His Pro Arg Met Ser Leu Ala Phe Glu Gln Asp
 35 40 45
 Asn Gly Glu Arg Arg Tyr Ile Thr Leu Trp Lys Asn Thr Thr Pro Lys
 50 55 60
 Asp Val Phe Thr Tyr Asp Tyr Ala Thr Gly Ser Thr Tyr Ile Phe Thr
 65 70 75 80
 Asn Ile Asp Tyr Glu Val Lys Asp Gly Tyr Glu Asn Leu Thr Ala Thr
 85 90 95
 Tyr Gln Thr Thr Val Glu Asn Ala Thr Ala Gln Glu Val Gly Thr Thr
 100 105 110
 Asp Glu Asp Glu Thr Phe Ala Gly Gly Glu Pro Leu Asp His His Leu
 115 120 125
 Asp Asp Ala Leu Asn Glu Thr Pro Asp Asp Ala Glu Thr Glu Ser Asp
 130 135 140
 Ser Gly His Val Met Thr Ser Phe Ala Ser Arg Asp Gln Leu Pro Glu
 145 150 155 160
 Trp Thr Leu His Thr Tyr Thr Leu Thr Ala Thr Asp Gly Ala Lys Thr
 165 170 175
 Asp Thr Glu Tyr Ala Arg Arg Thr Leu Ala Tyr Thr Val Arg Gln Glu
 180 185 190
 Leu Tyr Thr Asp His Asp Ala Ala Pro Val Ala Thr Asp Gly Leu Met
 195 200 205
 Leu Leu Thr Pro Glu Pro Leu Gly Glu Thr Pro Leu Asp Leu Asp Cys
 210 215 220
 Gly Val Arg Val Glu Ala Asp Glu Thr Arg Thr Leu Asp Tyr Thr Thr
 225 230 235 240

Ala Lys Asp Arg Leu Leu Ala Arg Glu Leu Val Glu Glu Gly Leu Lys
 245 250 255
 Arg Ser Leu Trp Asp Asp Tyr Leu Val Arg Gly Ile Asp Glu Val Leu
 260 265 270
 Ser Lys Glu Pro Val Leu Thr Cys Asp Glu Phe Asp Leu His Glu Arg
 275 280 285
 Tyr Asp Leu Ser Val Glu Val Gly His Ser Gly Arg Ala Tyr Leu His
 290 295 300
 Ile Asn Phe Arg His Arg Phe Val Pro Lys Leu Thr Leu Ala Asp Ile
 305 310 315 320
 Asp Asp Asp Asn Ile Tyr Pro Gly Leu Arg Val Lys Thr Thr Tyr Arg
 325 330 335
 Pro Arg Arg Gly His Ile Val Trp Gly Leu Arg Asp Glu Cys Ala Thr
 340 345 350
 Asp Ser Leu Asn Thr Leu Gly Asn Gln Ser Val Val Ala Tyr His Arg
 355 360 365
 Asn Asn Gln Thr Pro Ile Asn Thr Asp Leu Leu Asp Ala Ile Glu Ala
 370 375 380
 Ala Asp Arg Arg Val Val Glu Thr Arg Arg Gln Gly His Gly Asp Asp
 385 390 395 400
 Ala Val Ser Phe Pro Gln Glu Leu Leu Ala Val Glu Pro Asn Thr His
 405 410 415
 Gln Ile Lys Gln Phe Ala Ser Asp Gly Phe His Gln Gln Ala Arg Ser
 420 425 430
 Lys Thr Arg Leu Ser Ala Ser Arg Cys Ser Glu Lys Ala Gln Ala Phe
 435 440 445
 Ala Glu Arg Leu Asp Pro Val Arg Leu Asn Gly Ser Thr Val Glu Phe
 450 455 460
 Ser Ser Glu Phe Phe Thr Gly Asn Asn Glu Gln Gln Leu Arg Leu Leu
 465 470 475 480
 Tyr Glu Asn Gly Glu Ser Val Leu Thr Phe Arg Asp Gly Ala Arg Gly
 485 490 495
 Ala His Pro Asp Glu Thr Phe Ser Lys Gly Ile Val Asn Pro Pro Glu
 500 505 510
 Ser Phe Glu Val Ala Val Val Leu Pro Glu Gln Gln Ala Asp Thr Cys
 515 520 525
 Lys Ala Gln Trp Asp Thr Met Ala Asp Leu Leu Asn Gln Ala Gly Ala
 530 535 540
 Pro Pro Thr Arg Ser Glu Thr Val Gln Tyr Asp Ala Phe Ser Ser Pro

545	550	555	560
Glu Ser Ile Ser Leu Asn Val Ala Gly Ala Ile Asp Pro Ser Glu Val			
	565	570	575
Asp Ala Ala Phe Val Val Leu Pro Pro Asp Gln Glu Gly Phe Ala Asp			
	580	585	590
Leu Ala Ser Pro Thr Glu Thr Tyr Asp Glu Leu Lys Lys Ala Leu Ala			
	595	600	605
Asn Met Gly Ile Tyr Ser Gln Met Ala Tyr Phe Asp Arg Phe Arg Asp			
610	615	620	
Ala Lys Ile Phe Tyr Thr Arg Asn Val Ala Leu Gly Leu Leu Ala Ala			
625	630	635	640
Ala Gly Gly Val Ala Phe Thr Thr Glu His Ala Met Pro Gly Asp Ala			
	645	650	655
Asp Met Phe Ile Gly Ile Asp Val Ser Arg Ser Tyr Pro Glu Asp Gly			
	660	665	670
Ala Ser Gly Gln Ile Asn Ile Ala Ala Thr Ala Thr Ala Val Tyr Lys			
	675	680	685
Asp Gly Thr Ile Leu Gly His Ser Ser Thr Arg Pro Gln Leu Gly Glu			
690	695	700	
Lys Leu Gln Ser Thr Asp Val Arg Asp Ile Met Lys Asn Ala Ile Leu			
705	710	715	720
Gly Tyr Gln Gln Val Thr Gly Glu Ser Pro Thr His Ile Val Ile His			
	725	730	735
Arg Asp Gly Phe Met Asn Glu Asp Leu Asp Pro Ala Thr Glu Phe Leu			
	740	745	750
Asn Glu Gln Gly Val Glu Tyr Asp Ile Val Glu Ile Arg Lys Gln Pro			
	755	760	765
Gln Thr Arg Leu Leu Ala Val Ser Asp Val Gln Tyr Asp Thr Pro Val			
	770	775	780
Lys Ser Ile Ala Ala Ile Asn Gln Asn Glu Pro Arg Ala Thr Val Ala			
785	790	795	800
Thr Phe Gly Ala Pro Glu Tyr Leu Ala Thr Arg Asp Gly Gly Gly Leu			
	805	810	815
Pro Arg Pro Ile Gln Ile Glu Arg Val Ala Gly Glu Thr Asp Ile Glu			
	820	825	830
Thr Leu Thr Arg Gln Val Tyr Leu Leu Ser Gln Ser His Ile Gln Val			
	835	840	845
His Asn Ser Thr Ala Arg Leu Pro Ile Thr Thr Ala Tyr Ala Asp Gln			
850	855	860	

Ala Ser Thr His Ala Thr Lys Gly Tyr Leu Val Gln Thr Gly Ala Phe
865 870 875 880
Glu Ser Asn Val Gly Phe Leu
885

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<212> PRT
<213> 人工序列
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Leu Gln Arg Asn Ala Leu Val Cys Ala Gly Cys Glu Gln Ile Phe Glu
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Asp Lys Leu Ser Gly Thr Arg Thr Asp Arg Pro Gly Leu Lys Arg Ala
35 40 45
Leu Lys Arg Leu Gln Lys Gly Asp Thr Leu Val Val Trp Lys Leu Asp
50 55 60
Arg Leu Gly Arg Ser Met Lys His Leu Ile Ser Leu Val Gly Glu Leu
65 70 75 80
Arg Glu Arg Gly Ile Asn Phe Arg Ser Leu Thr Asp Ser Ile Asp Thr
85 90 95
Ser Ser Pro Met Gly Arg Phe Phe Phe Tyr Val Met Gly Ala Leu Ala
100 105 110
Glu Met Glu Arg Glu Leu Ile Ile Glu Arg Thr Met Ala Gly Leu Ala
115 120 125
Ala Ala Arg Asn Lys Gly Arg Arg Phe Gly Arg Pro Pro Lys Gly Gly
130 135 140
Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser
145 150 155 160
Gly Gly Ser Gly Gly Ser Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile
165 170 175
Gly Thr Asn Ser Val Gly Trp Ala Val Ile Thr Asp Glu Tyr Lys Val
180 185 190
Pro Ser Lys Lys Phe Lys Val Leu Gly Asn Thr Asp Arg His Ser Ile
195 200 205
Lys Lys Asn Leu Ile Gly Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala
210 215 220

Glu Ala Thr Arg Leu Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg																			
225					230					235									240
Lys Asn Arg Ile Cys Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala																			
					245					250									255
Lys Val Asp Asp Ser Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val																			
					260					265									270
Glu Glu Asp Lys Lys His Glu Arg His Pro Ile Phe Gly Asn Ile Val																			
					275					280									285
Asp Glu Val Ala Tyr His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg																			
					290					295									300
Lys Lys Leu Val Asp Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr																			
305					310					315									320
Leu Ala Leu Ala His Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu																			
					325					330									335
Gly Asp Leu Asn Pro Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln																			
					340					345									350
Leu Val Gln Thr Tyr Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala																			
					355					360									365
Ser Gly Val Asp Ala Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser																			
					370					375									380
Arg Arg Leu Glu Asn Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn																			
385					390					395									400
Gly Leu Phe Gly Asn Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn																			
					405					410									415
Phe Lys Ser Asn Phe Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser																			
					420					425									430
Lys Asp Thr Tyr Asp Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly																			
					435					440									445
Asp Gln Tyr Ala Asp Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala																			
					450					455									460
Ile Leu Leu Ser Asp Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala																			
465					470					475									480
Pro Leu Ser Ala Ser Met Ile Lys Arg Tyr Asp Glu His His Gln Asp																			
					485					490									495
Leu Thr Leu Leu Lys Ala Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr																			
					500					505									510
Lys Glu Ile Phe Phe Asp Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile																			
					515					520									525
Asp Gly Gly Ala Ser Gln Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile																			

Lys Ser Asp Gly Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp
 850 855 860
 Asp Ser Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly
 865 870 875 880
 Gln Gly Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro
 885 890 895
 Ala Ile Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu
 900 905 910
 Val Lys Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met
 915 920 925
 Ala Arg Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu
 930 935 940
 Arg Met Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile
 945 950 955 960
 Leu Lys Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu
 965 970 975
 Tyr Leu Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val Asp Gln Glu
 980 985 990
 Leu Asp Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp Ala Ile Val Pro
 995 1000 1005
 Gln Ser Phe Leu Lys Asp Asp Ser Ile Asp Asn Lys Val Leu Thr
 1010 1015 1020
 Arg Ser Asp Lys Asn Arg Gly Lys Ser Asp Asn Val Pro Ser Glu
 1025 1030 1035
 Glu Val Val Lys Lys Met Lys Asn Tyr Trp Arg Gln Leu Leu Asn
 1040 1045 1050
 Ala Lys Leu Ile Thr Gln Arg Lys Phe Asp Asn Leu Thr Lys Ala
 1055 1060 1065
 Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys Ala Gly Phe Ile Lys
 1070 1075 1080
 Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys His Val Ala Gln
 1085 1090 1095
 Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu Asn Asp Lys
 1100 1105 1110
 Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys Leu Val
 1115 1120 1125
 Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu Ile
 1130 1135 1140
 Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val

1145	1150	1155
Gly Thr Ala Leu Ile Lys	Lys Tyr Pro Lys Leu	Glu Ser Glu Phe
1160	1165	1170
Val Tyr Gly Asp Tyr Lys	Val Tyr Asp Val Arg	Lys Met Ile Ala
1175	1180	1185
Lys Ser Glu Gln Glu Ile	Gly Lys Ala Thr Ala	Lys Tyr Phe Phe
1190	1195	1200
Tyr Ser Asn Ile Met Asn	Phe Phe Lys Thr Glu	Ile Thr Leu Ala
1205	1210	1215
Asn Gly Glu Ile Arg Lys	Arg Pro Leu Ile Glu	Thr Asn Gly Glu
1220	1225	1230
Thr Gly Glu Ile Val Trp	Asp Lys Gly Arg Asp	Phe Ala Thr Val
1235	1240	1245
Arg Lys Val Leu Ser Met	Pro Gln Val Asn Ile	Val Lys Lys Thr
1250	1255	1260
Glu Val Gln Thr Gly Gly	Phe Ser Lys Glu Ser	Ile Leu Pro Lys
1265	1270	1275
Arg Asn Ser Asp Lys Leu	Ile Ala Arg Lys Lys	Asp Trp Asp Pro
1280	1285	1290
Lys Lys Tyr Gly Gly Phe	Asp Ser Pro Thr Val	Ala Tyr Ser Val
1295	1300	1305
Leu Val Val Ala Lys Val	Glu Lys Gly Lys Ser	Lys Lys Leu Lys
1310	1315	1320
Ser Val Lys Glu Leu Leu	Gly Ile Thr Ile Met	Glu Arg Ser Ser
1325	1330	1335
Phe Glu Lys Asn Pro Ile	Asp Phe Leu Glu Ala	Lys Gly Tyr Lys
1340	1345	1350
Glu Val Lys Lys Asp Leu	Ile Ile Lys Leu Pro	Lys Tyr Ser Leu
1355	1360	1365
Phe Glu Leu Glu Asn Gly	Arg Lys Arg Met Leu	Ala Ser Ala Gly
1370	1375	1380
Glu Leu Gln Lys Gly Asn	Glu Leu Ala Leu Pro	Ser Lys Tyr Val
1385	1390	1395
Asn Phe Leu Tyr Leu Ala	Ser His Tyr Glu Lys	Leu Lys Gly Ser
1400	1405	1410
Pro Glu Asp Asn Glu Gln	Lys Gln Leu Phe Val	Glu Gln His Lys
1415	1420	1425
His Tyr Leu Asp Glu Ile	Ile Glu Gln Ile Ser	Glu Phe Ser Lys
1430	1435	1440

Arg Val Ile Leu Ala Asp Ala Asn Leu Asp Lys Val Leu Ser Ala
 1445 1450 1455
 Tyr Asn Lys His Arg Asp Lys Pro Ile Arg Glu Gln Ala Glu Asn
 1460 1465 1470
 Ile Ile His Leu Phe Thr Leu Thr Asn Leu Gly Ala Pro Ala Ala
 1475 1480 1485
 Phe Lys Tyr Phe Asp Thr Thr Ile Asp Arg Lys Arg Tyr Thr Ser
 1490 1495 1500
 Thr Lys Glu Val Leu Asp Ala Thr Leu Ile His Gln Ser Ile Thr
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 35 40 45
 Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala Thr Arg Leu Lys
 50 55 60
 Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys Tyr
 65 70 75 80
 Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser Phe
 85 90 95
 Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys His
 100 105 110
 Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr His
 115 120 125
 Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp Ser
 130 135 140

Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His Met
 145 150 155 160
 Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro Asp
 165 170 175
 Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val Gln Thr Tyr Asn
 180 185 190
 Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala Lys
 195 200 205
 Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg Leu Glu Asn Leu
 210 215 220
 Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu Phe Gly Asn Leu
 225 230 235 240
 Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe Asp
 245 250 255
 Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp Asp
 260 265 270
 Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp Leu
 275 280 285
 Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp Ile
 290 295 300
 Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser Met
 305 310 315 320
 Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys Ala
 325 330 335
 Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe Asp
 340 345 350
 Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser Gln
 355 360 365
 Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp Gly
 370 375 380
 Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg Lys
 385 390 395 400
 Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu Gly
 405 410 415
 Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe Leu
 420 425 430
 Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile Pro
 435 440 445
 Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp Met

450	455	460
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465	470	475
Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr Ala		
	485	490
Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser Leu		
	500	505
Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys Tyr		
	515	520
Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln Lys		
	530	535
Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr Val		
545	550	555
Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp Ser		
	565	570
Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly Thr		
	580	585
Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp Asn		
	595	600
Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr Leu		
	610	615
Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys Thr Tyr Ala His		
625	630	635
Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr Thr		
	645	650
Gly Trp Gly Ala Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp Lys		
	660	665
Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe Ala		
	675	680
Asn Arg Asn Phe Met Ala Leu Ile His Asp Asp Ser Leu Thr Phe Lys		
	690	695
Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly Asp Ser Leu His		
705	710	715
Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly Ile		
	725	730
Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys Val Met Gly Arg		
	740	745
His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln Thr		
	755	760
		765

Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met Lys Arg Ile Glu			
770	775	780	
Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys Glu His Pro Val			
785	790	795	800
Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu Tyr Tyr Leu Gln			
	805	810	815
Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp Ile Asn Arg Leu			
	820	825	830
Ser Asp Tyr Asp Val Asp His Ile Val Pro Gln Ser Phe Leu Lys Asp			
	835	840	845
Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly			
	850	855	860
Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn			
865	870	875	880
Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe			
	885	890	895
Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys			
	900	905	910
Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Ala Ile Thr Lys			
	915	920	925
His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu			
	930	935	940
Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys			
945	950	955	960
Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu			
	965	970	975
Ile Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val			
	980	985	990
Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe Val			
	995	1000	1005
Tyr Gly Asp Tyr Lys Val Tyr Asp Val Arg Lys Met Ile Ala Lys			
	1010	1015	1020
Ser Glu Gln Glu Ile Gly Lys Ala Thr Ala Lys Tyr Phe Phe Tyr			
	1025	1030	1035
Ser Asn Ile Met Asn Phe Phe Lys Thr Glu Ile Thr Leu Ala Asn			
	1040	1045	1050
Gly Glu Ile Arg Lys Arg Pro Leu Ile Glu Thr Asn Gly Glu Thr			
	1055	1060	1065
Gly Glu Ile Val Trp Asp Lys Gly Arg Asp Phe Ala Thr Val Arg			

1070	1075	1080
Lys Val Leu Ser Met Pro Gln Val Asn Ile Val Lys Lys Thr Glu		
1085	1090	1095
Val Gln Thr Gly Gly Phe Ser Lys Glu Ser Ile Leu Pro Lys Arg		
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Asn Ser Asp Lys Leu Ile Ala Arg Lys Lys Asp Trp Asp Pro Lys		
1115	1120	1125
Lys Tyr Gly Gly Phe Asp Ser Pro Thr Val Ala Tyr Ser Val Leu		
1130	1135	1140
Val Val Ala Lys Val Glu Lys Gly Lys Ser Lys Lys Leu Lys Ser		
1145	1150	1155
Val Lys Glu Leu Leu Gly Ile Thr Ile Met Glu Arg Ser Ser Phe		
1160	1165	1170
Glu Lys Asn Pro Ile Asp Phe Leu Glu Ala Lys Gly Tyr Lys Glu		
1175	1180	1185
Val Lys Lys Asp Leu Ile Ile Lys Leu Pro Lys Tyr Ser Leu Phe		
1190	1195	1200
Glu Leu Glu Asn Gly Arg Lys Arg Met Leu Ala Ser Ala Gly Glu		
1205	1210	1215
Leu Gln Lys Gly Asn Glu Leu Ala Leu Pro Ser Lys Tyr Val Asn		
1220	1225	1230
Phe Leu Tyr Leu Ala Ser His Tyr Glu Lys Leu Lys Gly Ser Pro		
1235	1240	1245
Glu Asp Asn Glu Gln Lys Gln Leu Phe Val Glu Gln His Lys His		
1250	1255	1260
Tyr Leu Asp Glu Ile Ile Glu Gln Ile Ser Glu Phe Ser Lys Arg		
1265	1270	1275
Val Ile Leu Ala Asp Ala Asn Leu Asp Lys Val Leu Ser Ala Tyr		
1280	1285	1290
Asn Lys His Arg Asp Lys Pro Ile Arg Glu Gln Ala Glu Asn Ile		
1295	1300	1305
Ile His Leu Phe Thr Leu Thr Asn Leu Gly Ala Pro Ala Ala Phe		
1310	1315	1320
Lys Tyr Phe Asp Thr Thr Ile Asp Arg Lys Arg Tyr Thr Ser Thr		
1325	1330	1335
Lys Glu Val Leu Asp Ala Thr Leu Ile His Gln Ser Ile Thr Gly		
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Leu Tyr Glu Thr Arg Ile Asp Leu Ser Gln Leu Gly Gly Asp		
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 35 40 45
 Ser Ile Trp Arg His Thr Ser Gln Asn Thr Asn Lys His Val Glu Val
 50 55 60
 Asn Phe Ile Glu Lys Phe Thr Thr Glu Arg Tyr Phe Cys Pro Asn Thr
 65 70 75 80
 Arg Cys Ser Ile Thr Trp Phe Leu Ser Trp Ser Pro Cys Gly Glu Cys
 85 90 95
 Ser Arg Ala Ile Thr Glu Phe Leu Ser Arg Tyr Pro His Val Thr Leu
 100 105 110
 Phe Ile Tyr Ile Ala Arg Leu Tyr His His Ala Asp Pro Arg Asn Arg
 115 120 125
 Gln Gly Leu Arg Asp Leu Ile Ser Ser Gly Val Thr Ile Gln Ile Met
 130 135 140
 Thr Glu Gln Glu Ser Gly Tyr Cys Trp Arg Asn Phe Val Asn Tyr Ser
 145 150 155 160
 Pro Ser Asn Glu Ala His Trp Pro Arg Tyr Pro His Leu Trp Val Arg
 165 170 175
 Leu Tyr Val Leu Glu Leu Tyr Cys Ile Ile Leu Gly Leu Pro Pro Cys
 180 185 190
 Leu Asn Ile Leu Arg Arg Lys Gln Pro Gln Leu Thr Phe Phe Thr Ile
 195 200 205
 Ala Leu Gln Ser Cys His Tyr Gln Arg Leu Pro Pro His Ile Leu Trp
 210 215 220
 Ala Thr Gly Leu Lys Ser Gly Ser Glu Thr Pro Gly Thr Ser Glu Ser
 225 230 235 240
 Ala Thr Pro Glu Ser Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly
 245 250 255

Thr Asn Ser Val Gly Trp Ala Val Ile Thr Asp Glu Tyr Lys Val Pro
 260 265 270
 Ser Lys Lys Phe Lys Val Leu Gly Asn Thr Asp Arg His Ser Ile Lys
 275 280 285
 Lys Asn Leu Ile Gly Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu
 290 295 300
 Ala Thr Arg Leu Lys Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys
 305 310 315 320
 Asn Arg Ile Cys Tyr Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys
 325 330 335
 Val Asp Asp Ser Phe Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu
 340 345 350
 Glu Asp Lys Lys His Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp
 355 360 365
 Glu Val Ala Tyr His Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys
 370 375 380
 Lys Leu Val Asp Ser Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu
 385 390 395 400
 Ala Leu Ala His Met Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly
 405 410 415
 Asp Leu Asn Pro Asp Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu
 420 425 430
 Val Gln Thr Tyr Asn Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser
 435 440 445
 Gly Val Asp Ala Lys Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg
 450 455 460
 Arg Leu Glu Asn Leu Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly
 465 470 475 480
 Leu Phe Gly Asn Leu Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe
 485 490 495
 Lys Ser Asn Phe Asp Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys
 500 505 510
 Asp Thr Tyr Asp Asp Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp
 515 520 525
 Gln Tyr Ala Asp Leu Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile
 530 535 540
 Leu Leu Ser Asp Ile Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro
 545 550 555 560
 Leu Ser Ala Ser Met Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu

	565		570		575										
Thr	Leu	Leu	Lys	Ala	Leu	Val	Arg	Gln	Gln	Leu	Pro	Glu	Lys	Tyr	Lys
	580		585		590										
Glu	Ile	Phe	Phe	Asp	Gln	Ser	Lys	Asn	Gly	Tyr	Ala	Gly	Tyr	Ile	Asp
	595		600		605										
Gly	Gly	Ala	Ser	Gln	Glu	Glu	Phe	Tyr	Lys	Phe	Ile	Lys	Pro	Ile	Leu
	610		615		620										
Glu	Lys	Met	Asp	Gly	Thr	Glu	Glu	Leu	Leu	Val	Lys	Leu	Asn	Arg	Glu
625			630		635		640								
Asp	Leu	Leu	Arg	Lys	Gln	Arg	Thr	Phe	Asp	Asn	Gly	Ser	Ile	Pro	His
	645		650		655										
Gln	Ile	His	Leu	Gly	Glu	Leu	His	Ala	Ile	Leu	Arg	Arg	Gln	Glu	Asp
	660		665		670										
Phe	Tyr	Pro	Phe	Leu	Lys	Asp	Asn	Arg	Glu	Lys	Ile	Glu	Lys	Ile	Leu
	675		680		685										
Thr	Phe	Arg	Ile	Pro	Tyr	Tyr	Val	Gly	Pro	Leu	Ala	Arg	Gly	Asn	Ser
	690		695		700										
Arg	Phe	Ala	Trp	Met	Thr	Arg	Lys	Ser	Glu	Glu	Thr	Ile	Thr	Pro	Trp
705			710		715		720								
Asn	Phe	Glu	Glu	Val	Val	Asp	Lys	Gly	Ala	Ser	Ala	Gln	Ser	Phe	Ile
	725		730		735										
Glu	Arg	Met	Thr	Ala	Phe	Asp	Lys	Asn	Leu	Pro	Asn	Glu	Lys	Val	Leu
	740		745		750										
Pro	Lys	His	Ser	Leu	Leu	Tyr	Glu	Tyr	Phe	Thr	Val	Tyr	Asn	Glu	Leu
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Thr	Lys	Val	Lys	Tyr	Val	Thr	Glu	Gly	Met	Arg	Lys	Pro	Ala	Phe	Leu
	770		775		780										
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	820		825		830										
Ala	Ser	Leu	Gly	Thr	Tyr	His	Asp	Leu	Leu	Lys	Ile	Ile	Lys	Asp	Lys
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Asp	Phe	Leu	Asp	Asn	Glu	Glu	Asn	Glu	Asp	Ile	Leu	Glu	Asp	Ile	Val
	850		855		860										
Leu	Thr	Leu	Thr	Leu	Phe	Glu	Asp	Arg	Glu	Met	Ile	Glu	Glu	Arg	Leu
865			870		875		880								

Lys Thr Tyr Ala His Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys
 885 890 895
 Arg Arg Arg Tyr Thr Gly Trp Gly Ala Leu Ser Arg Lys Leu Ile Asn
 900 905 910
 Gly Ile Arg Asp Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys
 915 920 925
 Ser Asp Gly Phe Ala Asn Arg Asn Phe Met Ala Leu Ile His Asp Asp
 930 935 940
 Ser Leu Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln
 945 950 955 960
 Gly Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala
 965 970 975
 Ile Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu Val
 980 985 990
 Lys Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu Met Ala
 995 1000 1005
 Arg Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu
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 Arg Met Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly Ser Gln
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 Ile Leu Lys Glu His Pro Val Glu Asn Thr Gln Leu Gln Asn Glu
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 Lys Leu Tyr Leu Tyr Tyr Leu Gln Asn Gly Arg Asp Met Tyr Val
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 Asp Gln Glu Leu Asp Ile Asn Arg Leu Ser Asp Tyr Asp Val Asp
 1070 1075 1080
 His Ile Val Pro Gln Ser Phe Leu Lys Asp Asp Ser Ile Asp Asn
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 Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly Lys Ser Asp Asn
 1100 1105 1110
 Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn Tyr Trp Arg
 1115 1120 1125
 Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe Asp Asn
 1130 1135 1140
 Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys Ala
 1145 1150 1155
 Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Ala Ile Thr Lys
 1160 1165 1170
 His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp

1175	1180	1185
Glu Asn Asp Lys Leu Ile	Arg Glu Val Lys Val	Ile Thr Leu Lys
1190	1195	1200
Ser Lys Leu Val Ser Asp	Phe Arg Lys Asp Phe	Gln Phe Tyr Lys
1205	1210	1215
Val Arg Glu Ile Asn Asn	Tyr His His Ala His	Asp Ala Tyr Leu
1220	1225	1230
Asn Ala Val Val Gly Thr	Ala Leu Ile Lys Lys	Tyr Pro Lys Leu
1235	1240	1245
Glu Ser Glu Phe Val Tyr	Gly Asp Tyr Lys Val	Tyr Asp Val Arg
1250	1255	1260
Lys Met Ile Ala Lys Ser	Glu Gln Glu Ile Gly	Lys Ala Thr Ala
1265	1270	1275
Lys Tyr Phe Phe Tyr Ser	Asn Ile Met Asn Phe	Phe Lys Thr Glu
1280	1285	1290
Ile Thr Leu Ala Asn Gly	Glu Ile Arg Lys Arg	Pro Leu Ile Glu
1295	1300	1305
Thr Asn Gly Glu Thr Gly	Glu Ile Val Trp Asp	Lys Gly Arg Asp
1310	1315	1320
Phe Ala Thr Val Arg Lys	Val Leu Ser Met Pro	Gln Val Asn Ile
1325	1330	1335
Val Lys Lys Thr Glu Val	Gln Thr Gly Gly Phe	Ser Lys Glu Ser
1340	1345	1350
Ile Leu Pro Lys Arg Asn	Ser Asp Lys Leu Ile	Ala Arg Lys Lys
1355	1360	1365
Asp Trp Asp Pro Lys Lys	Tyr Gly Gly Phe Asp	Ser Pro Thr Val
1370	1375	1380
Ala Tyr Ser Val Leu Val	Val Ala Lys Val Glu	Lys Gly Lys Ser
1385	1390	1395
Lys Lys Leu Lys Ser Val	Lys Glu Leu Leu Gly	Ile Thr Ile Met
1400	1405	1410
Glu Arg Ser Ser Phe Glu	Lys Asn Pro Ile Asp	Phe Leu Glu Ala
1415	1420	1425
Lys Gly Tyr Lys Glu Val	Lys Lys Asp Leu Ile	Ile Lys Leu Pro
1430	1435	1440
Lys Tyr Ser Leu Phe Glu	Leu Glu Asn Gly Arg	Lys Arg Met Leu
1445	1450	1455
Ala Ser Ala Gly Glu Leu	Gln Lys Gly Asn Glu	Leu Ala Leu Pro
1460	1465	1470

Ser Lys Tyr Val Asn Phe Leu Tyr Leu Ala Ser His Tyr Glu Lys		
1475	1480	1485
Leu Lys Gly Ser Pro Glu Asp Asn Glu Gln Lys Gln Leu Phe Val		
1490	1495	1500
Glu Gln His Lys His Tyr Leu Asp Glu Ile Ile Glu Gln Ile Ser		
1505	1510	1515
Glu Phe Ser Lys Arg Val Ile Leu Ala Asp Ala Asn Leu Asp Lys		
1520	1525	1530
Val Leu Ser Ala Tyr Asn Lys His Arg Asp Lys Pro Ile Arg Glu		
1535	1540	1545
Gln Ala Glu Asn Ile Ile His Leu Phe Thr Leu Thr Asn Leu Gly		
1550	1555	1560
Ala Pro Ala Ala Phe Lys Tyr Phe Asp Thr Thr Ile Asp Arg Lys		
1565	1570	1575
Arg Tyr Thr Ser Thr Lys Glu Val Leu Asp Ala Thr Leu Ile His		
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Gln Ser Ile Thr Gly Leu Tyr Glu Thr Arg Ile Asp Leu Ser Gln		
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Leu Gly Gly Asp		
1610		
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Glu Ala Ala Ala Lys		
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 Val Glu Lys Ala Leu Ser Leu Gly Val Thr Lys Leu Val Glu Arg Trp
 210 215 220
 Ile Ser Val Ser Gly Val Ala Asp Asp Pro Asn Asn Tyr Leu Phe Cys
 225 230 235 240
 Arg Val Arg Lys Asn Gly Val Ala Ala Pro Ser Ala Thr Ser Gln Leu
 245 250 255
 Ser Thr Arg Ala Leu Glu Gly Ile Phe Glu Ala Thr His Arg Leu Ile
 260 265 270
 Tyr Gly Ala Lys Asp Asp Ser Gly Gln Arg Tyr Leu Ala Trp Ser Gly
 275 280 285
 His Ser Ala Arg Val Gly Ala Ala Arg Asp Met Ala Arg Ala Gly Val
 290 295 300
 Ser Ile Pro Glu Ile Met Gln Ala Gly Gly Trp Thr Asn Val Asn Ile
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 Val Met Asn Tyr Ile Arg Asn Leu Asp Ser Glu Thr Gly Ala Met Val
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 Arg Leu Leu Glu Asp Gly Asp
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 35 40 45
 Gly Thr Ala Ile Lys Arg Ala Thr Phe Met Ser Tyr Asn Thr Ile Ile
 50 55 60
 Ser Asn Ser Leu Ser Phe Asp Ile Val Asn Lys Ser Leu Gln Phe Lys
 65 70 75 80
 Tyr Lys Thr Gln Lys Ala Thr Ile Leu Glu Ala Ser Leu Lys Lys Leu
 85 90 95

Ile Pro Ala Trp Glu Phe Thr Ile Ile Pro Tyr Tyr Gly Gln Lys His			
100	105	110	
Gln Ser Asp Ile Thr Asp Ile Val Ser Ser Leu Gln Leu Gln Phe Glu			
115	120	125	
Ser Ser Glu Glu Ala Asp Lys Gly Asn Ser His Ser Lys Lys Met Leu			
130	135	140	
Lys Ala Leu Leu Ser Glu Gly Glu Ser Ile Trp Glu Ile Thr Glu Lys			
145	150	155	160
Ile Leu Asn Ser Phe Glu Tyr Thr Ser Arg Phe Thr Lys Thr Lys Thr			
165	170	175	
Leu Tyr Gln Phe Leu Phe Leu Ala Thr Phe Ile Asn Cys Gly Arg Phe			
180	185	190	
Ser Asp Ile Lys Asn Val Asp Pro Lys Ser Phe Lys Leu Val Gln Asn			
195	200	205	
Lys Tyr Leu Gly Val Ile Ile Gln Cys Leu Val Thr Glu Thr Lys Thr			
210	215	220	
Ser Val Ser Arg His Ile Tyr Phe Phe Ser Ala Arg Gly Arg Ile Asp			
225	230	235	240
Pro Leu Val Tyr Leu Asp Glu Phe Leu Arg Asn Ser Glu Pro Val Leu			
245	250	255	
Lys Arg Val Asn Arg Thr Gly Asn Ser Ser Ser Asn Lys Gln Glu Tyr			
260	265	270	
Gln Leu Leu Lys Asp Asn Leu Val Arg Ser Tyr Asn Lys Ala Leu Lys			
275	280	285	
Lys Asn Ala Pro Tyr Ser Ile Phe Ala Ile Lys Asn Gly Pro Lys Ser			
290	295	300	
His Ile Gly Arg His Leu Met Thr Ser Phe Leu Ser Met Lys Gly Leu			
305	310	315	320
Thr Glu Leu Thr Asn Val Val Gly Asn Trp Ser Asp Lys Arg Ala Ser			
325	330	335	
Ala Val Ala Arg Thr Thr Tyr Thr His Gln Ile Thr Ala Ile Pro Asp			
340	345	350	
His Tyr Phe Ala Leu Val Ser Arg Tyr Tyr Ala Tyr Asp Pro Ile Ser			
355	360	365	
Lys Glu Met Ile Ala Leu Lys Asp Glu Thr Asn Pro Ile Glu Glu Trp			
370	375	380	
Gln His Ile Glu Gln Leu Lys Gly Ser Ala Glu Gly Ser Ile Arg Tyr			
385	390	395	400
Pro Ala Trp Asn Gly Ile Ile Ser Gln Glu Val Leu Asp Tyr Leu Ser			

	405	410	415
Ser Tyr Ile Asn Arg Arg Ile			
	420		
<210> 727			
<211> 144			
<212> PRT			
<213> 人工序列			
<220>			
<223> 合成多肽			
<400> 727			
Met Arg Leu Phe Gly Tyr Ala Arg Val Ser Thr Ser Gln Gln Ser Leu			
1	5	10	15
Asp Ile Gln Val Arg Ala Leu Lys Asp Ala Gly Val Lys Ala Asn Arg			
	20	25	30
Ile Phe Thr Asp Lys Ala Ser Gly Ser Ser Ser Asp Arg Lys Gly Leu			
	35	40	45
Asp Leu Leu Arg Met Lys Val Glu Glu Gly Asp Val Ile Leu Val Lys			
	50	55	60
Lys Leu Asp Arg Leu Gly Arg Asp Thr Ala Asp Met Ile Gln Leu Ile			
65	70	75	80
Lys Glu Phe Asp Ala Gln Gly Val Ser Ile Arg Phe Ile Asp Asp Gly			
	85	90	95
Ile Ser Thr Asp Gly Glu Met Gly Lys Met Val Val Thr Ile Leu Ser			
	100	105	110
Ala Val Ala Gln Ala Glu Arg Gln Arg Ile Leu Glu Arg Thr Asn Glu			
	115	120	125
Gly Arg Gln Glu Ala Met Ala Lys Gly Val Val Phe Gly Arg Lys Arg			
	130	135	140
<210> 728			
<211> 144			
<212> PRT			
<213> 人工序列			
<220>			
<223> 合成多肽			
<400> 728			
Met Arg Leu Phe Gly Tyr Ala Arg Val Ser Thr Ser Gln Gln Ser Leu			
1	5	10	15
Asp Ile Gln Val Arg Ala Leu Lys Asp Ala Gly Val Lys Ala Asn Arg			
	20	25	30

Ile Phe Thr Asp Lys Ala Ser Gly Ser Ser Ser Asp Arg Lys Gly Leu
 35 40 45
 Asp Leu Leu Arg Met Lys Val Glu Glu Gly Asp Val Ile Leu Val Lys
 50 55 60
 Lys Leu Asp Arg Leu Gly Arg Asp Thr Ala Asp Met Ile Gln Leu Ile
 65 70 75 80
 Lys Glu Phe Asp Ala Gln Gly Val Ser Ile Arg Phe Ile Asp Asp Gly
 85 90 95
 Ile Ser Thr Asp Gly Glu Met Gly Lys Met Val Val Thr Ile Leu Ser
 100 105 110
 Ala Val Ala Gln Ala Glu Arg Gln Arg Ile Leu Glu Arg Thr Asn Glu
 115 120 125
 Gly Arg Gln Glu Ala Met Ala Lys Gly Val Val Phe Gly Arg Lys Arg
 130 135 140

<210> 729

<211> 144

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 729

Met Arg Leu Phe Gly Tyr Ala Arg Val Ser Thr Ser Gln Gln Ser Leu
 1 5 10 15
 Asp Ile Gln Val Arg Ala Leu Lys Asp Ala Gly Val Lys Ala Asn Arg
 20 25 30
 Ile Phe Thr Asp Lys Ala Ser Gly Ser Ser Ser Asp Arg Lys Gly Leu
 35 40 45
 Asp Leu Leu Arg Met Lys Val Glu Glu Gly Asp Val Ile Leu Val Lys
 50 55 60
 Lys Leu Asp Arg Leu Gly Arg Asp Thr Ala Asp Met Ile Gln Leu Ile
 65 70 75 80
 Lys Glu Phe Asp Ala Gln Gly Val Ser Ile Arg Phe Ile Asp Asp Gly
 85 90 95
 Ile Ser Thr Asp Gly Tyr Met Gly Lys Met Val Val Thr Ile Leu Ser
 100 105 110
 Ala Val Ala Gln Ala Glu Arg Gln Arg Ile Leu Gln Arg Thr Asn Glu
 115 120 125
 Gly Arg Gln Glu Ala Met Ala Lys Gly Val Val Phe Gly Arg Lys Arg
 130 135 140

<210> 730

<211> 147

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 730

Met Ala Lys Ile Gly Tyr Ala Arg Val Ser Ser Lys Glu Gln Asn Leu
 1 5 10 15
 Asp Arg Gln Leu Gln Ala Leu Gln Gly Val Ser Lys Val Phe Ser Asp
 20 25 30
 Lys Leu Ser Gly Gln Ser Val Glu Arg Pro Gln Leu Gln Ala Met Leu
 35 40 45
 Asn Tyr Ile Arg Glu Gly Asp Ile Val Val Val Thr Glu Leu Asp Arg
 50 55 60
 Leu Gly Arg Asn Asn Lys Glu Leu Thr Glu Leu Met Asn Ala Ile Gln
 65 70 75 80
 Gln Lys Gly Ala Thr Leu Glu Val Leu Asp Leu Pro Ser Met Asn Gly
 85 90 95
 Ile Glu Asp Glu Asn Leu Arg Arg Leu Ile Asn Asn Leu Val Ile Glu
 100 105 110
 Leu Tyr Lys Tyr Gln Ala Glu Ser Glu Arg Lys Arg Ile Lys Glu Arg
 115 120 125
 Gln Ala Gln Gly Ile Glu Ile Ala Lys Ser Lys Gly Lys Phe Lys Gly
 130 135 140
 Arg Gln His
 145

<210> 731

<211> 147

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 731

Met Ala Lys Ile Gly Tyr Ala Arg Val Ser Ser Lys Glu Gln Asn Leu
 1 5 10 15
 Asp Arg Gln Leu Gln Ala Leu Gln Gly Val Ser Lys Val Phe Ser Asp
 20 25 30
 Lys Leu Ser Gly Gln Ser Val Glu Arg Pro Gln Leu Gln Ala Met Leu

35	40	45
Asn Tyr Ile Arg Glu Gly Asp	Ile Val Val Val Thr	Glu Leu Asp Arg
50	55	60
Leu Gly Arg Asn Asn Lys Glu	Leu Thr Glu Leu Met	Asn Ala Ile Gln
65	70	75
Gln Lys Gly Ala Thr Leu Glu	Val Leu Asp Leu Pro	Ser Met Asp Gly
85	90	95
Ile Glu Asp Glu Asn Leu Arg	Arg Leu Ile Asn Asn	Leu Val Ile Glu
100	105	110
Leu Tyr Lys Tyr Gln Ala Glu	Ser Glu Arg Lys Arg	Ile Lys Glu Arg
115	120	125
Gln Ala Gln Gly Ile Glu Ile	Ala Lys Ser Lys Gly	Lys Phe Lys Gly
130	135	140
Arg Gln His		
145		
<210> 732		
<211> 150		
<212> PRT		
<213> 人工序列		
<220>		
<223> 合成多肽		
<400> 732		
Met Ile Ile Gly Tyr Ala Arg	Val Ser Ser Leu Asp	Gln Asn Leu Glu
1	5	10
Arg Gln Leu Glu Asn Leu Lys	Thr Phe Gly Ala Glu	Lys Ile Phe Thr
20	25	30
Glu Lys Gln Ser Gly Lys Ser	Ile Glu Asn Arg Pro	Ile Leu Gln Lys
35	40	45
Ala Leu Asn Phe Val Arg Met	Gly Asp Arg Phe Ile	Val Glu Ser Ile
50	55	60
Asp Arg Leu Gly Arg Asn Tyr	Asn Glu Val Ile His	Thr Val Asn Tyr
65	70	75
Leu Lys Asp Lys Glu Val Gln	Leu Met Ile Thr Ser	Leu Pro Met Met
85	90	95
Asn Glu Val Ile Gly Asn Pro	Leu Leu Asp Lys Phe	Met Lys Asp Leu
100	105	110
Ile Ile Gln Ile Leu Ala Met	Val Ser Glu Gln Glu	Arg Asn Glu Ser
115	120	125
Lys Arg Arg Gln Ala Gln Gly	Ile Gln Val Ala Lys	Glu Lys Gly Val

130	135	140
Tyr Lys Gly Arg Pro Leu		
145	150	
<210> 733		
<211> 150		
<212> PRT		
<213> 人工序列		
<220>		
<223> 合成多肽		
<400> 733		
Met Ile Ile Gly Tyr Ala Arg Val Ser Ser Leu Asp Gln Asn Leu Glu		
1	5	10
Arg Gln Leu Glu Asn Leu Lys Thr Phe Gly Ala Glu Lys Ile Phe Thr		
	20	25
Glu Lys Gln Ser Gly Lys Ser Ile Glu Asn Arg Pro Ile Leu Gln Lys		
	35	40
Ala Leu Asn Phe Val Arg Met Gly Asp Arg Phe Ile Val Glu Ser Ile		
	50	55
Asp Arg Leu Gly Arg Asn Tyr Asn Glu Val Ile His Thr Val Asn Tyr		
65	70	75
Leu Lys Asp Lys Glu Val Arg Leu Met Ile Thr Ser Leu Pro Met Met		
	85	90
Asn Glu Val Ile Gly Asn Pro Leu Leu Asp Lys Phe Met Lys Asp Leu		
	100	105
Ile Ile Arg Ile Leu Ala Met Val Ser Glu Gln Glu Arg Asn Glu Ser		
	115	120
Lys Arg Arg Gln Ala Gln Gly Ile Gln Val Ala Lys Glu Lys Gly Val		
	130	135
Tyr Lys Gly Arg Pro Leu		
145	150	
<210> 734		
<211> 144		
<212> PRT		
<213> 人工序列		
<220>		
<223> 合成多肽		
<400> 734		
Met Arg Leu Phe Gly Tyr Ala Arg Val Ser Thr Ser Gln Gln Ser Leu		
1	5	10
		15

Asp Leu Gln Val Arg Ala Leu Lys Asp Ala Gly Val Lys Ala Asn Arg
 20 25 30
 Ile Phe Thr Asp Lys Ala Ser Gly Ser Ser Thr Asp Arg Glu Gly Leu
 35 40 45
 Asp Leu Leu Arg Met Lys Val Lys Glu Gly Asp Val Ile Leu Val Lys
 50 55 60
 Lys Leu Asp Arg Leu Gly Arg Asp Thr Ala Asp Met Leu Gln Leu Ile
 65 70 75 80
 Lys Glu Phe Asp Ala Gln Gly Val Ala Val Arg Phe Ile Asp Asp Gly
 85 90 95
 Ile Ser Thr Asp Gly Asp Met Gly Gln Met Val Val Thr Ile Leu Ser
 100 105 110
 Ala Val Ala Gln Ala Glu Arg Arg Arg Ile Leu Glu Arg Thr Asn Glu
 115 120 125
 Gly Arg Gln Glu Ala Lys Leu Lys Gly Ile Lys Phe Gly Arg Arg Arg
 130 135 140

<210> 735

<211> 144

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 735

Met Arg Leu Phe Gly Tyr Ala Arg Val Ser Thr Ser Gln Gln Ser Leu
 1 5 10 15
 Asp Leu Gln Val Arg Ala Leu Lys Asp Ala Gly Val Lys Ala Asn Arg
 20 25 30
 Ile Phe Thr Asp Lys Ala Ser Gly Ser Ser Thr Asp Arg Glu Gly Leu
 35 40 45
 Asp Leu Leu Arg Met Lys Val Lys Glu Gly Asp Val Ile Leu Val Lys
 50 55 60
 Lys Leu Asp Arg Leu Ser Arg Asp Thr Ala Asp Met Leu Gln Leu Ile
 65 70 75 80
 Lys Glu Phe Asp Ala Gln Gly Val Ala Val Arg Phe Ile Asp Asp Gly
 85 90 95
 Ile Ser Thr Asp Gly Tyr Met Gly Gln Met Val Val Thr Ile Leu Ser
 100 105 110
 Ala Val Ala Gln Ala Glu Arg Arg Arg Ile Leu Gln Arg Thr Asn Glu
 115 120 125

Gly Arg Gln Glu Ala Lys Leu Lys Gly Ile Lys Phe Gly Arg Arg Arg
 130 135 140

<210> 736

<211> 142

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 736

Met Ala Thr Ile Gly Tyr Ile Arg Val Ser Thr Ile Asp Gln Asn Ile
 1 5 10 15

Asp Leu Gln Arg Asn Ala Leu Thr Ser Ala Asn Cys Asp Arg Ile Phe
 20 25 30

Glu Asp Arg Ile Ser Gly Lys Ile Ala Asn Arg Pro Gly Leu Lys Arg
 35 40 45

Ala Leu Lys Tyr Val Asn Lys Gly Asp Thr Leu Val Val Trp Lys Leu
 50 55 60

Asp Arg Leu Gly Arg Ser Val Lys Asn Leu Val Ala Leu Ile Ser Glu
 65 70 75 80

Leu His Glu Arg Gly Ala His Phe His Ser Leu Thr Asp Ser Ile Asp
 85 90 95

Thr Ser Ser Ala Met Gly Arg Phe Phe Phe His Val Met Ser Ala Leu
 100 105 110

Ala Glu Met Glu Arg Glu Leu Ile Val Glu Arg Thr Leu Ala Gly Leu
 115 120 125

Ala Ala Ala Arg Ala Gln Gly Arg Leu Gly Gly Arg Pro Val
 130 135 140

<210> 737

<211> 142

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 737

Met Ala Thr Ile Gly Tyr Ile Arg Val Ser Thr Ile Asp Gln Asn Ile
 1 5 10 15

Asp Leu Gln Arg Asn Ala Leu Thr Ser Ala Asn Cys Asp Arg Ile Phe
 20 25 30

Glu Asp Arg Ile Ser Gly Lys Ile Ala Asn Arg Pro Gly Leu Lys Arg

35	40	45
Ala Leu Lys Tyr Val Asn Lys Gly Asp Thr Leu Val Val Trp Lys Leu		
50	55	60
Asp Arg Leu Gly Arg Ser Val Lys Asn Leu Val Ala Leu Ile Ser Glu		
65	70	75
Leu His Glu Arg Gly Ala His Phe His Ser Leu Thr Asp Ser Ile Asp		
85	90	95
Thr Ser Ser Ala Met Gly Arg Phe Phe Phe Tyr Val Met Ser Ala Leu		
100	105	110
Ala Glu Met Glu Arg Glu Leu Ile Val Glu Arg Thr Leu Ala Gly Leu		
115	120	125
Ala Ala Ala Arg Ala Gln Gly Arg Leu Gly Gly Arg Pro Val		
130	135	140
<210> 738		
<211> 608		
<212> PRT		
<213> 人工序列		
<220>		
<223> 合成多肽		
<400> 738		
Met Asp Thr Tyr Ala Gly Ala Tyr Asp Arg Gln Ser Arg Glu Arg Glu		
1	5	10
Asn Ser Ser Ala Ala Ser Pro Ala Thr Gln Arg Ser Ala Asn Glu Asp		
20	25	30
Lys Ala Ala Asp Leu Gln Arg Glu Val Glu Arg Asp Gly Gly Arg Phe		
35	40	45
Arg Phe Val Gly His Phe Ser Glu Ala Pro Gly Thr Ser Ala Phe Gly		
50	55	60
Thr Ala Glu Arg Pro Glu Phe Glu Arg Ile Leu Asn Glu Cys Arg Ala		
65	70	75
Gly Arg Leu Asn Met Ile Ile Val Tyr Asp Val Ser Arg Phe Ser Arg		
85	90	95
Leu Lys Val Met Asp Ala Ile Pro Ile Val Ser Glu Leu Leu Ala Leu		
100	105	110
Gly Val Thr Ile Val Ser Thr Gln Glu Gly Val Phe Arg Gln Gly Asn		
115	120	125
Val Met Asp Leu Ile His Leu Ile Met Arg Leu Asp Ala Ser His Lys		
130	135	140
Glu Ser Ser Leu Lys Ser Ala Lys Ile Leu Asp Thr Lys Asn Leu Gln		

145	150	155	160
Arg Glu Leu Gly Gly Tyr Val Gly Gly Lys Ala Pro Tyr Gly Phe Glu			
	165	170	175
Leu Val Ser Glu Thr Lys Glu Ile Thr Arg Asn Gly Arg Met Val Asn			
	180	185	190
Val Val Ile Asn Lys Leu Ala His Ser Thr Thr Pro Leu Thr Gly Pro			
	195	200	205
Phe Glu Phe Glu Pro Asp Val Ile Arg Trp Trp Trp Arg Glu Ile Lys			
	210	215	220
Thr His Lys His Leu Pro Phe Lys Pro Gly Ser Gln Ala Ala Ile His			
225	230	235	240
Pro Gly Ser Ile Thr Gly Leu Cys Lys Arg Met Asp Ala Asp Ala Val			
	245	250	255
Pro Thr Arg Gly Glu Thr Ile Gly Lys Lys Thr Ala Ser Ser Ala Trp			
	260	265	270
Asp Pro Ala Thr Val Met Arg Ile Leu Arg Asp Pro Arg Ile Ala Gly			
	275	280	285
Phe Ala Ala Glu Val Ile Tyr Lys Lys Lys Pro Asp Gly Thr Pro Thr			
	290	295	300
Thr Lys Ile Glu Gly Tyr Arg Ile Gln Arg Asp Pro Ile Thr Leu Arg			
305	310	315	320
Pro Val Glu Leu Asp Cys Gly Pro Ile Ile Glu Pro Ala Glu Trp Tyr			
	325	330	335
Glu Leu Gln Ala Trp Leu Asp Gly Arg Gly Arg Gly Lys Gly Leu Ser			
	340	345	350
Arg Gly Gln Ala Ile Leu Ser Ala Met Asp Lys Leu Tyr Cys Glu Cys			
	355	360	365
Gly Ala Val Met Thr Ser Lys Arg Gly Glu Glu Ser Ile Lys Asp Ser			
	370	375	380
Tyr Arg Cys Arg Arg Arg Lys Val Val Asp Pro Ser Ala Pro Gly Gln			
385	390	395	400
His Glu Gly Thr Cys Asn Val Ser Met Ala Ala Leu Asp Lys Phe Val			
	405	410	415
Ala Glu Arg Ile Phe Asn Lys Ile Arg His Ala Glu Gly Asp Glu Glu			
	420	425	430
Thr Leu Ala Leu Leu Trp Glu Ala Ala Arg Arg Phe Gly Lys Leu Thr			
	435	440	445
Glu Ala Pro Glu Lys Ser Gly Glu Arg Ala Asn Leu Val Ala Glu Arg			
450	455	460	

<223> 合成多肽
<400> 741
Asn Gly Ala Asn
1
<210> 742
<211> 4
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<400> 742
Asn Gly Asn Gly
1
<210> 743
<211> 4
<212> PRT
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<400> 743
Asn Gly Ala Gly
1
<210> 744
<211> 4
<212> PRT
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<400> 744
Asn Gly Cys Gly
1
<210> 745
<211> 6
<212> PRT
<213> 人工序列
<220>
<223> 合成多肽
<400> 745
Asn Asn Gly Arg Arg Thr

1 5
<210> 746
<211> 5
<212> PRT
<213> 人工序列
<220>
<223> 合成多肽
<400> 746
Asn Gly Arg Arg Asn
1 5
<210> 747
<211> 6
<212> PRT
<213> 人工序列
<220>
<223> 合成多肽
<400> 747
Asn Asn Asn Arg Arg Thr
1 5
<210> 748
<211> 7
<212> PRT
<213> 人工序列
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<400> 748
Asn Asn Asn Gly Ala Thr Thr
1 5
<210> 749
<211> 7
<212> PRT
<213> 人工序列
<220>
<223> 合成多肽
<400> 749
Asn Asn Ala Gly Ala Ala Trp
1 5
<210> 750
<211> 5

<212> PRT
 <213> 人工序列
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 <223> 合成多肽
 <400> 750
 Asn Ala Ala Ala Cys
 1 5
 <210> 751
 <211> 4
 <212> PRT
 <213> 人工序列
 <220>
 <223> 合成多肽
 <400> 751
 Thr Thr Thr Asn
 1
 <210> 752
 <211> 1367
 <212> PRT
 <213> 人工序列
 <220>
 <223> 合成多肽
 <400> 752
 Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly Thr Asn Ser Val Gly
 1 5 10 15
 Trp Ala Val Ile Thr Asp Glu Tyr Lys Val Pro Ser Lys Lys Phe Lys
 20 25 30
 Val Leu Gly Asn Thr Asp Arg His Ser Ile Lys Lys Asn Leu Ile Gly
 35 40 45
 Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala Thr Arg Leu Lys
 50 55 60
 Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys Tyr
 65 70 75 80
 Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser Phe
 85 90 95
 Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys His
 100 105 110
 Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr His
 115 120 125

Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp Ser
 130 135 140
 Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His Met
 145 150 155 160
 Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro Asp
 165 170 175
 Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val Gln Thr Tyr Asn
 180 185 190
 Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala Lys
 195 200 205
 Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg Leu Glu Asn Leu
 210 215 220
 Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu Phe Gly Asn Leu
 225 230 235 240
 Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe Asp
 245 250 255
 Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp Asp
 260 265 270
 Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp Leu
 275 280 285
 Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp Ile
 290 295 300
 Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser Met
 305 310 315 320
 Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys Ala
 325 330 335
 Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe Asp
 340 345 350
 Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser Gln
 355 360 365
 Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp Gly
 370 375 380
 Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg Lys
 385 390 395 400
 Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu Gly
 405 410 415
 Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe Leu
 420 425 430
 Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile Pro

435	440	445
Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp Met		
450	455	460
Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu Val		
465	470	475
Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr Asn		
485	490	495
Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser Leu		
500	505	510
Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys Tyr		
515	520	525
Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln Lys		
530	535	540
Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr Val		
545	550	555
Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp Ser		
565	570	575
Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly Thr		
580	585	590
Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp Asn		
595	600	605
Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr Leu		
610	615	620
Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys Thr Tyr Ala His		
625	630	635
Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr Thr		
645	650	655
Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp Lys		
660	665	670
Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe Ala		
675	680	685
Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe Lys		
690	695	700
Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly Asp Ser Leu His		
705	710	715
Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly Ile		
725	730	735
Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys Val Met Gly Arg		
740	745	750

His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln Thr
 755 760 765
 Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met Lys Arg Ile Glu
 770 775 780
 Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys Glu His Pro Val
 785 790 795 800
 Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu Tyr Tyr Leu Gln
 805 810 815
 Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp Ile Asn Arg Leu
 820 825 830
 Ser Asp Tyr Asp Val Asp Ala Ile Val Pro Gln Ser Phe Leu Lys Asp
 835 840 845
 Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly
 850 855 860
 Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn
 865 870 875 880
 Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe
 885 890 895
 Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys
 900 905 910
 Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys
 915 920 925
 His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu
 930 935 940
 Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys
 945 950 955 960
 Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu
 965 970 975
 Ile Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val
 980 985 990
 Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe Val
 995 1000 1005
 Tyr Gly Asp Tyr Lys Val Tyr Asp Val Arg Lys Met Ile Ala Lys
 1010 1015 1020
 Ser Glu Gln Glu Ile Gly Lys Ala Thr Ala Lys Tyr Phe Phe Tyr
 1025 1030 1035
 Ser Asn Ile Met Asn Phe Phe Lys Thr Glu Ile Thr Leu Ala Asn
 1040 1045 1050
 Gly Glu Ile Arg Lys Arg Pro Leu Ile Glu Thr Asn Gly Glu Thr

1055	1060	1065
Gly Glu Ile Val Trp Asp	Lys Gly Arg Asp Phe	Ala Thr Val Arg
1070	1075	1080
Lys Val Leu Ser Met Pro	Gln Val Asn Ile Val	Lys Lys Thr Glu
1085	1090	1095
Val Gln Thr Gly Gly Phe	Ser Lys Glu Ser Ile	Leu Pro Lys Arg
1100	1105	1110
Asn Ser Asp Lys Leu Ile	Ala Arg Lys Lys Asp	Trp Asp Pro Lys
1115	1120	1125
Lys Tyr Gly Gly Phe Asp	Ser Pro Thr Val Ala	Tyr Ser Val Leu
1130	1135	1140
Val Val Ala Lys Val Glu	Lys Gly Lys Ser Lys	Lys Leu Lys Ser
1145	1150	1155
Val Lys Glu Leu Leu Gly	Ile Thr Ile Met Glu	Arg Ser Ser Phe
1160	1165	1170
Glu Lys Asn Pro Ile Asp	Phe Leu Glu Ala Lys	Gly Tyr Lys Glu
1175	1180	1185
Val Lys Lys Asp Leu Ile	Ile Lys Leu Pro Lys	Tyr Ser Leu Phe
1190	1195	1200
Glu Leu Glu Asn Gly Arg	Lys Arg Met Leu Ala	Ser Ala Gly Glu
1205	1210	1215
Leu Gln Lys Gly Asn Glu	Leu Ala Leu Pro Ser	Lys Tyr Val Asn
1220	1225	1230
Phe Leu Tyr Leu Ala Ser	His Tyr Glu Lys Leu	Lys Gly Ser Pro
1235	1240	1245
Glu Asp Asn Glu Gln Lys	Gln Leu Phe Val Glu	Gln His Lys His
1250	1255	1260
Tyr Leu Asp Glu Ile Ile	Glu Gln Ile Ser Glu	Phe Ser Lys Arg
1265	1270	1275
Val Ile Leu Ala Asp Ala	Asn Leu Asp Lys Val	Leu Ser Ala Tyr
1280	1285	1290
Asn Lys His Arg Asp Lys	Pro Ile Arg Glu Gln	Ala Glu Asn Ile
1295	1300	1305
Ile His Leu Phe Thr Leu	Thr Asn Leu Gly Ala	Pro Ala Ala Phe
1310	1315	1320
Lys Tyr Phe Asp Thr Thr	Ile Asp Arg Lys Arg	Tyr Thr Ser Thr
1325	1330	1335
Lys Glu Val Leu Asp Ala	Thr Leu Ile His Gln	Ser Ile Thr Gly
1340	1345	1350

Leu Tyr Glu Thr Arg Ile Asp Leu Ser Gln Leu Gly Gly Asp
 1355 1360 1365
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 <213> 人工序列
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 Asp Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly Thr Asn Ser Val Gly
 1 5 10 15
 Trp Ala Val Ile Thr Asp Glu Tyr Lys Val Pro Ser Lys Lys Phe Lys
 20 25 30
 Val Leu Gly Asn Thr Asp Arg His Ser Ile Lys Lys Asn Leu Ile Gly
 35 40 45
 Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala Thr Arg Leu Lys
 50 55 60
 Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys Tyr
 65 70 75 80
 Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser Phe
 85 90 95
 Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys His
 100 105 110
 Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr His
 115 120 125
 Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp Ser
 130 135 140
 Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His Met
 145 150 155 160
 Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro Asp
 165 170 175
 Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val Gln Thr Tyr Asn
 180 185 190
 Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala Lys
 195 200 205
 Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg Leu Glu Asn Leu
 210 215 220
 Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu Phe Gly Asn Leu
 225 230 235 240

Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe Asp
 245 250 255
 Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp Asp
 260 265 270
 Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp Leu
 275 280 285
 Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp Ile
 290 295 300
 Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser Met
 305 310 315 320
 Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys Ala
 325 330 335
 Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe Asp
 340 345 350
 Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser Gln
 355 360 365
 Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp Gly
 370 375 380
 Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg Lys
 385 390 395 400
 Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu Gly
 405 410 415
 Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe Leu
 420 425 430
 Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile Pro
 435 440 445
 Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp Met
 450 455 460
 Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu Val
 465 470 475 480
 Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr Asn
 485 490 495
 Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser Leu
 500 505 510
 Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys Tyr
 515 520 525
 Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln Lys
 530 535 540
 Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr Val

545	550	555	560
Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp Ser			
	565	570	575
Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly Thr			
	580	585	590
Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp Asn			
	595	600	605
Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr Leu			
	610	615	620
Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys Thr Tyr Ala His			
625	630	635	640
Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr Thr			
	645	650	655
Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp Lys			
	660	665	670
Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe Ala			
	675	680	685
Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe Lys			
	690	695	700
Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly Asp Ser Leu His			
705	710	715	720
Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly Ile			
	725	730	735
Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys Val Met Gly Arg			
	740	745	750
His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln Thr			
	755	760	765
Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met Lys Arg Ile Glu			
	770	775	780
Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys Glu His Pro Val			
785	790	795	800
Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu Tyr Tyr Leu Gln			
	805	810	815
Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp Ile Asn Arg Leu			
	820	825	830
Ser Asp Tyr Asp Val Asp His Ile Val Pro Gln Ser Phe Leu Lys Asp			
	835	840	845
Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly			
	850	855	860

Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn
 865 870 875 880
 Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe
 885 890 895
 Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys
 900 905 910
 Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys
 915 920 925
 His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu
 930 935 940
 Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys
 945 950 955 960
 Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu
 965 970 975
 Ile Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val
 980 985 990
 Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe Val
 995 1000 1005
 Tyr Gly Asp Tyr Lys Val Tyr Asp Val Arg Lys Met Ile Ala Lys
 1010 1015 1020
 Ser Glu Gln Glu Ile Gly Lys Ala Thr Ala Lys Tyr Phe Phe Tyr
 1025 1030 1035
 Ser Asn Ile Met Asn Phe Phe Lys Thr Glu Ile Thr Leu Ala Asn
 1040 1045 1050
 Gly Glu Ile Arg Lys Arg Pro Leu Ile Glu Thr Asn Gly Glu Thr
 1055 1060 1065
 Gly Glu Ile Val Trp Asp Lys Gly Arg Asp Phe Ala Thr Val Arg
 1070 1075 1080
 Lys Val Leu Ser Met Pro Gln Val Asn Ile Val Lys Lys Thr Glu
 1085 1090 1095
 Val Gln Thr Gly Gly Phe Ser Lys Glu Ser Ile Leu Pro Lys Arg
 1100 1105 1110
 Asn Ser Asp Lys Leu Ile Ala Arg Lys Lys Asp Trp Asp Pro Lys
 1115 1120 1125
 Lys Tyr Gly Gly Phe Asp Ser Pro Thr Val Ala Tyr Ser Val Leu
 1130 1135 1140
 Val Val Ala Lys Val Glu Lys Gly Lys Ser Lys Lys Leu Lys Ser
 1145 1150 1155
 Val Lys Glu Leu Leu Gly Ile Thr Ile Met Glu Arg Ser Ser Phe

1160	1165	1170
Glu Lys Asn Pro Ile Asp	Phe Leu Glu Ala Lys	Gly Tyr Lys Glu
1175	1180	1185
Val Lys Lys Asp Leu Ile	Ile Lys Leu Pro Lys	Tyr Ser Leu Phe
1190	1195	1200
Glu Leu Glu Asn Gly Arg	Lys Arg Met Leu Ala	Ser Ala Gly Glu
1205	1210	1215
Leu Gln Lys Gly Asn Glu	Leu Ala Leu Pro Ser	Lys Tyr Val Asn
1220	1225	1230
Phe Leu Tyr Leu Ala Ser	His Tyr Glu Lys Leu	Lys Gly Ser Pro
1235	1240	1245
Glu Asp Asn Glu Gln Lys	Gln Leu Phe Val Glu	Gln His Lys His
1250	1255	1260
Tyr Leu Asp Glu Ile Ile	Glu Gln Ile Ser Glu	Phe Ser Lys Arg
1265	1270	1275
Val Ile Leu Ala Asp Ala	Asn Leu Asp Lys Val	Leu Ser Ala Tyr
1280	1285	1290
Asn Lys His Arg Asp Lys	Pro Ile Arg Glu Gln	Ala Glu Asn Ile
1295	1300	1305
Ile His Leu Phe Thr Leu	Thr Asn Leu Gly Ala	Pro Ala Ala Phe
1310	1315	1320
Lys Tyr Phe Asp Thr Thr	Ile Asp Arg Lys Arg	Tyr Thr Ser Thr
1325	1330	1335
Lys Glu Val Leu Asp Ala	Thr Leu Ile His Gln	Ser Ile Thr Gly
1340	1345	1350
Leu Tyr Glu Thr Arg Ile	Asp Leu Ser Gln Leu	Gly Gly Asp
1355	1360	1365
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Asp Lys Lys Tyr Ser Ile	Gly Leu Asp Ile Gly	Thr Asn Ser Val Gly
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Trp Ala Val Ile Thr Asp	Glu Tyr Lys Val Pro	Ser Lys Lys Phe Lys
	20	25
		30
Val Leu Gly Asn Thr Asp	Arg His Ser Ile Lys	Lys Asn Leu Ile Gly

35	40	45
Ala Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala Thr Arg Leu Lys		
50	55	60
Arg Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys Tyr		
65	70	75
Leu Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser Phe		
85	90	95
Phe His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys His		
100	105	110
Glu Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr His		
115	120	125
Glu Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp Ser		
130	135	140
Thr Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His Met		
145	150	155
Ile Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro Asp		
165	170	175
Asn Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val Gln Thr Tyr Asn		
180	185	190
Gln Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala Lys		
195	200	205
Ala Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg Leu Glu Asn Leu		
210	215	220
Ile Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu Phe Gly Asn Leu		
225	230	235
Ile Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe Asp		
245	250	255
Leu Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp Asp		
260	265	270
Asp Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp Leu		
275	280	285
Phe Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp Ile		
290	295	300
Leu Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser Met		
305	310	315
Ile Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys Ala		
325	330	335
Leu Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe Asp		
340	345	350

Gln Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser Gln
 355 360 365
 Glu Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp Gly
 370 375 380
 Thr Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg Lys
 385 390 395 400
 Gln Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu Gly
 405 410 415
 Glu Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe Leu
 420 425 430
 Lys Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile Pro
 435 440 445
 Tyr Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp Met
 450 455 460
 Thr Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu Val
 465 470 475 480
 Val Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr Asn
 485 490 495
 Phe Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser Leu
 500 505 510
 Leu Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys Tyr
 515 520 525
 Val Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln Lys
 530 535 540
 Lys Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr Val
 545 550 555 560
 Lys Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp Ser
 565 570 575
 Val Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly Thr
 580 585 590
 Tyr His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp Asn
 595 600 605
 Glu Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr Leu
 610 615 620
 Phe Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys Thr Tyr Ala His
 625 630 635 640
 Leu Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr Thr
 645 650 655
 Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp Lys

660	665	670
Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly Phe Ala		
675	680	685
Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu Thr Phe Lys		
690	695	700
Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly Asp Ser Leu His		
705	710	715
Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala Ile Lys Lys Gly Ile		
725	730	735
Leu Gln Thr Val Lys Val Val Asp Glu Leu Val Lys Val Met Gly Arg		
740	745	750
His Lys Pro Glu Asn Ile Val Ile Glu Met Ala Arg Glu Asn Gln Thr		
755	760	765
Thr Gln Lys Gly Gln Lys Asn Ser Arg Glu Arg Met Lys Arg Ile Glu		
770	775	780
Glu Gly Ile Lys Glu Leu Gly Ser Gln Ile Leu Lys Glu His Pro Val		
785	790	795
Glu Asn Thr Gln Leu Gln Asn Glu Lys Leu Tyr Leu Tyr Tyr Leu Gln		
805	810	815
Asn Gly Arg Asp Met Tyr Val Asp Gln Glu Leu Asp Ile Asn Arg Leu		
820	825	830
Ser Asp Tyr Asp Val Asp His Ile Val Pro Gln Ser Phe Leu Lys Asp		
835	840	845
Asp Ser Ile Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly		
850	855	860
Lys Ser Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn		
865	870	875
Tyr Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe		
885	890	895
Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp Lys		
900	905	910
Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Gln Ile Thr Lys		
915	920	925
His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys Tyr Asp Glu		
930	935	940
Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr Leu Lys Ser Lys		
945	950	955
Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe Tyr Lys Val Arg Glu		
965	970	975

Ile Asn Asn Tyr His His Ala His Asp Ala Tyr Leu Asn Ala Val Val		
980	985	990
Gly Thr Ala Leu Ile Lys Lys Tyr Pro Lys Leu Glu Ser Glu Phe Val		
995	1000	1005
Tyr Gly Asp Tyr Lys Val Tyr Asp Val Arg Lys Met Ile Ala Lys		
1010	1015	1020
Ser Glu Gln Glu Ile Gly Lys Ala Thr Ala Lys Tyr Phe Phe Tyr		
1025	1030	1035
Ser Asn Ile Met Asn Phe Phe Lys Thr Glu Ile Thr Leu Ala Asn		
1040	1045	1050
Gly Glu Ile Arg Lys Arg Pro Leu Ile Glu Thr Asn Gly Glu Thr		
1055	1060	1065
Gly Glu Ile Val Trp Asp Lys Gly Arg Asp Phe Ala Thr Val Arg		
1070	1075	1080
Lys Val Leu Ser Met Pro Gln Val Asn Ile Val Lys Lys Thr Glu		
1085	1090	1095
Val Gln Thr Gly Gly Phe Ser Lys Glu Ser Ile Leu Pro Lys Arg		
1100	1105	1110
Asn Ser Asp Lys Leu Ile Ala Arg Lys Lys Asp Trp Asp Pro Lys		
1115	1120	1125
Lys Tyr Gly Gly Phe Asp Ser Pro Thr Val Ala Tyr Ser Val Leu		
1130	1135	1140
Val Val Ala Lys Val Glu Lys Gly Lys Ser Lys Lys Leu Lys Ser		
1145	1150	1155
Val Lys Glu Leu Leu Gly Ile Thr Ile Met Glu Arg Ser Ser Phe		
1160	1165	1170
Glu Lys Asn Pro Ile Asp Phe Leu Glu Ala Lys Gly Tyr Lys Glu		
1175	1180	1185
Val Lys Lys Asp Leu Ile Ile Lys Leu Pro Lys Tyr Ser Leu Phe		
1190	1195	1200
Glu Leu Glu Asn Gly Arg Lys Arg Met Leu Ala Ser Ala Gly Glu		
1205	1210	1215
Leu Gln Lys Gly Asn Glu Leu Ala Leu Pro Ser Lys Tyr Val Asn		
1220	1225	1230
Phe Leu Tyr Leu Ala Ser His Tyr Glu Lys Leu Lys Gly Ser Pro		
1235	1240	1245
Glu Asp Asn Glu Gln Lys Gln Leu Phe Val Glu Gln His Lys His		
1250	1255	1260
Tyr Leu Asp Glu Ile Ile Glu Gln Ile Ser Glu Phe Ser Lys Arg		

1265	1270	1275
Val Ile Leu Ala Asp Ala	Asn Leu Asp Lys Val	Leu Ser Ala Tyr
1280	1285	1290
Asn Lys His Arg Asp Lys	Pro Ile Arg Glu Gln	Ala Glu Asn Ile
1295	1300	1305
Ile His Leu Phe Thr Leu	Thr Asn Leu Gly Ala	Pro Ala Ala Phe
1310	1315	1320
Lys Tyr Phe Asp Thr Thr	Ile Asp Arg Lys Arg	Tyr Thr Ser Thr
1325	1330	1335
Lys Glu Val Leu Asp Ala	Thr Leu Ile His Gln	Ser Ile Thr Gly
1340	1345	1350
Leu Tyr Glu Thr Arg Ile	Asp Leu Ser Gln Leu	Gly Gly Asp
1355	1360	1365
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<213> 冰岛硫化叶菌		
<400> 755		
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Val Ala Thr Glu Ala Glu	Asn Ser Thr Ile Tyr	Asn Asn Lys Val Glu
	20	25
Ile Asp Asp Glu Glu Leu	Arg Asn Val Leu Asn	Leu Ala Tyr Lys Ile
	35	40
Ala Lys Asn Asn Glu Asp	Ala Ala Ala Glu Arg	Arg Gly Lys Ala Lys
	50	55
Lys Lys Lys Gly Glu Glu	Gly Glu Thr Thr Thr	Ser Asn Ile Ile Leu
65	70	75
Pro Leu Ser Gly Asn Asp	Lys Asn Pro Trp Thr	Glu Thr Leu Lys Cys
	85	90
Tyr Asn Phe Pro Thr Thr	Val Ala Leu Ser Glu	Val Phe Lys Asn Phe
	100	105
Ser Gln Val Lys Glu Cys	Glu Glu Val Ser Ala	Pro Ser Phe Val Lys
	115	120
Pro Glu Phe Tyr Glu Phe	Gly Arg Ser Pro Gly	Met Val Glu Arg Thr
	130	135
Arg Arg Val Lys Leu Glu	Val Glu Pro His Tyr	Leu Ile Ile Ala Ala
145	150	155
Ala Gly Trp Val Leu Thr	Arg Leu Gly Lys Ala	Lys Val Ser Glu Gly

	165	170	175
Asp Tyr Val Gly Val Asn Val Phe Thr Pro Thr Arg Gly Ile Leu Tyr			
	180	185	190
Ser Leu Ile Gln Asn Val Asn Gly Ile Val Pro Gly Ile Lys Pro Glu			
	195	200	205
Thr Ala Phe Gly Leu Trp Ile Ala Arg Lys Val Val Ser Ser Val Thr			
	210	215	220
Asn Pro Asn Val Ser Val Val Arg Ile Tyr Thr Ile Ser Asp Ala Val			
225	230	235	240
Gly Gln Asn Pro Thr Thr Ile Asn Gly Gly Phe Ser Ile Asp Leu Thr			
	245	250	255
Lys Leu Leu Glu Lys Arg Tyr Leu Leu Ser Glu Arg Leu Glu Ala Ile			
	260	265	270
Ala Arg Asn Ala Leu Ser Ile Ser Ser Asn Met Arg Glu Arg Tyr Ile			
	275	280	285
Val Leu Ala Asn Tyr Ile Tyr Glu Tyr Leu Thr Gly Ser Lys Arg Leu			
	290	295	300
Glu Asp Leu Leu Tyr Phe Ala Asn Arg Asp Leu Ile Met Asn Leu Asn			
305	310	315	320
Ser Asp Asp Gly Lys Val Arg Asp Leu Lys Leu Ile Ser Ala Tyr Val			
	325	330	335
Asn Gly Glu Leu Ile Arg Gly Glu Gly			
	340	345	
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Met Glu Val Pro Leu Tyr Asn Ile Phe Gly Asp Asn Tyr Ile Ile Gln			
1	5	10	15
Val Ala Thr Glu Ala Glu Asn Ser Thr Ile Tyr Asn Asn Lys Val Glu			
	20	25	30
Ile Asp Asp Glu Glu Leu Arg Asn Val Leu Asn Leu Ala Tyr Lys Ile			
	35	40	45
Ala Lys Asn Asn Glu Asp Ala Ala Ala Glu Arg Arg Gly Lys Ala Lys			
	50	55	60
Lys Lys Lys Gly Glu Glu Gly Glu Thr Thr Thr Ser Asn Ile Ile Leu			
65	70	75	80
Pro Leu Ser Gly Asn Asp Lys Asn Pro Trp Thr Glu Thr Leu Lys Cys			

	85	90	95
Tyr Asn Phe Pro Thr Thr Val Ala Leu Ser Glu Val Phe Lys Asn Phe			
	100	105	110
Ser Gln Val Lys Glu Cys Glu Glu Val Ser Ala Pro Ser Phe Val Lys			
	115	120	125
Pro Glu Phe Tyr Lys Phe Gly Arg Ser Pro Gly Met Val Glu Arg Thr			
	130	135	140
Arg Arg Val Lys Leu Glu Val Glu Pro His Tyr Leu Ile Met Ala Ala			
145	150	155	160
Ala Gly Trp Val Leu Thr Arg Leu Gly Lys Ala Lys Val Ser Glu Gly			
	165	170	175
Asp Tyr Val Gly Val Asn Val Phe Thr Pro Thr Arg Gly Ile Leu Tyr			
	180	185	190
Ser Leu Ile Gln Asn Val Asn Gly Ile Val Pro Gly Ile Lys Pro Glu			
	195	200	205
Thr Ala Phe Gly Leu Trp Ile Ala Arg Lys Val Val Ser Ser Val Thr			
	210	215	220
Asn Pro Asn Val Ser Val Val Ser Ile Tyr Thr Ile Ser Asp Ala Val			
225	230	235	240
Gly Gln Asn Pro Thr Thr Ile Asn Gly Gly Phe Ser Ile Asp Leu Thr			
	245	250	255
Lys Leu Leu Glu Lys Arg Asp Leu Leu Ser Glu Arg Leu Glu Ala Ile			
	260	265	270
Ala Arg Asn Ala Leu Ser Ile Ser Ser Asn Met Arg Glu Arg Tyr Ile			
	275	280	285
Val Leu Ala Asn Tyr Ile Tyr Glu Tyr Leu Thr Gly Ser Lys Arg Leu			
	290	295	300
Glu Asp Leu Leu Tyr Phe Ala Asn Arg Asp Leu Ile Met Asn Leu Asn			
305	310	315	320
Ser Asp Asp Gly Lys Val Arg Asp Leu Lys Leu Ile Ser Ala Tyr Val			
	325	330	335
Asn Gly Glu Leu Ile Arg Gly Glu Gly			
	340	345	
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Met Ser Lys Arg His Pro Arg Ile Ser Gly Val Lys Gly Tyr Arg Leu			

1	5	10	15
His Ala Gln Arg Leu Glu Tyr Thr Gly Lys Ser Gly Ala Met Arg Thr			
	20	25	30
Ile Lys Tyr Pro Leu Tyr Ser Ser Pro Ser Gly Gly Arg Thr Val Pro			
	35	40	45
Arg Glu Ile Val Ser Ala Ile Asn Asp Asp Tyr Val Gly Leu Tyr Gly			
	50	55	60
Leu Ser Asn Phe Asp Asp Leu Tyr Asn Ala Glu Lys Arg Asn Glu Glu			
65	70	75	80
Lys Val Tyr Ser Val Leu Asp Phe Trp Tyr Asp Cys Val Gln Tyr Gly			
	85	90	95
Ala Val Phe Ser Tyr Thr Ala Pro Gly Leu Leu Lys Asn Val Ala Glu			
	100	105	110
Val Arg Gly Gly Ser Tyr Glu Leu Thr Lys Thr Leu Lys Gly Ser His			
	115	120	125
Leu Tyr Asp Glu Leu Gln Ile Asp Lys Val Ile Lys Phe Leu Asn Lys			
	130	135	140
Lys Glu Ile Ser Arg Ala Asn Gly Ser Leu Asp Lys Leu Lys Lys Asp			
145	150	155	160
Ile Ile Asp Cys Phe Lys Ala Glu Tyr Arg Glu Arg His Lys Asp Gln			
	165	170	175
Cys Asn Lys Leu Ala Asp Asp Ile Lys Asn Ala Lys Lys Asp Ala Gly			
	180	185	190
Ala Ser Leu Gly Glu Arg Gln Lys Lys Leu Phe Arg Asp Phe Phe Gly			
	195	200	205
Ile Ser Glu Gln Ser Glu Asn Asp Lys Pro Ser Phe Thr Asn Pro Leu			
	210	215	220
Asn Leu Thr Cys Cys Leu Leu Pro Phe Asp Thr Val Asn Asn Asn Arg			
225	230	235	240
Asn Arg Gly Glu Val Leu Phe Asn Lys Leu Lys Glu Tyr Ala Gln Lys			
	245	250	255
Leu Asp Lys Asn Glu Gly Ser Leu Glu Met Trp Glu Tyr Ile Gly Ile			
	260	265	270
Gly Asn Ser Gly Thr Ala Phe Ser Asn Phe Leu Gly Glu Gly Phe Leu			
	275	280	285
Gly Arg Leu Arg Glu Asn Lys Ile Thr Glu Leu Lys Lys Ala Met Met			
	290	295	300
Asp Ile Thr Asp Ala Trp Arg Gly Gln Glu Gln Glu Glu Glu Leu Glu			
305	310	315	320

Lys Arg Leu Arg Ile Leu Ala Ala Leu Thr Ile Lys Leu Arg Glu Pro
 325 330 335
 Lys Phe Asp Asn His Trp Gly Gly Tyr Arg Ser Asp Ile Asn Gly Lys
 340 345 350
 Leu Ser Ser Trp Leu Gln Asn Tyr Ile Asn Gln Thr Val Lys Ile Lys
 355 360 365
 Glu Asp Leu Lys Gly His Lys Lys Asp Leu Lys Lys Ala Lys Glu Met
 370 375 380
 Ile Asn Arg Phe Gly Glu Ser Asp Thr Lys Glu Glu Ala Val Val Ser
 385 390 395 400
 Ser Leu Leu Glu Ser Ile Glu Lys Ile Val Pro Asp Asp Ser Ala Asp
 405 410 415
 Asp Glu Lys Pro Asp Ile Pro Ala Ile Ala Ile Tyr Arg Arg Phe Leu
 420 425 430
 Ser Asp Gly Arg Leu Thr Leu Asn Arg Phe Val Gln Arg Glu Asp Val
 435 440 445
 Gln Glu Ala Leu Ile Lys Glu Arg Leu Glu Ala Glu Lys Lys Lys Lys
 450 455 460
 Pro Lys Lys Arg Lys Lys Lys Ser Asp Ala Glu Asp Glu Lys Glu Thr
 465 470 475 480
 Ile Asp Phe Lys Glu Leu Phe Pro His Leu Ala Lys Pro Leu Lys Leu
 485 490 495
 Val Pro Asn Phe Tyr Gly Asp Ser Lys Arg Glu Leu Tyr Lys Lys Tyr
 500 505 510
 Lys Asn Ala Ala Ile Tyr Thr Asp Ala Leu Trp Lys Ala Val Glu Lys
 515 520 525
 Ile Tyr Lys Ser Ala Phe Ser Ser Ser Leu Lys Asn Ser Phe Phe Asp
 530 535 540
 Thr Asp Phe Asp Lys Asp Phe Phe Ile Lys Arg Leu Gln Lys Ile Phe
 545 550 555 560
 Ser Val Tyr Arg Arg Phe Asn Thr Asp Lys Trp Lys Pro Ile Val Lys
 565 570 575
 Asn Ser Phe Ala Pro Tyr Cys Asp Ile Val Ser Leu Ala Glu Asn Glu
 580 585 590
 Val Leu Tyr Lys Pro Lys Gln Ser Arg Ser Arg Lys Ser Ala Ala Ile
 595 600 605
 Asp Lys Asn Arg Val Arg Leu Pro Ser Thr Glu Asn Ile Ala Lys Ala
 610 615 620
 Gly Ile Ala Leu Ala Arg Glu Leu Ser Val Ala Gly Phe Asp Trp Lys

625	630	635	640
Asp Leu Leu Lys Lys Glu Glu His Glu Glu Tyr Ile Asp Leu Ile Glu			
	645	650	655
Leu His Lys Thr Ala Leu Ala Leu Leu Leu Ala Val Thr Glu Thr Gln			
	660	665	670
Leu Asp Ile Ser Ala Leu Asp Phe Val Glu Asn Gly Thr Val Lys Asp			
	675	680	685
Phe Met Lys Thr Arg Asp Gly Asn Leu Val Leu Glu Gly Arg Phe Leu			
	690	695	700
Glu Met Phe Ser Gln Ser Ile Val Phe Ser Glu Leu Arg Gly Leu Ala			
705	710	715	720
Gly Leu Met Ser Arg Lys Glu Phe Ile Thr Arg Ser Ala Ile Gln Thr			
	725	730	735
Met Asn Gly Lys Gln Ala Glu Leu Leu Tyr Ile Pro His Glu Phe Gln			
	740	745	750
Ser Ala Lys Ile Thr Thr Pro Lys Glu Met Ser Arg Ala Phe Leu Asp			
	755	760	765
Leu Ala Pro Ala Glu Phe Ala Thr Ser Leu Glu Pro Glu Ser Leu Ser			
	770	775	780
Glu Lys Ser Leu Leu Lys Leu Lys Gln Met Arg Tyr Tyr Pro His Tyr			
785	790	795	800
Phe Gly Tyr Glu Leu Thr Arg Thr Gly Gln Gly Ile Asp Gly Gly Val			
	805	810	815
Ala Glu Asn Ala Leu Arg Leu Glu Lys Ser Pro Val Lys Lys Arg Glu			
	820	825	830
Ile Lys Cys Lys Gln Tyr Lys Thr Leu Gly Arg Gly Gln Asn Lys Ile			
	835	840	845
Val Leu Tyr Val Arg Ser Ser Tyr Tyr Gln Thr Gln Phe Leu Glu Trp			
	850	855	860
Phe Leu His Arg Pro Lys Asn Val Gln Thr Asp Val Ala Val Ser Gly			
865	870	875	880
Ser Phe Leu Ile Asp Glu Lys Lys Val Lys Thr Arg Trp Asn Tyr Asp			
	885	890	895
Ala Leu Thr Val Ala Leu Glu Pro Val Ser Gly Ser Glu Arg Val Phe			
	900	905	910
Val Ser Gln Pro Phe Thr Ile Phe Pro Glu Lys Ser Ala Glu Glu Glu			
	915	920	925
Gly Gln Arg Tyr Leu Gly Ile Asp Ile Gly Glu Tyr Gly Ile Ala Tyr			
	930	935	940

Thr Ala Leu Glu Ile Thr Gly Asp Ser Ala Lys Ile Leu Asp Gln Asn			
945	950	955	960
Phe Ile Ser Asp Pro Gln Leu Lys Thr Leu Arg Glu Glu Val Lys Gly			
	965	970	975
Leu Lys Leu Asp Gln Arg Arg Gly Thr Phe Ala Met Pro Ser Thr Lys			
	980	985	990
Ile Ala Arg Ile Arg Glu Ser Leu Val His Ser Leu Arg Asn Arg Ile			
	995	1000	1005
His His Leu Ala Leu Lys His Lys Ala Lys Ile Val Tyr Glu Leu			
	1010	1015	1020
Glu Val Ser Arg Phe Glu Glu Gly Lys Gln Lys Ile Lys Lys Val			
	1025	1030	1035
Tyr Ala Thr Leu Lys Lys Ala Asp Val Tyr Ser Glu Ile Asp Ala			
	1040	1045	1050
Asp Lys Asn Leu Gln Thr Thr Val Trp Gly Lys Leu Ala Val Ala			
	1055	1060	1065
Ser Glu Ile Ser Ala Ser Tyr Thr Ser Gln Phe Cys Gly Ala Cys			
	1070	1075	1080
Lys Lys Leu Trp Arg Ala Glu Met Gln Val Asp Glu Thr Ile Thr			
	1085	1090	1095
Thr Gln Glu Leu Ile Gly Thr Val Arg Val Ile Lys Gly Gly Thr			
	1100	1105	1110
Leu Ile Asp Ala Ile Lys Asp Phe Met Arg Pro Pro Ile Phe Asp			
	1115	1120	1125
Glu Asn Asp Thr Pro Phe Pro Lys Tyr Arg Asp Phe Cys Asp Lys			
	1130	1135	1140
His His Ile Ser Lys Lys Met Arg Gly Asn Ser Cys Leu Phe Ile			
	1145	1150	1155
Cys Pro Phe Cys Arg Ala Asn Ala Asp Ala Asp Ile Gln Ala Ser			
	1160	1165	1170
Gln Thr Ile Ala Leu Leu Arg Tyr Val Lys Glu Glu Lys Lys Val			
	1175	1180	1185
Glu Asp Tyr Phe Glu Arg Phe Arg Lys Leu Lys Asn Ile Lys Val			
	1190	1195	1200
Leu Gly Gln Met Lys Lys Ile			
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<212> PRT			

<213> 人工序列

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<223> 合成多肽

<400> 758

Ser Gly Gly Ser

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<210> 759

<211> 4

<212> PRT

<213> 人工序列

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<223> 合成多肽

<400> 759

Gly Gly Gly Ser

1

<210> 760

<211> 91

<212> DNA

<213> 人工序列

<220>

<223> 合成多核苷酸

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catgttgagg agacttaagt ccaaaacctg g 91

<210> 761

<211> 30

<212> PRT

<213> 人工序列

<220>

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<400> 761

Met Asp Ser Leu Leu Met Asn Arg Arg Lys Phe Leu Tyr Gln Phe Lys

1

5

10

15

Asn Val Arg Trp Ala Lys Gly Arg Arg Glu Thr Tyr Leu Cys

20

25

30

<210> 762

<211> 1129

<212> PRT

<213> 酸土脂环酸芽孢杆菌

<400> 762

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 20 25 30
 Val Arg Tyr Tyr Thr Glu Trp Leu Ser Leu Leu Arg Gln Glu Asn Leu
 35 40 45
 Tyr Arg Arg Ser Pro Asn Gly Asp Gly Glu Gln Glu Cys Asp Lys Thr
 50 55 60
 Ala Glu Glu Cys Lys Ala Glu Leu Leu Glu Arg Leu Arg Ala Arg Gln
 65 70 75 80
 Val Glu Asn Gly His Arg Gly Pro Ala Gly Ser Asp Asp Glu Leu Leu
 85 90 95
 Gln Leu Ala Arg Gln Leu Tyr Glu Leu Leu Val Pro Gln Ala Ile Gly
 100 105 110
 Ala Lys Gly Asp Ala Gln Gln Ile Ala Arg Lys Phe Leu Ser Pro Leu
 115 120 125
 Ala Asp Lys Asp Ala Val Gly Gly Leu Gly Ile Ala Lys Ala Gly Asn
 130 135 140
 Lys Pro Arg Trp Val Arg Met Arg Glu Ala Gly Glu Pro Gly Trp Glu
 145 150 155 160
 Glu Glu Lys Glu Lys Ala Glu Thr Arg Lys Ser Ala Asp Arg Thr Ala
 165 170 175
 Asp Val Leu Arg Ala Leu Ala Asp Phe Gly Leu Lys Pro Leu Met Arg
 180 185 190
 Val Tyr Thr Asp Ser Glu Met Ser Ser Val Glu Trp Lys Pro Leu Arg
 195 200 205
 Lys Gly Gln Ala Val Arg Thr Trp Asp Arg Asp Met Phe Gln Gln Ala
 210 215 220
 Ile Glu Arg Met Met Ser Trp Glu Ser Trp Asn Gln Arg Val Gly Gln
 225 230 235 240
 Glu Tyr Ala Lys Leu Val Glu Gln Lys Asn Arg Phe Glu Gln Lys Asn
 245 250 255
 Phe Val Gly Gln Glu His Leu Val His Leu Val Asn Gln Leu Gln Gln
 260 265 270
 Asp Met Lys Glu Ala Ser Pro Gly Leu Glu Ser Lys Glu Gln Thr Ala
 275 280 285
 His Tyr Val Thr Gly Arg Ala Leu Arg Gly Ser Asp Lys Val Phe Glu
 290 295 300

Lys Trp Gly Lys Leu Ala Pro Asp Ala Pro Phe Asp Leu Tyr Asp Ala
305 310 315 320
Glu Ile Lys Asn Val Gln Arg Arg Asn Thr Arg Arg Phe Gly Ser His
325 330 335
Asp Leu Phe Ala Lys Leu Ala Glu Pro Glu Tyr Gln Ala Leu Trp Arg
340 345 350
Glu Asp Ala Ser Phe Leu Thr Arg Tyr Ala Val Tyr Asn Ser Ile Leu
355 360 365
Arg Lys Leu Asn His Ala Lys Met Phe Ala Thr Phe Thr Leu Pro Asp
370 375 380
Ala Thr Ala His Pro Ile Trp Thr Arg Phe Asp Lys Leu Gly Gly Asn
385 390 395 400
Leu His Gln Tyr Thr Phe Leu Phe Asn Glu Phe Gly Glu Arg Arg His
405 410 415
Ala Ile Arg Phe His Lys Leu Leu Lys Val Glu Asn Gly Val Ala Arg
420 425 430
Glu Val Asp Asp Val Thr Val Pro Ile Ser Met Ser Glu Gln Leu Asp
435 440 445
Asn Leu Leu Pro Arg Asp Pro Asn Glu Pro Ile Ala Leu Tyr Phe Arg
450 455 460
Asp Tyr Gly Ala Glu Gln His Phe Thr Gly Glu Phe Gly Gly Ala Lys
465 470 475 480
Ile Gln Cys Arg Arg Asp Gln Leu Ala His Met His Arg Arg Arg Gly
485 490 495
Ala Arg Asp Val Tyr Leu Asn Val Ser Val Arg Val Gln Ser Gln Ser
500 505 510
Glu Ala Arg Gly Glu Arg Arg Pro Pro Tyr Ala Ala Val Phe Arg Leu
515 520 525
Val Gly Asp Asn His Arg Ala Phe Val His Phe Asp Lys Leu Ser Asp
530 535 540
Tyr Leu Ala Glu His Pro Asp Asp Gly Lys Leu Gly Ser Glu Gly Leu
545 550 555 560
Leu Ser Gly Leu Arg Val Met Ser Val Asp Leu Gly Leu Arg Thr Ser
565 570 575
Ala Ser Ile Ser Val Phe Arg Val Ala Arg Lys Asp Glu Leu Lys Pro
580 585 590
Asn Ser Lys Gly Arg Val Pro Phe Phe Phe Pro Ile Lys Gly Asn Asp
595 600 605
Asn Leu Val Ala Val His Glu Arg Ser Gln Leu Leu Lys Leu Pro Gly

610	615	620
Glu Thr Glu Ser Lys Asp	Leu Arg Ala Ile Arg	Glu Glu Arg Gln Arg
625	630	635
Thr Leu Arg Gln Leu Arg	Thr Gln Leu Ala Tyr	Leu Arg Leu Leu Val
	645	650
Arg Cys Gly Ser Glu Asp	Val Gly Arg Arg	Glu Arg Ser Trp Ala Lys
	660	665
Leu Ile Glu Gln Pro Val	Asp Ala Ala Asn His	Met Thr Pro Asp Trp
	675	680
Arg Glu Ala Phe Glu Asn	Glu Leu Gln Lys Leu	Lys Ser Leu His Gly
690	695	700
Ile Cys Ser Asp Lys Glu	Trp Met Asp Ala Val	Tyr Glu Ser Val Arg
705	710	715
Arg Val Trp Arg His Met	Gly Lys Gln Val Arg	Asp Trp Arg Lys Asp
	725	730
Val Arg Ser Gly Glu Arg	Pro Lys Ile Arg	Gly Tyr Ala Lys Asp Val
	740	745
Val Gly Gly Asn Ser Ile	Glu Gln Ile Glu Tyr	Leu Glu Arg Gln Tyr
	755	760
Lys Phe Leu Lys Ser Trp	Ser Phe Phe Gly Lys	Val Ser Gly Gln Val
770	775	780
Ile Arg Ala Glu Lys Gly	Ser Arg Phe Ala Ile	Thr Leu Arg Glu His
785	790	795
Ile Asp His Ala Lys Glu	Asp Arg Leu Lys Lys	Leu Ala Asp Arg Ile
	805	810
Ile Met Glu Ala Leu Gly	Tyr Val Tyr Ala Leu	Asp Glu Arg Gly Lys
	820	825
Gly Lys Trp Val Ala Lys	Tyr Pro Pro Cys Gln	Leu Ile Leu Leu Glu
	835	840
Glu Leu Ser Glu Tyr Gln	Phe Asn Asn Asp Arg	Pro Pro Ser Glu Asn
	850	855
Asn Gln Leu Met Gln Trp	Ser His Arg Gly Val	Phe Gln Glu Leu Ile
865	870	875
Asn Gln Ala Gln Val His	Asp Leu Leu Val Gly	Thr Met Tyr Ala Ala
	885	890
Phe Ser Ser Arg Phe Asp	Ala Arg Thr Gly Ala	Pro Gly Ile Arg Cys
	900	905
Arg Arg Val Pro Ala Arg	Cys Thr Gln Glu His	Asn Pro Glu Pro Phe
	915	920
		925

Pro Trp Trp Leu Asn Lys Phe Val Val Glu His Thr Leu Asp Ala Cys
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 Pro Leu Arg Ala Asp Asp Leu Ile Pro Thr Gly Glu Gly Glu Ile Phe
 945 950 955 960
 Val Ser Pro Phe Ser Ala Glu Glu Gly Asp Phe His Gln Ile His Ala
 965 970 975
 Asp Leu Asn Ala Ala Gln Asn Leu Gln Gln Arg Leu Trp Ser Asp Phe
 980 985 990
 Asp Ile Ser Gln Ile Arg Leu Arg Cys Asp Trp Gly Glu Val Asp Gly
 995 1000 1005
 Glu Leu Val Leu Ile Pro Arg Leu Thr Gly Lys Arg Thr Ala Asp
 1010 1015 1020
 Ser Tyr Ser Asn Lys Val Phe Tyr Thr Asn Thr Gly Val Thr Tyr
 1025 1030 1035
 Tyr Glu Arg Glu Arg Gly Lys Lys Arg Arg Lys Val Phe Ala Gln
 1040 1045 1050
 Glu Lys Leu Ser Glu Glu Glu Ala Glu Leu Leu Val Glu Ala Asp
 1055 1060 1065
 Glu Ala Arg Glu Lys Ser Val Val Leu Met Arg Asp Pro Ser Gly
 1070 1075 1080
 Ile Ile Asn Arg Gly Asn Trp Thr Arg Gln Lys Glu Phe Trp Ser
 1085 1090 1095
 Met Val Asn Gln Arg Ile Glu Gly Tyr Leu Val Lys Gln Ile Arg
 1100 1105 1110
 Ser Arg Val Pro Leu Gln Asp Ser Ala Cys Glu Asn Thr Gly Asp
 1115 1120 1125

Ile

<210> 763

<211> 5

<212> PRT

<213> 人工序列

<220>

<223> 合成多肽

<400> 763

Asn Gly Gly Asn Gly

1 5

<210> 764

<211> 1389

<212> PRT

<213> *Leptotrichia shahii*

<400> 764

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	20	25	30
Gly Asn Lys Tyr Ile Leu Asn Ile Asn Glu Asn Asn Asn Lys Glu Lys			
	35	40	45
Ile Asp Asn Asn Lys Phe Ile Arg Lys Tyr Ile Asn Tyr Lys Lys Asn			
	50	55	60
Asp Asn Ile Leu Lys Glu Phe Thr Arg Lys Phe His Ala Gly Asn Ile			
65	70	75	80
Leu Phe Lys Leu Lys Gly Lys Glu Gly Ile Ile Arg Ile Glu Asn Asn			
	85	90	95
Asp Asp Phe Leu Glu Thr Glu Glu Val Val Leu Tyr Ile Glu Ala Tyr			
	100	105	110
Gly Lys Ser Glu Lys Leu Lys Ala Leu Gly Ile Thr Lys Lys Lys Ile			
	115	120	125
Ile Asp Glu Ala Ile Arg Gln Gly Ile Thr Lys Asp Asp Lys Lys Ile			
	130	135	140
Glu Ile Lys Arg Gln Glu Asn Glu Glu Glu Ile Glu Ile Asp Ile Arg			
145	150	155	160
Asp Glu Tyr Thr Asn Lys Thr Leu Asn Asp Cys Ser Ile Ile Leu Arg			
	165	170	175
Ile Ile Glu Asn Asp Glu Leu Glu Thr Lys Lys Ser Ile Tyr Glu Ile			
	180	185	190
Phe Lys Asn Ile Asn Met Ser Leu Tyr Lys Ile Ile Glu Lys Ile Ile			
	195	200	205
Glu Asn Glu Thr Glu Lys Val Phe Glu Asn Arg Tyr Tyr Glu Glu His			
	210	215	220
Leu Arg Glu Lys Leu Leu Lys Asp Asp Lys Ile Asp Val Ile Leu Thr			
225	230	235	240
Asn Phe Met Glu Ile Arg Glu Lys Ile Lys Ser Asn Leu Glu Ile Leu			
	245	250	255
Gly Phe Val Lys Phe Tyr Leu Asn Val Gly Gly Asp Lys Lys Lys Ser			
	260	265	270
Lys Asn Lys Lys Met Leu Val Glu Lys Ile Leu Asn Ile Asn Val Asp			
	275	280	285
Leu Thr Val Glu Asp Ile Ala Asp Phe Val Ile Lys Glu Leu Glu Phe			

290	295	300
Trp Asn Ile Thr Lys Arg	Ile Glu Lys Val Lys Lys Val	Asn Asn Glu
305	310	315
Phe Leu Glu Lys Arg Arg	Asn Arg Thr Tyr Ile Lys Ser Tyr	Val Leu
	325	330
Leu Asp Lys His Glu Lys	Phe Lys Ile Glu Arg Glu	Asn Lys Lys Asp
	340	345
Lys Ile Val Lys Phe Phe	Val Glu Asn Ile Lys Asn	Asn Ser Ile Lys
	355	360
Glu Lys Ile Glu Lys Ile	Leu Ala Glu Phe Lys Ile	Asp Glu Leu Ile
	370	380
Lys Lys Leu Glu Lys Glu	Leu Lys Lys Gly Asn Cys	Asp Thr Glu Ile
385	390	395
Phe Gly Ile Phe Lys Lys	His Tyr Lys Val Asn Phe	Asp Ser Lys Lys
	405	410
Phe Ser Lys Lys Ser Asp	Glu Glu Lys Glu Leu Tyr	Lys Ile Ile Tyr
	420	425
Arg Tyr Leu Lys Gly Arg	Ile Glu Lys Ile Leu Val	Asn Glu Gln Lys
	435	440
Val Arg Leu Lys Lys Met	Glu Lys Ile Glu Ile Glu	Lys Ile Leu Asn
	450	455
Glu Ser Ile Leu Ser Glu	Lys Ile Leu Lys Arg Val	Lys Gln Tyr Thr
465	470	475
Leu Glu His Ile Met Tyr	Leu Gly Lys Leu Arg His	Asn Asp Ile Asp
	485	490
Met Thr Thr Val Asn Thr	Asp Asp Phe Ser Arg Leu	His Ala Lys Glu
	500	505
Glu Leu Asp Leu Glu Leu	Ile Thr Phe Phe Ala Ser	Thr Asn Met Glu
	515	520
Leu Asn Lys Ile Phe Ser	Arg Glu Asn Ile Asn Asn	Asp Glu Asn Ile
	530	535
Asp Phe Phe Gly Gly Asp	Arg Glu Lys Asn Tyr Val	Leu Asp Lys Lys
545	550	555
Ile Leu Asn Ser Lys Ile	Lys Ile Ile Arg Asp Leu	Asp Phe Ile Asp
	565	570
Asn Lys Asn Asn Ile Thr	Asn Asn Phe Ile Arg Lys	Phe Thr Lys Ile
	580	585
Gly Thr Asn Glu Arg Asn	Arg Ile Leu His Ala Ile	Ser Lys Glu Arg
	595	600
		605

Asp Leu Gln Gly Thr Gln Asp Asp Tyr Asn Lys Val Ile Asn Ile Ile
 610 615 620
 Gln Asn Leu Lys Ile Ser Asp Glu Glu Val Ser Lys Ala Leu Asn Leu
 625 630 635 640
 Asp Val Val Phe Lys Asp Lys Lys Asn Ile Ile Thr Lys Ile Asn Asp
 645 650 655
 Ile Lys Ile Ser Glu Glu Asn Asn Asn Asp Ile Lys Tyr Leu Pro Ser
 660 665 670
 Phe Ser Lys Val Leu Pro Glu Ile Leu Asn Leu Tyr Arg Asn Asn Pro
 675 680 685
 Lys Asn Glu Pro Phe Asp Thr Ile Glu Thr Glu Lys Ile Val Leu Asn
 690 695 700
 Ala Leu Ile Tyr Val Asn Lys Glu Leu Tyr Lys Lys Leu Ile Leu Glu
 705 710 715 720
 Asp Asp Leu Glu Glu Asn Glu Ser Lys Asn Ile Phe Leu Gln Glu Leu
 725 730 735
 Lys Lys Thr Leu Gly Asn Ile Asp Glu Ile Asp Glu Asn Ile Ile Glu
 740 745 750
 Asn Tyr Tyr Lys Asn Ala Gln Ile Ser Ala Ser Lys Gly Asn Asn Lys
 755 760 765
 Ala Ile Lys Lys Tyr Gln Lys Lys Val Ile Glu Cys Tyr Ile Gly Tyr
 770 775 780
 Leu Arg Lys Asn Tyr Glu Glu Leu Phe Asp Phe Ser Asp Phe Lys Met
 785 790 795 800
 Asn Ile Gln Glu Ile Lys Lys Gln Ile Lys Asp Ile Asn Asp Asn Lys
 805 810 815
 Thr Tyr Glu Arg Ile Thr Val Lys Thr Ser Asp Lys Thr Ile Val Ile
 820 825 830
 Asn Asp Asp Phe Glu Tyr Ile Ile Ser Ile Phe Ala Leu Leu Asn Ser
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 865 870 875 880
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 900 905 910
 Asp Phe Asp Asp Phe Lys Ile Gln Thr Lys Lys Glu Ile Phe Asn Asn

915	920	925
Tyr Tyr Glu Asp Ile Lys	Asn Asn Ile Leu Thr	Glu Phe Lys Asp Asp
930	935	940
Ile Asn Gly Cys Asp Val	Leu Glu Lys Lys Leu	Glu Lys Ile Val Ile
945	950	955
Phe Asp Asp Glu Thr Lys	Phe Glu Ile Asp Lys	Lys Ser Asn Ile Leu
	965	970
Gln Asp Glu Gln Arg Lys	Leu Ser Asn Ile Asn	Lys Lys Asp Leu Lys
	980	990
Lys Lys Val Asp Gln Tyr	Ile Lys Asp Lys Asp	Gln Glu Ile Lys Ser
995	1000	1005
Lys Ile Leu Cys Arg Ile	Ile Phe Asn Ser Asp	Phe Leu Lys Lys
1010	1015	1020
Tyr Lys Lys Glu Ile Asp	Asn Leu Ile Glu Asp	Met Glu Ser Glu
1025	1030	1035
Asn Glu Asn Lys Phe Gln	Glu Ile Tyr Tyr Pro	Lys Glu Arg Lys
1040	1045	1050
Asn Glu Leu Tyr Ile Tyr	Lys Lys Asn Leu Phe	Leu Asn Ile Gly
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Asn Pro Asn Phe Asp Lys	Ile Tyr Gly Leu Ile	Ser Asn Asp Ile
1070	1075	1080
Lys Met Ala Asp Ala Lys	Phe Leu Phe Asn Ile	Asp Gly Lys Asn
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Ile Arg Lys Asn Lys Ile	Ser Glu Ile Asp Ala	Ile Leu Lys Asn
1100	1105	1110
Leu Asn Asp Lys Leu Asn	Gly Tyr Ser Lys Glu	Tyr Lys Glu Lys
1115	1120	1125
Tyr Ile Lys Lys Leu Lys	Glu Asn Asp Asp Phe	Phe Ala Lys Asn
1130	1135	1140
Ile Gln Asn Lys Asn Tyr	Lys Ser Phe Glu Lys	Asp Tyr Asn Arg
1145	1150	1155
Val Ser Glu Tyr Lys Lys	Ile Arg Asp Leu Val	Glu Phe Asn Tyr
1160	1165	1170
Leu Asn Lys Ile Glu Ser	Tyr Leu Ile Asp Ile	Asn Trp Lys Leu
1175	1180	1185
Ala Ile Gln Met Ala Arg	Phe Glu Arg Asp Met	His Tyr Ile Val
1190	1195	1200
Asn Gly Leu Arg Glu Leu	Gly Ile Ile Lys Leu	Ser Gly Tyr Asn
1205	1210	1215

Thr Gly Ile Ser Arg Ala Tyr Pro Lys Arg Asn Gly Ser Asp Gly		
1220	1225	1230
Phe Tyr Thr Thr Thr Ala Tyr Tyr Lys Phe Phe Asp Glu Glu Ser		
1235	1240	1245
Tyr Lys Lys Phe Glu Lys Ile Cys Tyr Gly Phe Gly Ile Asp Leu		
1250	1255	1260
Ser Glu Asn Ser Glu Ile Asn Lys Pro Glu Asn Glu Ser Ile Arg		
1265	1270	1275
Asn Tyr Ile Ser His Phe Tyr Ile Val Arg Asn Pro Phe Ala Asp		
1280	1285	1290
Tyr Ser Ile Ala Glu Gln Ile Asp Arg Val Ser Asn Leu Leu Ser		
1295	1300	1305
Tyr Ser Thr Arg Tyr Asn Asn Ser Thr Tyr Ala Ser Val Phe Glu		
1310	1315	1320
Val Phe Lys Lys Asp Val Asn Leu Asp Tyr Asp Glu Leu Lys Lys		
1325	1330	1335
Lys Phe Lys Leu Ile Gly Asn Asn Asp Ile Leu Glu Arg Leu Met		
1340	1345	1350
Lys Pro Lys Lys Val Ser Val Leu Glu Leu Glu Ser Tyr Asn Ser		
1355	1360	1365
Asp Tyr Ile Lys Asn Leu Ile Ile Glu Leu Leu Thr Lys Ile Glu		
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Asn Thr Asn Asp Thr Leu		
1385		

<210> 765

<211> 5268

<212> DNA

<213> 人工序列

<220>

<223> 合成多核苷酸

<400> 765

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acgtggaaga tgctcctgtc agtgtgtaga agctgggcag cttggtgcaa gttgaacaac 180
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gacttcgatc aggtgctggag tcttatggag aatagtgaca gatgccagga cattcggaac 480

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 20 25 30
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 35 40 45
 Cys Arg Ser Trp Ala Ala Trp Cys Lys Leu Asn Asn Arg Lys Trp Phe
 50 55 60
 Pro Ala Glu Pro Glu Asp Val Arg Asp Tyr Leu Leu Tyr Leu Gln Ala
 65 70 75 80
 Arg Gly Leu Ala Val Lys Thr Ile Gln Gln His Leu Gly Gln Leu Asn
 85 90 95
 Met Leu His Arg Arg Ser Gly Leu Pro Arg Pro Ser Asp Ser Asn Ala
 100 105 110
 Val Ser Leu Val Met Arg Arg Ile Arg Lys Glu Asn Val Asp Ala Gly
 115 120 125
 Glu Arg Ala Lys Gln Ala Leu Ala Phe Glu Arg Thr Asp Phe Asp Gln
 130 135 140
 Val Arg Ser Leu Met Glu Asn Ser Asp Arg Cys Gln Asp Ile Arg Asn
 145 150 155 160
 Leu Ala Phe Leu Gly Ile Ala Tyr Asn Thr Leu Leu Arg Ile Ala Glu
 165 170 175
 Ile Ala Arg Ile Arg Val Lys Asp Ile Ser Arg Thr Asp Gly Gly Arg
 180 185 190
 Met Leu Ile His Ile Gly Arg Thr Lys Thr Leu Val Ser Thr Ala Gly
 195 200 205
 Val Glu Lys Ala Leu Ser Leu Gly Val Thr Lys Leu Val Glu Arg Trp
 210 215 220
 Ile Ser Val Ser Gly Val Ala Asp Asp Pro Asn Asn Tyr Leu Phe Cys
 225 230 235 240

Arg Val Arg Lys Asn Gly Val Ala Ala Pro Ser Ala Thr Ser Gln Leu
 245 250 255
 Ser Thr Arg Ala Leu Glu Gly Ile Phe Glu Ala Thr His Arg Leu Ile
 260 265 270
 Tyr Gly Ala Lys Asp Asp Ser Gly Gln Arg Tyr Leu Ala Trp Ser Gly
 275 280 285
 His Ser Ala Arg Val Gly Ala Ala Arg Asp Met Ala Arg Ala Gly Val
 290 295 300
 Ser Ile Pro Glu Ile Met Gln Ala Gly Gly Trp Thr Asn Val Asn Ile
 305 310 315 320
 Val Met Asn Tyr Ile Arg Asn Leu Asp Ser Glu Thr Gly Ala Met Val
 325 330 335
 Arg Leu Leu Glu Asp Gly Asp Gly Gly Ser Gly Gly Ser Gly Gly Ser
 340 345 350
 Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Asp
 355 360 365
 Lys Lys Tyr Ser Ile Gly Leu Ala Ile Gly Thr Asn Ser Val Gly Trp
 370 375 380
 Ala Val Ile Thr Asp Glu Tyr Lys Val Pro Ser Lys Lys Phe Lys Val
 385 390 395 400
 Leu Gly Asn Thr Asp Arg His Ser Ile Lys Lys Asn Leu Ile Gly Ala
 405 410 415
 Leu Leu Phe Asp Ser Gly Glu Thr Ala Glu Ala Thr Arg Leu Lys Arg
 420 425 430
 Thr Ala Arg Arg Arg Tyr Thr Arg Arg Lys Asn Arg Ile Cys Tyr Leu
 435 440 445
 Gln Glu Ile Phe Ser Asn Glu Met Ala Lys Val Asp Asp Ser Phe Phe
 450 455 460
 His Arg Leu Glu Glu Ser Phe Leu Val Glu Glu Asp Lys Lys His Glu
 465 470 475 480
 Arg His Pro Ile Phe Gly Asn Ile Val Asp Glu Val Ala Tyr His Glu
 485 490 495
 Lys Tyr Pro Thr Ile Tyr His Leu Arg Lys Lys Leu Val Asp Ser Thr
 500 505 510
 Asp Lys Ala Asp Leu Arg Leu Ile Tyr Leu Ala Leu Ala His Met Ile
 515 520 525
 Lys Phe Arg Gly His Phe Leu Ile Glu Gly Asp Leu Asn Pro Asp Asn
 530 535 540
 Ser Asp Val Asp Lys Leu Phe Ile Gln Leu Val Gln Thr Tyr Asn Gln

545	550	555	560
Leu Phe Glu Glu Asn Pro Ile Asn Ala Ser Gly Val Asp Ala Lys Ala			
	565	570	575
Ile Leu Ser Ala Arg Leu Ser Lys Ser Arg Arg Leu Glu Asn Leu Ile			
	580	585	590
Ala Gln Leu Pro Gly Glu Lys Lys Asn Gly Leu Phe Gly Asn Leu Ile			
	595	600	605
Ala Leu Ser Leu Gly Leu Thr Pro Asn Phe Lys Ser Asn Phe Asp Leu			
	610	615	620
Ala Glu Asp Ala Lys Leu Gln Leu Ser Lys Asp Thr Tyr Asp Asp Asp			
625	630	635	640
Leu Asp Asn Leu Leu Ala Gln Ile Gly Asp Gln Tyr Ala Asp Leu Phe			
	645	650	655
Leu Ala Ala Lys Asn Leu Ser Asp Ala Ile Leu Leu Ser Asp Ile Leu			
	660	665	670
Arg Val Asn Thr Glu Ile Thr Lys Ala Pro Leu Ser Ala Ser Met Ile			
	675	680	685
Lys Arg Tyr Asp Glu His His Gln Asp Leu Thr Leu Leu Lys Ala Leu			
	690	695	700
Val Arg Gln Gln Leu Pro Glu Lys Tyr Lys Glu Ile Phe Phe Asp Gln			
705	710	715	720
Ser Lys Asn Gly Tyr Ala Gly Tyr Ile Asp Gly Gly Ala Ser Gln Glu			
	725	730	735
Glu Phe Tyr Lys Phe Ile Lys Pro Ile Leu Glu Lys Met Asp Gly Thr			
	740	745	750
Glu Glu Leu Leu Val Lys Leu Asn Arg Glu Asp Leu Leu Arg Lys Gln			
	755	760	765
Arg Thr Phe Asp Asn Gly Ser Ile Pro His Gln Ile His Leu Gly Glu			
	770	775	780
Leu His Ala Ile Leu Arg Arg Gln Glu Asp Phe Tyr Pro Phe Leu Lys			
785	790	795	800
Asp Asn Arg Glu Lys Ile Glu Lys Ile Leu Thr Phe Arg Ile Pro Tyr			
	805	810	815
Tyr Val Gly Pro Leu Ala Arg Gly Asn Ser Arg Phe Ala Trp Met Thr			
	820	825	830
Arg Lys Ser Glu Glu Thr Ile Thr Pro Trp Asn Phe Glu Glu Val Val			
	835	840	845
Asp Lys Gly Ala Ser Ala Gln Ser Phe Ile Glu Arg Met Thr Asn Phe			
	850	855	860

Asp Lys Asn Leu Pro Asn Glu Lys Val Leu Pro Lys His Ser Leu Leu
 865 870 875 880
 Tyr Glu Tyr Phe Thr Val Tyr Asn Glu Leu Thr Lys Val Lys Tyr Val
 885 890 895
 Thr Glu Gly Met Arg Lys Pro Ala Phe Leu Ser Gly Glu Gln Lys Lys
 900 905 910
 Ala Ile Val Asp Leu Leu Phe Lys Thr Asn Arg Lys Val Thr Val Lys
 915 920 925
 Gln Leu Lys Glu Asp Tyr Phe Lys Lys Ile Glu Cys Phe Asp Ser Val
 930 935 940
 Glu Ile Ser Gly Val Glu Asp Arg Phe Asn Ala Ser Leu Gly Thr Tyr
 945 950 955 960
 His Asp Leu Leu Lys Ile Ile Lys Asp Lys Asp Phe Leu Asp Asn Glu
 965 970 975
 Glu Asn Glu Asp Ile Leu Glu Asp Ile Val Leu Thr Leu Thr Leu Phe
 980 985 990
 Glu Asp Arg Glu Met Ile Glu Glu Arg Leu Lys Thr Tyr Ala His Leu
 995 1000 1005
 Phe Asp Asp Lys Val Met Lys Gln Leu Lys Arg Arg Arg Tyr Thr
 1010 1015 1020
 Gly Trp Gly Arg Leu Ser Arg Lys Leu Ile Asn Gly Ile Arg Asp
 1025 1030 1035
 Lys Gln Ser Gly Lys Thr Ile Leu Asp Phe Leu Lys Ser Asp Gly
 1040 1045 1050
 Phe Ala Asn Arg Asn Phe Met Gln Leu Ile His Asp Asp Ser Leu
 1055 1060 1065
 Thr Phe Lys Glu Asp Ile Gln Lys Ala Gln Val Ser Gly Gln Gly
 1070 1075 1080
 Asp Ser Leu His Glu His Ile Ala Asn Leu Ala Gly Ser Pro Ala
 1085 1090 1095
 Ile Lys Lys Gly Ile Leu Gln Thr Val Lys Val Val Asp Glu Leu
 1100 1105 1110
 Val Lys Val Met Gly Arg His Lys Pro Glu Asn Ile Val Ile Glu
 1115 1120 1125
 Met Ala Arg Glu Asn Gln Thr Thr Gln Lys Gly Gln Lys Asn Ser
 1130 1135 1140
 Arg Glu Arg Met Lys Arg Ile Glu Glu Gly Ile Lys Glu Leu Gly
 1145 1150 1155
 Ser Gln Ile Leu Lys Glu His Pro Val Glu Asn Thr Gln Leu Gln

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Asn Glu Lys Leu Tyr Leu Tyr Tyr Leu Gln Asn Gly Arg Asp Met		
1175	1180	1185
Tyr Val Asp Gln Glu Leu Asp Ile Asn Arg Leu Ser Asp Tyr Asp		
1190	1195	1200
Val Asp Ala Ile Val Pro Gln Ser Phe Leu Lys Asp Asp Ser Ile		
1205	1210	1215
Asp Asn Lys Val Leu Thr Arg Ser Asp Lys Asn Arg Gly Lys Ser		
1220	1225	1230
Asp Asn Val Pro Ser Glu Glu Val Val Lys Lys Met Lys Asn Tyr		
1235	1240	1245
Trp Arg Gln Leu Leu Asn Ala Lys Leu Ile Thr Gln Arg Lys Phe		
1250	1255	1260
Asp Asn Leu Thr Lys Ala Glu Arg Gly Gly Leu Ser Glu Leu Asp		
1265	1270	1275
Lys Ala Gly Phe Ile Lys Arg Gln Leu Val Glu Thr Arg Gln Ile		
1280	1285	1290
Thr Lys His Val Ala Gln Ile Leu Asp Ser Arg Met Asn Thr Lys		
1295	1300	1305
Tyr Asp Glu Asn Asp Lys Leu Ile Arg Glu Val Lys Val Ile Thr		
1310	1315	1320
Leu Lys Ser Lys Leu Val Ser Asp Phe Arg Lys Asp Phe Gln Phe		
1325	1330	1335
Tyr Lys Val Arg Glu Ile Asn Asn Tyr His His Ala His Asp Ala		
1340	1345	1350
Tyr Leu Asn Ala Val Val Gly Thr Ala Leu Ile Lys Lys Tyr Pro		
1355	1360	1365
Lys Leu Glu Ser Glu Phe Val Tyr Gly Asp Tyr Lys Val Tyr Asp		
1370	1375	1380
Val Arg Lys Met Ile Ala Lys Ser Glu Gln Glu Ile Gly Lys Ala		
1385	1390	1395
Thr Ala Lys Tyr Phe Phe Tyr Ser Asn Ile Met Asn Phe Phe Lys		
1400	1405	1410
Thr Glu Ile Thr Leu Ala Asn Gly Glu Ile Arg Lys Arg Pro Leu		
1415	1420	1425
Ile Glu Thr Asn Gly Glu Thr Gly Glu Ile Val Trp Asp Lys Gly		
1430	1435	1440
Arg Asp Phe Ala Thr Val Arg Lys Val Leu Ser Met Pro Gln Val		
1445	1450	1455

Asn Ile Val Lys Lys Thr Glu Val Gln Thr Gly Gly Phe Ser Lys 1460	1465	1470
Glu Ser Ile Leu Pro Lys Arg Asn Ser Asp Lys Leu Ile Ala Arg 1475	1480	1485
Lys Lys Asp Trp Asp Pro Lys Lys Tyr Gly Gly Phe Asp Ser Pro 1490	1495	1500
Thr Val Ala Tyr Ser Val Leu Val Val Ala Lys Val Glu Lys Gly 1505	1510	1515
Lys Ser Lys Lys Leu Lys Ser Val Lys Glu Leu Leu Gly Ile Thr 1520	1525	1530
Ile Met Glu Arg Ser Ser Phe Glu Lys Asn Pro Ile Asp Phe Leu 1535	1540	1545
Glu Ala Lys Gly Tyr Lys Glu Val Lys Lys Asp Leu Ile Ile Lys 1550	1555	1560
Leu Pro Lys Tyr Ser Leu Phe Glu Leu Glu Asn Gly Arg Lys Arg 1565	1570	1575
Met Leu Ala Ser Ala Gly Glu Leu Gln Lys Gly Asn Glu Leu Ala 1580	1585	1590
Leu Pro Ser Lys Tyr Val Asn Phe Leu Tyr Leu Ala Ser His Tyr 1595	1600	1605
Glu Lys Leu Lys Gly Ser Pro Glu Asp Asn Glu Gln Lys Gln Leu 1610	1615	1620
Phe Val Glu Gln His Lys His Tyr Leu Asp Glu Ile Ile Glu Gln 1625	1630	1635
Ile Ser Glu Phe Ser Lys Arg Val Ile Leu Ala Asp Ala Asn Leu 1640	1645	1650
Asp Lys Val Leu Ser Ala Tyr Asn Lys His Arg Asp Lys Pro Ile 1655	1660	1665
Arg Glu Gln Ala Glu Asn Ile Ile His Leu Phe Thr Leu Thr Asn 1670	1675	1680
Leu Gly Ala Pro Ala Ala Phe Lys Tyr Phe Asp Thr Thr Ile Asp 1685	1690	1695
Arg Lys Arg Tyr Thr Ser Thr Lys Glu Val Leu Asp Ala Thr Leu 1700	1705	1710
Ile His Gln Ser Ile Thr Gly Leu Tyr Glu Thr Arg Ile Asp Leu 1715	1720	1725
Ser Gln Leu Gly Gly Asp Gly Gly Ser Asp Tyr Lys Asp Asp Asp 1730	1735	1740
Asp Lys Gly Gly Ser Pro Lys Lys Lys Arg Lys Val		

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Arg Xaa Lys

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<211> 5

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5



图1A

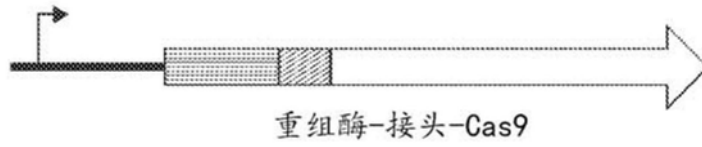


图1B

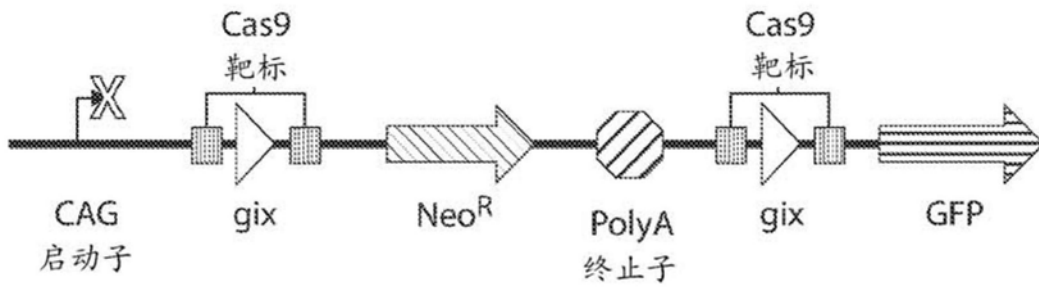


图1C

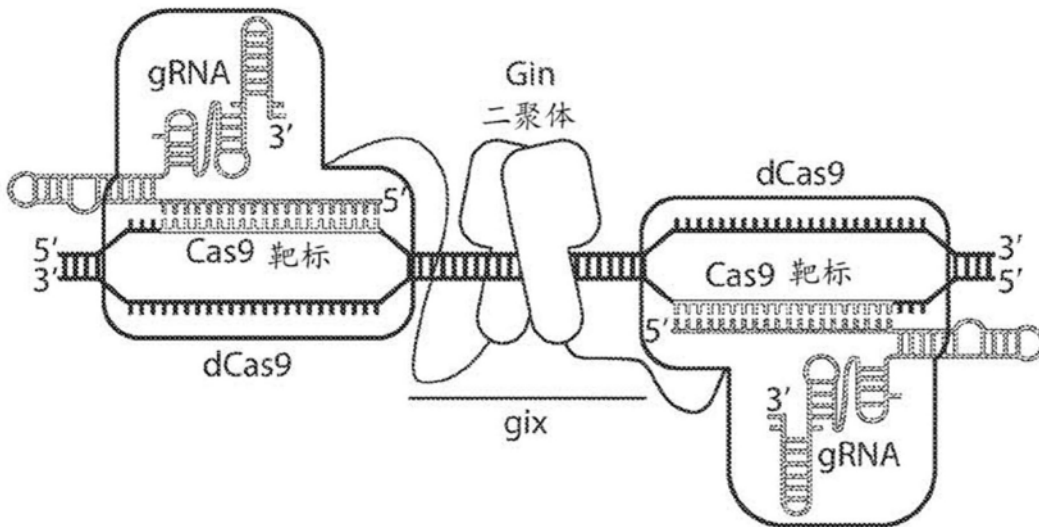


图1D



图2B

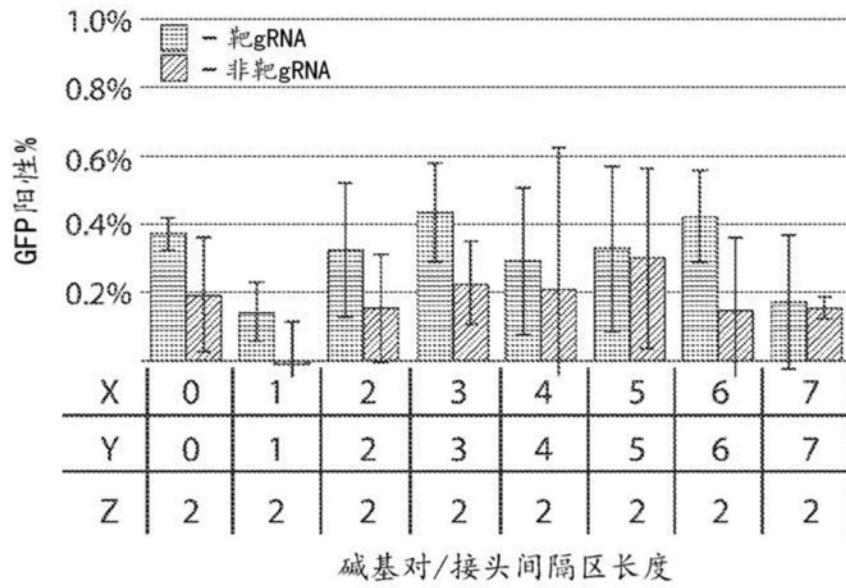


图2C

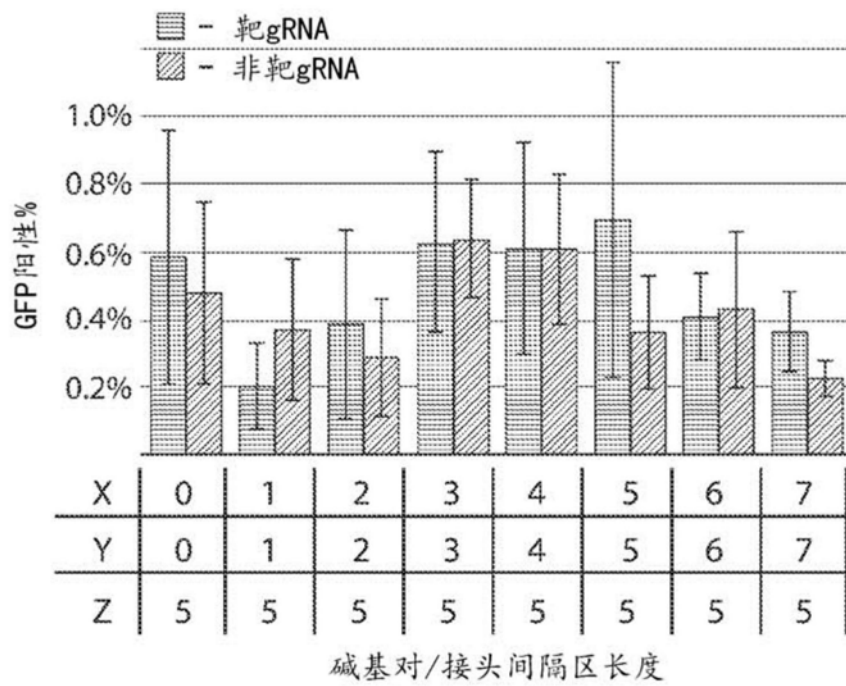


图2D

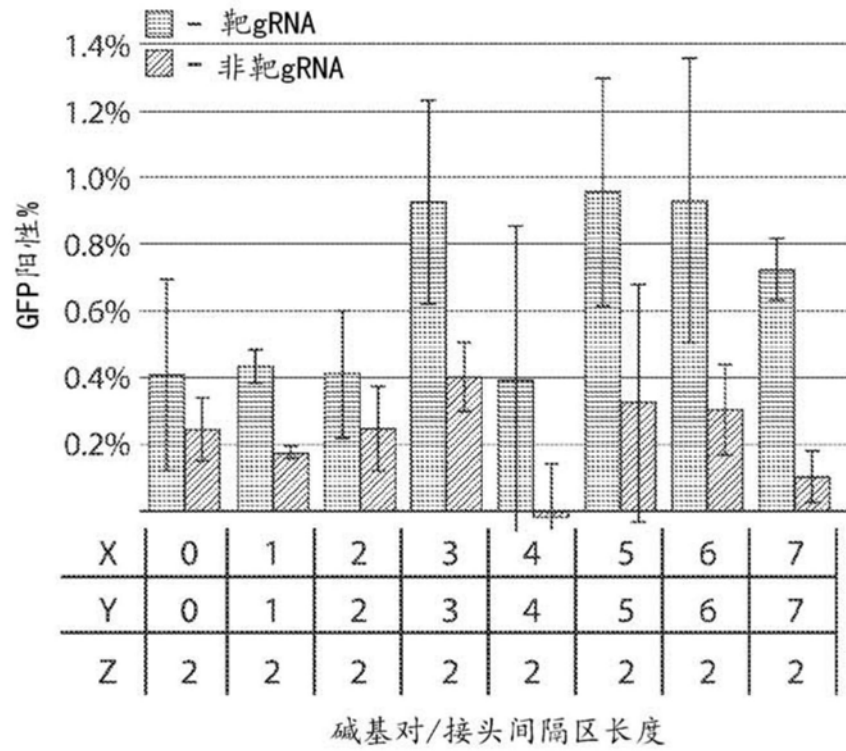


图2E

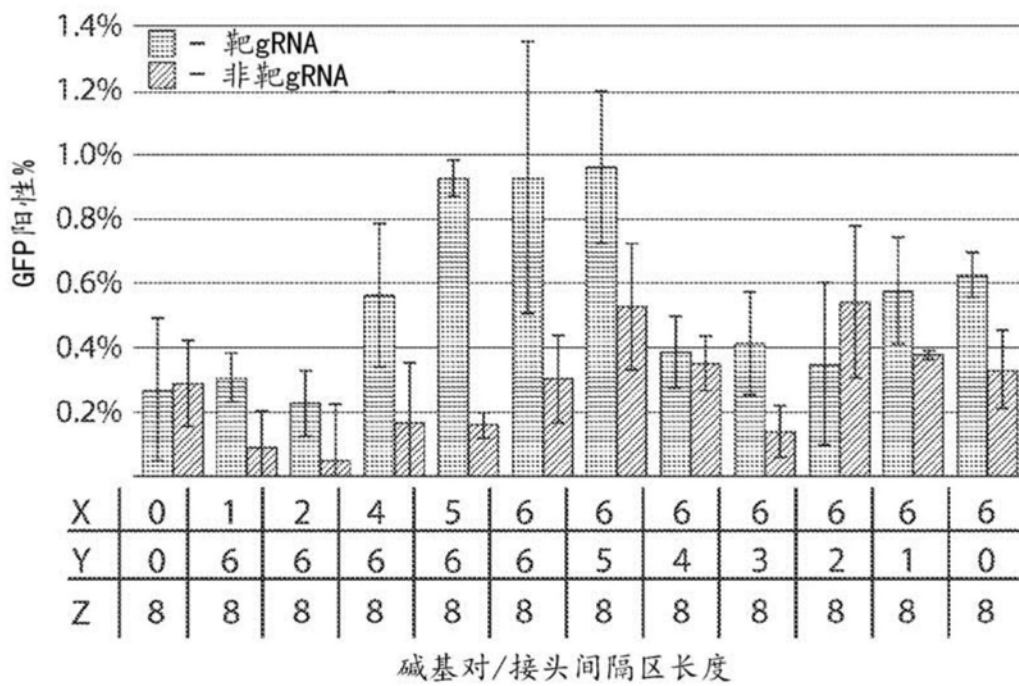


图2F



图3A

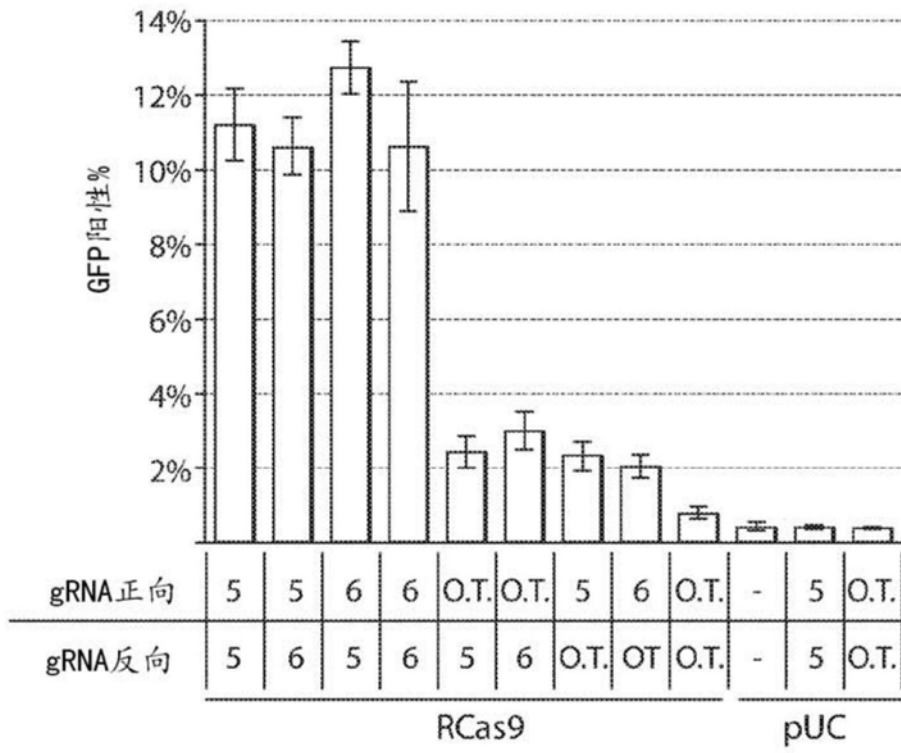


图3B

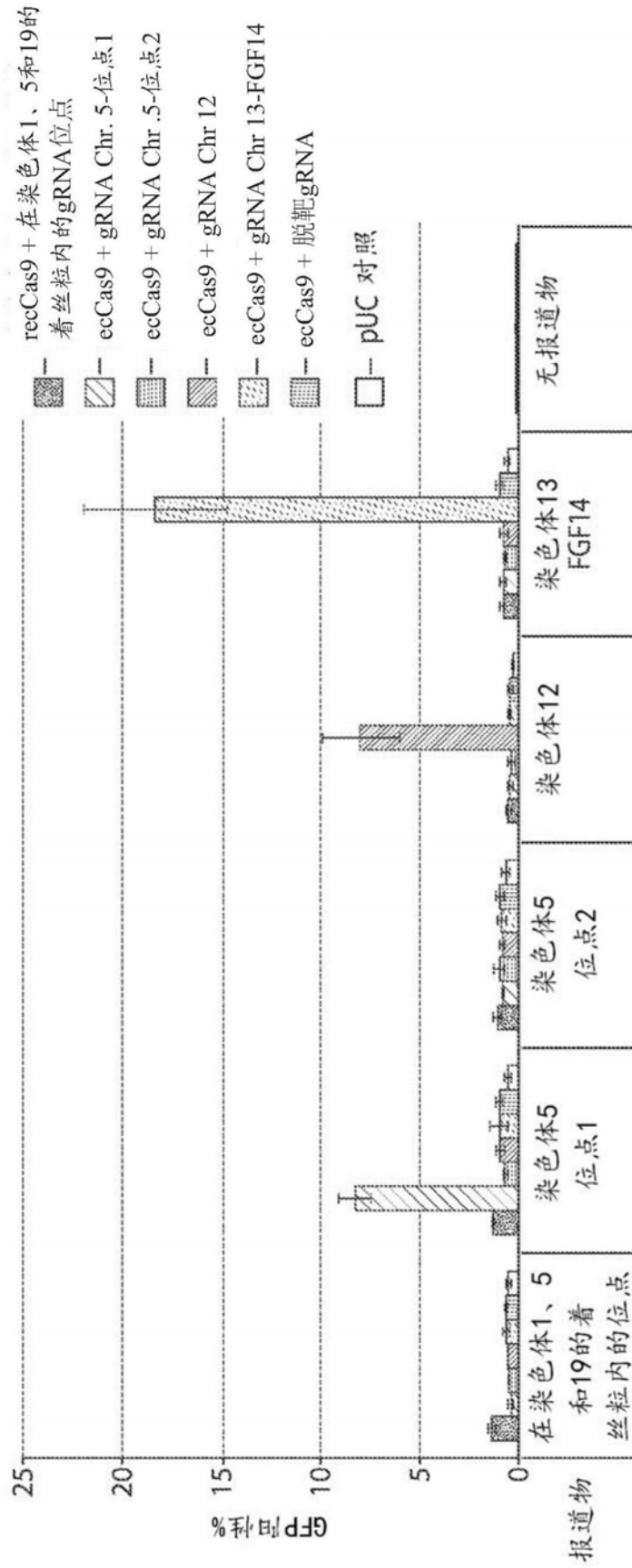


图4A



图4B

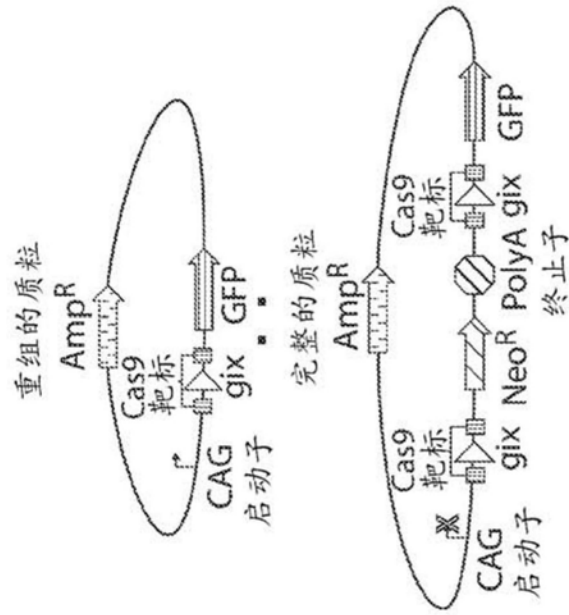


图4C

gRNA/RCas9组合

	gRNAs 染色体5位点1	gRNAs 染色体12	gRNAs 染色体13 FGF14	pUC 对照
	GinB-8GGS- -dCas9	GinB-8GGS- -dCas9	GinB-8GGS- -dCas9	pUC 对照
报道物: 染色体5位点1	11.96±0.54%	0.00%	0.00%	0.00%
报道物: 染色体12	0.00%	23.49±0.41%	0.00%	0.00%
报道物: 染色体13 FGF14	0.00%	0.00%	31.73±4.27%	0.00%

报道物

图4D

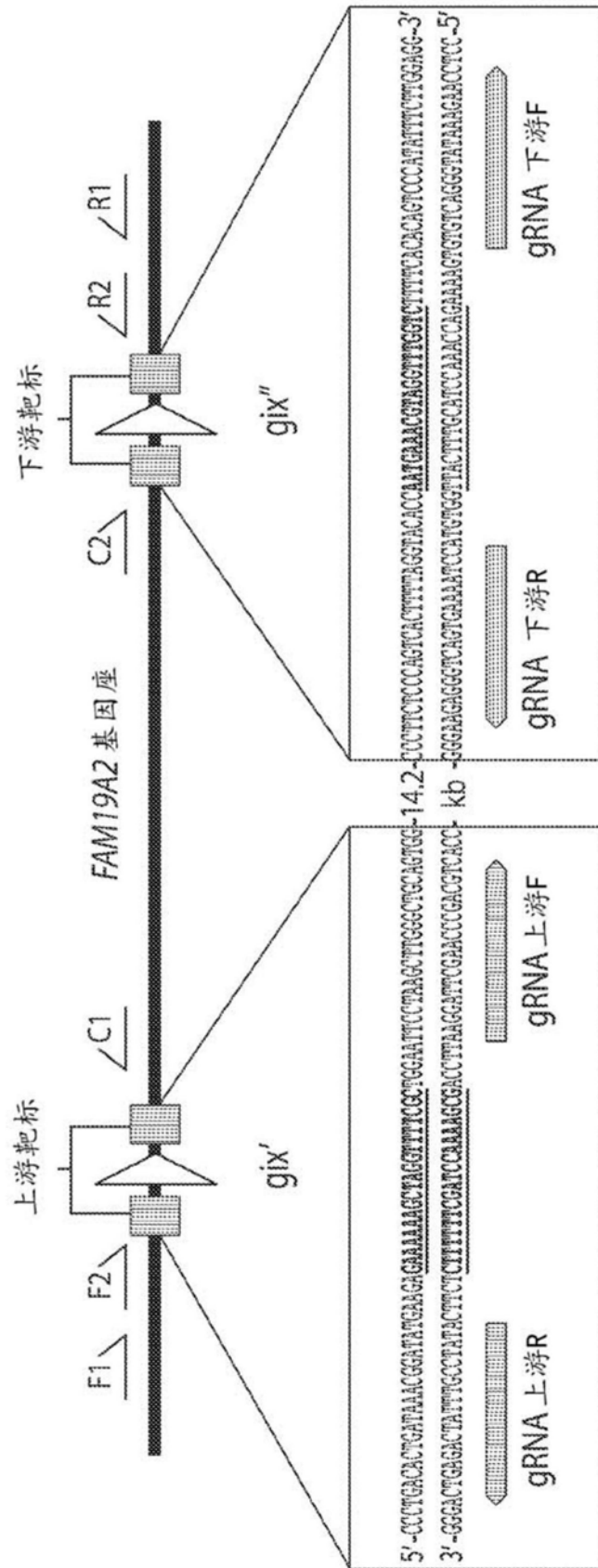


图5A

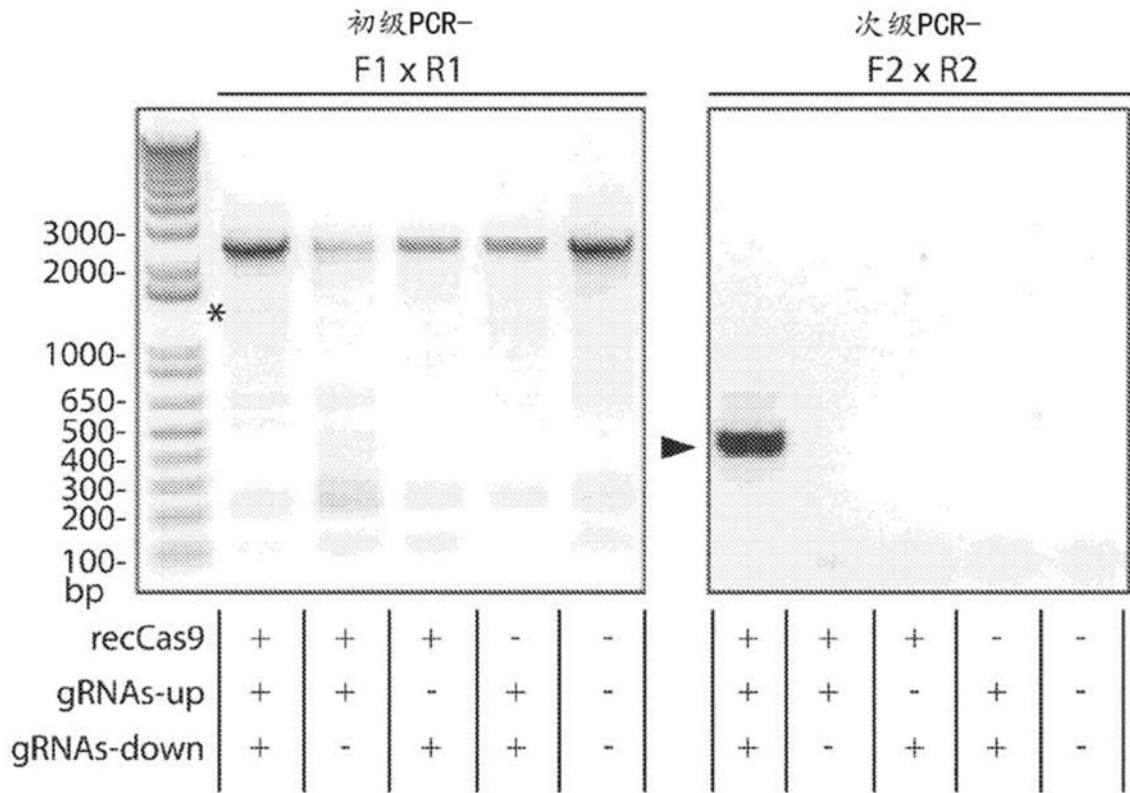


图5B

重组后接合

5' .. CCCTGACACTGATAAACGGGATATGAAGAGAAAGCTAGGTTGGTCTTTTCACACAGTCCCATATTTCTTTGGAGG... 3'
3' .. GGGACTGAGACTATTTGCCCTATCTCTCTTTTCGATCCAAACCAGAAAAGTGTGCAGGGTATAAAGAACCCTCC... 5'

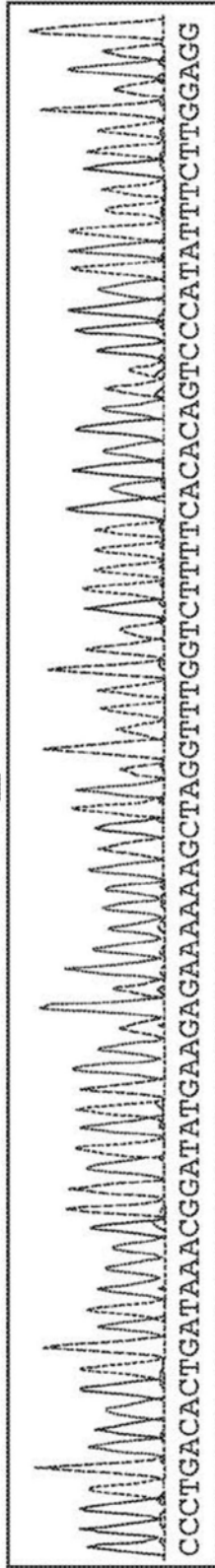


图5C

巢式PCR模板

最小缺失					
	样品1	样品2	样品3	未转染的对照	
	0.036±0.0233%	0.011±0.0072%	0.021±0.0091%	<0.0072%	

图5D

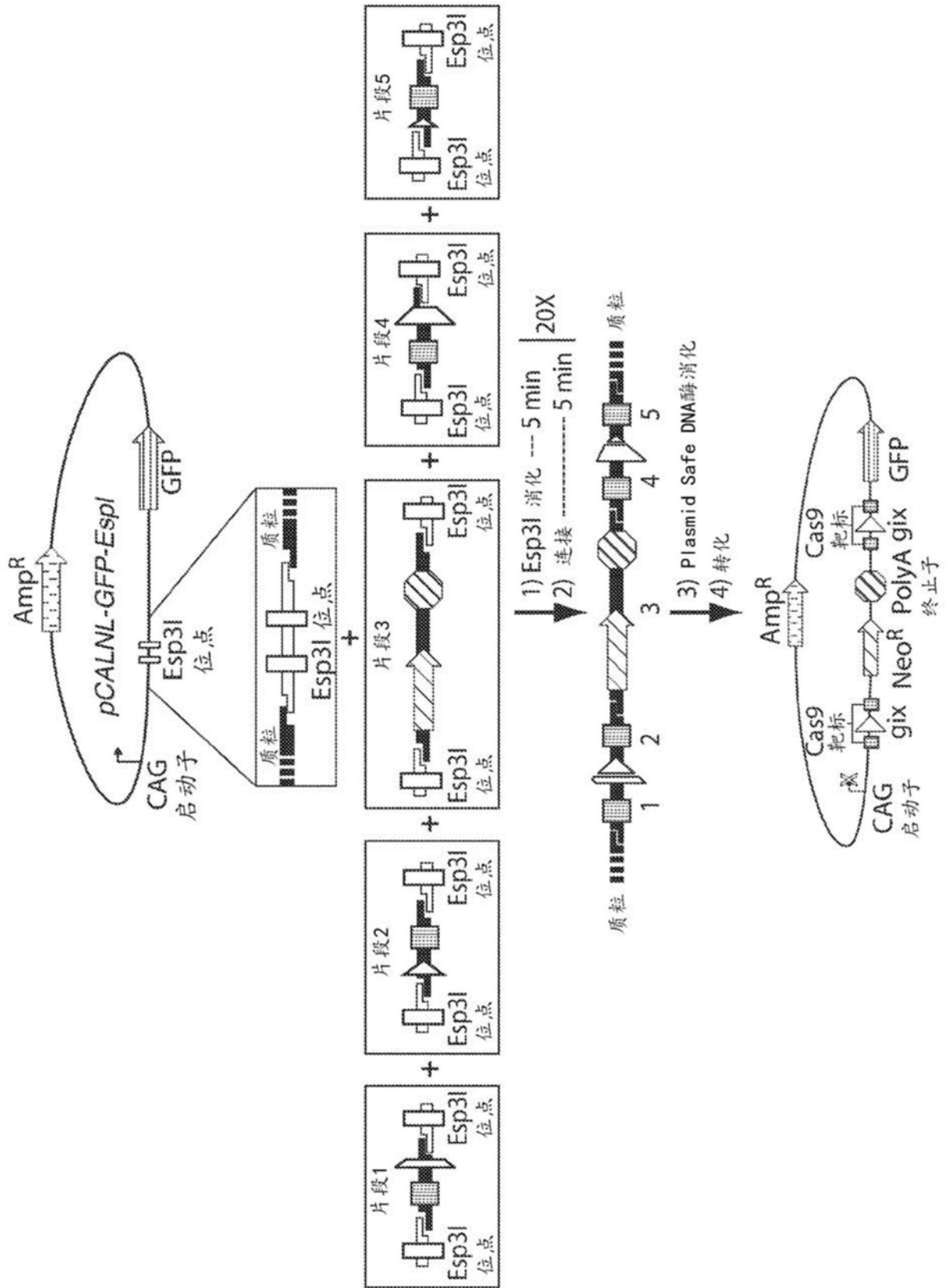


图6

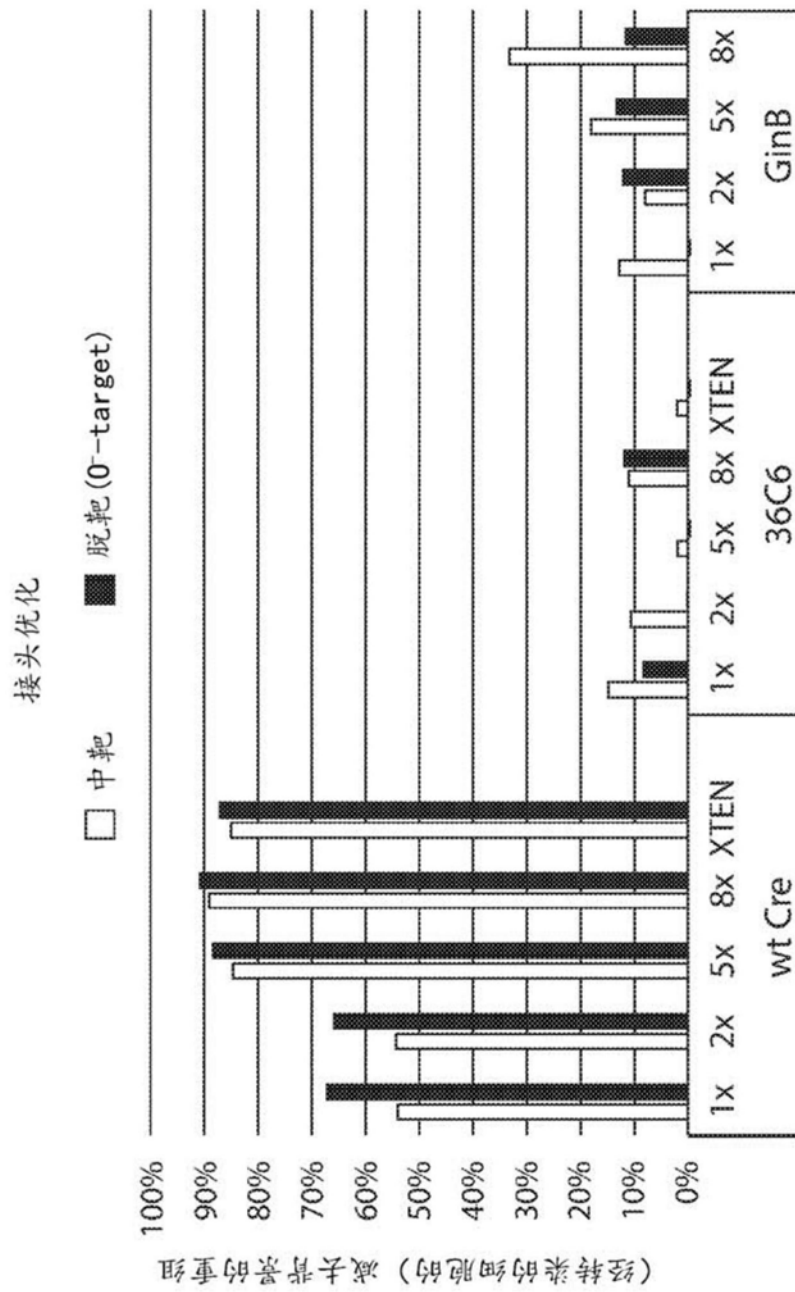


图7A

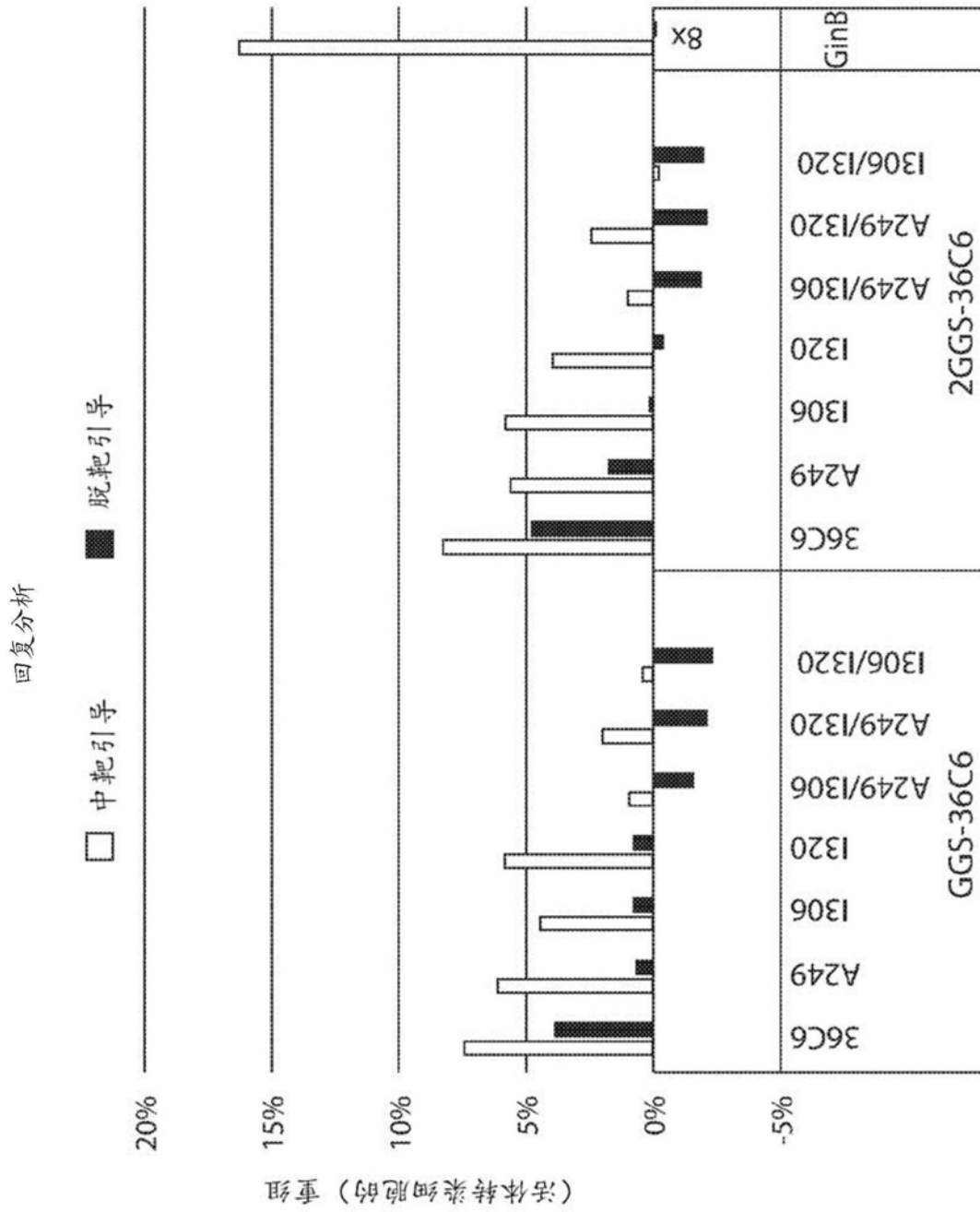


图7B

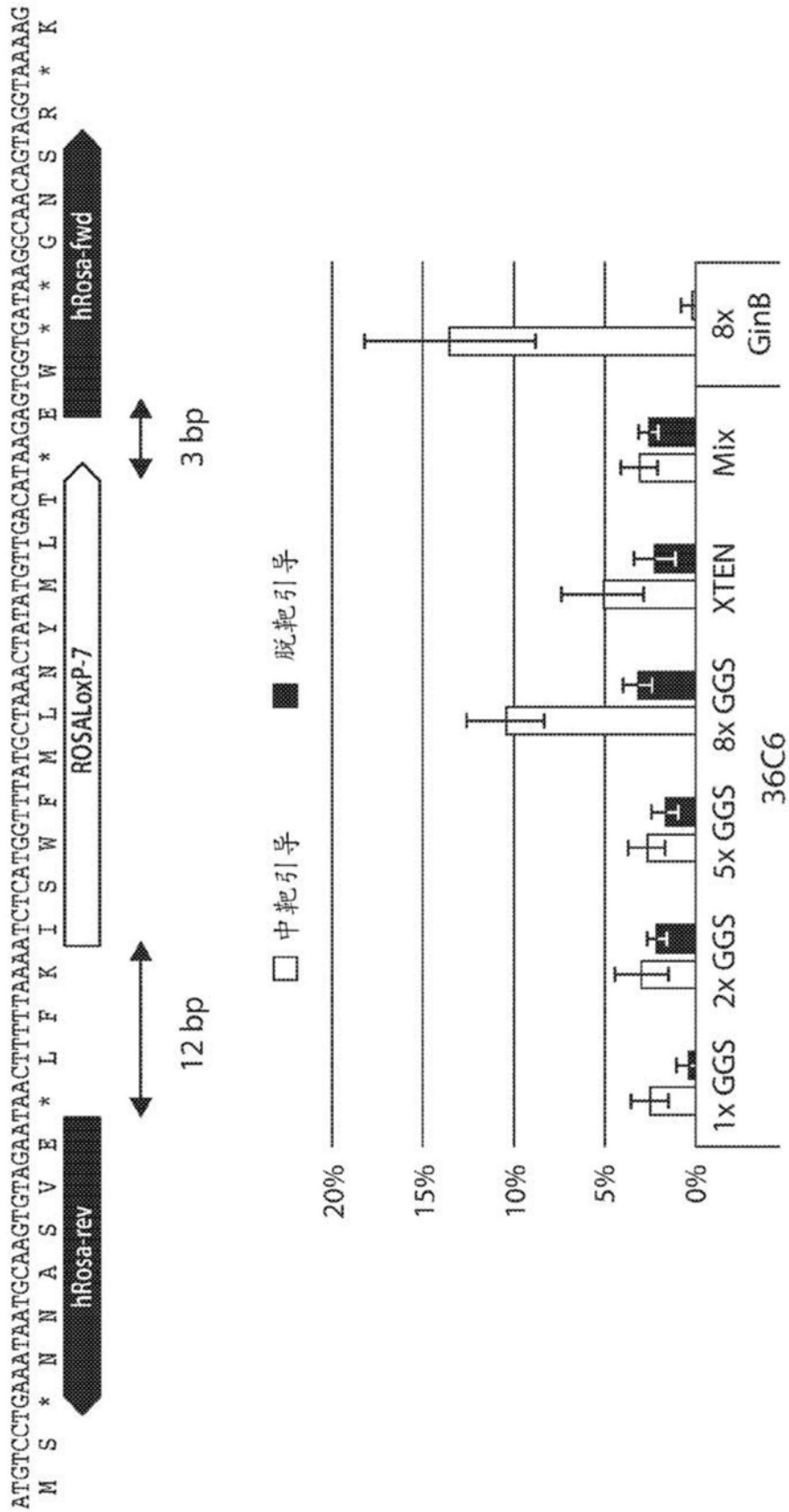


图8

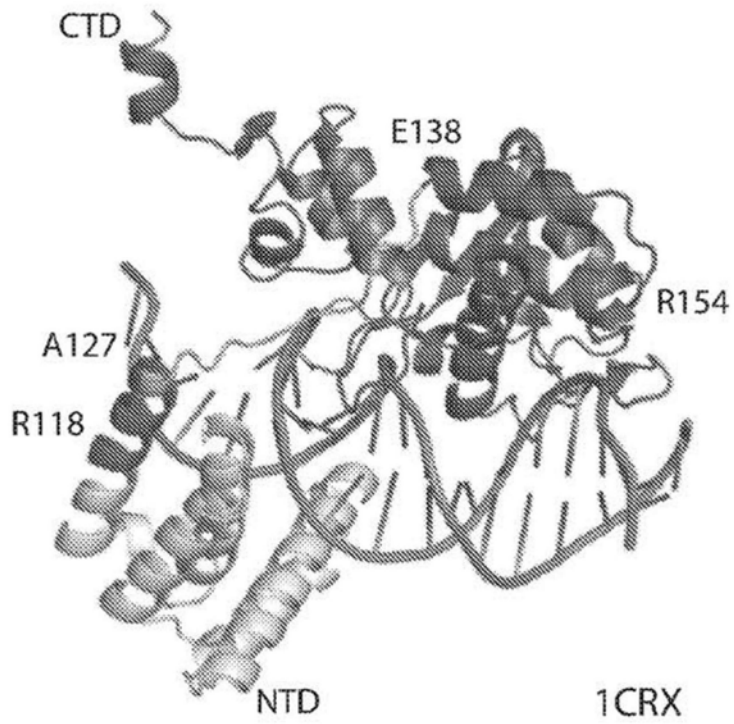


图9A

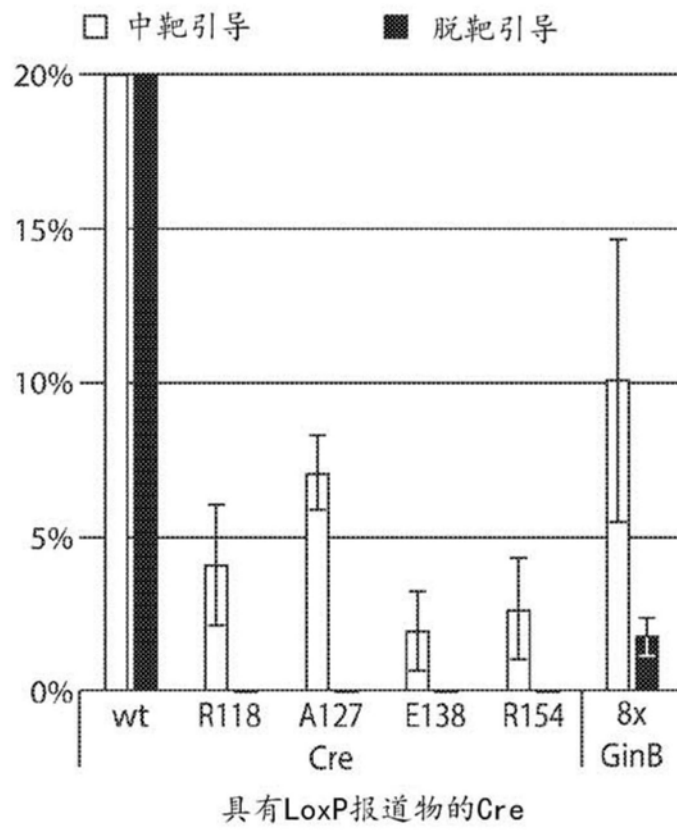


图9B