

TABLE 1—ESTIMATED REPORTING BURDEN <sup>1</sup>—Continued

Surveys of pharmacists and patients on variations in the physical characteristics of generic drug pills and patients' perceptions	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total Hours
Survey of patients #2 .....	1,000	1	1,000	0.3 (18 minutes)	300
Total .....					1,017

<sup>1</sup> There are no capital costs or operating and maintenance costs associated with this collection of information.

<sup>2</sup> Eligibility is determined prior to mailing the surveys; screening is not required.

**References**

The following references have been placed on display in the Division of Dockets Management (see **ADDRESSES**) and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday, and are available electronically at <http://www.regulations.gov>.

1. Woodward, C.A., "Questionnaire Construction and Question Writing for Research in Medical Education," *Medical Education*, 22, pp. 345–363 (1988).
2. Fitzpatrick, R., "Surveys of Patient Satisfaction: II—Designing a Questionnaire and Conducting a Survey," *British Medical Journal*, 302(6785), pp. 1129–1132 (1991).

Dated: May 8, 2015.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2015–11623 Filed 5–13–15; 8:45 am]

**BILLING CODE 4164–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2014–N–1031]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; FDA Recall Regulations**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled, "FDA Recall Regulations" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, [PRAStaff@fda.hhs.gov](mailto:PRAStaff@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** On February 27, 2015, the Agency submitted a proposed collection of information entitled, "FDA Recall Regulations" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0249. The approval expires on March 31, 2018. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: May 8, 2015.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2015–11624 Filed 5–13–15; 8:45 am]

**BILLING CODE 4164–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2014–N–1076]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry: Formal Dispute Resolution; Scientific and Technical Issues Related to Pharmaceutical Current Good Manufacturing Practice**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled, "Guidance for Industry: Formal Dispute Resolution; Scientific and Technical Issues Related to Pharmaceutical Current Good Manufacturing Practice" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** FDA PRA Staff, Office of Operations, Food

and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, [PRAStaff@fda.hhs.gov](mailto:PRAStaff@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** On January 8, 2015, the Agency submitted a proposed collection of information entitled, "Guidance for Industry: Formal Dispute Resolution; Scientific and Technical Issues Related to Pharmaceutical Current Good Manufacturing Practice" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0563. The approval expires on April 30, 2018. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: May 8, 2015.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2015–11609 Filed 5–13–15; 8:45 am]

**BILLING CODE 4164–01–P**

**DEPARTMENT OF HOMELAND SECURITY**

**U.S. Customs and Border Protection**

[Docket No. USCBP–2015–0020]

**The U.S. Customs and Border Protection Airport and Seaport Inspections User Fee Advisory Committee (UFAC)**

**AGENCY:** U.S. Customs and Border Protection, Department of Homeland Security (DHS).

**ACTION:** Committee Management; Notice of Federal Advisory Public Committee Meeting.

**SUMMARY:** The U.S. Customs and Border Protection Airport and Seaport Inspections User Fee Advisory Committee (UFAC) will meet on Tuesday, June 2, 2015, in Washington,

DC. The meeting will be open to the public.

**DATES:** The UFAC will meet on Tuesday, June 2, 2015, from 1:00 p.m. to 2:30 p.m. EST. Please note that the meeting is scheduled for one and a half hours and that the meeting may close early if the committee completes its business.

*Pre-Registration:* Meeting participants may attend either in person or via webinar after pre-registering using a method indicated below:

—For members of the public who plan to attend the meeting in person, please register either online at [https://apps.cbp.gov/te\\_reg/index.asp?w=43](https://apps.cbp.gov/te_reg/index.asp?w=43), by email to [tradeevents@dhs.gov](mailto:tradeevents@dhs.gov); or by fax to (202) 325-4290 by 5:00 p.m. EST on May 29, 2015.

—For members of the public who plan to participate via webinar, please register online at [https://apps.cbp.gov/te\\_reg/index.asp?w=44](https://apps.cbp.gov/te_reg/index.asp?w=44), by 5:00 p.m. EST on May 29, 2015.

Feel free to share this information with other interested members of your organization or association.

Members of the public who are pre-registered and later require cancellation, please do so in advance of the meeting by accessing one (1) of the following links: [https://apps.cbp.gov/te\\_reg/cancel.asp?w=43](https://apps.cbp.gov/te_reg/cancel.asp?w=43) to cancel an in person registration, or [https://apps.cbp.gov/te\\_reg/cancel.asp?w=44](https://apps.cbp.gov/te_reg/cancel.asp?w=44) to cancel a webinar registration.

**ADDRESSES:** The meeting will be held at the U.S. International Trade Commission, 500 E Street SW., Courtroom A, Washington, DC 20436.

All visitors to the International Trade Commission Building must show a state-issued ID or Passport to proceed through the security checkpoint for admittance to the building. There will be signage posted directing visitors to the location of Courtroom A.

For information on facilities or services for individuals with disabilities or to request special assistance at the meeting, contact Ms. Wanda Tate, Office of Trade Relations, U.S. Customs and Border Protection at (202) 344-1661 as soon as possible.

To facilitate public participation, we are inviting public comment on the issues to be considered by the committee prior to the formulation of recommendations as listed in the "Agenda" section below.

Comments must be submitted in writing no later than May 25, 2015, and must be identified by Docket No. USCBP-2015-0020, and may be submitted by one of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments.

- *Email:* [Tradeevents@dhs.gov](mailto:Tradeevents@dhs.gov). Include the docket number in the subject line of the message.

- *Fax:* (202) 325-4290.

- *Mail:* Ms. Wanda Tate, Office of Trade Relations, U.S. Customs and Border Protection, 1300 Pennsylvania Avenue NW., Room 3.5A, Washington, DC 20229.

*Instructions:* All submissions received must include the words "Department of Homeland Security" and the docket number for this action. Comments received will be posted without alteration at <http://www.regulations.gov>, including any personal information provided. Do not submit personal information to this docket.

*Docket:* For access to the docket to read background documents or comments, go to <http://www.regulations.gov> and search for Docket Number USCBP-2015-0020. To submit a comment, see the link on the Regulations.gov Web site for "How do I submit a comment?" located on the right hand side of the main site page.

There will be two (2) public comment periods held during the meeting on June 2, 2015. Speakers are requested to limit their comments to two (2) minutes or less to facilitate greater participation. Contact the individual listed below to register as a speaker. Please note that the public comment periods for speakers may end before the times indicated on the schedule that is posted on the CBP Web page, <http://www.cbp.gov/trade/stakeholder-engagement/user-fee-advisory-committee>.

**FOR FURTHER INFORMATION CONTACT:** Ms. Wanda Tate, Office of Trade Relations, U.S. Customs and Border Protection, 1300 Pennsylvania Avenue NW., Room 3.5A, Washington, DC 20229; telephone (202) 344-1440; facsimile (202) 325-4290.

**SUPPLEMENTARY INFORMATION:** Pursuant to the Federal Advisory Committee Act (5 U.S.C. Appendix), the Department of Homeland Security (DHS) hereby announces the meeting of the U.S. Customs and Border Protection Airport and Seaport Inspections User Fee Advisory Committee (UFAC). The UFAC is tasked with providing advice to the Secretary of Homeland Security (DHS) through the Commissioner of U.S. Customs and Border Protection (CBP) on matters related to the performance of airport and seaport inspections coinciding with the assessment of an agriculture, customs, or immigration user fee. The UFAC meeting will be

held on the date and time specified above.

### Agenda

The UFAC will meet to discuss and report the work completed by the Financial Assessment and Options Subcommittee and the Processes Subcommittee:

1. The Financial Assessment and Options Subcommittee will discuss an overview of current worldwide user fees being paid by industry, and mapping how industry collects and transmits user fees to U.S. Customs and Border Protection (CBP).

2. The Processes Subcommittee will discuss developing advice that would enhance U.S. Customs and Border Protection (CBP) operational efficiencies.

Dated: May 8, 2015.

**Maria Luisa Boyce,**

*Senior Advisor for Private Sector Engagement, Office of Trade Relations, U.S. Customs and Border Protection.*

[FR Doc. 2015-11619 Filed 5-13-15; 8:45 am]

**BILLING CODE 9111-14-P**

## DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS-2015-0022]

### Agency Information Collection Activities: Post-Award Contract Information

**AGENCY:** Office of the Chief Procurement Officer, DHS.

**ACTION:** 60-Day Notice and request for comments; Extension without change, 1600-0003.

**SUMMARY:** The Department of Homeland Security, Office of the Chief Procurement Officer, will submit the following Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. Chapter 35).

**DATES:** Comments are encouraged and will be accepted until July 13, 2015. This process is conducted in accordance with 5 CFR 1320.1

**ADDRESSES:** You may submit comments, identified by docket number DHS-2014-0022, by one of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Please follow the instructions for submitting comments.

- *Email:* [dhs.pra@hq.dhs.gov](mailto:dhs.pra@hq.dhs.gov) Please include docket number DHS-2015-0022 in the subject line of the message.