TABLE 1—ESTIMATED REPORTING BURDEN [1]—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

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.
[2] 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.*

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.*

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,