three requests for documentation of successful completion of staff training using the CFP Training Plan and Log for a total of 1,500 annual responses. Each submission is estimated to take 0.1 hour (or 6 minutes) per response for a total of 150 hours. Thus, the total reporting burden for this information collection is 200 hours.

Based on a review of the information collection since our last request for OMB approval, we have made no adjustments to our burden estimate.

Dated: July 6, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020–14879 Filed 7–9–20; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2018–N–2434, FDA–2016–N–3535, FDA–2013–N–1619, FDA–2016–N–0736, FDA–2019–N–3885, FDA–2013–N–1423, FDA–2013–N–0804, FDA–2016–N–3995, FDA–2018–D–1592, FDA–2016–N–2066, and FDA–2017–N–0366]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approvals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of information collections that have been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–7726, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The following is a list of FDA information collections recently approved by OMB under section 3507 of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507). The OMB control number and expiration date of OMB approval for each information collection are shown in table 1. Copies of the supporting statements for the information collections are available on the internet at *https://www.reginfo.gov/public/do/PRAMain.* 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.

TABLE 1—LIST OF INFORMATION COLLECTIONS APPROVED BY OMB

Title of collection	OMB control No.	Date approval expires
Formal Meetings Between the Food and Drug Administration and Sponsors and Applicants of Prescription Drug User Fee Act Products	0910–0429	5/31/2023
Special Protocol Assessments	0910–0470	5/31/2023
Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements	0910–0606	5/31/2023
Tracking Network for PETNet, LivestockNet, and SampleNet	0910–0680	5/31/2023
Center for Tobacco Products, Food and Drug Administration Funded Trainee/Scholar Survey	0910–0887	5/31/2023
Importer's Entry Notice	0910–0046	6/30/2023
Premarket Notification Submission 510(k), Subpart E	0910–0120	6/30/2023
Medical Devices; Pediatric Uses of Devices; Requirement for Submission of Information on Pediatric Subpopulations	0910–0748	6/30/2023
Controlled Correspondence Related to Generic Drug Development	0910–0797	6/30/2023
Certification of Identity for Freedom of Information Act and Privacy Act Requests	0910–0832	6/30/2023
FDA Advisory Committee Membership Nominations	0910–0833	6/30/2023

Dated: July 6, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020–14875 Filed 7–9–20; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–N–1391]

Office of Women's Health Strategic Priorities; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA or Agency) is opening a public docket to solicit input and comments from stakeholders interested in informing strategic priorities for the Office of Women's Health (OWH). This will help the Agency ensure that important health concerns are carefully considered in establishing OWH's scientific, educational, and outreach priorities.

DATES: Submit either electronic or written comments by September 8, 2020.

ADDRESSES: You may submit comments as follows. Please note that untimely comments will not be considered. Electronic comments must be submitted on or before September 8, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of September 8, 2020. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- Federal eRulemaking Portal: *https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted,