DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2016–N–0628; FDA–2012–N–0306; FDA–2002–N–0323; FDA–2012–N–0427; FDA–2012–N–0536; FDA–2012–N–0560; FDA–2015–N–3662; FDA–2012–N–0976; FDA–2013–N–0297; FDA–2012–N–1203; FDA–2011–D–0893; FDA–2014–N–0189; FDA–2012–N–1210]

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: FDA PRA Staff, Office of Operations, Food and Drug Administration, Three White Flint North, 11601 Landsdown St., North Bethesda, MD 20852, *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 *http://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
Reporting Associated with New Animal Drug Applications	0910–0032	8/31/2019
Administrative Detention and Banned Medical Devices	0910–0114	8/31/2019
Registration of Food Facilities	0910–0502	8/31/2019
Inspection by Accredited Persons Program Under the Medical Device User Fee and Modernization Act of 2002	0910–0510	8/31/2019
Medical Device User Fee Cover Sheet—FDA Form 3601	0910–0511	8/31/2019
Guidance on Informed Consent for in Vitro Diagnostic Studies Using Leftover Human Specimens That Are Not Individually Identifiable	0910–0582	8/31/2019
Guidance for Reagents for Detection of Specific Novel Influenza A Viruses	0910–0584	8/31/2019
Guidance: Emergency Use Authorization of Medical Products	0910–0595	8/31/2019
Prevention of *Salmonella* Enteritidis in Shell Eggs During Production—Recordkeeping and Registration Provisions	0910–0660	8/31/2019
Information to Accompany Humanitarian Device Exemption Applications and Annual Distribution Number Reporting Requirements	0910–0661	8/31/2019
Guidance for Center for Devices and Radiological Health Appeals Processes	0910–0738	8/31/2019
Deeming Tobacco Products To Be Subject to the FD&C Act	0910–0768	8/31/2019
Food Labeling: Revision of the Nutrition Facts Label and Supplement Facts Label	0910–0813	7/31/2019

Dated: September 6, 2016.

Leslie Kux,

Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–0001]

Request for Nominations for Voting Members on a Public Advisory Committee; Tobacco Products Scientific Advisory Committee

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is requesting nominations for voting members to serve on the Tobacco Products Scientific Advisory Committee, Office of Science, Center for Tobacco Products.

FDA seeks to include the views of women and men, members of all racial and ethnic groups, and individuals with and without disabilities on its advisory committees and, therefore, encourages nominations of appropriately qualified candidates from these groups.

DATES: Nominations received on or before November 14, 2016 will be given first consideration for membership on the Tobacco Products Scientific Advisory Committee. Nominations received after November 14, 2016 will be considered for nomination to the committee as later vacancies occur.

ADDRESSES: All nominations for membership should be sent electronically by logging into the FDA Advisory Nomination Portal: *http://www.accessdata.fda.gov/scripts/FACTRSPortal/FACTRS/index.cfm* or by mail to Advisory Committee Oversight and Management Staff, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 5103, Silver Spring, MD 20993–0002.

FOR FURTHER INFORMATION CONTACT: *Regarding all nomination questions for membership, the primary contact is:* Caryn Cohen, Office of Science, Center for Tobacco Products, Food and Drug Administration, Document Control Center, 10903 New Hampshire Ave., Bldg. 71, Rm. G335, Silver Spring, MD 20993–0002, 1–877–287–1373 (choose Option 5), *TPSAC@fda.hhs.gov.*

Information about becoming a member on an FDA advisory committee can also be obtained by visiting FDA's Web site by using the following link: *http://www.fda.gov/AdvisoryCommittees/default.htm.*

SUPPLEMENTARY INFORMATION: FDA is requesting nominations for voting members on the Tobacco Products Scientific Advisory Committee.

I. General Description of the Committee Duties

The Tobacco Products Scientific Advisory Committee (the Committee)