4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA-2018-N-3442; FDA-2013-N-0557; FDA-2013-N-0514; FDA-2013-N-0190; FDA-2013-N-1428; FDA-2019-N-0075; FDA-2016-N-2544; FDA-2019-N-2778; FDA-2012-N-0977; FDA-2013-N-0823; FDA-2009-N-0380; FDA-2013-N-1147; FDA-2010-N-0117; FDA-2010-D-0350; FDA-2010-D-0319; FDA-2012-D-0530; and FDA-2016-N-2683]

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 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 Number	Date Approval Expires
Web-Based Pilot Survey to Assess Allergy to Cosmetics in the United States	0910-0881	1/31/2021
Postmarket Surveillance of Medical Devices	0910-0449	11/30/2022
Administrative Procedures for Clinical Laboratory Improvement Amendments Categorization	0910-0607	11/30/2022
Requirements Under the Comprehensive Smokeless Tobacco Health Education Act of 1986, as Amended by the Family Smoking Prevention and Tobacco Control Act	0910-0671	11/30/2022
Electronic Drug Product Reporting for Human Drug Compounding Outsourcing Facilities Under Section 503B of the Federal Food, Drug, and Cosmetic Act	0910-0827	11/30/2022
Experimental Study on Measuring Consumer Comprehension of Displays of Harmful and Potentially Harmful Constituents in Tobacco Products and Tobacco Smoke	0910-0880	11/30/2022
Medical Devices; Current Good Manufacturing Practice Quality System Regulation	0910-0073	12/31/2022
Threshold of Regulation for Substances Used in Food-Contact Articles	0910-0298	12/31/2022
Regulations Restricting the Sale and Distribution of Cigarettes and Smokeless Tobacco to Protect Children and Adolescents	0910-0312	12/31/2022
Format and Content Requirements for Over-the-Counter Drug Product Labeling	0910-0340	12/31/2022
Product Jurisdiction: Assignment of Agency Component for Review of Premarket Applications	0910-0523	12/31/2022
Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition	0910-0541	12/31/2022
Hypertension Indication: Drug Labeling for Cardiovascular Outcome Claims	0910-0670	12/31/2022
Guidance for Tobacco Retailers on Tobacco Retailer Training Programs	0910-0745	12/31/2022
Dear Health Care Provider Letters: Improving Communication of Important Safety Information	0910-0754	12/31/2022
Requests for Feedback on Medical Device Submissions	0910-0756	12/31/2022
Data to Support Social and Behavioral Research as Used by the Food and Drug Administration	0910-0847	12/31/2022

Dated: January 24, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

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