4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA-2022-N-1894; FDA-2018-N-3303; FDA-2022-N-0576; FDA-2022-N-1794; FDA-2011-N-0902; FDA-2009-N-0545; FDA-2016-N-2474; FDA-2010-D-0350; FDA-2012-D-0530; FDA-2016-N-2683; FDA-2013-N-0403; FDA-2013-N-0134; FDA-2022-N-2440; FDA-2013-N-0879; and FDA-2014-N-1048]

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: JonnaLynn Capezzuto, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-3794, 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 Number	Date Approval Expires
Yale-Mayo Clinic Centers of Excellence in Regulatory Science and Innovation B12 Pediatric Device Survey	0910-0912	3/31/2024
Electronic Products Requirements	0910-0025	2/28/2026
Investigational Device Exemptions	0910-0078	2/28/2026
General Drug Labeling Provisions and OTC Monograph Drug User Fee Submissions	0910-0340	2/28/2026
Prescription Drug Product Labeling; Medication Guide Requirements	0910-0393	2/28/2026
Reporting of Biological Product Deviations and Human Cells, Tissues, and Cellular and Tissue-Based Product Deviations in Manufacturing	0910-0458	2/28/2026
Designation of New Animal Drugs for Minor Use or Minor Species	0910-0605	2/28/2026
Tobacco Retailer Training Programs	0910-0745	2/28/2026
Q-Submission and Early Payor Feedback Request Programs for Medical Devices	0910-0756	2/28/2026
Data To Support Social and Behavioral Research as Used by the Food and Drug Administration	0910-0847	2/28/2026
Protection of Human Subjects and Institutional Review Boards	0910-0130	3/31/2026
Mammography Standards Quality Act Requirements	0910-0309	3/31/2026
Biologics License Applications; Procedures & Requirements	0910-0338	3/31/2026
Procedures for the Safe Processing and Importing of Fish and Fishery Products	0910-0354	3/31/2026
Medical Device Labeling Regulations; Unique Device Identification	0910-0485	3/31/2026

Dated: April 28, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

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