4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA-2016-N-2976; FDA-2016-N-3535; FDA-2013-N-1089; FDA-2013-N-1619; FDA-2013-N-0719; FDA-2016-N-3586; FDA-2013-N-0796; FDA-2016-N-0736; FDA-2016-N-3995; FDA-2013-D-0575; FDA-2016-N-0735]

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, 10A63, 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 Number	Date Approval Expires
Request for Information From U.S. Processors That Export to the European Community	0910-0320	5/31/2020
Guidance for Industry: Special Protocol Assessment	0910-0470	5/31/2020
Use of Symbols on Labels and in Labeling of In Vitro Diagnostic Devices Intended for Professional Use	0910-0553	5/31/2020
Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements	0910-0606	5/31/2020
Guidance for Industry: Planning for the Effects of High Absenteeism to Ensure Availability of Medically Necessary Drug Products	0910-0675	5/31/2020
Focus Groups About Drug Products as Used by the Food and Drug Administration	0910-0677	5/31/2020
Testing Communication on Medical Devices and Radiation-Emitting Products	0910-0678	5/31/2020
Tracking Network for PETNet, LivestockNet, and SampleNet	0910-0680	5/31/2020
Medical Devices: Pediatric Uses of Devices; Requirements for Submission of Information on Pediatric Subpopulations That Suffer From a Disease or Condition That a Device is Intended to Treat, Diagnose, or Cure	0910-0748	5/31/2020
Guidance for Industry: Expedited Programs for Serious Conditions--Drugs and Biologics	0910-0765	5/31/2020
Superimposed Text in Direct-to-Consumer Promotion of Prescription Drugs	0910-0831	5/31/2020

Dated: June 27, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

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