Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Transmittal #1—Initial Request	54	14,216	0.17	130,503
Transmittal #1—Initial Request Acknowledgement	54	14,216	0.05	38,383
Transmittal #2—Subsequent Action	54	10,662	0.08	46,060
Transmittal #3—Request for Assistance/Discovery	54	2,132	0.08	9,210
Uniform Support Petition	54	5,686	0.05	15,352
General Testimony	54	5,686	0.33	101,325
Declaration in Support of Establishing Parentage	54	2,132	0.15	17,269
Child Support Locate Request	54	142	0.05	383
Notice of Determination of Controlling Order	54	1	0.25	14
Letter of Transmittal Requesting Registration	54	8,529	0.08	36,845
Personal Information Form for UIFSA § 311	54	5,686	0.05	15,352
Child Support Agency Confidential Information Form	54	17,059	0.05	46,059
Request for Change of Support Payment Location Pursuant to UIFSA 319(b)	54	71	0.05	192
Estimated Total Annual Burden Hours				456,947

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: 45 CFR 303.7.

Mary C. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2024–N–2844; FDA–2024–N–4687; FDA–2024–N–5581; FDA–2024–N–2931; FDA–2024–N–4470; FDA–2024–N–2865]

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: Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, *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
Reclassification Petitions for Medical Devices	0910–0138	7/31/2028
Medicated Feed Mill License Application	0910–0337	6/30/2028
Additives in Animal Food	0910–0546	6/30/2028
Microbiological Testing and Corrective Measures for Bottled Water	0910–0658	6/30/2028
Antimicrobial Animal Drug Sales and Distribution	0910–0659	6/30/2028
Generic Clearance for Quantitative Testing for the Development of FDA Communications	0910–0865	6/30/2028

Dated: July 23, 2025.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2025–14224 Filed 7–28–25; 8:45 am]

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