

*Estimated Total Annual Burden Hours:* 3,883.

*Authority:* 42 U.S.C. 652(a)(7); 42 U.S.C. 666(c)(1); and 45 CFR 303.7(a)(5).

**Mary B. Jones,**  
*ACF/OPRE Certifying Officer.*

[FR Doc. 2022-08770 Filed 4-22-22; 8:45 am]

**BILLING CODE 4184-41-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for OMB Review; Comprehensive Child Welfare Information System (CCWIS) Automated Function Checklist and Data Quality Plan (OMB #0970-0463)**

**AGENCY:** Children’s Bureau, Administration for Children and Families, HHS.

**ACTION:** Request for public comment.

**SUMMARY:** The Administration for Children and Families (ACF) is requesting a 3-year extension of the Comprehensive Child Welfare Information System (CCWIS) information collection (OMB #0970-0463, expiration 8/31/2022). The CCWIS information collection includes the Automated Function List and the Data Quality Plan. There are no required instruments associated with the data collection and no changes to the data collection.

**DATES:** *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/](http://www.reginfo.gov/public/do/)

*PRAMain.* Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. You can also obtain copies of the proposed collection of information by emailing [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). Identify all emailed requests by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* The CCWIS information collection includes two components:

- The Automated Function List update required pursuant to section 1355.52(i)(2); and
- The Data Quality Plan update required pursuant to section 1355.52(d)(5).

The CCWIS regulations require updates of this information to confirm that the project meets CCWIS requirements and that project costs are appropriately allocated to benefiting programs.

*Respondents:* Title IV–E agencies under the Social Security Act.

**ANNUAL BURDEN ESTIMATES**

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Automated Function List section 1355.52(i)(2) .....	55	1	10	550
Data Quality Plan section 1355.52(d)(5) .....	55	1	40	2,200

*Estimated Annual Burden Hours:* 2,750.

*Authority:* 42 U.S.C. 620 *et seq.*, 42 U.S.C. 670 *et seq.*, 42 U.S.C. 1301 and 1302.

**Mary B. Jones,**  
*ACF/OPRE Certifying Officer.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2018-N-4206]

**Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Device User Fee Small Business Qualification and Certification**

**AGENCY:** Food and Drug Administration, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing

that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Fax written comments on the collection of information by May 25, 2022.

**ADDRESSES:** To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to <https://www.reginfo.gov/public/do/PRAMain>. Find this particular information collection by selecting “Currently under Review—Open for Public Comments” or by using the search function. All comments should be identified with the OMB control number 0910-0508. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-45, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-5733, [PRAStaff@fda.hhs.gov](mailto:PRAStaff@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

**Medical Device User Fee Small Business Qualification and Certification**

*OMB Control Number 0910-0508—Extension*

This information collection helps support implementation of the Medical Device User Fee and Modernization Act of 2002 (MDUFMA) (Pub. L. 107-250) and FDA’s Medical Device User Fee program. Current authorization for medical device user fees will be in place from October 1, 2017, until September 30, 2022.

Section 738(d)(2)(A) and (e)(2)(A) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379j(d)(2)(A) and (e)(2)(A)) define a “small business” as an entity that reported \$100 million or less of gross receipts or sales in its most recent Federal income tax return, including such returns of its affiliates, partners, and parent firms. If a firm’s gross receipts or sales are no more than