DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Federal Case Registry (FCR) (OMB #0970–0421)

AGENCY: Office of Child Support Enforcement, Administration for Children and Families, Health and Human Services (HHS).

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting a 3-year extension of the Federal Case Registry (FCR). There are no changes to the collection instruments used for the FCR (current Office of Management and Budget (OMB) approval expires January 31, 2021).

DATES: *Comments due within 30 days of publication.* OMB is required to make a decision concerning 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/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

SUPPLEMENTARY INFORMATION:

Description: ACF implemented the FCR within the Federal Parent Locator Service (FPLS) on October 1, 1998, pursuant to federal law. The FCR is a national database of information pertaining to child support cases processed by state child support agencies, referred to as ''IV–D'' cases, and non-IV–D support orders privately established or modified by courts or tribunals on or after October 1, 1998. FCR information is submitted by each State Case Registry (SCR), which is a central registry of child support orders and cases. The FCR automatically compares new SCR submissions to existing FCR information and notifies state agencies if an IV–D case participant in the state appears as a participant in an IV–D or non-IV-case in another state.

Respondents: State child support enforcement agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Annual burden hours
Appendix G: Input Record Layout	54	151	0.0333	272

Estimated Total Annual Burden Hours: 272.

Authority: The information collection activities pertaining to the FCR are authorized by: 42 U.S.C. 653(h), which requires the establishment of the FCR within the FPLS; 42 U.S.C. 654a(e), which requires state child support agencies to include a SCR in the state's automated system; and 42 U.S.C. 654a(f)(1), which requires states to conduct information comparison activities between the SCR and the FCR.

John M. Sweet Jr.,

ACF/OPRE Certifying Officer.

[FR Doc. 2020–19840 Filed 9–4–20; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Generic for Administration for Children and Families (ACF) Program Monitoring Activities (New Collection)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families (ACF), Health and Human Services (HHS).

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) intends to request approval from the Office of Management and Budget (OMB) for a new generic clearance for information collections related to ACF program office monitoring activities. ACF programs promote the economic and social well-being of families, children, individuals, and communities. The proposed Generic for ACF Program Monitoring Activities would allow ACF program offices to collect standardized information from recipients that receive federal funds to ensure oversight, evaluation, support purposes, and stewardship of federal funds.

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/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

SUPPLEMENTARY INFORMATION:

Description: Program monitoring is a post-award process through which ACF assesses a recipient's programmatic performance and business management performance. Monitoring activities are necessary to ensure timely action by ACF to support grantees and protect federal interests.

Program offices would use information collected under this generic clearance to monitor funding recipient activities and to provide support or take appropriate action, as needed. The information gathered will be used primarily for internal purposes, but aggregate data may be included in public materials such as Reports to Congress or program office documents. Following standard OMB requirements, ACF will submit a request for each individual data collection activity under this generic clearance. Each request will include the individual form(s) or instrument(s), a justification specific to the individual information collection, and any supplementary documents. OMB is requested to review requests within 10 days of submission.

Respondents: ACF funding recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hour per response	Total burden hours
Program Monitoring Forms	1500	3	10	45,000

John M. Sweet Jr.,

ACF/OPRE Certifying Officer.

[FR Doc. 2020–19811 Filed 9–4–20; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2020–N–0907]

Medical Device User Fee Amendments for Fiscal Years 2023 Through 2027; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is announcing a virtual public meeting to discuss proposed recommendations for the reauthorization of the Medical Device User Fee Amendments (MDUFA) for fiscal years (FYs) 2023 through 2027 (MDUFA V). MDUFA authorizes FDA to collect user fees to support the process for the review of medical device applications. The current legislative authority for MDUFA expires after September 30, 2022, and new legislation will be required for FDA to continue collecting user fees for the medical device program in future fiscal years. The Federal Food, Drug, and Cosmetic Act (FD&C Act) directs that FDA begin MDUFA reauthorization by publishing a notice in the **Federal Register** requesting public input and holding a public meeting where the public may present its views on the reauthorization, providing a period of 30 days after the public meeting to obtain written comments from the public suggesting changes to MDUFA, and publishing the comments on FDA's website.

DATES: The public meeting will be held on October 27, 2020, from 9 a.m. Eastern Time to 2 p.m. Eastern Time. Submit either electronic or written comments on the medical device user fee program and suggestions regarding the commitments FDA should propose for the next reauthorized program by November 27, 2020. Registration to view the meeting must be received by October 23, 2020.

ADDRESSES: Registration to attend this virtual public meeting and other information can be found at *https://www.fda.gov/medical-devices/workshops-conferences-medical-devices/2020-medical-device-meetings-and-workshops.* (Select this meeting from the posted events list.) See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before November 27, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time on November 27, 2020. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2020–N–0907 for ''Medical Device User Fee Amendments for Fiscal Years 2023 Through 2027.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting