ESTIMATED TOTAL ANNUAL BURDEN HOURS

Data collection activity	Total number of respondents	Number of responses per respondent (each year)	Average burden hours per response (in hours)	Total annual burden hours
Site Visit and Key Informant Data Collection:				
Program director individual interview	8	0.33	2	5
Program manager/supervisor individual interviews	8	0.33	1	3
Frontline staff interviews	16	0.33	1	5
Partner representative interviews	24	0.33	1	8
Partner survey	40	0.33	0.42	6
Sustainability survey	126	0.42	0.33	18
Enrollment, client and service data:				
Semi-annual progress reports	18	2	16.5	594
Case enrollment data	54	33	0.25	446
Case closure	54	33	0.0167	30
Case closure—prenatal	18	10	0.0167	3
Service log entries	108	1,560	0.033	5,560
Outcome and impact data:				
Administrative Data:				
Obtain access to administrative data	18	1	41	738
Report administrative data	18	2	144	5,184
Standardized instruments:				
Enter data into local database	18	100	.625	1,125
Review records and submit	18	2	25	900
Data entry for comparison study sites (16 grantees)	16	100	.625	1,000
Estimated Total Burden Hours	.....	.....	.....	15,625

Authority: The Child and Family Services Improvement Act of 2006 (Pub. L. 109–288) created the competitive RPG program. The September 30, 2011, passage of the Child and Family Services Improvement and Innovation Act (Pub. L. 112–34) extended funding for the RPG program from federal fiscal year (FFY) 2012 to FFY 2016. In 2018, the president signed the Bipartisan Budget Act of 2018 (Pub. L. 115–123) into law reauthorizing the RPG program through FFY 2021 and added a focus on opioid abuse.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–03520 Filed 2–17–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Office of Community Services Data Collection for the Low Income Household Water Assistance Program Reports (0970–0578)

AGENCY: Office of Community Services; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Community Services (OCS), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is requesting an extension of approval for an information request to collect data from Low Income Household Water Assistance Program (LIHWAP) grant recipients. This information collection was originally approved for 6 months as an emergency approval. OCS is proposing revisions to the information collection based on feedback received to date.

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: The LIHWAP Quarterly Performance and Management Report and the LIHWAP Annual Report provide ACF and Congress information necessary for oversight of recipients' performance in administering the LIHWAP program. The LIHWAP Quarterly Performance and Management Report solicits information on total households assisted, type of assistance provided, LIHWAP implementation information, performance management, and ongoing training/technical assistance needs. The LIHWAP Annual Report is modeled after the Low Income Home Energy Assistance Annual Report and has been streamlined to reduce recipient burden. The LIHWAP Annual Report collects data in three distinct modules: (1) Use of Funds, (2) Household Report, (3) Performance Measures.

This information collection package also includes a burden estimate related to the information collected from households. While grant recipients will collect necessary information from households using a variety of intake systems and local forms, OCS is providing technical assistance in this area and has included a sample application template in supplementary materials. This is a sample template; there will be no mandated household application format, and OCS will not receive or analyze copies of individual household application materials. OCS is proposing changes based on feedback received, including comments in response to a request for comments in the **Federal Register** (86 FR 59166). The currently approved versions of the LIHWAP Quarterly and Annual Reports and the sample application can be found here *https://www.reginfo.gov/*

public/do/PRAViewICR?ref_nbr=202110-0970-011. Updated materials can be found by following the directions to submit a comment in the **ADDRESSES** section of this notice.

Respondents: LIHWAP grant recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Annual burden hours
Quarterly Report for FY 2022	157	4	13	8,164
Quarterly Report for FY 2023	157	4	10	6,280
Annual Report	157	2	211	33,127
Household Application	1,200,000	1	.5	200,000

Estimated Total Annual Burden Hours: 241,291 (for FY 2022), 239,407 (for FY 2023).

Authority: Public Law 116–260 and LIHWAP Terms and Conditions Section 10 (*https://www.acf.hhs.gov/sites/default/files/documents/LIHWAP%20Terms%20and%20Conditions%20for%20States.pdf*).

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–03610 Filed 2–17–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–D–0893]

Agency Information Collection Activities; Proposed Collection; Comment Request; Center for Devices and Radiological Health Appeals Processes

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection associated with certain Center for Devices and Radiological Health (CDRH) appeals processes.

DATES: Submit either electronic or written comments on the collection of information by April 19, 2022.

ADDRESSES: 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 April 19, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of April 19, 2022. 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–2011–D–0893 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Center for Devices and Radiological Health Appeals Processes.'' 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