payment, specifying: The amounts withheld; and the dates that the withholding began and ended.

Paragraph (g) of FAR clause 52.232–27 requires contractors to issue a written notice of any withholding to a subcontractor (with copy to the contracting officer), specifying—

- The amount to be withheld;
- The specific causes for the withholding under the terms of the subcontract; and
- The remedial actions to be taken by the subcontractor in order to receive payment of the amounts withheld.

Paragraph (l) of FAR clause 52.232–27 requires contractors to remit overpayments to the payment office cited in the contract along with a description that includes the following:

- Circumstances of the overpayment (*e.g.,* duplicate payment, erroneous payment, liquidation errors, date(s) of overpayment);
- Affected contract number and delivery order number if applicable;
- Affected line item or subline item, if applicable; and
- Contractor point of contact.

Contractors are required to provide a copy of the remittance and supporting documentation to the contracting officer.

The information is used to understand when the contractor withholds amounts from subcontractors and suppliers after the Government has already paid the contractor the amounts withheld.

FAR 52.232–34, Payment by Electronic Funds Transfer—Other than System for Award Management. This clause requires contractors to provide the following information to enable the Government to make payments under the contract by EFT:

- The contract number (or other procurement identification number).
- The contractor's name and remittance address.
- The signature, title, and telephone number of the contractor official authorized to provide this information.
- The name, address, and 9-digit Routing Transit Number of the contractor's financial agent.
- The contractor's account number and the type of account.
- If applicable, the Fedwire Transfer System telegraphic abbreviation of the contractor's financial agent.
- If applicable, the contractor must provide the name, address, telegraphic abbreviation, and 9-digit Routing Transit Number of the correspondent financial institution receiving the wire transfer payment if the contractor's financial agent is not directly on-line to the Fedwire Transfer System.

The burden to provide the information required by the FAR clause at 52.232–33, Payment by Electronic Funds Transfer—System for Award Management, is covered by OMB Control Number 9000–0189, Certain Federal Acquisition Regulation Part 4 Requirements. OMB Control Number 9000–0189 accounts for new registrations and renewals in the System for Award Management, which includes providing the EFT information.

The information is used to enable the Government to make contract payments by EFT.

C. Annual Burden

Respondents: 275,319.

Total Annual Responses: 1,817,432.

Total Burden Hours: 471,947.

D. Public Comment

A 60-day notice was published in the **Federal Register** at 87 FR 1599, on March 21, 2022. Two comments were received; however, they did not change the estimate of the burden.

Comments: One of the comments is not related to the information collection. The other comment is a vendor's presentation of their products and services regarding payment solutions.

Response: The commenters did not express an opinion on whether the estimated number of burden hours is accurate; or ways to minimize the burden of the collection of information. The information collection revision does not reflect any changes to the FAR requirements. Adjustments are made to the public and Government burden estimates based on the most recent data available.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division, by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.* Please cite OMB Control No. 9000–0073, Certain Federal Acquisition Regulation Part 32 Requirements.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2022–11389 Filed 5–26–22; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Office of Refugee Resettlement Cash and Medical Assistance Program Quarterly Report on Expenditures and Obligations (ORR–2) (OMB #0970–0407)

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Refugee Resettlement (ORR), Administration for Children and Families (ACF) is requesting a 3-year extension of the form ORR–2, Cash and Medical Assistance Program Quarterly Report on Expenditures and Obligations (OMB #0970–0407, expiration 9/30/2022). There are no changes requested to the form.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing *infocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: ORR reimburses, to the extent of available appropriations, certain non-federal costs for the provision of cash and medical assistance (CMA) to refugees, along with allowable expenses for the administration of the refugee resettlement program at the state level. States and Replacement Designees currently submit the ORR–2 Quarterly Report on Expenditures and Obligations, which provides aggregate expenditure and obligation data. The ORR–2 collects expenditures and obligations data separately for each of the four following CMA program components: Refugee cash assistance, refugee medical assistance, CMA administration, and services for unaccompanied minors. This breakdown of financial status data allows ORR to track program expenditures in greater detail to anticipate any funding issues and to meet the requirements of ORR regulations at CFR 400.211 to collect these data for use in estimating future

costs of the refugee resettlement program. ORR must implement the methodology at CFR 400.211 each year after receipt of its annual appropriation to ensure that appropriated funds will be adequate for reimbursement to states of the costs for assistance provided to entering refugees. The estimating methodology prescribed in the regulations requires the use of actual past costs by program component. If the methodology indicates that appropriated funds are inadequate, ORR must take steps to reduce federal expenses, such as by limiting the number of months of eligibility for Refugee Cash Assistance and Refugee Medical Assistance. The ORR–2 is a single-page financial report that allows ORR to collect the necessary data to ensure that funds are adequate for the projected need and thereby meet the requirements of both the Refugee Act and ORR regulations.

Respondents: State governments and Replacement Designees.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Annual burden hours
ORR–2, Cash and Medical Assistance Program Quarterly Report on Expenditures and Obligations	66	4	1.5	396

Estimated Total Annual Burden Hours: 396.

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: 8 U.S.C. 1522 Sec. 412 and 8 U.S.C. 524 (Title IV), Sec. 414.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–11409 Filed 5–26–22; 8:45 am]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0557]

Agency Information Collection Activities; Proposed Collection; Comment Request; Postmarket Surveillance of Medical Devices

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 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 requirements for postmarket surveillance of medical devices.

DATES: Submit either electronic or written comments on the collection of information by July 26, 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 July 26, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of July 26, 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–2013–N–0557 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Postmarket Surveillance of Medical Devices.'' 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