

date. *Form Number*: CMS–10241 (OMB control number 0938–1041); *Frequency*: Monthly; *Affected Public*: Private sector (Business or other for-profits); *Number of Respondents*: 30,000; *Total Annual Responses*: 30,000; *Total Annual Hours*: 15,000. (For policy questions regarding this collection contact: Lisa Shochet at 410–786–5445).

Dated: May 15, 2018.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2018–10673 Filed 5–17–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10540]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by July 17, 2018.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically*. You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail*. You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10540 Quality Improvement Strategy Implementation Plan and Progress Form

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for

approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request*: Revision of a currently approved collection; *Title of Information Collection*: Quality Improvement Strategy Implementation Plan and Progress Form; *Use*: Section 1311(c)(1)(E) of the Affordable Care Act requires qualified health plans (QHPs) offered through an Exchange must implement a quality improvement strategy (QIS) as described in section 1311(g)(1). Section 1311(g)(3) of the Affordable Care Act specifies the guidelines under Section 1311(g)(2) shall require the periodic reporting to the applicable Exchange the activities that a qualified health plan has conducted to implement a strategy as described in section 1311(g)(1). CMS intends to have QHP issuers complete the QIS Plan and Reporting Template annually for initial certification and subsequent annual updates of progress in implementation of their strategy. The template will include topics to assess an issuer's compliance in creating a payment structure that provides increased reimbursement or other incentives to improve the health outcomes of plan enrollees, prevent hospital readmissions, improve patient safety and reduce medical errors, promote wellness and health, and reduce health and health care disparities, as described in Section 1311(g)(1) of the Affordable Care Act.

The Quality Improvement Strategy Plan and Reporting Template will allow: (1) The Department of Health & Human Services (HHS) to evaluate the compliance and adequacy of QHP issuers' quality improvement efforts, as required by Section 1311(c) of the Affordable Care Act, and (2) HHS will use the issuers' validated information to evaluate the issuers' quality improvement strategies for compliance with the requirements of Section 1311(g) of the Affordable Care Act. *Form Number*: CMS–10540 (OMB control number: 0938–1286); *Frequency*: Annually; *Affected Public*: Public sector (Individuals and Households); Private sector (Business or other for-profits and Not-for-profit institutions); *Number of Respondents*: 250 respondents; *Total Annual Responses*: 250 responses; *Total Annual Hours*: 12,000 hours. (For policy questions regarding this collection, contact Nidhi Singh Shah at 301–492–5110.)

Dated: May 15, 2018.
William N. Parham, III,
Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.
 [FR Doc. 2018-10672 Filed 5-17-18; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Preventing and Addressing Intimate Violence When Engaging Dads.
OMB No.: New Collection.
Description: The Administration for Children and Families (ACF), Office of Planning, Research, and Evaluation (OPRE) proposes to collect information

as part of the Preventing and Addressing Intimate Violence when Engaging Dads (PAIVED) study. Since 2006, the Healthy Marriage and Responsible Fatherhood (HMRF) initiative has funded programs that play a key role in helping the Office of Family Assistance (OFA) achieve its goals to foster economically secure households and communities for the well-being and long-term success of children and families. The purpose of the PAIVED study is to better understand the prevalence of intimate partner violence (IPV) experienced by the population of fathers served by Responsible Fatherhood (RF) programs, and the services that federally- and non-federally funded RF programs are providing to address and contribute to the prevention of IPV among its participants.

The proposed data collection will include whether IPV content is included

in RF programs, the types of activities they are using to address IPV, and related successes and challenges. Other collected data will include barriers to addressing IPV in RF programs, the relevance of addressing IPV with fathers, fathers' reactions to this programming, and what types of partnerships RF programs have with other agencies to address IPV. This information will be collected through interviews conducted over the phone and in-person with RF program staff and community partners. This information will be critical to inform future efforts to address and contribute to the prevention of IPV through RF programming.

Respondents: Responsible Fatherhood program staff (e.g., program directors and facilitators) and community partners.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
RF program/community partner screening and participant recruitment	50	1	1	50
RF program staff semi-structured interview	25	1	1.5	38
Community partner semi-structured interview	15	1	1.5	23

Estimated Total Annual Burden Hours: 111.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: OPREinfocollection@acf.hhs.gov.

OMB Comment: 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. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: OIRA_SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families.

Mary Jones,
ACF/OPRE Certifying Officer.
 [FR Doc. 2018-10656 Filed 5-17-18; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

U.S. National Authority for the WHO Global Code of Practice on the International Recruitment of Health Personnel; Notice of Public Meeting

Time and date: Monday, July 2, 2018, 2:00 p.m.–4:00 p.m. EST.

Place: Hubert H. Humphrey Building, Room 325A, 200 Independence Ave. SW, Washington, District of Columbia 20201.

Status: Open, but requiring RSVP to us.who.irhp@hhs.gov by Monday, June 25, 2018.

Purpose: The purpose of the World Health Organization (WHO) Global Code of Practice on International Recruitment of Health Personnel (Global Code) is “to establish and promote voluntary principles and practices for the ethical international recruitment of health personnel and to facilitate the strengthening of health systems.” The United States Government has designated the Office of Global Affairs (OGA) and the Health Resources and Services Administration (HRSA) as co-National Authorities to be the point of contact for implementation activities. The Global Code encourages WHO Member States to cooperate with all

relevant stakeholders in their implementation efforts. This meeting is intended to provide an update to all interested stakeholders on U.S. Global Code implementation efforts to date and to provide a forum for questions on activities related to implementation of the Global Code.

The meeting will be open to the public as indicated above, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify within their RSVP at least 10 business days prior to the meeting. Foreign nationals planning to attend the session in person will require additional paperwork for security clearance and that this clearance process requires a minimum of 10 business days.

RSVP: Due to security restrictions for entry into the HHS Humphrey Federal Building, we will need to receive RSVPs for this event. Please send your full name and organization to us.who.irhp@hhs.gov. If you are not a U.S. citizen, you must RSVP no later than Monday, June 18, 2018. Please note this in the subject line of your RSVP, and our office will contact you to gain additional biographical information for your