security, resulting from a suspected or confirmed breach.

j. To the Government Publishing Office (GPO), when *Login.gov* needs to mail a user an address confirmation form or if a user requests mailed notifications of account changes or of proofing attempts.

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**HISTORY:**

This notice modifies the routine use section of the system of records notice that is published in full at 82 FR 6552, January 19, 2017. The comments GSA received on that notice, and its responses to them, may be searched for and viewed on *regulations.gov* using Docket ID ''GSA–GSA–2017–0002''.

[FR Doc. 2017–16852 Filed 8–9–17; 8:45 am]

**BILLING CODE 6820–34–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

**[Document Identifier: CMS–10147]**

### Agency Information Collection Activities: Submission for OMB Review; 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 (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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by September 11, 2017.

**ADDRESSES:** When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 *OR*, Email: *OIRA_submission@omb.eop.gov*.

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' Web site address at Web site 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:** Under the Paperwork Reduction Act of 1995 (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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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. No comments were received in response to the 60-day comment period. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Medicare Prescription Drug Coverage and Your Rights; *Use:* Through the delivery of this standardized notice, Part D plan sponsors' network pharmacies are in the best position to inform enrollees (at the point of sale) about how to contact their Part D plan if their prescription cannot be filled and how to request an exception to the Part D plan's formulary. The notice restates certain rights and protections related to the enrollees Medicare prescription drug benefits, including the right to receive a written explanation from the drug plan about why a prescription drug is not covered. *Form Number:* CMS–10147 (OMB control number: 0938–0975); *Frequency:* Occasionally; *Affected Public:* Private sector (business or other for-profits); *Number of Respondents:* 62,000; *Total Annual Responses:* 40,100,000; *Total Annual Hours:* 668,066. (For policy questions regarding this collection contact Sabrina Sparkman at 410–786–3209.)

Dated: August 7, 2017.

**William N. Parham, III,**

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

[FR Doc. 2017–16892 Filed 8–9–17; 8:45 am]

**BILLING CODE 4120–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Proposed Information Collection Activity; Comment Request

*Title:* Personal Responsibility Education Program (PREP) Multi-Component Evaluation Extension.

*OMB No.:* 0970–0398.

*Description:* The Family and Youth Services Bureau (FYSB) and the Office of Planning, Research, Evaluation (OPRE) in the Administration for Children and Families (ACF) are requesting an extension without change of a currently approved information collection (OMB No. 0970–0398). The purpose of the extension is to complete the ongoing follow-up data collection for the Personal Responsibility Education Program (PREP) Multi-Component Evaluation, which was designed to document how PREP programs are designed and implemented in the field, collect performance measure data for PREP programs, and assess the effectiveness of selected PREP-funded programs.

The PREP Multi-Component Evaluation contains three components: A Design and Implementation Study, a Performance Analysis Study, and an Impact and In-Depth Implementation Study. Data collection related to the

Design and Implementation Study is complete; data collection related to the Performance Analysis Study will be complete in late summer 2017. This notice is specific to data collection activities for the Impact and In-Depth Implementation Study, which is being conducted in four sites. The proposed extension is necessary to complete ongoing follow-up data collection. The resulting data will be used in a rigorous program impact analysis to assess the effectiveness of each program in reducing teen sexual activity and associated risk behaviors.

*Respondents:* Youth participants who agreed to participate in the study upon enrollment in the four impact study sites.

ANNUAL BURDEN ESTIMATES

| Instrument | Total/annual number of respondents | Number of responses per respondents | Average burden hours per response | Total/annual burden hours |
|---|---|---|---|---|
| Second follow-up survey | 325 | 1 | .75 | 244 |

*Estimated Total/Annual Burden Hours:* 244

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded 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. Email address: *OPREinfocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

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.

**Mary Jones,**

*ACF/OPRE Reports Clearance Officer.*

[FR Doc. 2017–16843 Filed 8–9–17; 8:45 am]

**BILLING CODE 4184–37–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Proposed Information Collection Activity; Comment Request

*Title:* Variations in Implementation of Quality Interventions (VIQI) Project: Data Collection.

*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 Variations in Implementation of Quality Interventions (VIQI): Examining the Quality-Child Outcomes Relationship in Child Care and Early Education Project.

The VIQI Project will inform policymakers, practitioners, and stakeholders about effective ways to support the quality and effectiveness of early care and education (ECE) centers for promoting young children's learning and development. In partnership with ECE centers across the United States that serve young children with diverse economic backgrounds, the VIQI Project aims to (1) identify dimensions of quality within ECE settings that are key levers for promoting children's outcomes; (2) inform what levels of quality are necessary to successfully support children's developmental gains; (3) identify drivers that facilitate and inhibit successful implementation of interventions aimed at strengthening quality; and (4) understand how these relations vary across different ECE settings, staff, and children. To achieve these aims, the VIQI Project will include a year-long pilot study that will pilot up to three curricular and professional development models, followed by a year-long impact evaluation and process study that involve testing the effectiveness of two curricular and professional development models that aim to strengthen teacher practices, the quality of classroom processes, and children's outcomes. The study will include up to 189 community-based and Head Start ECE centers spread across seven different metropolitan areas in the United States.

To test the effectiveness of the curricular and professional development models, the VIQI project will consist of a 3- or 4-group experimental design in the pilot study and a 3-group experimental design in the impact evaluation and the process study in which the initial quality and other characteristics of ECE centers are measured. The centers then will be stratified based upon select information collected—by setting type (*e.g.,* Head Start and community-based ECE centers) and initial levels of quality—and randomly assigned to one of the intervention conditions where they will be offered curricular and professional development supports aimed at strengthening the quality of classroom and teacher practices, or to a business-as-usual comparison condition.

In the pilot study, 24 centers in one metropolitan area will participate in the VIQI Project. Information about center and staff characteristics and classroom and teacher practices will be collected (1) to stratify and randomly assign centers; (2) to describe how the different interventions are implemented and are experienced by centers and teachers; and (3) to document the treatment differentials across research conditions. The information will then be used to adjust and to refine the research design and measures that will be used in the impact evaluation and process study.

In the impact evaluation and process study, 165 centers in seven metropolitan areas will participate in the VIQI Project. Information about center and staff characteristics and classroom and teacher practices will be collected (1) to stratify and randomly assign centers; (2) to identify subgroups of interest; (3) to describe how the interventions are implemented and are experienced by