

evaluation findings and lessons learned to provide data-driven technical assistance to the grantees and other organizations delivering the National DPP lifestyle change program. The data and lessons learned from DP12–1212 were also used to inform decision-making and policy, including the development of the Centers for Medicare & Medicaid Services (CMS) Medicare Diabetes Prevention Program (MDPP). As of April 1, 2018, the MDPP Expanded Model provides coverage for the National DPP lifestyle change program for eligible Medicare beneficiaries.

Despite the fact that over 1800 CDC-recognized organizations in 50 states, the District of Columbia, Puerto Rico, and the Virgin Islands offer the National DPP lifestyle change program, there are still many geographic areas with few or no in-person delivery programs. In addition, some populations, including Medicare beneficiaries, men, African-Americans, Asian-Americans, Hispanics, American Indians, Alaska Natives, Pacific Islanders, and people with visual impairment or physical disabilities, are under-enrolled relative to their estimated numbers and disease burden. To address these gaps, CDC funded a new, five-year cooperative agreement with ten new national organizations in September 2017, “Scaling the National DPP in Underserved Areas” (DP17–1705). CDC funded ten national organizations with affiliate program delivery sites in at least three states, each to start new CDC-

recognized organizations in underserved areas and to enroll both general and priority populations in new or existing CDC-recognized organizations. The DP17–1705 grantees will work on activities designed to accomplish three main goals:

(1) Build the infrastructure in underserved areas necessary to deliver the National DPP lifestyle change program to the general population and to priority populations, including Medicare beneficiaries, men, African-Americans, Asian-Americans, Hispanics, American Indians, Alaska Natives, Pacific Islanders, and non-institutionalized people with visual or physical disabilities;

(2) Tailor and adapt the program to address the unique needs and challenges of the enrolled participants; and

(3) Provide participants with specialized support needed to successfully complete the program and achieve 5–7% weight loss. Through this new cooperative agreement, it is anticipated that enrollment, retention, and achievement of 5–7% weight loss goals for the targeted populations will increase.

At this time, CDC requests an additional three years of OMB approval to continue collecting information needed to evaluate the effectiveness of CDC’s funding for the new grantees. The data collection will allow CDC to continue to provide data-driven, tailored programmatic technical assistance to ensure continuous quality

improvement for each year of the cooperative agreement. A number of changes to the evaluation forms are proposed to ensure that reporting and evaluation requirements are consistent with the aims of the new cooperative agreement and reflect the lessons learned from the original funded national organizations and their affiliate delivery sites. Evaluation data elements have been added or modified accordingly. Also, the method of data collection has converted from using an Excel spreadsheet to a web-based data system to allow for real-time feedback and technical assistance.

The reporting burden of this collection of information is estimated to vary between two and four hours with an average of three hours per grantee response (decreased from 12 hours), and between four and six hours with an average of five hours per affiliate delivery site response (increased from an average of 45 minutes per response). These estimated burden hours include the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information by the grantee and affiliate delivery site to submit information directly to the web-based data system. The number of respondents will increase with the increased number of grantees. These changes result in a net increase of 368 annualized burden hours. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
National DPP Affiliate Delivery Sites	Evaluation Form for Sites	100	1	5	500
National DPP Grantees	Evaluation Form for Grantees	10	1	3	30
Total	530

Jeffrey M. Zirger,

Acting Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10391]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Department of Health and Human Services.

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 December 3, 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/Paperwork-ReductionActof1995/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) 786-1326.

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-10391 Methods for Assuring Access to Covered Medicaid Services Under 42 CFR 447.203 and 447.204

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. *Title of Information Collection:* Methods for Assuring Access to Covered Medicaid Services Under 42 CFR 447.203 and 447.204; *Type of Information Collection Request:*

Extension of a currently approved collection; *Use:* Current regulations at 42 CFR 447.203(b) require states to develop an access monitoring review plan (AMRP) that is updated at least every three years for: Primary care services, physician specialist services, behavioral health services, pre and post-natal obstetric services (including labor and delivery), and home health services. When states reduce rates for other Medicaid services, they must add those services to the AMRP and monitor the effects of the rate reductions for 3 years. If access issues are detected, a state must submit a corrective action plan to CMS within 90 days and work to address the issues within 12 months. Section 447.203(b)(7) requires that states have mechanisms to obtain ongoing beneficiary and provider feedback. A state is also required to maintain a record of data on public input and how the state responded to the input. Prior to submitting proposals to reduce or restructure Medicaid service payment rates, states must receive input from beneficiaries, providers, and other affected stakeholders on the extent of beneficiary access to the affected services.

The information is used by states to document that access to care is in compliance with section 1902(a)(30)(A) of the Social Security Act, to identify issues with access within a state's

Medicaid program, and to inform any necessary programmatic changes to address issues with access to care. CMS uses the information to make informed approval decisions on State plan amendments that propose to make Medicaid rate reductions or restructure payment rates and to provide the necessary information for CMS to monitor ongoing compliance with section 1902(a)(30)(A). Beneficiaries, providers and other affected stakeholders may use the information to raise access issues to state Medicaid agencies and work with agencies to address those issues. *Form Number:* CMS-10391 (OMB control number: 0938-1134); *Frequency:* Annually; *Affected Public:* State, Local, or Tribal Governments); *Number of Respondents:* 51; *Number of Responses:* 212; *Total Annual Hours:* 12,262. (For questions regarding this collection contact Jeremy Silanskis at 410-786-1592.)

Dated: September 28, 2018.

William N. Parham, III,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10008, CMS-R-234, and CMS-R-194]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Department of Health and Human Services.

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