

pricing information for each plan. It is an Excel workbook with multiple worksheets and special functions through which bidders present to CMS their plan pricing information. Bidders enter information, such as plan experience, projected enrollment, and risk profile, and the BPT calculates the plan premiums and other values that drive the bidding process. CMS maintains and updates each BPT file and releases new versions every April.

The BPT files may be downloaded from the Health Plan Management System website (or HPMS), which is a restricted-access website, so users must obtain approval from CMS before using it. From HPMS, the BPT files may be downloaded as part of the Plan Benefit Package (or PBPP) software, or they may be downloaded as stand-alone blank files. These files are made available to users on the first Monday of April every year and an HPMS memo is released announcing the software availability. Plan sponsors are required to upload the completed BPTs to HPMS by the first Monday in June each year.

MAOs and PDPs use the Bid Pricing Tool (BPT) software to develop their actuarial pricing bid. The information provided in the BPT is the basis for the plan's enrollee premiums and CMS payments for each contract year. The tool collects data such as medical expense development (from claims data and/or manual rating), administrative expenses, profit levels, and projected plan enrollment information. By statute, completed BPTs are due to CMS by the first Monday of June each year. *Form Number:* CMS-10142 (OMB control number: 0938-0944); *Frequency:* Yearly; *Affected Public:* Private Sector, Business or other for-profits and Not-for-profit institution; *Number of Respondents:* 555; *Total Annual Responses:* 4,995; *Total Annual Hours:* 149,850. (For policy questions regarding this collection contact Rachel Shevland at 410-786-3026.)

Dated: December 11, 2018.

**William N. Parham, III,**

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

[FR Doc. 2018-27104 Filed 12-13-18; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS-3070G-I and CMS-10692]

#### 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 February 12, 2019.

**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 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](mailto:Paperwork@cms.hhs.gov).

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

**FOR FURTHER INFORMATION CONTACT:** Reports Clearance Officer, 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-3070G-I ICF/IID Survey Report Form and Supporting Regulations  
CMS-10692 Home and Community Based Services (HCBS) Incident Management Survey

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:* Extension of a currently approved collection; *Title of Information Collection:* ICF/IID Survey Report Form and Supporting Regulations; *Use:* The information collected with forms 3070G-I is used to determine the level of compliance with Intermediate Care Facilities for Individuals with Intellectual Disabilities

(ICF/IID) CoPs necessary to participate in the Medicare/Medicaid program. Information needed to monitor the State's performance as well as the ICF/IID program in general, is available to CMS only through the use of information abstracted from the survey report form. The form serves as a coding worksheet designed to facilitate data entry and retrieval into the Automated Survey Processing Environment Suite (ASPEN) in the State and at the CMS regional offices. *Form Number:* CMS-3070G-I (OMB control number: 0938-0062); *Frequency:* Reporting—Yearly; *Affected Public:* Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 6,100; *Total Annual Responses:* 6,100; *Total Annual Hours:* 18,300. (For policy questions regarding this collection contact Melissa Rice at 410-786-3270.)

2. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Home and Community Based Services (HCBS) Incident Management Survey; *Use:* The Survey will be disseminated to all 51 state Medicaid agencies (including the District of Columbia) to assess incident management systems in 1915(c) waivers. States will be surveyed to identify methods and promising practices for identifying, reporting, tracking, and resolving incidents of abuse, neglect, and exploitation. The survey results will also be used to review the strengths and weaknesses of each state's incident management system and will inform guidance to help ensure compliance with sections 1902(a)(30)(A) and 1915(c)(2)(A) of the Social Security Act. *Form Number:* CMS-10692 (OMB control number: 0938-TBD); *Frequency:* Once and on occasion; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 102; *Total Annual Hours:* 153. (For policy questions regarding this collection contact Ryan Shannahan at 410-786-0295.)

Dated: December 11, 2018.

**William N. Parham, III,**  
 Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2018-27101 Filed 12-13-18; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Proposed Collection; 60-Day Comment Request; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (NIAID)**

**AGENCY:** National Institutes of Health, HHS.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Institute of Allergy and Infectious Diseases (NIAID) will publish periodic summaries of propose projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

**DATES:** Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

**FOR FURTHER INFORMATION CONTACT:** To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Ms. Dione Washington, Health Science Policy Analyst, Office of Strategic Planning, Initiative Development and Analysis, 5601 Fishers Lane, Rockville, Maryland 20892 or call non-toll-free number (240) 669-2100 or email your request, including your address to: [washingtondi@niaid.nih.gov](mailto:washingtondi@niaid.nih.gov). Formal requests for additional plans and instruments must be requested in writing.

**SUPPLEMENTARY INFORMATION:** Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the

proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

*Proposed Collection Title:* Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery, 0925-0668, Expiration Date 2/28/2019, EXTENSION, National Institute of Allergy and Infectious Diseases (NIAID).

*Need and Use of Information Collection:* There are no changes being requested for this submission. The proposed information collection activity provides a means to garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide information about the NIAID's customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the NIAID and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 2511.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Type of collection	Number of respondents	Annual frequency per response	Hours per response	Total hours
Customer satisfaction surveys .....	4000	1	30/60	2000
In-Depth Interviews (IDIs) or Small Discussion Groups .....	50	1	90/60	75