

GENERAL SERVICES ADMINISTRATION

[Notice MV–2020–02; Docket No. 2020–0002; Sequence No. 27]

Notice of GSA Live Webinar regarding GSA's Implementation of Section 889 of the FY 2019 National Defense Authorization Act (NDAA); Correction

AGENCY: Office of Governmentwide Policy (OGP), General Services Administration (GSA).

ACTION: Notice; correction.

SUMMARY: On July 22, 2020, GSA published a notice regarding the hosting of a live and recorded virtual webinar on August 12, 2020 at 1:00 p.m. Eastern Standard Time (EST). This notice is to list the correct website for the meeting registration.

FOR FURTHER INFORMATION CONTACT: Patricia Richardson at patricia.m.richardson@gsa.gov or Maria Swaby at 202–208–0291.

SUPPLEMENTARY INFORMATION:

Correction

In FR Doc. 2020–15846, published on July 22, 2020 at 85 FR 44302, make the following correction:

On page 44302, third column, in the **ADDRESSES** section, remove “HERE” and add “https://gsa.zoomgov.com/webinar/register/WN_hQ6tHTRDR-mMNnRRxJy22Q” in its place.

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2020–16242 Filed 7–24–20; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–116]

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 September 28, 2020.

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

FOR FURTHER INFORMATION CONTACT: William N. 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–116 Clinical Laboratory Improvement Amendments (CLIA) Application Form and Supporting Regulations

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:* Clinical Laboratory Improvement Amendments (CLIA) Application Form and Supporting Regulations; *Use:* Section 353 (b) of the Public Health Service Act specifies that the laboratory must submit an application in such form and manner as the Secretary shall prescribe that describes the characteristics of the laboratory and examinations and procedures performed by the laboratory. The application must be completed by entities performing laboratory's testing specimens for diagnostic or treatment purposes. This information is vital to the certification process. In this revision, the majority of changes were minor changes to the form and accompanying instructions to facilitate the completion and data entry of the form. We anticipate that the changes will not increase the time to complete the form. *Form Number:* CMS–116 (OMB control number: 0938–0581); *Frequency:* Biennially and Occasionally; *Affected Public:* Private Sector—Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 52,140; *Total Annual Responses:* 52,140; *Total Annual Hours:* 52,140. (For policy questions regarding this collection contact Kathleen Todd at 410–786–3385.)

Dated: July 22, 2020.

William N. Parham, III,

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

[FR Doc. 2020-16243 Filed 7-24-20; 8:45 am]

BILLING CODE 4120-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Application Requirements for the Low Income Home Energy Assistance Program (LIHEAP) Model Plan Application (OMB #0970-0075)

AGENCY: Office of Community Services, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Community Services (OCS), Administration for Children and Families (ACF), U.S. Department of Health and Human

Services (HHS), is requesting a 3-year extension of the form OCS-0024: Low Income Home Energy Assistance Program (LIHEAP) Model Plan Application (OMB #0970-0075, expiration 9/30/2020). There are no changes requested to the form.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing infocollection@acf.hhs.gov. Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation (OPRE), 330 C Street SW, Washington, DC 20201, Attn: ACF Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: States, including the

District of Columbia, tribes, tribal organizations, and U.S. territories applying for LIHEAP block grant funds must, prior to receiving federal funds, submit an annual application (Model Plan, ACF-122) that meets the LIHEAP statutory and regulatory requirements. In addition to the Model Plan, grantees are also required to complete the Mandatory Grant Application, SF-424—Mandatory, which is included as the first section of the Model Plan.

The LIHEAP Model Plan is an electronic form and is submitted to OCS/ACF through the On-Line Data Collection (OLDC) system within GrantSolutions, which is currently being used by all LIHEAP grantees to submit other required LIHEAP reporting forms. In order to reduce the reporting burden, all data entries from each grantee's prior year's submission of the Model Plan in OLDC are saved and re-populated into the form for the following fiscal year's application.

Respondents: States, the District of Columbia, U.S. territories, and tribal governments.

ANNUAL BURDEN ESTIMATES

Instrument	Total annual number of respondents	Total annual number of responses per respondent	Average burden hours per response	Total annual burden hours
LIHEAP Detailed Model Plan	210	1	.50	105

Estimated Total Annual Burden Hours: 105.

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: 42 U.S.C. 8621)

John M. Sweet Jr.,

ACF/OPRE Certifying Officer.

[FR Doc. 2020-16197 Filed 7-24-20; 8:45 am]

BILLING CODE 4184-80-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Allergy and Infectious Diseases Special Emphasis Panel Emergency Awards: Rapid Investigation of Severe Acute Respiratory

Syndrome Coronavirus 2 (SARS-CoV-2) and Coronavirus Disease 2019 (COVID-19) (R21/R01 Clinical Trial Not Allowed).

Date: August 13, 2020.

Time: 11:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3F52, Rockville, MD 20892, (Telephone Conference Call).

Contact Person: Margaret A. Morris Fears, Ph.D., Scientific Review Officer, Scientific Review Program, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3F52, Rockville, MD 20852, maggie.morrisfears@nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)