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.

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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:

Contonte

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 Notfor-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.)