SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning submission of a Professional Employee Compensation Plan.

DATES: Submit comments on or before August 18, 2015.

ADDRESSES: Submit comments identified by Information Collection 9000–0066, Professional Employee Compensation Plan by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.*

Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 9000–0066, Professional Employee Compensation Plan''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 9000–0066, Professional Employee Compensation Plan'' on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0066, Professional Employee Compensation Plan.

Instructions: Please submit comments only and cite Information Collection 9000–0066, Professional Employee Compensation Plan, in all correspondence related to this collection. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Mr. Edward Loeb, Procurement Analyst, Office of Acquisition Policy, GSA, 202–501–3775 or email *edward.loeb@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

FAR 22.1103 requires that all professional employees are compensated fairly and properly. Accordingly, FAR 52.222–46, Evaluation of Compensation for Professional Employees, is required to be inserted in solicitations for negotiated service contracts when the contract amount is expected to exceed $650,000 and the service to be provided will require meaningful numbers of professional employees. The purpose of the provision at FAR 52.222–46 is to require offerors to submit for evaluation a total compensation plan setting forth proposed salaries and fringe benefits for professional employees working on the contract. Plans indicating unrealistically low professional employees' compensation may be assessed adversely as one of the factors considered in making a contract award.

B. Annual Reporting and Recordkeeping Burden

Respondents: 12,921.

Responses per Respondent: 3.

Total Responses: 38,763.

Hours per Response: 1.333333.

Total Burden Hours: 51,684.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary; whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0066, Professional Employee Compensation Plan, in all correspondence.

Dated: June 15, 2015.

Edward Loeb,

Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2015–15132 Filed 6–18–15; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[Docket 2015–0076; Sequence 19; [OMB Control No. 9000–0080]

Federal Acquisition Regulation; Information Collection; Integrity of Unit Prices

AGENCIES: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice of request for public comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement concerning Integrity of Unit Prices.

DATES: Submit comments on or before August 18, 2015.

ADDRESSES: Submit comments identified by Information Collection 9000–0080, Integrity of Unit Prices by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.*

Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 9000–0080, Integrity of Unit Prices''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 9000–0080, Integrity of Unit Prices'' on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NE, Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0080, Integrity of Unit Prices.

Instructions: Please submit comments only and cite Information Collection 9000–0080, Integrity of Unit Prices, in all correspondence related to this collection. All comments received will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Mr. Edward Loeb, Procurement Analyst,

Office of Acquisition Policy, GSA, 202–501–0650 or email *edward.loeb@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The clause at FAR 52.215–14, Integrity of Unit Prices, requires offerors and contractors under Federal contracts that are to be awarded without adequate price competition to identify in their proposals those supplies which they will not manufacture or to which they will not contribute significant value. The policies included in the FAR are required by 41 U.S.C. 3503 (a)(1)(A)(for the civilian agencies) and 10.U.S.C 2306a(b)(1)(A)(i)(for DOD and NASA). The rule contains no reporting requirements on contracts below the simplified acquisition threshold, construction and architect-engineering services, utility services, service contracts where supplies are not required, commercial items, and contracts for petroleum products.

B. Annual Reporting Burden

Respondents: 950.
Responses per Respondent: 10.
Annual Responses: 9,500.
Hours per Response: 1.
Total Burden Hours: 9,500.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary; whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street, NW., Washington, DC, 20405, telephone 202501–4755. Please cite OMB Control No. 9000–0080, Integrity of Unit Prices.

Dated: June 15, 2015.

Edward Loeb,

Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2015–15131 Filed 6–18–15; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–15–15GE]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Improving the Impact of Laboratory Practice Guidelines: A New Paradigm for Metrics—Clinical and Laboratory Standards Institute—NEW —Center for Surveillance, Epidemiology and Laboratory Services (CSELS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) is funding three 5-year projects collectively entitled ''Improving the Impact of Laboratory Practice Guidelines: A New Paradigm for Metrics''. An ''LPG'' is defined as written recommendations for voluntary, standardized approaches for medical laboratory testing that takes into account processes for test selection, sample procurement and processing, analytical methods, and results reporting for effective diagnosis and management of disease and health conditions. LPGs may be disseminated to, and used by, laboratorians and clinicians to assist with test selection and test result interpretation. The overall purpose of these cooperative agreements is to increase the effectiveness of LPGs by defining measures and collecting information to inform better LPG creation, revision, dissemination, promotion, uptake, and impact on clinical testing and public health. The project will explore how these processes and their impediments and facilitators differ among various intended users of LPGs. Through this demonstration project, CDC seeks to understand how to customize LPG creation and promotion to better serve these intended users of LPGs. An important goal is to help organizations that sponsor the development of LPGs create a sustainable approach for continuous quality improvement to evaluate and improve an LPG's impact through better collection of information.

The CDC selected three organizations that currently create and disseminate LPGs to support activities under a cooperative agreement funding mechanism to improve the impact of their LPGs. The American Society for Microbiology (ASM), the Clinical and Laboratory Standards Institute (CLSI), and the College of American Pathologists (CAP), will each use their LPGs as models to better understand how to improve uptake and impact of these and future LPGs. Only the CLSI submission will be described in this notice.

Specifically, the CLSI project will address two LPGs that are important to clinical testing and have a high public health impact: *POCT12, Point-of-Care Blood Glucose Testing in Acute and Chronic Care Facilities* and *POCT13, Glucose Monitoring in Settings without Laboratory Support.* These LPGs provide guidance and recommendations for personnel monitoring patient glucose levels at sites that have access to a hospital laboratory and at locations, such as physician offices or nursing