collects and processes this report on behalf of all three agencies.

Request for Comment

Public comment is requested on all aspects of this notice. Comment is also specifically invited on:

a. Whether the information collection is necessary for the proper performance of the agencies' functions, including whether the information has practical utility;

b. The accuracy of the agencies' estimate of the burden of the information collection, including the validity of the methodology and assumptions used;

c. Ways to enhance the quality, utility, and clarity of the information to be collected;

d. Ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology; and

e. Estimates of capital or start up costs and costs of operation, maintenance, and purchase of services to provide information.

Comments submitted to the Board in response to this notice will be shared with the other agencies. All comments will become a matter of public record.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2024–04398 Filed 3–1–24; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0205; Docket No. 2023–0053; Sequence No. 11]

Submission for OMB Review; Implementation of Federal Acquisition Supply Chain Security Act (FASCSA) Orders

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

ACTION: Notice.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division has submitted to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement regarding implementation of Federal Acquisition Supply Chain Security Act (FASCSA) Orders.

DATES: Submit comments on or before April 3, 2024.

ADDRESSES: Written comments and recommendations for this information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Marissa Ryba, Procurement Analyst, at telephone 314–586–1280, or *marissa.ryba@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. OMB Control Number, Title, and Any Associated Form(s)

9000–0205, Implementation of Federal Acquisition Supply Chain Security Act (FASCSA) Orders.

B. Need and Uses

This clearance covers the information that offerors and contractors must submit to comply with the following FAR requirements:

a. FAR 52.204–29, Federal Acquisition Supply Chain Security Act Orders—Representation and Disclosures. This provision prohibits contractors from providing or using as part of the performance of the contract any covered article, or any products or services produced or provided by a source, if the covered article or the source is subject to an applicable FASCSA order identified in the clause at FAR 52.204–30(b)(1).

By submitting an offer, offerors are representing compliance with the prohibition. If an offeror cannot represent compliance with the prohibition, then the offeror must disclose the following information in accordance with 52.204–29(e):

(1) Name of the product or service provided to the Government;

(2) Name of the covered article or source subject to an FASCA order;

(3) If applicable, name of the vendor, including the Commercial and Government Entity code and unique entity identifier (if known), that supplied the covered article or the product or service to the Offeror;

(4) Brand;

(5) Model number (original equipment manufacturer number, manufacturer part number, or wholesaler number);

(6) Item description;

(7) Reason why the applicable covered article or the product or service is being provided;

b. FAR 52.204–30, Federal Acquisition Supply Chain Security Act Orders—Prohibition. This clause requires contractors to provide a report to the Government within 3 business days if the contractor identifies that the contractor or any-tier subcontractor, delivered or used a covered article or product or service subject to a FASCSA order. The report requires the following information:

(1) Contract number;

(2) Order number(s), if applicable;

(3) Name of the product or service provided to the Government;

(4) Name of the covered article or source subject to a FASCSA order;

(5) If applicable, name of the vendor, including the Commercial and Government Entity code and unique entity identifier (if known), that supplied the covered article or the product or service to the Contractor;

(6) Brand;

(7) Model number (original equipment manufacturer number, manufacturer part number, or wholesaler number);

(8) Item description; and

(9) Any readily available information about mitigation actions undertaken or recommended.

The contractor must also submit additional information within 10 days of submitting the first report identifying any further available information about mitigation actions undertaken or recommended. Additionally, the contractor shall describe the efforts it undertook to prevent submission and any additional efforts to prevent future submission of the covered article or the product or service produced or provided by a source subject to an applicable FASCSA order.

FAR provision 52.204–29. Information collected under will be by the government to determine whether to seek a waiver from a FASCSA order issued under the authority of the Federal Acquisition Supply Chain Security Act of 2018.

FAR clause 52.204–30 will Information collected will be used by the contracting officer working with the requirement activity to determine whether it is necessary to take further action and modify the contract.

C. Annual Burden

Respondents: 6,113.
Total Annual Responses: 1.
Total Burden Hours: 12,226.

D. Public Comment

A 60-day notice was published in the **Federal Register** at 88 FR 88923, on

December 26, 2023. No comments were received.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division, by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.* Please cite OMB Control No. 9000–0205, Implementation of Federal Acquisition Supply Chain Security Act (FASCSA) Orders.

Janet Fry,

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

[FR Doc. 2024–04505 Filed 3–1–24; 8:45 am]

BILLING CODE 6820–EP–P

UNITED STATES AGENCY FOR GLOBAL MEDIA

Fiscal Year (FY) 2022 Service Contract Inventory Report & FY 2023 Planned Analysis

AGENCY: United States Agency for Global Media.

ACTION: Notice.

SUMMARY: The United States Agency for Global Media (USAGM) announces the members of its FY 2022 Service Contract Inventory Report and FY 2023 Planned Analysis.

ADDRESSES: USAGM Office of Contracts, 330 Independence Ave. SW, Washington, DC 20237.

FOR FURTHER INFORMATION CONTACT: Khilena Adhin, Acquisition Policy Branch Chief, at *conpolicy@usagm.gov,* 202–920–2302.

SUPPLEMENTARY INFORMATION: In accordance with section 743 of division C of the Consolidated Appropriations Act of 2010, the U.S. Agency for Global Media (USAGM) is publishing this notice to advise the public of the availability of its FY 2022 Service Contract Inventory Report and FY 2023 Planned Analysis. They are available on the USAGM website, through the following link: *https://www.usagm.gov/our-work/strategy-and-results/strategic-priorities/research-reports/service-contract-inventory/.* The service contract inventory provides information on service contract actions over $25,000 made in FY 2022. The information is organized by function to show how contracted resources are distributed throughout the Agency. The inventory has been developed in accordance with guidance on service contract inventories issued on November 5, 2010 and on December 19, 2011 by the Office of Management and Budget, Office of Federal Procurement Policy (OFPP).

Dated: February 28, 2024.

Armanda Matthews,

Program Support Specialist, U.S. Agency for Global Media.

[FR Doc. 2024–04475 Filed 3–1–24; 8:45 am]

BILLING CODE 8610–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting of the Clinical Laboratory Improvement Advisory Committee

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC) announces the following meeting for the Clinical Laboratory Improvement Advisory Committee (CLIAC). This is a virtual meeting. It is open to the public, limited only by the number of webcast lines available. Time will be available for public comment, and the public is also welcome to submit written comments in advance of the meeting (see the public participation section below).

DATES: The meeting will be held on April 10, 2024, from 10 a.m. to 6 p.m., EDT.

ADDRESSES: This is a virtual meeting. Meeting times are tentative and subject to change. The confirmed meeting times, agenda items, and meeting materials, including instructions for accessing the live meeting broadcast, will be available on the CLIAC website at *https://www.cdc.gov/cliac.* Check the website on the day of the meeting for the web conference link.

FOR FURTHER INFORMATION CONTACT: Heather Stang, M.S., Senior Advisor for Clinical Laboratories, Division of Laboratory Systems, Center for Laboratory Systems and Response, Office of Laboratory Science and Safety, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop V24–3, Atlanta, Georgia 30329–4027. Telephone: (404) 498–2769; Email: *HStang@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: The Clinical Laboratory Improvement Advisory Committee (CLIAC) is charged with providing scientific and technical advice and guidance to the Secretary, Department of Health and Human Services; the Assistant Secretary for Health; the Director, Centers for Disease Control and Prevention (CDC); the Commissioner, Food and Drug Administration (FDA); and the Administrator, Centers for Medicare & Medicaid Services (CMS). The advice and guidance pertain to general issues related to improvement in clinical laboratory quality and laboratory medicine and specific questions related to possible revision of the Clinical Laboratory Improvement Amendments of 1988 (CLIA) standards. Examples include providing guidance on studies designed to improve quality, safety, effectiveness, efficiency, timeliness, equity, and patient-centeredness of laboratory services; revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards on medical and laboratory practice; and the modification of the standards and provision of non-regulatory guidelines to accommodate technological advances, such as new test methods, the electronic transmission of laboratory information, and mechanisms to improve the integration of public health and clinical laboratory practices.

Matters to be Considered: The agenda will include agency updates from CDC, CMS, and FDA. Presentations and CLIAC discussions will focus on the applicability of CLIA personnel requirements to preanalytic testing, the role of artificial intelligence and machine learning in the clinical laboratory, and the use of clinical standards to improve laboratory quality. Agenda items are subject to change as priorities dictate.

Public Participation

It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments pertinent to agenda items.

Oral Public Comment: Public comment periods for each agenda item are scheduled immediately prior to the Committee discussion period for that item. In general, each individual or group requesting to present an oral comment will be limited to a total time of five minutes (unless otherwise indicated). Speakers should email *CLIAC@cdc.gov* or notify the contact person above (see **FOR FURTHER INFORMATION CONTACT**) at least five business days prior to the meeting date.

Written Public Comment: CLIAC accepts written comments until the date of the meeting (unless otherwise stated). However, it is requested that comments be submitted at least five business days prior to the meeting date so that the