with the following Federal Acquisition Regulation (FAR) requirements:

FAR 52.212–5, Contract Terms and Conditions Required to Implement Statutes or Executive Orders—Commercial Products and Commercial Services. Paragraph (d) of this clause requires contractors to make available at its offices at all reasonable times the records, materials, and other evidence for examination, audit, or reproduction by the Comptroller General of the United States, or an authorized representative. As used in this clause, records include books, documents, accounting procedures and practices, and other data, regardless of type and regardless of form.

FAR 52.214–26, Audit and Records—Sealed Bidding. This clause requires contractors required to submit certified cost or pricing data in connection with the pricing of a modification under a contract to make all records available to the contracting officer, or its authorized representative, including computations and projections related to the proposal for the modification; the discussions conducted on the proposal(s), including those related to negotiating; pricing of the modification; or performance of the modification. This clause requires contractors to make all records available to the Comptroller General of the United States, or an authorized representative, in the case of pricing a modification. This clause allows the Comptroller General to interview any current employee regarding such transactions.

FAR 52.215–2, Audit and Records—Negotiation. This clause requires contractors to maintain records for cost-reimbursement, incentive, time-and-materials, labor-hour, or price redeterminable contracts, or any combination of these, for contracting officers, or an authorized representative, to examine and audit all records and other evidence sufficient to reflect properly all costs claimed to have been incurred or anticipated to be incurred directly or indirectly in performance of a contract. The right of examination includes inspection at all reasonable times of contractor's plants, or parts of them, engaged in performing the pertinent contract. Contractors required to submit certified cost or pricing data in connection with a pricing action under a contract must make all records available to the contracting officer, or its authorized representative, including computations and projections related to the proposal for the contract, subcontract, or modification; the discussions conducted on the proposal(s), including those related to negotiating; pricing of the contract, subcontract, or modification; or performance of the contract, subcontract or modification. Also, this clause requires contractors to make all records available to the Comptroller General of the United States, or an authorized representative, to examine any of the contractor's directly pertinent records involving transactions under the pertinent contract or subcontract. This clause allows the Comptroller General to interview any current employee regarding such transactions.

The information must be retained so that audits necessary for contract surveillance, verification of contract pricing, and reimbursement of contractor costs can be performed. This information collection does not require contractors to create or maintain any record that the contractor does not maintain in its ordinary course of business.

C. Annual Burden

Respondents: 19,033.
Total Annual Responses: 93,578.
Total Burden Hours: 93,578.
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. 9000–0034, Examination of Records by Comptroller General and Contract Audit, in all correspondence.

Janet Fry,

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

[FR Doc. 2022–27433 Filed 12–16–22; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0161; Docket No. 2022–0053; Sequence No. 23]

Information Collection; Reporting Purchases From Sources Outside the United States

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

ACTION: Notice and request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, and the Office of Management and Budget (OMB) regulations, DoD, GSA, and NASA invite the public to comment on an extension concerning reporting purchases from sources outside the United States. DoD, GSA, and NASA invite comments on: whether the proposed collection of information is necessary for the proper performance of the functions of Federal Government acquisitions, including whether the information will have practical utility; the accuracy of the estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including the use of automated collection techniques or other forms of information technology. OMB has approved this information collection for use through May 31, 2023. DoD, GSA, and NASA propose that OMB extend its approval for use for three additional years beyond the current expiration date.

DATES: DoD, GSA, and NASA will consider all comments received by February 17, 2023.

ADDRESSES: DoD, GSA, and NASA invite interested persons to submit comments on this collection through *https://www.regulations.gov* and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at 202–501–4755 or *GSARegSec@gsa.gov*.

Instructions: All items submitted must cite OMB Control No. 9000–0161, Reporting Purchases from Sources Outside the United States. Comments received generally will be posted without change to *https://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Zenaida Delgado, Procurement Analyst, at telephone 202–969–7207, or *zenaida.delgado@gsa.gov*.

SUPPLEMENTARY INFORMATION:

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

9000–0161, Reporting Purchases from Sources Outside the United States.

B. Need and Uses

This clearance covers the information that offerors must submit to comply

with the FAR provision 52.225–18, Place of Manufacture. This provision requires offerors of manufactured end products to indicate in response to a solicitation, by checking a box, whether the place of manufacture of the end products it expects to provide is predominantly manufactured in the United States or outside the United States. Contracting officers use the information as the basis for entry into the Federal Procurement Data System for further data on the rationale for purchasing foreign manufactured items. The data is necessary for analysis of the application of the Buy American statute and the trade agreements.

C. Annual Burden

Respondents: 50,106.
Total Annual Responses: 2,600,361.
Total Burden Hours: 26,004.
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. 9000–0161, Reporting Purchases from Sources Outside the United States, in all correspondence.

Janet Fry,

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

[FR Doc. 2022–27430 Filed 12–16–22; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

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, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. 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: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—RFA–OH–22–002, NIOSH Centers for Agricultural Safety and Health.

Date: March 7, 2023.

Time: 11:00 a.m.–6:00 p.m., EST.

Place: Video-Assisted Meeting.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Marilyn Ridenour, B.S.N., M.P.H., Scientific Review Officer, Office of Extramural Programs, National Institute for Occupational Safety and Health, CDC, 1095 Willowdale Road, Morgantown, West Virginia, 26505; Telephone: (304) 285–5879; Email: *MRidenour@cdc.gov.*

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2022–27454 Filed 12–16–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–3429–FN]

Medicare and Medicaid Programs: Application From the Center for Improvement in Healthcare Quality for Continued Approval of Its Hospital Accreditation Program

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice.

SUMMARY: This notice announces our decision to approve the Center for Improvement in Healthcare Quality (CIHQ) for continued recognition as a national accrediting organization for hospitals that wish to participate in the Medicare or Medicaid programs.

DATES: The decision announced in this notice is applicable January 1, 2023 through January 1, 2028.

FOR FURTHER INFORMATION CONTACT: Erin Imhoff, (410) 786–2337. Caecilia Blondiaux, (410) 786–2190.

SUPPLEMENTARY INFORMATION:

I. Background

Under the Medicare program, eligible beneficiaries may receive covered services from a hospital, provided certain requirements are met. Section 1861(e) of the Social Security Act (the Act) establishes statutory authority for the Secretary of the Department of Health and Human Services (Secretary) to set distinct criteria for facilities seeking designation as a hospital. Regulations concerning provider agreements are at 42 CFR part 489 and those pertaining to activities relating to the survey and certification of facilities are at 42 CFR part 488. The regulations at 42 CFR part 482 specify the minimum conditions of participation that a hospital must meet to participate in the Medicare program.

Generally, to enter into an agreement, a hospital must first be certified by a state survey agency (SA) as complying with the conditions or requirements set forth in part 482 of our regulations. Thereafter, the hospital is subject to regular surveys by a SA to determine whether it continues to meet these requirements.

Section 1865(a)(1) of the Act provides that, if a provider entity demonstrates through accreditation by a Centers for Medicare & Medicaid Services (CMS) approved national accrediting organization (AO) that all applicable Medicare requirements are met or exceeded, we will deem those provider entities as having met such requirements. Accreditation by an AO is voluntary and is not required for Medicare participation.

If an AO is recognized by the Secretary of the Department of Health and Human Services (the Secretary) as having standards for accreditation that meet or exceed Medicare requirements, any provider entity accredited by the national accrediting body's approved program would be deemed to meet the Medicare requirements. A national AO applying for approval of its accreditation program under part 488, subpart A, must provide CMS with reasonable assurance that the AO requires the accredited provider entities to meet requirements that are at least as stringent as the Medicare requirements. Our regulations concerning the approval of AOs are set forth at §§ 488.4, 488.5 and 488.5(e)(2)(i). The regulations at § 488.5(e)(2)(i) require AOs to reapply for continued approval of its accreditation program every 6 years or sooner, as determined by CMS.