establish, manage, and terminate user accounts limited to their own agency.

The GSA system administrator accounts are an additional level of security and management in that they oversee all partner agency accounts, including both designated partner agency account managers and agency users. The GSA system administrator accounts require additional tokens that meet multi-factor authentication standards in accordance with National Institute of Standards and Technology (NIST) standards. The controls assist in restricting access to authorized users who require it for official business purposes. Records in FDMS are maintained in a secure, password protected electronic system that utilizes security hardware and software to include multiple firewalls, active intrusion detection, encryption, identification and authentication of users.

RECORD ACCESS PROCEDURES:

Partner agency users can access and manage their user credentials through their designated partner agency account manager. If an access inquiry is not resolved by the designated partner agency account manager, the partner agency user may contact the GSA system manager listed above. Procedures for requesting access from GSA can be found at 41 CFR part 105–64.4.

CONTESTING RECORD PROCEDURES:

If partner agency users have questions or concerns about their account records, they can contact their designated partner agency account manager. If a question or concern is not resolved by the designated partner agency account manager, a partner agency user may contact the GSA system manager listed above. Procedures for contesting records stored by GSA can be found at 41 CFR part 105–64.4.

NOTIFICATION PROCEDURES:

If partner agency users wish to receive notice about their account records, they can contact their designated partner agency account manager. If not resolved by the designated partner agency account manager, the partner agency user may contact the GSA system manager listed above. Procedures for requesting notice of records stored by GSA can be found at 41 CFR part 105–64.4.

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

N/A.

[FR Doc. 2019–21885 Filed 10–7–19; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0032; Docket No. 2019–0003; Sequence No. 30]

Information Collection; Contractor Use of Interagency Fleet Management System Vehicles

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 section 3506(c)(2)(A) of 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 a revision and renewal concerning contractor use of Interagency Fleet Management System Vehicles. 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 January 31, 2020. 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 December 9, 2019.

ADDRESSES: DoD, GSA, and NASA invite interested persons to submit comments on this collection by either of the following methods:

- *Federal eRulemaking Portal:* This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. Go to *http://www.regulations.gov* and follow the instructions on the site.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405. ATTN: Lois Mandell/IC 9000–0032, Contractor Use of Interagency Fleet Management System Vehicles.

Instructions: All items submitted must cite Information Collection 9000–0032, Contractor Use of Interagency Fleet Management System Vehicles. Comments received generally will be posted without change to *http://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 (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Mr. Michael O. Jackson, Procurement Analyst, at telephone 202–208–4949, or email at *michaelo.jackson@gsa.gov.*

SUPPLEMENTARY INFORMATION:

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

9000–0032, Contractor Use of Interagency Fleet Management System Vehicles.

B. Needs and Uses

Federal Acquisition Regulation (FAR) 51.203 and the clause at FAR 52.251–2, Interagency Fleet Management System (IFMS) Vehicles and Related Services, are to be used in solicitations and contracts when a cost-reimbursement contract is contemplated and the contracting officer may authorize, if in the best interest of the Government, the contractor to use IFMS vehicles and related services. Before such an authorization, the contracting officer must have, among other requirements: (1) A written statement that the contractor will assume, without the right of reimbursement from the Government, the cost or expense of any use of the IFMS vehicles and services not related to the performance of the contract; (2) Evidence that the contractor has obtained motor vehicle liability insurance covering bodily injury and property damage, with limits of liability as required or approved by the agency, protecting the contractor and the Government against third-party claims arising from the ownership, maintenance, or use of an IFMS vehicle; and (3) Considered any recommendations of the contractor.

Authorized contractors shall submit requests for IFMS vehicles and related services in writing to the appropriate GSA point of contact in accordance with the FAR. Contractors' requests must include: (1) Two copies of the agency authorization; (2) The number of vehicles and related services required and period of use; (3) A list of employees who are authorized to request the vehicles or related services; (4) A listing of equipment authorized to be serviced; and (5) Billing instructions and address.

C. Annual Burden

Respondents: 132.
Total Annual Responses: 132.
Total Burden Hours: 132.
Obtaining Copies: 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. 9000–0032, Contractor Use of Interagency Fleet Management System Vehicles, in all correspondence.

Dated: October 2, 2019.

Janet Fry,

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

[FR Doc. 2019–21886 Filed 10–7–19; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Inventory for Poliovirus Containment: Minimizing Risk of Poliovirus Release From Laboratories in the United States; Availability

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

ACTION: Notice.

SUMMARY: The United States National Authority for Containment of Poliovirus (NAC), Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS), announces the availability of the National Inventory for Poliovirus Containment survey. This survey is designed to collect relevant laboratory inventory data to ensure facilities throughout the United States are in compliance with requirements established in the World Health Organization (WHO) Global Action Plan (GAPIII), as adapted for the WHO Region of the Americas. Per GAPIII, each country is required to complete a national inventory of poliovirus-containing materials, including poliovirus potentially infectious materials (PIM).

DATES: The deadline for completion of the survey is December 31, 2019.

FOR FURTHER INFORMATION CONTACT: Lia Haynes Smith, Director, National Authority for Containment of Poliovirus, Centers for Disease Control and Prevention, 1600 Clifton Road, NE, MS H21–6, Atlanta, GA 30329. Telephone: (404)718–5160.

SUPPLEMENTARY INFORMATION: The survey should be completed by laboratories, storage sites, or other facilities that test, extract, handle, or store biological samples from humans, experimentally infected animals, sewage, or environmental waters. The survey questions are intended to identify facilities that possess any materials that may contain poliovirus. The questions seek to distinguish between potentially infectious materials (PIM) containing wild poliovirus (WPV), circulating vaccine-derived poliovirus (cVDPV), and oral poliovirus vaccine (OPV). PIM includes historical domestic and international specimens, human respiratory secretions, fecal specimens and environmental samples collected for non-polio related work in a time and place where wild poliovirus (WPV) or vaccine-derived poliovirus (cVDPV) was circulating or where oral polio vaccine (OPV) was in use. A table of country-specific poliovirus data can be found at *http://polioeradication.org/wp-content/uploads/2018/11/PIM-Annex-2-16-Nov-18.pdf.* Additionally, PIM cultured in some common cell lines in order to isolate other viruses of interest may have unintentionally amplified poliovirus, so respiratory or enteric viral isolates obtained from PIM specimens using these cell lines are also considered PIM. With the release of the WHO PIM guidance in April 2018, nucleic acid extracted using a validated method and specimens that potentially contain only OPV (OPV PIM), are no longer subject to containment under WHO GAP III. However, they are still considered part of the U.S. inventory and should be reported.

For the purpose of this survey, PIM should be identified based on where and when the specimens were collected, not based on any test results.

If a facility intends to destroy any of the potentially infectious poliovirus material or infectious material it possesses, it must submit material destruction attestation to the NAC. The NAC will send this attestation form to the facility once the completed survey is received.

Although the U.S. no longer immunizes with OPV, poliovirus materials are still present within a limited number of U.S. facilities for public health and virologic research, as well as diagnostic and manufacturing-related purposes. In these essential facilities [poliovirus-essential facilities; PEFs], poliovirus materials will continue to be retained, post-eradication, to serve critical national and international functions. It is crucial that poliovirus materials are appropriately contained under strict biosafety and biosecurity handling and storage conditions to ensure that the virus is not released into the environment, either accidentally or intentionally, to cause outbreaks of the disease in susceptible populations. The risk from a poliovirus reintroduction can be minimized, in part, by ensuring that facilities retaining poliovirus are located in areas with high levels of vaccination coverage. The data collected from this survey will be used to identify facilities with poliovirus materials, to inform poliovirus immunization activities at PEFs including the potential need to immunize particular facility staff, and to identify vaccination coverage estimates for communities surrounding these facilities.

Survey Overview

An overview of the survey questions can be found at *https://www.cdc.gov/cpr/polioviruscontainment/00_docs/SurveyGuidance.pdf.* This overview document is provided to help facilities prepare their survey responses and is not intended to be completed as a paper-based format. The survey must be completed online.

Access to the survey, including appendices and other references, can be found at *https://www.cdc.gov/cpr/polioviruscontainment/NIPC.htm* The time needed to complete the online survey will vary depending on the complexity of a facility and the availability of needed information.

Paperwork Reduction Act

CDC has determined that the information collection activities conducted under this project are exempt from the requirements of the Paperwork Reduction Act (PRA) as they fall under the activities authorized under the National Childhood Vaccine Injury Act (NCVIA) at section 2102(a)(6)–(a)(7) of the Public Health Service Act (42 U.S.C. 300aa–2(a)(6)–(a)(7).