GSARegSec@gsa.gov. Please cite OMB Control No. 9000–0047, Place of Performance.

Janet Fry,

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

[FR Doc. 2024–07822 Filed 4–11–24; 8:45 am]

BILLING CODE 6820–EP–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0293; Docket No. 2024–0001; Sequence No. 5]

Information Collection; Reporting and Use of Information Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements

AGENCY: Office of Technology Strategy/Office of Government-wide Policy, General Services Administration (GSA).

ACTION: Notice and request for comments.

SUMMARY: Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division (MVCB) will be submitting to the Office of Management and Budget (OMB) a request to review and approve a revision and renewal of the currently approved information collection requirement on the reporting and use of information concerning integrity and performance of recipients of grants and cooperative agreements.

DATES: Submit comments on or before June 11, 2024.

ADDRESSES: Submit comments identified by Information Collection 3090–0293; Reporting and Use of Information Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements to *http://www.regulations.gov.*

Submit comments via the Federal eRulemaking portal by searching the OMB control number 3090–0293. Select the link ''Comment Now'' that corresponds with ''Information Collection 3090–0293, Reporting and Use of Information Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements. Follow the instructions provided on the screen. Please include your name, company name (if any), and ''Information Collection 3090–0293, Reporting and Use of Information Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements'' on your attached document. If your comment cannot be submitted using *regulations.gov,* call or email the points of contact in the **FOR FURTHER INFORMATION CONTACT** section of this document for alternate instructions.

Instructions: Please submit comments only and cite Information Collection 3090–0293, Reporting and Use of Information Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements, in all correspondence related to this collection. Comments received generally will be posted without change to *regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *regulations.gov,* approximately two-to-three business days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Ms. Salomeh Ghorbani, Director, IAE Outreach and Stakeholder Engagement Division, at 703–605–3467 or *IAE_Admin@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

This information collection requirement, OMB Control No. 3090–0293, currently titled ''Reporting and Use of Information Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements'' is necessary in order to comply with section 872 of the Duncan Hunter National Defense Authorization Act of 2009, Public Law 110–417, as amended by Public Law 111–212, hereafter referred to as ''the Act.'' The Duncan Hunter National Defense Authorization Act of 2009 (Pub. L. 110–417) was enacted on October 14, 2008. Section 872 of this Act required the development and maintenance of an information system that contains specific information on the integrity and performance of covered Federal agency contractors and grantees.

The Federal Awardee Performance and Integrity Information System (FAPIIS) was developed to address these requirements and has been superseded by the System for Award Management (SAM) at *SAM.gov.* SAM provides users access to integrity information from the FAPIIS reporting module in the Contractor Performance Assessment Reporting System (CPARS), proceedings information from the Entity Management section of the SAM database, and suspension/debarment information from the Exclusions section of SAM.

As required by 2 CFR part 200 Uniform Administrative Requirements, Cost Principles, and Audit Requirements for Federal Awards, effective January 1, 2016, Federal agencies are required to review and consider any information about the applicant that is in SAM before making any award in excess of the simplified acquisition threshold, defined in 41 U.S.C. 134, over the period of performance.

Non-Federal entities (NFEs) are required to disclose any information about criminal, civil, and administrative proceedings, and/or affirm that there is no new information to provide. This applies to NFEs that receive Federal awards (currently active grants, cooperative agreements, and procurement contracts), as required by 45 CFR part 75, Appendix XII, of the Uniform Guidance, for any period of time during the period of performance of an award/project.

B. Annual Reporting Burden

Proceedings Screening Question #1

Respondents: 19,152.
Responses per Respondent: 1.
Total annual responses: 19,152.
Hours per response: .1.
Total response burden hours: 1,915.

Proceedings Screening Question #2

Respondents: 141.
Responded per Respondent: 1.
Total annual responses: 141.
Hours per response: .1.
Total response burden hours: 14.

Proceedings Details

Respondents: 141.
Responses per respondent: 2.
Total annual responses: 282.
Hours per response: .5.
Total response burden hours: 141.

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 theinformation 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 appropriatetechnological collection techniques or other forms of information technology.

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. OMB Control No. 3090–0293, Reporting and Use of Information

Concerning Integrity and Performance of Recipients of Grants and Cooperative Agreements, in all correspondence.

Lois Mandell,

Director, Regulatory Secretariat Division, General Services Administration.

[FR Doc. 2024–07804 Filed 4–11–24; 8:45 am]

BILLING CODE 6820–WY–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–24–24EK; Docket No. CDC–2024–0026]

Proposed Data Collection Submitted for Public Comment and Recommendations

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

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other federal agencies the opportunity to comment on a proposed information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled *B. multivorans* Ice Machine Multistate Investigation. This is an outbreak investigation which aims to evaluate the associations between Burkholderia multivorans infections among hospitalized patients and potential exposures to nonsterile ice and water from ice machines to help inform measures to prevent ongoing transmission.

DATES: CDC must receive written comments on or before June 11, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0026 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (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. In addition, the PRA also requires federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected;

4. 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 submissions of responses; and

5. Assess information collection costs.

Proposed Project

B. multivorans Ice Machine Multistate Investigation—New—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC has been assisting state and local jurisdictions investigate clusters of *Burkholderia multivorans* infections among patients admitted across four hospitals in two non-contiguous states. The outbreak strain of the bacteria has been identified in environmental samples from ice machines. Molecular analysis has shown that the bacterial strain identified in ice machines is genetically highly similar to the patient isolates. Further investigation revealed that the same brand of ice machine and the same filters, descaling/cleaning, and sanitizing products were used by the four hospitals. Epidemiologic and laboratory evidence suggest the possibility of contaminated nonsterile ice and water from the same brand of ice machines as a common source of exposure.

Further investigation is needed to identify the scope of the outbreak and the source of the ice machine contamination. CDC has deemed it necessary to conduct a national call for cases requesting that public health authorities report cases and clusters of *B. multivorans.* A case report form (CRF) was developed by CDC to assist jurisdictions in this effort. Jurisdictions will gather information using this case report form to assist in determining epidemiologic characteristics and risk factors of patients with *B. multivorans* as well as potential source(s) of *B. multivorans,* including ice machines and ice machine-related products (*e.g.,* cleaning solutions).

The Centers for Disease Control and Prevention will share findings and recommendations with public health and healthcare partners to prevent further spread of *B. multivorans* infections; findings may also be shared with other relevant stakeholders and/or published in scientific journals to disseminate investigation outcomes. CDC requests OMB approval for an estimated 120 annual burden hours. There are no costs to respondents other than their time.