rate for moving purposes and the POA rate when a Government-furnished automobile is authorized.

DATES: *Applicability date:* This notice applies to travel and relocation performed on or after July 1, 2022 through December 31, 2022.

FOR FURTHER INFORMATION CONTACT: For clarification of content, please contact Ms. Cheryl D. McClain-Barnes, Program Analyst, Office of Government-wide Policy, Office of Asset and Transportation Management, at 202–208–4334, or by email at *travelpolicy@gsa.gov.* Please cite Notice of FTR Bulletin 22–06.

SUPPLEMENTARY INFORMATION: GSA is required by statute to set the mileage reimbursement rate for privately owned automobiles (POA) as the single standard mileage rate established by the IRS. On June 9, 2022, the IRS announced a midyear mileage rate adjustment to reflect the rising cost of fuel. Therefore, in line with the IRS, GSA adjusted the POV mileage reimbursement rates starting July 1, 2022 through the remainder of calendar year (CY) 2022.

FTR Bulletin 22–06 establishes and announces the newly adjusted CY 2022 POV mileage reimbursement rates for official temporary duty and relocation travel. This notice is the only notification to agencies, in addition to the changes posted on GSA's website at *https://gsa.gov/ftrbulletins* and *https://gsa.gov/mileage.*

Krystal J. Brumfield,

Associate Administrator, Office of Government-wide Policy.

[FR Doc. 2022–14264 Filed 7–1–22; 8:45 am]

BILLING CODE 6820–14–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0308; Docket No. 2022–0001; Sequence No. 5]

Submission for OMB Review; General Services Administration Acquisition Regulation (GSAR); Construction Contract Administration

AGENCY: Office of Acquisition Policy, General Services Administration (GSA).

ACTION: Notice and request for comments.

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 regarding OMB Control No. 3090–0308, Construction Contract Administration.

DATES: *Submit comments on or before:* August 4, 2022.

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: Mr. Marten Wallace, General Services Acquisition Policy Division, GSA, by phone at 202–286–5807 or by email at *marten.wallace@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The information collected is used by PBS to evaluate a contractor's proposals, negotiate contract modifications, evaluate a contractor's progress, and review payment requests during contract administration. The clause was previously GSAR 552.236–78 Shop Drawings, Coordination Drawings, and Schedules. The clause is simplified, including removing the requirement for a specific number of prints and copies of various submittals. This simplification will ease the compliance burden for the contractor during contract administration from the current state.

B. Annual Reporting Burden

Public reporting burden for GSAR 552.236–72 Submittals is estimated to average .25 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

The annual reporting burden is estimated as follows:

Respondents: 890.

Responses per respondent: 5.

Total annual responses: 4,452.

Preparation hours per response: .25.

Total response burden hours: 1,113.

C. Public Comments

A 60-day notice published in the **Federal Register** at 87 FR 24303 on April 25, 2022. No comments were received.

Obtaining Copies of Proposals: 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.*

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2022–14187 Filed 7–1–22; 8:45 am]

BILLING CODE 6820–61–P

GENERAL SERVICES ADMINISTRATION

[Notice–PBS–2022–02; Docket No. 2022–0002; Sequence No. 11]

Federal Management Regulation; Designation of a Federal Building

AGENCY: Public Buildings Service (PBS), General Services Administration.

ACTION: Notice of a bulletin.

SUMMARY: The attached bulletin announces the redesignation of a federal building.

DATES: This bulletin expires January 5, 2023. The building designation remains in effect until canceled or superseded by another bulletin.

FOR FURTHER INFORMATION CONTACT: General Services Administration, PBS, Office of Portfolio Management, Attn: Chandra Kelley, 77 Forsyth Street SW, Atlanta, GA 30303, at 404–562–2763, or by email at *chandra.kelley@gsa.gov.*

SUPPLEMENTARY INFORMATION: This bulletin announces the designation of a federal building. Public Law 117–103, dated March 15, 2022, designated the Federal Building located at 2005 University Boulevard in Tuscaloosa, Alabama, as the ''Richard Shelby Federal Building and Courthouse''.

Robin Carnahan,

Administrator of General Services.

[FR Doc. 2022–14238 Filed 7–1–22; 8:45 am]

BILLING CODE 6820–Y1–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Invitation to Manufacturers of Platforms for Nucleic Acid Amplification or Detection Suitable for Assay Development and Molecular Diagnostics for Detection of Agents That Cause Infectious Diseases

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

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS), is interested in obtaining information on available platforms for nucleic acid amplification or detection that meet criteria outlined below in the **SUPPLEMENTARY INFORMATION** section below.

DATES: Manufacturers are asked to contact CDC at the address below by August 19, 2022.

FOR FURTHER INFORMATION CONTACT: Laura Hughes-Baker, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H24–12, Atlanta, GA 30329–4027. Telephone: (404) 639–1402; Email: *eocevent521@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Background: Nucleic acid amplification or detection is used in many diagnostic tests. Rapid and accurate results that can specifically detect small amounts of pathogen material are essential to identifying and tracking diseases. The recent pandemic has demonstrated the need for tests that can be used in public health laboratories across the United States and internationally.

Many CDC laboratories across the agency use a particular diagnostic platform for nucleic acid detection. Because this current platform will be retired in the future, CDC is interested in hearing from manufacturers regarding the availability of current and potential platforms that could support CDC's overall diagnostics and surveillance.

Criteria: Ideally, the replacement platform should:

- Be suitable for research, surveillance, or assay development, and in vitro diagnostic purposes;
- Have Food and Drug Administration (FDA) clearance for diagnostic use or a research platform capable of obtaining FDA clearance;
- Be compatible with a 96 well format;
- Be compatible with diagnostic, surveillance, or characterization tests targeting a variety of pathogens; and
- Have software that allows for flexibility in analysis.

Manufactures who may have a platform that meets these criteria should submit information to CDC at *eocevent521@cdc.gov* or the address provided in the **FOR FURTHER INFORMATION** section above.

All information submitted to CDC will be kept confidential as allowed by relevant federal law, including the Freedom of Information Act (5 U.S.C. 552) and the Trade Secrets Act (18 U.S.C. 1905).

Disclaimer and Important Notes

This notice is for planning purposes; it does not constitute a formal announcement for comprehensive applications. In accordance with Federal Acquisition Regulation 48 CFR 15.201(e), responses to this notice are not offers and cannot be accepted by the Government to form a binding award. CDC will not provide reimbursement for costs incurred in responding to this notice.

Dated: June 29, 2022.

Angela K. Oliver,

Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2022–14211 Filed 7–1–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–22–1273; Docket No. CDC–2022–0080]

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 continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Pregnancy Risk Assessment Monitoring System (PRAMS). PRAMS is a surveillance project of the Centers for Disease Control and Prevention (CDC) and state health departments that collects jurisdiction-specific, population-based data on maternal attitudes and experiences before, during, and shortly after pregnancy.

DATES: CDC must receive written comments on or before September 6, 2022.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2022–0080 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: 404–639–7118; 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,