Type of Review: Extension of currently approved collection.

Affected Public: Businesses and other for-profit entities.

Estimated Annual Burden Hours: 7,880.

Estimated Annual Labor Costs: $429,838.

Estimated Annual Non-Labor Costs: de minimis.

Abstract: The Affiliate Marketing Rule, 16 CFR part 680 (Affiliate Marketing Rule or Rule) requires covered entities to provide consumers with notice and an opportunity to opt out of the use of certain information before sending marketing solicitations. The Rule has no recordkeeping or reporting requirements. On December 23, 2025, the Commission sought comment on the disclosure requirements associated with the Rule. 90 FR 60101. No relevant comments were received. For more details about the Rule requirements, the background behind these information collection provisions, and the basis for these calculations, see 90 FR 60101 (Dec. 23, 2025).

Pursuant to the OMB regulations, 5 CFR part 1320, that implement the PRA, 44 U.S.C. 3501 *et seq.,* the FTC is providing this second opportunity for public comment while seeking OMB approval to renew the pre-existing clearance for those information collection requirements.

Your comment—including your name and your state—will be placed on the public record of this proceeding. Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, such as anyone's Social Security number; date of birth; driver's license number or other state identification number, or foreign country equivalent; passport number; financial account number; or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, your comment should not include any ''trade secret or any commercial or financial information which . . . is privileged or confidential'' —as provided by Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2)—including in particular competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Josephine Liu,

Assistant General Counsel for Legal Counsel.

[FR Doc. 2026–06057 Filed 3–27–26; 8:45 am]

BILLING CODE 6750–01–P

OFFICE OF MANAGEMENT AND BUDGET

Office of Federal Procurement Policy

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0161; Docket No. 2025–0121; Sequence No. 1]

Submission for OMB Review; Reporting Purchases From Sources Outside the United States

AGENCY: Office of Federal Procurement Policy (OFPP), Office of Management and Budget (OMB); 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 OMB a request to review and approve an extension of a previously approved information collection requirement regarding reporting purchases from sources outside the United States.

DATES: Submit comments on or before April 29, 2026.

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: *FARPolicy@gsa.gov* or call 202–969–4075.

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: 23,134.
Total Annual Responses: 1,295,504.
Total Burden Hours: 12,955.

D. Public Comment

A 60-day notice was published in the **Federal Register** at 90 FR 59122, on December 18, 2025. A respondent provided several comments; however, they did not result in changes to the estimated burden.

Summary of Comments

The respondent supports the extension but expressed concern over the use, quality, and consistency of the collected information. The respondent recommended the following:

- Adopt standardized governmentwide collection of place-of-manufacture data.
- Improve processes by preventing unnecessary duplication in existing Buy American certification frameworks and providing clearer and more consistent definitions.
- Update burden estimates to reflect technological advancements and administrative efficiencies, recognizing the cumulative paperwork impact.

Response: The FAR Council acknowledges the comments received. Regarding the comments addressing duplication in existing Buy American certification frameworks, the respondent will have the opportunity to provide feedback when proposed changes to FAR Part 25 are published for comment as part of FAR Case 2026–004, Revolutionary Federal Acquisition Regulation Overhaul parts 19, 22, 23, and 25.

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.

Janet Fry,

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

[FR Doc. 2026–06072 Filed 3–27–26; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting of the Advisory Board on Radiation and Worker Health, Subcommittee for Procedure Reviews, National Institute for Occupational Safety and Health

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 Subcommittee on Procedures Reviews (SPR) of the Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board). This meeting is open to the public, but without a public comment period. The public is welcome to submit written comments in advance of the meeting, to the contact person below. The public is also welcome to listen to the meeting by joining the audio conference (information below). The audio conference line has 150 ports for callers.

DATES: The meeting will be held on June 5, 2026, from 11 a.m. to 4:30 p.m., EDT.

Written comments must be received on or before May 29, 2026.

ADDRESSES: You may submit comments by mail to: Rashaun Roberts, Ph.D., Designated Federal Officer, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1090 Tusculum Avenue, Mailstop C–24, Cincinnati, Ohio 45226. Email: *ocas@cdc.gov.*

Written comments received in advance of the meeting will be included in the official record of the meeting.

Meeting Information: Audio Conference Call via FTS Conferencing. The USA toll-free dial-in number is 1–866–659–0537; the passcode is 9933701.

FOR FURTHER INFORMATION CONTACT: Rashaun Roberts, Ph.D., Designated Federal Officer, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1090 Tusculum Avenue, Mailstop C–24, Cincinnati, Ohio 45226, Telephone: (513) 533–6800, Email: *ocas@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines, which have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction, which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC). In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to the CDC. NIOSH implements this responsibility for CDC.

The charter was issued on August 3, 2001, renewed at appropriate intervals, and rechartered under Executive Order 14109 (September 29, 2023) on March 22, 2024. Unless continued by the President, the Advisory Board will terminate on September 30, 2027, consistent with Executive Order 14354 of September 29, 2025.

Purpose: The Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179 ; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advising the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class. The ABRWH Subcommittee on Procedure Reviews (SPR) is responsible for overseeing, tracking, and participating in the reviews of all procedures used in the dose reconstruction process by the NIOSH Division of Compensation Analysis and Support (DCAS) and its dose reconstruction contractor (Oak Ridge Associated Universities—ORAU).

Matters to be Considered: The agenda will include discussions on the following:

1. Administrative items, including: a. SPR-approved documents for August Board meeting, b. SC&A revised PER protocol presentation, and c. Updated list of documents awaiting NIOSH responses; 2. Carry-over items from January 28, 2026, SPR Meeting including a. ORAUT–RPRT–0071, rev. 00 ''External Dose Coworker Methodology'' including: i. SC&A memo, ii. BRS update, iii. Closeout discussion, b. DCAS–PER–079 rev. 0 ''Pinellas TBD Revision'' presentation, c.DCAS–PER–036, rev. 0 ''Blockson TBD Revision'' presentation, d. DCAS–PER–039, rev. 0, ''Baker Perkins TBD Revision'' presentation; 3. NIOSH responses to SC&A reviews including: a. Battelle-TIB–5000, rev. 00 ''Default Assumptions and Methods for Atomic Weapons Employer Dose Reconstructions,'' and b. ORAUT–RPRT–0087, rev. 00, ''Applications of Regression in External Dose Reconstruction''; 4. SC&A issued reviews: a. ORAUT–OTIB–0092, rev. 00 ''Correction Factors for Neutron Dose Measured with Nuclear Track Emulsion, Type A Film'' presentation including: i. NIOSH response to OTIB–0092 observation, b. DCAS–PER–090, rev. 0 ''Grand Junction Operations Office'' presentation, c. ORAUT–OTIB–0034, rev. 04 ''Internal Dosimetry Co-Exposure Data for Oak Ridge National Laboratory'' presentation, d. DCAS–PER–069, rev. 0 ''Jessop Steel Company'' presentation, e. DCAS–PER–078, rev. 0 ''Extrusion Plant'' presentation, f. ORAUT–TKBS–0020, rev. 00 ''Technical Basis Document: Basis for the Development of an Exposure Matrix for Tennessee Valley Authority, Muscle Shoals, Alabama, Period of Operations: 1951–1955'' presentation, and g. ORAUT–TKBS–0016–4, rev. 02 ''Mound Plant—Occupational Environmental Dose'' presentation, and h. ORAUT–TKBS–0016–5, rev. 03 ''Mound Plant—Occupational Internal Dose'' presentation; 5. Newly-Issued Guidance and Supplemental Topics. Agenda items are subject to change as priorities dictate. For additional information, please contact Toll Free 1(800) 232–4636.

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to