B. Need and Uses

DoD, GSA, and NASA are combining OMB Control Nos. for the Federal Acquisition Regulation (FAR) by FAR part. This consolidation is expected to improve industry's ability to easily and efficiently identify burdens associated with a given FAR part. The review of the information collections by FAR part allows improved oversight to ensure there is no redundant or unaccounted for burden placed on industry. Lastly, combining information collections in a given FAR part is also expected to reduce the administrative burden associated with processing multiple information collections.

This justification supports the revision of OMB Control No. 9000–0077 and combines it with the previously approved information collections under OMB Control No. 9000–0187, with the new title ''Federal Acquisition Regulation Part 46 Requirements''. Upon approval of this consolidated information collection, OMB Control No. 9000–0187 will be discontinued. The burden requirements previously approved under the discontinued number will be covered under OMB Control No. 9000–0077.

This clearance covers the information that contractors may be required to submit to comply with the following FAR clauses:

- FAR Inspection Clauses
- 52.246–2, Inspection of Supplies—Fixed-Price
- 52.246–3, Inspection of Supplies—Cost-Reimbursement
- 52.246–4, Inspection of Services—Fixed-Price
- 52.246–5, Inspection of Services—Cost-Reimbursement
- 52.246–6, Inspection—Time-and-Material and Labor-Hour
- 52.246–7, Inspection of Research and Development—Fixed-Price
- 52.246–8, Inspection of Research and Development—Cost-Reimbursement
- 52.246–12, Inspection of Construction

These FAR clauses require the contractor to provide and maintain an inspection system that is acceptable to the Government, and to keep complete records of all inspection work performed and make it available to the Government. These clauses give the Government the right to inspect and test all work.

Records required under these clauses are kept as a part of a contractor's normal business operations. To ensure they provide a quality product or service, every business must have standards and methods for reviewing or inspecting the quality of their product or service. These standards will differ by industry and the complexity of the product or service provided.

The Government relies on a contractor's existing quality assurance system for contracts for commercial products. The Government relies on the contractor to accomplish all inspection and testing needed to ensure that acquired commercial services conform to contract requirements before they are tendered to the Government. See FAR 12.208 and 46.202–1. Likewise, when the contract amount is expected to be less than the simplified acquisition threshold, these clauses do not apply.

The FAR ''inspection clauses'' are used for quality assurance depending on the type of contract, or the product or service being provided. These clauses do not require the transmittal or sending of documentation to the Government, but they have record keeping requirements. The Government may review these records to confirm the contract quality requirements are being met. This review is risk-based and may or may not include the review of all quality assurance records. Generally, the records are more likely to be reviewed when the contractor is not meeting quality standards or as part of Government Contract quality assurance surveillance for complex requirements. Subject matter experts estimate these records are requested from 10 percent or fewer of contractors.

The information is used to assure that supplies and services provided under Government contracts conform to contract requirements.

- FAR 52.246–15, Certificate of Conformance. This clause requires the contractor to complete and sign a certificate of conformance (CoC). This clause is used in solicitations and contracts for supplies or services at the discretion of the contracting officer when it is in the Government's interest, small losses would be incurred in the event of a defect; or because of the contractor's reputation or past performance, or when it is likely that the supplies or services furnished will be acceptable and any defective work would be replaced, corrected, or repaired without contest.
- FAR 52.246–26, Reporting Nonconforming Items. This clause requires contractors to provide written notification to the contracting officer within 60 days of becoming aware or having reason to suspect, such as through inspection, testing, record review, or notification from another source (*e.g.,* seller, customer, third party) that any end item, component, subassembly, part, or material contained in supplies purchased by the contractor for delivery to, or for, the Government is counterfeit or suspect counterfeit. This clause requires certain contractors to submit a report to the Government-Industry Data Exchange Program (GIDEP) system at *www.gidep.org* within 60 days of becoming aware or having reason to suspect, such as through inspection, testing, record review, or notification from another source (*e.g.,* seller, customer, third party) that an item purchased by the contractor for delivery to, or for, the Government is a counterfeit or suspect counterfeit item; or a common item that has a major or critical nonconformance.

This information will be used by the Government to address and detect nonconforming and counterfeit items. Perhaps more important, this information will be available to businesses for searching prior to placing orders, thus enabling the avoidance of purchasing counterfeit items in the first place.

C. Annual Burden

Respondents: 7,859.
Total Annual Responses: 9,301.
Total Burden Hours: 33,015.
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. 9000–0077, Federal Acquisition Regulation Part 46 Requirements.

Janet Fry,

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

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BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[Docket No. ATSDR–2022–0005]

Proposed Substances To Be Evaluated for Toxicological Profile Development

AGENCY: Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Agency for Toxic Substances and Disease Registry (ATSDR) within the Department of Health and Human Services is initiating the development of another set of

Toxicological Profiles. This notice solicits public nominations of substances for ATSDR to evaluate for Toxicological Profile development.

DATES: All nominations, whether for substances on the Substance Priority List or for other substances, must be received by August 8, 2022.

ADDRESSES: You may submit nominations, identified by Docket No. ATSDR–2022–0005, by either of the following methods:

- Federal eRulemaking portal at *www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 4770 Buford Highway, Mail Stop S102–1, Atlanta, GA 30341–3717. Attn: Docket No. ATSDR–2022–0005.

Instructions: All submissions must include the agency name and docket number for this notice. All relevant comments will be posted without change to *http://www.regulations.gov,* including any personal information provided. Do not submit comments by email. ATSDR does not accept comments by email. This means that no confidential business information or other confidential information should be submitted in response to this notice. Refer to the Submission of Nominations section (below) for the specific information required to be included in a nomination.

FOR FURTHER INFORMATION CONTACT: Kambria Haire, Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 1600 Clifton Rd. NE, Mail Stop S102–1, Atlanta, GA 30329–4027; Email: *ATSDRToxProfileFRNs@cdc.gov;* Phone: 1–800–232–4636.

SUPPLEMENTARY INFORMATION: The Superfund Amendments and Reauthorization Act of 1986 (SARA) [42 U.S.C. 9601 *et seq.*] amended the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA or Superfund) [42 U.S.C. 9601 *et seq.*] by establishing certain requirements for ATSDR and the U.S. Environmental Protection Agency (EPA) concerning hazardous substances most commonly found at facilities on the CERCLA National Priorities List (NPL) (*https://www.epa.gov/superfund/superfund-national-priorities-list-npl*). Among these statutory requirements is a mandate for the Administrator of ATSDR to prepare Toxicological Profiles for each substance included on the Priority List of Hazardous Substances, also known as the Substance Priority list (SPL). This list identifies 275 hazardous substances found at NPL sites that ATSDR and EPA have determined currently pose the most significant potential threat to human health.

Substances To Be Evaluated for Toxicological Profile Development

Each year, ATSDR develops a list of substances to be considered for Toxicological Profile development. The nomination process includes consideration of all substances on ATSDR's SPL, as well as other substances nominated by the public. For more information on ATSDR's SPL, visit *https://www.atsdr.cdc.gov/SPL/.*

Submission of Nominations for Toxicological Profile Development

Today's notice invites voluntary public nominations of substances for toxicological profile development. If nominating a substance that is not on the SPL, please include the rationale for the nomination and any supporting data. ATSDR will evaluate data and information associated with nominated substances and will determine the final list of substances to be chosen for Toxicological Profile development. Substances will be chosen according to ATSDR's specific guidelines for selection. These guidelines can be found in the *Selection Criteria,* which may be accessed at *https://www.atsdr.cdc.gov/toxprofiles/guidance/ATSDR_TP_Selection%20Criteria.pdf.*

Pamela I. Protzel Berman,

Associate Director, Office of Policy, Planning and Partnerships, Agency for Toxic Substances and Disease Registry.

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BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2022–0014]

Draft Supplemental Environmental Impact Statement; Notice of Public Meeting and Comment Period

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) within the Department of Health and Human Services (HHS) announces the opening of a docket and public meeting to obtain comments on the Draft Supplemental Environmental Impact Statement (SEIS) for CDC's Roybal Campus in Atlanta, Georgia. The Draft SEIS was prepared to address changes proposed since completing the 2014 Final Environmental Impact Statement (EIS) for the CDC Roybal Campus 2025 Master Plan (2014 Final EIS) and issuing the Record of Decision dated November 7, 2014. The 2014 Final EIS analyzed the potential impacts associated with implementing a new long-range Master Plan to guide the future physical development of the Roybal Campus for the planning horizon of 2015 to 2025.

DATES: Written comments must be received on or before August 22, 2022.

A virtual public meeting will be held on July 27, 2022, from 6:00 p.m. EST to 8:00 p.m. EST. This meeting will occur via the Zoom platform.

Please register at *https://us06web.zoom.us/meeting/register/tZ0vduiqrT8oEtfyyzvqDUn_oUl5nS-LvfUE.*

Registration is required prior to the meeting. Once registered, you will receive an email with the meeting link and call-in number. The meeting will be recorded using the Zoom platform and a stenographer will transcribe the public meeting. The transcript will be posted on the Docket and included in the Final SEIS.

ADDRESSES: You may submit comments, identified by Docket Number CDC–2022–0014, by either of the following methods:

- *Federal eRulemaking Portal: https://www.regulations.gov/.* Follow the instructions for submitting comments.
- *U.S. Mail:* Thayra Riley, NEPA Coordinator, Office of Safety, Security, and Asset Management, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H20–4, Atlanta, Georgia 30329.

Instructions: All submissions received must include the Agency name and Docket Number (CDC–2022–0014). CDC will post, without change, all relevant comments to *https://www.regulations.gov,* including any personal information provided. *Do not submit comments by email. CDC does not accept comments by email.* For access to the docket to read background documents or comments received, go to *https://www.regulations.gov.* Oral comments on the Draft SEIS will also be accepted during the virtual public meeting scheduled for July 27, 2022.

FOR FURTHER INFORMATION CONTACT: Thayra Riley, NEPA Coordinator, Office of Safety, Security, and Asset Management, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H20–4, Atlanta, Georgia 30329. Email: *cdc-roybalga-*