prosecuting, enforcing, or carrying out a statute, rule, regulation, or order when GSA becomes aware of a violation or potential violation of civil or criminal law or regulation.

c. To an appeal, grievance, hearing, or complaints examiner; an equal employment opportunity investigator, arbitrator, or mediator; and an exclusive representative or other person authorized to investigate or settle a grievance, complaint, or appeal filed by an individual who is the subject of the record.

d. To the Office of Personnel Management (OPM), the Office of Management and Budget (OMB), and the Government Accountability Office (GAO) in accordance with their responsibilities for evaluating Federal programs.

e. To a Member of Congress or his or her staff on behalf of and at the request of the individual who is the subject of the record.

f. To an expert, consultant, or contractor of GSA in the performance of a Federal duty to which the information is relevant.

g. To the National Archives and Records Administration (NARA) for records management purposes.

h. In connection with any litigation or settlement discussions regarding claims by or against the GSA, including public filing with a court, to the extent that GSA determines the disclosure of the information is relevant and necessary to the litigation or discussions.

i. To appropriate agencies, entities, and persons when (1) GSA suspects or has confirmed that there has been a breach of the system of records, (2) GSA has determined that as a result of the suspected or confirmed breach there is a risk of harm to individuals, GSA (including its information systems, programs, and operations), the Federal Government, or national security; and (3) the disclosure made to such agencies, entities, and persons is reasonably necessary to assist in connection with GSA's efforts to respond to the suspected or confirmed breach or to prevent, minimize, or remedy such harm.

j. To another Federal agency or Federal entity, when GSA determines that information from this system of records is reasonably necessary to assist the recipient agency or entity in (1) responding to a suspected or confirmed breach or (2) preventing, minimizing, or remedying the risk of harm to individuals, the recipient agency or entity (including its information systems, programs, and operations), the Federal Government, or national security, resulting from a suspected or confirmed breach.

k. To compare such records to other agencies' systems of records or to non-Federal records, in coordination with an OIG in conducting an audit, investigation, inspection, evaluation, or some other review as authorized by the IG Act.

POLICIES AND PRACTICES FOR STORAGE OF RECORDS:

Electronic records are stored on a secure server with access limited to staff who may access the records only by means of a lawful government purpose. Information is encrypted in transit and at rest.

POLICIES AND PRACTICES FOR RETRIEVAL OF RECORDS:

Records are retrievable by a personal identifier or by other appropriate type of designation approved by GSA.

POLICIES AND PRACTICES FOR RETENTION AND DISPOSAL OF RECORDS:

Content in this system will be disposed according to the following GSA schedule:

121.1/050 Property Disposal Case Records. This series contains records related to the process of appraising federally-owned real property (both developed and undeveloped), and the disposal activities associated with closing, selling, destroying, transferring, or otherwise removing from the Federal Government's real property inventory. Such records include those used in the determination of excess real property, disposal of excess and surplus real property case files, correspondence, and related records.

Retention Instructions:

Permanent. Cut off at the end of the fiscal year following case completion and fulfillment of all restrictions on the disposed property. Transfer to NARA 15 years after cutoff.

Legal Disposition Authority: DAA–0121–2015–0001–0009 (121.1/050).

ADMINISTRATIVE, TECHNICAL, AND PHYSICAL SAFEGUARDS:

Records in the system are protected from unauthorized access and misuse through a combination of administrative and technical measures. Administrative measures include but are not limited to policies that limit system access to individuals within an agency with a legitimate business need, and regular review of security procedures and best practices to enhance security. Technical measures include but are not limited to system design that allows authorized system users access only to data for which they are responsible; AWS security tools; required use of *Login.gov* for application user account authorization and identity verification for real property bidders; required use of GSA SecureAuth for internal GSA users; and use of encryption for certain data transfers. Physical security measures are provided by the hosting service and ensure that unauthorized access to physical systems is not permitted.

RECORD ACCESS PROCEDURES:

If an individual wishes to access any data or record pertaining to him or her in the system after it has been submitted, that individual should consult the GSA's Privacy Act implementation rules available at 41 CFR part 105–64.2.

CONTESTING RECORD PROCEDURES:

If an individual wishes to contest the content of any record pertaining to him or her in the system after it has been submitted, that individual should consult the GSA's Privacy Act implementation rules available at 41 CFR part 105–64.4.

NOTIFICATION PROCEDURES:

If an individual wishes to be notified at his or her request if the system contains a record pertaining to him or her after it has been submitted, that individual should consult the GSA's Privacy Act implementation rules available at 41 CFR part 105–64.4.

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

None.

Richard Speidel,

Chief Privacy Officer, Office of Enterprise Data & Privacy Management, General Services Administration.

[FR Doc. 2024–25577 Filed 11–1–24; 8:45 am]

BILLING CODE 6820–AB–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[60Day–25–0048; Docket No. ATSDR–2024–0005]

Proposed Data Collection Submitted for Public Comment and Recommendations

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), 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 ATSDR Exposure Investigations (EIs). ATSDR EIs are deigned to fill data gaps by conducting environmental and biological sampling and to evaluate public health issues at a site resulting from environmental exposures.

DATES: ATSDR must receive written comments on or before January 3, 2025.

ADDRESSES: You may submit comments, identified by Docket No. ATSDR–2024–0005 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. ATSDR 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

ATSDR Exposure Investigations (EIs) (OMB Control No. 0923–0048, Exp. 6/30/2025)—Extension—Agency for Toxic Substances and Disease Registry (ATSDR).

Background and Brief Description

The Agency for Toxic Substances and Disease Registry (ATSDR) is requesting a three-year Extension of this Generic Clearance to allow the agency to conduct Exposure Investigations (EIs), through methods developed by ATSDR. After a chemical release or suspected release into the environment, EIs are usually requested by officials of a state health agency, county health departments, the Environmental Protection Agency (EPA), the general public, and ATSDR staff.

EI results are used by public health professionals, environmental risk managers, and other decision makers to determine if current conditions warrant intervention strategies to minimize or eliminate human exposure. For example, an EI that ATSDR previously conducted during included environmental sampling to evaluate non-drinking water exposure to per- and polyfluoroalkyl substances (PFAS) in two communities that were shown to be exposed to PFAS in their drinking water.

During the most recent clearance period (4/30/2022–present) a single Generic Exposure Investigation information collection request (ICR) was submitted. The EI conducted under this clearance period was an EI in Jasper and Newton Counties, Missouri to evaluate exposure to lead in a former mining community. ATSDR collected blood samples from community members most vulnerable to the impacts of lead exposure: children five years old and younger along with pregnant women and women of childbearing age. ATSDR partnered with the U.S. Environmental Protection Agency (EPA) and the Missouri Department of Health and Senior Services (MDHSS) who collected environmental samples, including soil, dust wipes, drinking water and paint, along with the results of the blood sampling. Appropriate EI procedures, including use of consent forms and questionnaires were used in the EI. The environmental sampling was submitted under this OMB Control Number with a burden of 426 hours.

All of ATSDR's targeted biological assessments (*e.g.,* urine, blood) and some of the environmental investigations (*e.g.,* air, water, soil, dust, or food sampling) involve participants to determine whether they are or have been exposed to unusual levels of pollutants at specific locations (*e.g.,* where people live, spend leisure time, or anywhere they might come into contact with contaminants under investigation). Questionnaires, appropriate to the specific contaminant, are generally needed in about half of the EIs (at most approximately 12 per year) to assist in interpreting the biological or environmental sampling results. ATSDR collects contact information (*e.g.,* name, address, phone number) to provide the participant with their individual results. ATSDR also collects information on other possible confounding sources of chemical(s) exposure such as medicines taken, foods eaten, hobbies, jobs, etc. In addition, ATSDR asks questions on recreational or occupational activities that could increase a participant's exposure potential. That information represents an individual's exposure history.

The number of questions can vary depending on the number of chemicals being investigated, the route of exposure (*e.g.,* breathing, eating, touching), and number of other sources of the chemical(s) (*e.g.,* products used, jobs). We use approximately 12–20 questions about the pertinent environmental exposures per investigation. A question bank is available for health assessors to use as a basis of questions to be asked during the EI, but EI-specific questions may be included as appropriate. Typically, the number of participants in an individual EI ranges from 10 to 100. Participation is completely voluntary, and there are no costs to participants

other than their time. Based on a maximum of 12 EIs per year and 100 participants each, the estimated annualized burden hours are 600.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Exposure Investigation Participants ..	Chemical Exposure Questions	1,200	1	30/60	600
Total	.....	.....	.....	.....	600

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2024–25554 Filed 11–1–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–25–1424; Docket No. CDC–2024–0087]

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 U.S. National Authority for Containment of Poliovirus Data Collection Tools. Data collection will capture information relating to a poliovirus containment breach or incident at a U.S. facility and will assist the U.S. National Authority for Containment of Poliovirus (U.S. NAC) in the initial stages of the investigation into the breach, and also gather information regarding personal protective equipment and practices used for work and/or storage of infectious materials or potentially infectious at laboratory facilities.

DATES: CDC must receive written comments on or before January 3, 2025.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0087 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

U.S. National Authority for Containment of Poliovirus Data Collection Tools (OMB Control No. 0920–1424, Exp. 12/31/2026)—Revision—Office of Readiness and Response (ORR), Centers for Disease Control and Prevention CDC)

Background and Brief Description

The role of the National Authority for Containment of Poliovirus (U.S. NAC) is to ensure that the requirements established in the World Health Organization (WHO) Global Action Plan (GAP) III/IV standard are effectively implemented and maintained in facilities working with or storing infectious poliovirus or potentially infectious materials.

Risk assessments following an incident are a critical component for adequate application of the GAP standard. To support risk assessment activities, The Facility Incident Reporting Form for Poliovirus Release and Potential Exposure and the Facility Incident Reporting Form for Poliovirus Theft or Loss was created for facilities to capture and submit incident