Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551.

You may also submit your Privacy Act request electronically by filling out the required information at: *https://foia.federalreserve.gov/.*

CONTESTING RECORD PROCEDURES:

The Privacy Act allows individuals to seek amendment of information that is erroneous, irrelevant, untimely, or incomplete and is maintained in a system of records that pertains to them. To request an amendment to your record, you should clearly mark the request as a ''Privacy Act Amendment Request.'' You have the burden of proof for demonstrating the appropriateness of the requested amendment and you must provide relevant and convincing evidence in support of your request.

Your request for amendment must: (1) provide the name of the specific Board system of records containing the record you seek to amend; (2) identify the specific portion of the record you seek to amend; (3) describe the nature of and reasons for each requested amendment; (4) explain why you believe the record is not accurate, relevant, timely, or complete; and (5) unless you have already done so in a related Privacy Act request for access or amendment, provide the necessary information to verify your identity.

NOTIFICATION PROCEDURES:

Same as ''Access procedures'' above. You may also follow this procedure in order to request an accounting of previous disclosures of records pertaining to you as provided for by 5 U.S.C. 552a(c).

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

Certain portions of this system of records may be exempt from 5 U.S.C. 552a(c)(3), (d), (e)(1), (e)(4)(G), (H), and (I), and (f) of the Privacy Act pursuant to 5 U.S.C. 552a(k)(5).

HISTORY:

This SORN was previously published in the **Federal Register** at 73 FR 24984 at 24997 (May 6, 2008). The SORN was also amended to incorporate two new routine uses required by OMB at 83 FR 43872 (August 28, 2018).

Board of Governors of the Federal Reserve System.

Benjamin W. McDonough,

Secretary of the Board.

[FR Doc. 2026–05156 Filed 3–16–26; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[Docket No. ATSDR–2026–0001]

Nominations for 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 (HHS), announces that it is soliciting nominations of substances to be evaluated for an upcoming set of toxicological profiles. ATSDR is opening a docket for the public to submit nominations and provide comments on which toxicological profiles are developed next. Members of the public, government agencies, or private organizations may comment on which substances they are concerned about so that ATSDR may take this information into consideration when developing future toxicological profiles.

DATES: Written comments must be received on or before May 18, 2026.

ADDRESSES: You may submit comments, identified by Docket No. ATSDR–2026–0001 by either of the methods listed below. Do not submit comments by email. ATSDR does not accept comments by email.

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Agency for Toxic Substances and Disease Registry, Office of Innovation and Analytics, 4770 Buford Highway, Mail Stop S106–5, Atlanta, GA, 30341–3717. Attn: Docket No. ATSDR–2026–0001.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov*.

FOR FURTHER INFORMATION CONTACT: Farhana Rahman, Agency for Toxic Substances and Disease Registry, Office of Innovation and Analytics, 4770 Buford Highway, Mail Stop S106–5, Atlanta, GA, 30341–3717; 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). 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 has determined currently pose the most significant potential threat to human health. For more information on ATSDR's SPL, visit *https://www.atsdr.cdc.gov/programs/substance-priority-list.html*.

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.

Submission of Nominations for Toxicological Profile Development

This notice invites 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.

Public Participation

Interested persons or organizations are invited to participate by submitting nominations for substances. These submissions may include written views and data to support the nomination. Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov*. Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If

you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. ATSDR will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign related to substances being nominated. Do not submit comments by email. ATSDR does not accept comments by email.

Donata Green,

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

[FR Doc. 2026–05164 Filed 3–16–26; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–26–0134]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled ''Land travel-related Public Health Activities'' to the Office of Management and Budget (OMB) for review and approval. CDC previously published a ''Proposed Data Collection Submitted for Public Comment and Recommendations'' notice on January 13, 2026 to obtain comments from the public and affected agencies. CDC received one comment related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) 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;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) 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 submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570. Comments and recommendations for the proposed 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 30-day Review—Open for Public Comments'' or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Land travel-related Public Health Activities (OMB Control No. 0920–0134, Exp. 3/31/2026)—Revision—National Center for Emerging and Zoonotic Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The goal of this information collection is to ensure that, consistent with the authorities in the Public Health Service Act and CFR parts 70 and 71, Centers for Disease Control and Prevention (CDC) is able to prevent the introduction, transmission and spread of communicable diseases from foreign countries into the United States or from one State or possession into any other State or possession.

This information collection focuses on collecting necessary information needed for CDC to conduct public health response and follow up in the event an individual with a confirmed or suspected communicable disease is known to have traveled via land conveyance (*e.g.,* bus, train, other) across an international land border or state borders while infectious or potentially infectious, presenting a risk of disease spread to others on the conveyance.

This information collection includes collection of conveyance, passenger and crew contact information from land conveyance operators (for example, via manifests) for contact investigations. Additionally, this information collection includes forms to obtain information on the outcomes of the contact investigations carried out by international, state, local, or territorial public health professionals to assess the impacts of CDC public health activities. With this current submission, CDC is requesting a Revision that will consolidate all land travel activity-related information collections under one OMB Control Number, thereby improving efficiency of CDC's land travel activities PRA submission process. In this Revision, CDC plans to keep one form in the previously approved version of 0920–0134, and add two that are currently approved under other OMB Control Numbers. CDC is also requesting to add four new information collection tools related to land travel that are based on similar tools for air travel. All other information collection tools currently approved in 0920–0134 are undergoing other requests to be moved into other approved OMB packages. With the aggregation of these land activity tools to simplify the PRA process, CDC is recommending revising the previous title from ''*Foreign Quarantine Regulations (42 CFR 71)*'' to ''*Land travel-related Public Health Activities.*''

CDC requests OMB approval for a total estimated 154 annual burden hours. There are no costs to respondents other than their time to participate.