the advisory committee's recommendations; (iii) The types of specific perspectives required, for example, such as those of consumers, technical experts, the public at-large, academia, business, or other sectors; (iv) The need to obtain divergent points of view on the issues before the advisory committee; and (v) The relevance of State, local, or tribal governments to the development of the advisory committee's recommendations.

*Member Selection Criteria*

The following selection criteria will be used to evaluate nominees.

**Committee Members**

a. Educational background (*e.g.,* degree in business, Information Technology, law, public policy, or engineering);

b. Professional experiences and accomplishments (*e.g.,* projects, nature of work, or publications);

c. Current employment and membership in associations or other activities (*e.g.,* manufacturers, academia, and civil society organizations); and

d. Subject matter expertise in the key issue the GAP FAC is examining for the current period.

e. Willingness to commit time to the Committee and demonstrated ability to work constructively and effectively on committees;

Members will serve one (1) to three (3) year terms.

*Miscellaneous*

The GAP FAC will meet approximately four times per year. Such meetings will be open to the public unless an appropriate authority determines, in accordance with the FACA, that a meeting shall be closed or partially closed. The Committee will meet virtually with the potential exception of one in person meeting per year.

Committee members (including the Committee Chair) will not be compensated for their services but will be allowed travel expenses, including per diem, in accordance with 5 U.S.C. 5703. Regardless of the type of committee membership appointment, any travel expenses shall be paid at rates equivalent to that allowable to Federal employees.

*Nomination Submissions*

Any interested person and/or organization may nominate qualified individuals for membership. Individuals are also encouraged to self-nominate. The following items must be submitted in a nomination package:

(1) A letter of nomination stating the nominee's name and organizational affiliation(s), nominee's field of expertise, specific qualifications to serve on the Committee, and a brief statement of interest;

(2) A professional resume or curriculum vitae (CV); and

(3) A short biography (no more than two paragraphs) describing the nominee's professional and educational qualifications, including a list of relevant activities and any current or previous service on advisory committees.

The letter of nomination, resume or CV, and a short biography should include the candidate's full name, address of the current organization, position title, email address, and daytime telephone number(s) of the nominee and nominator.

In preparing the letter of nomination, please describe how the nominee's background, knowledge, and experience will bring value to the work of the Committee and how these qualifications would contribute to the overall diversity of the Committee. Also, describe any previous involvement with GSA through employment, grant funding, and/or contracting sources, if applicable.

Nominations are due by August 15, 2024 and must be submitted via email to: *gapfac@gsa.gov.*

**Jeffrey A. Koses,**

*Senior Procurement Executive and Acting Chief Acquisition Officer, Office of Government-wide Policy, General Services Administration.*

[FR Doc. 2024–15596 Filed 7–15–24; 8:45 am]

**BILLING CODE 6820–61–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[30Day–24–0199]**

### 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 ''Import Permit Applications (42 CFR 71.54)'' 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 February 5, 2024 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**

Import Permit Applications (42 CFR 71.54) (OMB Control No. 0920–0199, Exp. 8/31/2024)—Revision—Office of Readiness and Response (ORR), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

Section 361 of the Public Health Service Act (42 U.S.C. 264), as amended, authorizes the Secretary of Health and Human Services to make and enforce such regulations as are

necessary to prevent the introduction, transmission, or spread of communicable diseases from foreign countries into the States or possessions, or from one State or possession into any other State or possession. Part 71 of title 42, Code of Federal Regulations (Foreign Quarantine) sets forth provisions to prevent the introduction, transmission, and spread of communicable disease from foreign countries into the United States. Subpart F—Importations—contains provisions for the importation of infectious biological agents, infectious substances, and vectors (42 CFR 71.54); requiring persons that import these materials to obtain a permit issued by the CDC.

The Application for Permit to Import Biological Agents, Infectious Substances and Vectors of Human Disease into the United States form is used by laboratory facilities, such as those operated by government agencies, universities, and research institutions to request a permit for the importation of biological agents, infectious substances, or vectors of human disease. This form currently requests applicant and sender contact information; description of material for importation; facility isolation and containment information; and personnel qualifications. Based on questions we have received from prior applicants, CDC is proposing to reduce open text questions and replace them with more streamlined check boxes. The goal is that this will clarify what is being asked of applicants and will increase efficiency and speed of processing by reducing back and forth communication necessary to clarify to applicants.

On February 5, 2024, CDC published in the **Federal Register** a 60-day notice (89 FR 7712) seeking public comments on ''Application for Permit to Import Biological Agents and Vectors of Human Disease into the United States,'' ''Application for Permit to Import or Transport Live Bats,'' ''Application for Permit to Import Infectious Human Remains into the United States,'' and ''Importer Certification Statement'' to initiate the revision of the information collection. As a result of this notice, CDC received one comment that was not related to the notice or CDC's Import Permit Program's regulatory authority, therefore, no changes to the data collection instruments were made.

The Application for Permit to Import or Transport Live Bats form is used by laboratory facilities such as those operated by government agencies, universities, research institutions, and for educational, exhibition, or scientific purposes to request a permit for the importation, and any subsequent distribution after importation, of live bats. This form currently requests the applicant and sender contact information; a description and intended use of bats to be imported; and facility isolation and containment information. CDC does not plan to revise this application.

The Application for Permit to Import Infectious Human Remains into the United States is used by facilities that will bury/cremate the imported cadaver and educational facilities to request a permit for the importation and subsequent transfers throughout the U.S. of human remains or body parts that contains biological agents, infectious substances, or vectors of human disease. This form will request applicant and sender contact information; facility processing human remains; cause of death; biosafety and containment information; and final destination(s) of imported infectious human remains. CDC does not plan to revise this application.

The Importer Certification Statement is a new form and will be used as an attestation by an importer stating that they are importing only noninfectious biological agent(s) or biological substance(s). The noninfectious, imported agent or substance must be accompanied by an importer certification statement confirming that the material is not known to contain or suspected of containing an infectious biological agent or has been rendered noninfectious. This form requests a detailed description of the material, statements affirming that the material is not known or suspected to contain an infectious biological agent, and one of the following: (1) How the person knows that the material does not contain an infectious biological agent; (2) Why there is no reason to suspect that the material contains an infectious biological agent; or (3) A detailed description of how the material was rendered noninfectious.

Annualized burden hours were calculated based on updated data obtained from the CDC import permit database on the number of permits issued on annual basis since 2021. There is an increase in burden from 764 hours to 2,044 hours which reflects the new, proposed form (Importer Certification Statement) and the increase in the number of respondents, to this project. There was no change due to program changes or adjustments.

ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondents | Form name | Number of respondents | Number of responses per respondent | Average burden per response (in hours) |
|---|---|---|---|---|
| Applicants Requesting to Import Biological Agents, Infectious Substances and Vectors. | Application for Permit to Import Biological Agents, Infectious Substances and Vectors of Human Disease into the United States. | 3,300 | 1 | 20/60 |
| Applicants Requesting to Import Biological Agents, Infectious Substances and Vectors. | Application for Permit to Import Biological Agents, Infectious Substances and Vectors of Human Disease into the United States Subsequent Transfer. | 650 | 1 | 10/60 |
| Applicants Requesting to Import Live Bats | Application for a Permit to Import Live Bats | 3 | 1 | 20/60 |
| Applicants Requesting to Import Infectious Human Remains into the United States. | Application for Permit to Import Infectious Human Remains into the United States. | 3 | 1 | 20/60 |
| Importer Attestation that the Imported Biological Agent or Substance is Noninfectious. | Importer Certification Statement | 5,000 | 1 | 10/60 |

**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. 2024–15535 Filed 7–15–24; 8:45 am]
**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[30Day–24–0493]**

### 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 ''2025 and 2027 National Youth Risk Behavior Survey (YRBS)'' 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 February 9, 2024 to obtain comments from the public and affected agencies. CDC received one non-substantive 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**

2025 and 2027 National Youth Risk Behavior Survey (OMB Control No. 0920–0493)—Reinstatement with Change—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

The purpose of this request is to obtain OMB approval for a Reinstatement with Change, of the data collection titled National Youth Risk Behavior Survey (YRBS), a school-based survey that has been conducted biennially since 1991. OMB approval for the 2021 YRBS and 2023 YRBS expired November 30, 2023 (OMB Control No. 0920–0493). CDC seeks a three-year approval to conduct the YRBS in Spring 2025 and Spring 2027. Changes incorporated into this Reinstatement request include the addition of a validation study of fruit and vegetable intake, the results of which will be used to inform changes to the 2027 YRBS questionnaire. Additional changes include an updated title for the information collection to accurately reflect the years in which the survey will be conducted and minor changes to the data collection instrument.

The YRBS assesses priority health risk behaviors related to the major preventable causes of mortality, morbidity, and social problems among both youth and young adults in the United States. Data on health risk behaviors of adolescents are the focus of approximately 65 national health objectives in Healthy People 2030, an initiative of the U.S. Department of Health and Human Services (HHS). The YRBS provides data to measure 14 Healthy People 2030 objectives. In addition, the YRBS can identify racial and ethnic disparities in health risk behaviors. No other national source of data measures as many of the Healthy People 2030 objectives addressing adolescent health risk behaviors as the YRBS. The data also will have significant implications for policy and program development for school health programs nationwide.

In Spring 2025 and Spring 2027, the YRBS will be conducted among nationally representative samples of students attending public and private schools in grades 9–12, and in 2025, the validation study will be conducted among a convenience sample of schools and students. Information supporting the YRBS also will be collected from state-, district-, and school-level administrators and teachers. The table below reports the number of respondents annualized over the three-year project period. The total estimated annualized burden hours are 4,389. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondent | Form name | Number of respondents | Number of responses per respondent | Average burden per response (in hr) |
|---|---|---|---|---|
| State Administrators | State-level Recruitment Script for the Youth Risk Behavior Survey. | 17 | 1 | 30/60 |
| District Administrators | District-level Recruitment Script for the Youth Risk Behavior Survey. | 80 | 1 | 30/60 |
| School Administrators | School-level Recruitment Script for the Youth Risk Behavior Survey. | 133 | 1 | 30/60 |
| School Administrators | School-level Recruitment Script for the Validation Study | 6 | 1 | 30/60 |