

on the basis of race, religion, color, national origin, age, disability, or sex. Nominees must be U.S. citizens and cannot be full-time employees of the U.S. Government. Current participation on federal workgroups or prior experience serving on a Federal advisory committee does not disqualify a candidate; however, HHS policy is to avoid excessive individual service on advisory committees and multiple committee memberships. Committee members are Special Government Employees, requiring the filing of financial disclosure reports at the beginning and annually during their terms. The Centers for Disease Control and Prevention (CDC) reviews potential candidates for LEPAC membership each year and provides a slate of nominees for consideration to the Secretary of HHS for final selection. HHS notifies selected candidates of their appointment as soon as the HHS selection process is completed. Note that the need for different expertise varies from year to year and a candidate who is not selected in one year may be reconsidered in a subsequent year.

Candidates should submit the following items:

- Current curriculum vitae, including complete contact information (telephone numbers, mailing address, email address).
- At least one letter of recommendation from person(s) not employed by HHS. Candidates may submit letter(s) from current HHS employees if they wish, but at least one letter must be submitted by a person not employed by an HHS agency (e.g., CDC, National Institutes of Health, Food and Drug Administration)

Nominations may be submitted by the candidate or by the person/organization recommending the candidate.

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 announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-25-1378]

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 “Assessing Respirator Perceptions, Experiences, and Maintenance” 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 July 18, 2025, to obtain comments from the public and affected agencies. CDC did not receive comments 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

Assessing Respirator Perceptions, Experiences, and Maintenance (OMB Control No. 0920-1378, Exp. 11/30/2025)—Extension—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC), National Institute for Occupational Safety and Health (NIOSH), is requesting approval of the Extension of a Generic Information Collection Request (ICR) for a period of three years under the project titled Assessing Respirator Perceptions, Experiences, and Maintenance. The National Personal Protective Technology Laboratory (NPPTL) is a division of NIOSH, which operates within the CDC. NPPTL was established in 2001, at the request of Congress, with the mission of preventing disease, injury, and death for the millions of working men and women relying on personal protective technology (PPT). As the nation’s respirator approver for all workplaces (42 CFR part 84), the development of NPPTL filled a need for improved personal protective equipment (PPE) and focused research into PPT. To this end, NPPTL conducts respiratory protection research to examine exposures to inhalation hazards, dermal hazards, and any other hazardous environmental threats within an occupational setting.

Federal regulations exist regarding the use of respirators in the workplace. The Occupational Safety and Health Administration (OSHA) requires employers whose hazard management includes the use of respirators to have a respiratory protection program, which has specified components. Thus, the information collected from human subjects about their use of respirators is generally consistent across NPPTL studies with only the use conditions changing (e.g., respirator type or management implementation practices related to cleaning/decontamination, fit testing, and training). NPPTL requests a Generic ICR package for information collected from individual workers and managers related to the perceptions,

maintenance, and evaluation of respirator use on the job.

Different types of data collection including surveys, focus groups, interviews, and physiological monitoring will be used to: (1) assess workers' health and safety knowledge, attitudes, skills, and other personal attributes as they relate to their respiratory protection use and maintenance; (2) identify and overcome barriers that workers face while using respiratory protection to prevent exposure to contaminants and other hazards; (3) understand organizations' maintenance of respiratory protection

programs (RPP), directives, and guidelines that support worker best practices; and (4) determine appropriate training, interventions, and programs that support activities around respirator use and maintenance. Data collection may focus on respirator types ubiquitous to the industry being studied, new to the industry being studied, or novel to any industry. These data collection efforts may occur either electronically or in the field.

Respondents are expected to include a variety of employees from occupations such as public safety and emergency response, healthcare, and social

assistance occupations who wear or manage respirator use on the job. Expected respondent job roles include industrial hygienists, occupational health professionals, infection control professionals, physicians, nurse practitioners, nurses, infection preventionists, fire department chiefs, battalion chiefs, sheriffs, shift supervisors, firefighters, police officers, and paramedics.

The total estimated burden hours are 643,626 with an estimated annual burden of 214,542 hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form Name	Number of respondents	Number of responses per respondent	Average Hours per response
Individuals who wear respirators in any occupational setting or oversee/advise on respirator use.	Informed consent	110,000	1	5/60
	Demographics standardized survey with decision logic allowing some questions to be omitted.	110,000	1	15/60
	Qualitative fit testing survey measurements	675	20	15/60
	Perceptions-based survey instrument	105,000	2	15/60
	Knowledge-based survey instrument	105,000	2	30/60
	Interview/Focus group	4,000	2	1
	Physiological Monitoring: Heart rate, blood pressure, blood oxygen saturation, breathing rate, etc.	1,000	1	9

Jeffrey M. Zirger,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–25–0213]

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 “National Vital Statistics Report Form” 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 June 16, 2025, to obtain comments from the public and affected agencies. CDC received three comments 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

National Vital Statistics Report Form (OMB Control No. 0920–0213)—Reinstatement—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The compilation of national vital statistics by the federal government dates to the beginning of the 20th century. To administer these functions, the National Office of Vital Statistics was established in the Public Health