

Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

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for Disease Control and Prevention.*

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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

[60Day-20-0109; Docket No. CDC-2020-0080]

**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 proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Respiratory Protective Devices—42 CFR part 84—Regulation. The purpose of the data collection is to enable 42 CFR part 84 respirator approval certification activities.

DATES: CDC must receive written comments on or before September 18, 2020.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2020-0080 by any of the following methods:

- *Federal eRulemaking Portal:* [Regulations.gov](https://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-D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to [Regulations.gov](https://www.regulations.gov).

Please note: Submit all comments through the Federal eRulemaking portal ([regulations.gov](https://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-D74, Atlanta, Georgia 30329; phone: 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; and
 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.
5. Assess information collection costs.

Proposed Project

Respiratory Protective Devices—42 CFR part 84—Regulation (OMB Control No. 0920-0109, Exp. 10/31/20)—Revision—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The regulatory authority for the National Institute for Occupational

Safety and Health (NIOSH) certification program for respiratory protective devices is found in the Mine Safety and Health Amendments Act of 1977 (30 U.S.C. 577a, 651 *et seq.*, and 657(g)) and the Occupational Safety and Health Act of 1970 (30 U.S.C. 3, 5, 7, 811, 842(h), 844). These regulations have, as their basis, the performance tests and criteria for approval of respirators used by millions of American construction workers, miners, painters, asbestos removal workers, fabric mill workers, and fire fighters.

Regulations of the Environmental Protection Agency (EPA) and the Nuclear Regulatory Commission (NRC) also require the use of NIOSH-approved respirators. These regulations also establish methods for respirator manufacturers to submit respirators for testing under the regulation and have them certified as NIOSH-approved if they meet the criteria given in the above regulation. This data collection was formerly named Respiratory Protective Devices 30 CFR part 11 but in 1995, the respirator standard was moved to 42 CFR part 84.

NIOSH, in accordance with 42 CFR part 84: (1) Issues certificates of approval for respirators which have met specified construction, performance, and protection requirements; (2) establishes procedures and requirements to be met in filing applications for approval; (3) specifies minimum requirements and methods to be employed by NIOSH and by applicants in conducting inspections, examinations, and tests to determine effectiveness of respirators; (4) establishes a schedule of fees to be charged applicants for testing and certification, and (5) establishes approval labeling requirements. Information is collected from those who request services under 42 CFR part 84 in order to properly establish the scope and intent of request.

Information collected from requests for respirator approval functions includes contact information and information about factors likely to affect respirator performance and use. Such information includes, but is not necessarily limited to, respirator design, manufacturing methods and materials, quality assurance plans and procedures, and user instruction and draft labels, as specified in the regulation.

The main instrument for data collection for respirator approval functions is the Standard Application Form for the Approval of Respirators (SAF), currently Version 9. Respirator manufacturers are the respondents (estimated to average 140 each year over the years 2020-2023) and upon

completion of the SAF, their requests for approval are evaluated. A total of 375 applications were submitted in CY2019. To date, 300 applications have been submitted in CY2020. The increased submission rate is due to the publication of a new respirator class, PAPR100, as well certification requests due to COVID-19. No survey was conducted to more thoroughly analyze the reasons for the change in number of respondents. The applications are submitted at will, and taking into account both historical conditions, as well as the current situation, our prediction of the number of respondents each year between CY2020 and CY2022 is 140. A \$200 fee is required for each application. Respondents requesting respirator approval or certain extensions of approval are required to submit

additional fees for necessary testing and evaluation as specified in 42 CFR parts 84.20-22, 84.66, 84.258 and 84.1102.

Applicants are required to provide test data that shows the manufacturer is capable of ensuring the respirator is capable of meeting the specified requirements in 42 CFR part 84. The requirement for submitted test data is likely to be satisfied by standard testing performed by the manufacturer, and is not required to follow the relevant NIOSH Standard Test Procedures. As additional testing is not required, providing proof that an adequate test has been performed is limited to providing existing paperwork.

42 CFR part 84 approvals offer corroboration that approved respirators are produced to certain quality standards. Although 42 CFR part 84 Subpart E prescribes certain quality

standards, it is not expected that requiring approved quality standards will impose an additional cost burden over similarly effective quality standards that are not approved under 42 CFR part 84.

Manufacturers with current approvals are subject to site audits by the Institute or its agents. Audits may occur periodically, typically every second year, or as a result of a reported issue. Sixty-four site audits from 90 respirator approval holders were scheduled for the 2020 fiscal year. There is an average fee of \$12,656 for each audit to align with fee collection provisions of the Independent Offices Appropriations Act of 1952 (31 U.S.C. 9701), and OMB Circular A-25 Revised. It is estimated that the average over the next three years (FY21-FY23) will be 70.

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)
Business or other for-profit	Standard Application Form for the Approval of Respirators.	140	4	229	128,240
Business or other for-profit	Audit	70	1	24	1,680
Total	129,920

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, 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

Advisory Committee on Breast Cancer in Young Women (ACBCYW); Notice of Charter Renewal

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of charter renewal.

SUMMARY: This gives notice under the Federal Advisory Committee Act of October 6, 1972, that the Advisory Committee on Breast Cancer in Young Women (ACBCYW), Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through June 17, 2022.

FOR FURTHER INFORMATION CONTACT: Jeremy McCallister, Designated Federal

Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway NE, Mailstop S107-4, Atlanta, Georgia 30341, Telephone (404) 639-7989, Fax (770) 488-4760; Email: acbcyw@cdc.gov.

The Director, Strategic Business Initiatives Unit, 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, Strategic Business Initiatives Unit, 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

[Docket No. CDC-2020-0083]

Advisory Committee on Immunization Practices (ACIP)

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting and request for comment.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC, announces the following meeting of the Advisory Committee on Immunization Practices (ACIP). This meeting is open to the public. Time will be available for public comment. The meeting will be webcast live via the World Wide Web.

DATES: The meeting will be held on August 26, 2020 from 10:00 a.m. to 4:00 p.m., EDT (times subject to change).

Written comments must be received on or before August 27, 2020.

ADDRESSES: For more information on ACIP please visit the ACIP website: