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: