OMB approval is requested for three years. Participation is based on previous Emergency Epidemic Investigations. There are no costs to respondents.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Total number of responses per respondent	Average burden per response (in hours)	Total burden hours (in hours)
Emergency Epidemic Investigation Participants.	Emergency Epidemic Investigation Data Collection Instruments.	12,000	1	30/60	6,000
Total					6,000

Jeffrey M. Zirger,

Acting Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2016–16882 Filed 7–15–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 81 FR 30307–30308, dated May 16, 2016) is amended to reflect the reorganization of the Office of the Director, Centers for Disease Control and Prevention.

Section C–B, Organization and Functions, is hereby amended as follows:

After the title and the mission and function statements for the *Office of the Associate Director for Laboratory Science and Safety (CAC)* insert the following:

Office of the Director (CAC1). (1) Provides scientific, technical, and managerial expertise and leadership in the development and enhancement of laboratory science and safety programs; (2) oversees and monitors the development, implementation, and evaluation of the laboratory safety and quality management programs across CDC; (3) advises on policy, partnerships, and issues management matters; (4) advises on matters related to internal and external public health communications; (5) provides oversight to ensure CDC compliance with regulations for select agents and toxins, and the safe possession, use and transfer of select agents and toxins; and (6) leads responses to laboratory incidents and emergencies.

Office of Laboratory Science (CACB). (1) Provides high-level oversight and coordination of laboratory quality and safety training programs at all CDC campuses; (2) develops agency-level plans, policies, procedures and guidelines for implementation of quality management programs within Centers, Institute, and Offices (CIOs); (3) assures regulatory compliance and tracking for CDC's portfolio of laboratory developed tests; and (4) provides oversight of the catalog of laboratory safety training activities and tracking agency-wide progress and compliance with laboratory safety training requirements.

Office of Laboratory Safety (CACC). (1) Provides high-level oversight and coordination of laboratory safety at all CDC campuses; (2) develops and assures effectiveness of agency-level plans, policies, manuals and tools for implementation of laboratory safety standards; (3) assures regulatory compliance for biological safety, chemical safety, radiation safety and the possession, use and transport of select agents and toxins; and (4) provides expertise and consultation for biological safety, chemical safety and radiation safety.

Sherri Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2016–16884 Filed 7–15–16; 8:45 am]

BILLING CODE 4160–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day–16–16AVB]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (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 or send an email to *omb@cdc.gov.* Written comments and/or suggestions regarding the items contained in this notice