

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)	Total burden (in hr.)
Adult/Adolescent Facility Users	Eligibility Screening Script	110	1	5/60	9
	Adult and Adolescent Questionnaire	100	1	30/60	50
	Exposure Measurement Form	100	1	20/60	33
Parents/Guardians of Youth/Child Facility Users.	Eligibility Screening Script	110	1	5/60	9
	Youth and Child Questionnaire	100	1	30/60	50
Youth/Child Facility Users	Exposure Measurement Form	100	1	20/60	33
Total	184

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2021-02949 Filed 2-11-21; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control; Correction

Notice is hereby given of a change in the meeting of the Board of Scientific Counselors, National Center for Injury Prevention and Control; February 16, 2021, 10:00 a.m. to 4:15 p.m., EST which was published in the **Federal Register** on January 8, 2021, Volume 86, Number 5, page 1502.

The dates and addresses should read as follows:

DATES: The meeting will be held on February 16, 2021, from 10:00 a.m.–4:30 p.m., EST.

ADDRESSES: Zoom Virtual Meeting. If you would like to attend the virtual meeting, please pre-register by accessing the link at https://dceproductions.zoom.us/webinar/register/WN_AQ70-aWpTqKvPX9Ftap-UA. Instructions to access the Zoom virtual meeting will be provided in the link following your registration.

Meeting Information: There will be a public comment period at the end of the meeting; from 3:45 p.m.–4:15 p.m. The public is encouraged to register to provide public comment using the registration form available at the link provided: <https://www.surveymonkey.com/r/cbyh878>.

Individuals registered to provide public comment will be called upon first to speak based on the order of registration, followed by others from the public. All public comments will be limited to two (2) minutes per speaker.

Written comments may also be submitted for the meeting record and must be received on or before February 23, 2021; ncipcbssc@cdc.gov.

FOR FURTHER INFORMATION CONTACT:

Gwendolyn H. Cattledge, Ph.D., M.S.E.H., Deputy Associate Director for Science, NCIPC, CDC, 4770 Buford Highway NE, Mailstop S-106-9, Atlanta, Georgia 30341, Telephone: (770) 488-1430, Email: ncipcbssc@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

[30Day-21-1071]

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 Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery 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 October 21, 2020 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

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery (OMB Control No. 0920–1071, Exp. 2/28/2021)—Extension—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

CDC/NCEZID is seeking a three-year extension of OMB control No. 0920–1071 to continue collecting routine customer feedback on agency service delivery. Executive Order 12862 directs Federal agencies to provide service to the public that matches or exceeds the best service available in the private sector. In order to work continuously to

ensure that our programs are effective and meet our customers' needs, the National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention (CDC) (hereafter the "Agency") seeks to obtain OMB approval of a generic clearance to collect qualitative feedback on our service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study.

This collection of information is necessary to enable the Agency to garner customer and stakeholder feedback in an efficient, timely manner, in accordance with our commitment to improving service delivery. The information collected from our customers and stakeholders will help ensure that users have an effective, efficient, and satisfying experience with the Agency's programs. This feedback

will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Since getting approval in February 2018, NCEZID has utilized 0920–1071 ten separate times. The total number of responses was 15,585. The total number of burden hours was 2,525. Authorizing legislation for this collection comes from Section 301 of the Public Health Service Act (42 U.S.C. 241). The estimated annual burden hours requested for this Extension are 3,850. There is no cost to respondents other than the time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
General public	Online surveys	1500	1	30/60
	Focus groups	800	1	2
	In-person surveys	1000	1	30/60
	Usability testing	1500	1	30/60
	Customer comment cards	1000	1	15/60

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

[60Day–21–21CT; Docket No. CDC–2021–0006]

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 request for emergency clearance of the information collection titled Requirement for Negative Pre-Departure Covid–19 Test Result or Documentation of Recovery from Covid–19 for all Airline or other Aircraft Passengers arriving into the United States from any foreign country. This collection accompanies a CDC Order of the same name and is designed to prohibit the introduction into the United States of any airline passenger departing from the any foreign country unless the passenger:

(1) Has a negative pre-departure test result for COVID–19 (Qualifying Test), or (2) has written or electronic documentation of recovery from COVID–19 in the form of a positive viral test result and a letter from a licensed health care provider or public health

official stating that the passenger has been cleared for travel.

DATES: CDC must receive written comments on or before April 13, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2021–0006 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