

in detail the information that the EPA will be collecting, are available in the public docket for this ICR. The docket can be viewed online at www.regulations.gov or in person at the EPA Docket Center, WJC West, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The telephone number for the Docket Center is 202-566-1744. For additional information about EPA's public docket, visit <http://www.epa.gov/dockets>.

Abstract: The information is collected from applicants and recipients of EPA assistance to monitor adherence to the programmatic and administrative requirements of the Agency's financial assistance program, which includes the Agency's DBE program. The information collected is used to make awards, pay recipients, and collect information on how Federal funds are being spent. EPA needs this information to meet its Federal stewardship responsibilities. This ICR renewal requests authorization for the collection of information under EPA's General Regulation for Assistance Programs, which establishes minimum management requirements for all recipients of EPA grants or cooperative agreements (assistance agreements). This ICR combines all of these requirements under OMB Control Number 2030-0020. The information required by these regulations will be used by EPA award officials to make assistance awards and assistance payments and to verify that the recipient is using Federal funds appropriately.

Form Numbers:

EPA Form 190-F-04-001, "EPA Payment Request"
 EPA Form 190-F-05-001, "Fellowship Stipend Payment Enrollment Form"
 EPA Form 4700-4, "Preaward Compliance Review Report for All Applicants and Recipients Requesting Federal Financial Assistance"
 EPA Form 5700-52A, "MBE/WBE Utilization Under Federal Grants and Cooperative Agreements"
 EPA Form 5700-53, "Lobbying and Litigation Certification for Grants and Cooperative Agreements"
 EPA Form 5700-54, "Key Contacts Form," and EPA Form 5700-54-2, "Key Contacts Form for Multiple Principal Investigators"
 EPA Form 5770-2, "Fellowship Application"
 EPA Form 5770-3, "Fellowship Facilities and Commitment Statement"
 EPA Form 5770-5, "Agency Fellowship Certification"
 EPA Form 5770-7, "EPA Fellowship Activation Notice"

EPA Form 5770-8, "Fellowship Agreement"
 EPA Form 5770-9, "Completion of Studies Notice"
 EPA Form 6600-01, "EPA Administrative and Financial Onsite Review Questionnaire"
 EPA Form 6600-06, "Certification Regarding Lobbying"
 EPA Form 6600-08A, "Certificate of Indirect Costs for State & Local Governments"
 EPA Form 6600-08B, "Lobbying Indirect Cost Certificate for Non-Profit Organizations" and "Certificate of Indirect Costs for Indirect (F&A) Cost Rate for Non-Profit Organizations"
 EPA Form 6600-09, "EPA Administrative Capability Questionnaire"
 NCER Form 5, "EPA Office of Research and Development Current and Pending Support"

Respondents/affected entities: The primary recipients of EPA assistance agreements are State and local governments, Indian Tribes, educational institutions, and not-for-profit institutions.

Respondent's obligation to respond: Required to obtain an assistance agreement (40 CFR part 30, 40 CFR part 31, and 40 CFR part 33 for awards made prior to December 26, 2014, and 2 CFR 200, 2 CFR 1500, and 40 CFR part 33 for awards made after December 26, 2014).

Estimated number of respondents: 3,048 (total).

Frequency of response: On occasion, quarterly, and annually.

Total estimated burden: 94,606 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: \$6,054,791 (per year), includes \$0 annualized capital or operation & maintenance costs.

Changes in the Estimates: There is an increase of 4,482 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. The increase is partially due to the incorporation of burden into this ICR associated with the relocation of the DBE Program from OSDDBU to OGD. EPA also made adjustments to the estimated number of respondents for each of the requirements included in the ICR and to the burden hour estimates for three of the requirements.

Courtney Kerwin,

Director, Regulatory Support Division.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-21-21DA; Docket No. CDC-2021-0011]

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 Phased Approach to the Resumption of Passenger Operations. The proposed collection outlines a number of information collection activities required as part of the process to returning to passenger operations.

DATES: CDC must receive written comments on or before June 29, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2021-0011 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: 404-639-7118; 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;
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; and
5. Assess information collection costs.

Proposed Project

Phased Approach to the Resumption of Passenger Operations—Existing Collection in use without an OMB Control Number—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Framework for Conditional Sailing Order published in the **Federal Register** on November 4, 2020 prohibits a cruise ship operator from commencing or continuing any regular passenger operations without a COVID–19 Conditional Sailing Certificate issued by HHS/CDC. This information collection request outlines the reporting and document retention requirements that

are part of a phased approach to resuming passenger operations.

Per CDC's Framework for Conditional Sailing Order, cruise ship operators with ships that have not been in U.S. waters during the period of the No Sail Order (NSO) or voluntarily withdrew their ships, must have a NSO response plan deemed complete and accurate, including having submitted to CDC a signed Acknowledgment of No Sail Order Response Plan Completeness and Accuracy. In addition, cruise ship operators must continue to follow their cruise lines' complete, accurate, and acknowledged NSO response plans per the No Sail Order and Suspension of Further Embarkation; Notice of Modification and Extension and Other Measures Related to Operations published at 85 FR 21004 (April 15, 2020) (*i.e.*, “No Sail Order response plan”), as modified and extended July 16, 2020 (published at 85 FR 44085 (July 21, 2020)), and September 30, 2020 (published at 85 FR 62732 (October 5, 2020)).

The Framework for Conditional Sailing Order introduced a phased-in approach to the resumption of cruise ship passenger operations. This Framework Order details the requirements of the initial phase, which focuses on mass testing of crew and building the laboratory capacity needed to test both crew and future passengers.

The Second Phase of the Framework Order focuses on preparation for simulated voyages. As required under the CSO, a cruise ship operator's agreement with U.S. port authorities and local health authorities must include the following elements: (1) A port agreement between the cruise ship operator and port authority to determine the number of cruise ships operating out of any single port in order to not overburden the public health response resources of any single jurisdiction in the event of a COVID–19 outbreak; (2) medical care agreements between the cruise ship operator and health care entities, addressing evacuation and medical transport to onshore hospitals for passengers and crew in need of medical care, in accordance with CDC technical instructions and orders; and (3) housing agreements between the cruise ship operator and one or more shoreside facilities for isolation and quarantine of passengers or crew members with COVID–19 and close contacts, identified from the day of embarkation through disembarkation for each voyage. This Phase also includes a shift in reporting requirements using the

CDC (EDC) form previously approved in OMB Control Number 0920–0134 Foreign Quarantine Regulations. Starting in this phase, the form will be required from cruise ships on a daily, rather than weekly, rhythm.

Phase 2B of the Framework Order focuses on simulated voyages with volunteers playing the role of passengers to test cruise ship operators' ability to mitigate COVID–19 risk. A cruise ship operator must submit a Request for Approval to Conduct a Simulated Voyage Prior to Issuance of COVID–19 Conditional Sailing Certificate to conduct a simulated voyage at least 30 calendar days prior to the voyage. CDC will issue additional technical guidance outlining the specific areas that may be inspected and corresponding recommendations.

Following each simulated voyage, the cruise ship operator must document any deficiencies in its health and safety protocols through an “after-action” report and address how the cruise ship operator intends to address those deficiencies prior to applying for a COVID–19 Conditional Sailing Certificate. This after-action report must also include test results for any volunteer passengers or crew on the simulated voyage.

As a condition of applying for a COVID–19 Conditional Sailing Certificate (Phase 3), a cruise ship operator must have successfully conducted a simulated voyage or series of simulated voyages demonstrating the cruise ship operator's ability to mitigate the risks of COVID–19 onboard its cruise ship. The CDC COVID–19 Conditional Sailing Certificate Application form includes certain minimum requirements that must be met prior to a restricted voyage and burden for these requirements is outlined in section 12 below. These documents must be submitted 60 days prior to any proposed restricted voyage. If the Certificate is denied, revoked, or suspended, a cruise ship operator may submit a written appeal of a denial of its application for a COVID–19 Conditional Sailing Certificate or a revocation or suspension of its COVID–19 Conditional Sailing Certificate.

Compliance with the Framework for Conditional Sailing Order, beyond the information collections outline above, are primarily associated with the testing required, both onshore and onboard. This estimate includes the cost of onboard testing and lab equipment and maintenance on the ship.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total burden (in hrs.)
Cruise ship operator	No Sail Order Response Plan	5	1	2,400/60	200
Cruise ship operator	Request for Embarkation of Essential Crew and Contractors submitted to USCG.	5	1	10/60	1
Cruise ship operator	CDC an Attestation for Non-Commercial Travel of Disembarking Crew for Cruise Ship Operators During the No Sail Order.	5	5	15/60	7
Cruise ship operator	Virtual Implementation Checks	5	2	75/60	13
Cruise ship operator	Enhanced Data Collection (EDC) During COVID-19 Pandemic Form (Daily).	130	365	15/60	11,863
Cruise ship operator	CLIA Certified Laboratory Information—Onshore.	20	25	5/60	42
Cruise ship operator	Approval of Onboard COVID-19 Testing Instrument.	20	25	5/60	42
Cruise ship operator	Request for Embarkation of Essential Crew and Contractors.	130	5	10/60	109
Cruise ship operator	Attestation for Commercial Transportation of Disembarking Crew for Cruise Ship Operators During the Initial Phases of CDC’s Framework for Conditional Sailing Order (CSO).	130	5	15/60	163
Cruise ship operator	Agreement with Health Care Organization with signoff from Local Health Authorities.	130	1	600/60	1,300
Cruise ship operator	Agreement with Port of Entry with signoff from Local Health Authority.	130	1	600/60	1,300
Cruise ship operator	Agreement with Housing Facility with signoff from Local Health Authority.	130	1	600/60	1,300
Cruise ship operator	Request for Embarkation of Non-Essential Crew and contractors.	130	1	10/60	10
Cruise ship operator	Request for Approval to Conduct a Simulated Voyage Prior to Issuance of COVID-19 Conditional Sailing Certificate.	130	1	600/60	1,300
Passenger (3rd party disclosure)	Informed Consent and Medical Certification with no pre-existing conditions for Simulated Voyage.	39,000	1	75/60	48,750
Cruise ship operator	Remote and In-person Inspections ..	130	1	120/60	260
Cruise ship operator	After Action Report, Simulated Voyage.	130	1	600/60	1,300
Cruise ship operator	COVID-19 Conditional Sailing Certificate Application.	130	1	600/60	1,300
Cruise ship operator	Remote and In-person Inspections ..	130	1	120/60	260
Total	69,530

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-21-0891; Docket No. CDC-2021-0045]

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 World Trade Center Health Program Enrollment, Petitions, Designated Representative/HIPAA Authorization, and Member Satisfaction. Data collection is designed to provide healthcare monitoring and treatment to responders of the 9/11/2001 terrorist attacks at the World Trade Center in New York City, the Pentagon in Washington, DC, and Shanksville, Pennsylvania, as well as survivors in the New York City area.

DATES: CDC must receive written comments on or before June 29, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2021-0045 by any of the following methods:

Federal eRulemaking Portal: 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.

Please note: Submit all comments through the Federal eRulemaking portal

(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: 404-639-7118; 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;
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; and
5. Assess information collection costs.

Proposed Project

World Trade Center Health Program Enrollment, Petitions, Designated Representative/HIPAA Authorization, and Member Satisfaction. (OMB Control No. 0920-0891, Exp. 12/31/2021)—Revision—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

NIOSH seeks to request OMB approval to revise the currently approved information collection activities that support the World Trade Center (WTC) Health Program. The James Zadroga 9/11 Health and Compensation Act of 2010 (Pub. L. 111-347, as amended by Pub. L. 114-113) created the WTC Health Program to provide medical monitoring and treatment benefits to eligible firefighters and related personnel, law enforcement officers, and rescue, recovery, and cleanup workers who responded to the September 11, 2001, terrorist attacks in New York City, at the Pentagon, and in Shanksville, Pennsylvania (responders), and to eligible persons who were present in the dust or dust cloud on September 11, 2001, or who worked, resided, or attended school, childcare, or adult daycare in the New York City disaster area (survivors).

Since its inception in 2011, the WTC Health Program has been approved to collect information from applicants and program members (enrolled WTC responders and survivors) concerning enrollment, appointment of a designated representative or third party, and petitions regarding adding a new WTC-related health condition in order to determine coverage under the Program. The current approved total estimated burden is 14,063 hours annually (OMB Control No. 0920-0891, Exp. December 31, 2021).

The WTC Health Program has determined that some existing forms need to be updated (WTC Health Program Applications for Enrollment and Designated Representative Appointment/Designated Representative HIPAA Authorization Forms). For this revision, the burden hours on the WTC Health Program Applications for Enrollment increased due to an expected increase of application volume. The Program updated the enrollment applications for plain language and improved processing. We estimate 15,837 individuals will submit either a FDNY (+95 from previous package), General Responder (+3,740 from previous package), Pentagon/Shanksville Responder (-388 from previous package), or WTC Survivor (+7,881 from previous package) application annually. The applications will take approximately 0.5 hours to complete. The burden estimate for the applications is 7,919 hours. This is an increase from 2018 when the estimated annualized burden was 2,251.

Of the Applications for Enrollment we expect to receive each year, CDC estimates 3,830 (+1,355 from 2018) are