maintained by each of the contractors. As part of the base (minimum sample size), each round of data collection will collect 2,000 responses per quarter. The RSS can be expanded by increasing the number of completed responses per round or the number of rounds per year as needed up to a maximum of 28,000 responses per year per contractor or 56,000 total responses per year. Additionally, each data collection may include up to 2,000 additional responses per quarter (8,000 for the year) to improve representativeness. This increases the maximum burden by up to 16,000 responses per year. The RSS may also target individual surveys to collect data only from specific subgroups within existing survey panels and may supplement data collection for such groups with additional respondents from other probability or nonprobability samples. An additional 12,000 responses per year may be used for these developmental activities.

Each round's questionnaire will consist of four main components: (1)

basic demographic information on respondents to be used as covariates in analyses; (2) new, emerging, or supplemental content proposed by NCHS, other CDC Centers, Institute, and Offices, and other HHS agencies; (3) questions used for calibrating the survey weights; and (4) additional content selected by NCHS to evaluate against relevant benchmarks. NCHS will use questions from Components 1 and 2 to provide relevant, timely data on new, emerging, and priority health topics to be used for decision making. NCHS will use questions from Components 3 and 4 to weight and evaluate the quality of the estimates coming from questions in Components 1 and 2. NCHS submits a 30-day Federal Register Notice with information on the contents of each round of data collection.

NCHS calibrates survey weights from the RSS to gold standard surveys. Questions used for calibration in this round of RSS will include social and work limitation, employment, health information technology use, telephone use, marital status, language used at home and in other settings, and civic engagement. All of these questions have been on the National Health Interview Survey (NHIS) in prior years allowing calibration to these data.

Finally, all RSS rounds will include several questions that were previously on NHIS or other suitable federal surveys for benchmarking to evaluate data quality. Panelists in the RSS will be asked about health status, life satisfaction, chronic conditions, diabetes, veteran status, health care access and utilization, mental health, and body mass index. Female panelists will be asked about their pregnancy history and family planning.

The estimated total annual burden hours for the three-year approval period remains at 28,079 burden hours. The NCHS RSS Round 7 (2025) data collection is based on 8,000 complete surveys (2,664 hours). There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Adults 18+	Survey: NCHS RSS Round 7	8,000 20	1 1	20/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, 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

[30Day-25-1255]

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 "Emergency Cruise Ship Outbreak Investigations (CSOIs)" 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 November 25, 2024, 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

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

Emergency Cruise Ship Outbreak Investigations (CSOIs) (OMB Control No. 0920–1255 Exp. 3/31/2025)— Reinstatement—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Established in 1975 as a cooperative activity with the cruise ship industry, the Centers for Disease Control and Prevention (CDC) Vessel Sanitation Program (VSP) develops and implements comprehensive operational public health programs to minimize the risk of gastrointestinal illness. VSP coordinates and conducts public health inspections, ongoing surveillance of gastrointestinal illness, and outbreak investigations on cruise ships.

Under the authority of the Public Health Service Act (42 U.S.C. 264 and 269), VSP is requesting a three-year Extension for an existing Generic Clearance Information Collection Request (ICR). This ICR will provide for the quick turnaround necessary to conduct emergency Cruise Ship Outbreak Investigations (CSOIs) in response to acute gastroenteritis (AGE) outbreaks. CSOIs are used to determine causative agents and their sources, modes of transmission, or risk factors. VSP's jurisdiction includes passenger vessels carrying 13 or more people sailing from foreign ports and within 15 days of arriving at a U.S. port.

VSP uses its syndromic surveillance system called the "Maritime Illness and Death Reporting System (MIDRS)" (OMB Control No. 0920–1260, Exp. 3/31/2026) to collect aggregate data about the number of people onboard ships in VSP's jurisdiction who are experiencing AGE symptoms. When the levels of illness meet VSP's alert threshold (*i.e.*,

at least 2% in either the passenger or crew populations), a special report is made to VSP via MIDRS and VSP provides environmental health and epidemiologic assistance. VSP considers an outbreak to be \geq 3% of reportable AGE cases in either passenger or crew populations.

When a cruise ship has an AGE outbreak, VSP often must deploy a response team to meet the ship in port within 24 hours of reaching the outbreak threshold. In some cases, the response team must board the ship before its U.S. arrival and sail back to the U.S. port of disembarkation to conduct a more detailed and comprehensive epidemiologic and environmental health evaluation of the outbreak.

VSP can ascertain causative agent, sources of exposure, modes of transmission, and risk factors by gathering the following types of information from both the affected and (seemingly) unaffected populations:

- Demographic information,
- Pre-embarkation travel information,
- Symptoms, including type, onset, duration,
- Contact with people who were sick or their body fluids,
- Participation in ship and onshore activities,
- Locations of eating and drinking, and
- Foods and beverages consumed both on the ship and on shore. Rapid and flexible data collection is imperative given the mobile environment, the remaining duration of the voyage left for investigation, and the loss to follow-up if delays allow passengers to disembark and leave the

ship, including those returning to locations outside of the United States.

This Generic Clearance will cover investigations that meet all the following criteria:

- The investigation is urgent in nature (*i.e.*, timely data are needed to inform rapid public health action to prevent or reduce morbidity or mortality).
- The investigation is characterized by undetermined agents, undetermined sources, undetermined modes of transmission, or undetermined risk factors.
- One or more CDC staff (including trainees and fellows) will be deployed to the field.
- Data collection is completed in 30 days or less (most CSOIs involve two to five days of data collection).

This Generic Clearance excludes each of the following:

- Investigations related to non-urgent outbreaks or events.
- Investigations conducted for the primary purpose of program evaluation, surveillance, needs assessment, or research (e.g., to contribute to generalizable knowledge).
- Investigations with data collection expected for greater than 30 days.

VSP estimates 10 CSOIs annually in response to cruise ship AGE outbreaks. The estimated number of respondents is 1,300 per CSOI, for a total of 13,000 respondents per year. The average time burden is 15 minutes for each respondent. Therefore, the total estimated annual burden in hours is 4,063. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Cruise ship crew	Self-administered questionnaire	3,000 450	1 1	15/60 15/60
Cruise ship processor	Biospecimen collection	300	1	15/60 15/60
Cruise ship passenger Cruise ship passenger	Self-administered questionnaire	10,000 1,500	1	15/60
Cruise ship passenger	Biospecimen collection	1,000	1	15/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

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