online panels. The RSS is designed to have four rounds of data collection each year with data being collected by two contractors with probability panels. A cross-sectional nationally representative sample will be drawn from the online probability panel 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. Survey questions being asked of the panelists will be cognitively tested. This cognitive testing will help survey users interpret the findings by understanding how respondents answer each question.

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. Components 1 and 2 will contain different topics in each round of the survey. 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 marital status, employment, social and work limitations, use of the internet in general and for medical reasons, telephone use, civic engagement, and language used at home and in other settings. All 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 for benchmarking to evaluate data quality. Panelists in the RSS will be asked about health status; chronic conditions; disability; healthcare access and utilization; health behaviors; and food insecurity.

The estimated total annual burden hours for the three-year approval period remains at 28,079 burden hours. The NCHS RSS Round 3 (2024) data collection is based on 13,100 complete surveys (4,367 hours) and 20 cognitive interviews (20 hours) using the same survey instrument for a total of 4,387 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 3	13,100	1	20/60
Adult 18+	Cognitive Interviews	20	1	1

Jeffrey M. Zirger,

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

[FR Doc. 2023–25086 Filed 11–13–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–24–23BJ]

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 ''U.S. National Authority for Containment of Poliovirus Data Collection Tools'' 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 December 12, 2022 to obtain comments from the public and affected agencies. CDC received one comment 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

U.S. National Authority for Containment of Poliovirus Data Collection Tools—New—Office of Readiness and Response (ORR), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The role of the National Authority for Containment (U.S. NAC) of Poliovirus is to ensure that the requirements established in the World Health Organization (WHO) Global Action Plan (GAP) III/IV standard are effectively implemented and maintained in facilities working with or storing infectious poliovirus or potentially infectious materials. Risk assessments following an incident are a critical component for adequate application of the GAP standard. To support risk assessment activities, The ''Facility Incident Reporting Form for Poliovirus Release and Potential Exposure'' and the ''Facility Incident Reporting Form for Poliovirus Theft or Loss'' forms were created for facilities to capture and submit incident information to the U.S. NAC. These forms will not only address the biosafety and biosecurity containment emergency elements of the GAP standard but will also inform the U.S. NAC risk assessments and thereby, guide CDC's determination of the emergency response level and direction.

The information collected in the ''Personal Protective Equipment Survey for Laboratories'' will assist the Centers for Disease Control and Prevention (CDC), U.S. NAC and National Institute for Occupational Safety and Health (NIOSH) with developing guidance and recommendations for PPE selection and use in support of poliovirus containment, as well as identify laboratory PPE commonly used to evaluate laboratory PPE performance characteristics in testing studies.

Information collected in the ''Global Action Plan (GAP) Poliovirus Containment Poliovirus-Essential Facility Assessment Checklist'' will aid U.S. facilities in preparing for an audit to obtain a poliovirus certificate of containment. Data collected from this form will also collect additional information on poliovirus materials held by a U.S. facility, their work activities, and facility features.

The ''Poliovirus Containment Sampling Plan and Sanitation Assessment Form for Wastewater (WW) Systems Supporting a Poliovirus-Essential Facility (PEF) in the United States'' form will collect information to assess a poliovirus facility's essential WW system, the primary safeguards to reduce and control the release of poliovirus from the facility. In addition, it will verify the safeguards of local WW utilities that receive WW from the PEF.

OMB approval is sought for three years. The annualized time burden for this information collection is estimated to be 125 hours. 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)
Facility Staff/Leadership	Facility Incident Reporting Form for Poliovirus Release or Potential Exposure.	10	1	45/60
Facility Staff/Leadership	Facility Incident Reporting Form for Poliovirus Theft or Loss	10	1	45/60
Facility Staff/Leadership	Personal Protective Equipment Survey for Laboratories	20	1	90/60
Facility Staff/Leadership	GAP Poliovirus Containment Poliovirus-Essential Facility Questionnaire.	20	1	90/60
Facility Staff/Leadership	GAP Facility Assessment Checklist	20	1	1
Facility Staff/Leadership	The Poliovirus Containment Sampling Plan and Sanitation Assessment Form for Wastewater (WW) Systems Supporting a Poliovirus-Essential Facility (PEF) in the United States.	20	1	90/60

Jeffrey M. Zirger,

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

[FR Doc. 2023–25082 Filed 11–13–23; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–24–0214]

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 ''National Health Interview Survey (NHIS)'' 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 August 21, 2023 to obtain comments from the public and affected agencies. CDC receive three 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;