

compared among sites. Jurisdictions can implement the survey on an ongoing basis or as a point-in-time survey. In participating jurisdictions, a sample of women who have recently given birth to a live born or stillborn infant is selected from birth certificates or fetal death files. The sample is stratified based on the site's population of interest to ensure high-risk populations are adequately represented in the data.

The PRAMS survey instrument for live births is based on a core set of questions common across all jurisdictions that remain the same throughout each phase of data collection. In addition, CDC provides optional standardized modules (pre-grouped questions on a select topic) that jurisdictions may use to customize survey content at the beginning of each phase of data collection. Topics for both the core and standard modules include demographic and background characteristics; health conditions (which includes chronic conditions such as diabetes, hypertension, mental

health, oral health, cancer, as well as pregnancy-induced health conditions and family history of select conditions); health behaviors (including tobacco and alcohol use, substance use [licit and illicit], injury prevention and safety, nutrition, and physical activity); health care services (such as preconception care, prenatal care, postpartum care, contraceptive care, vaccinations, access to care, insurance coverage, receipt of recommended services, and provider counseling received); infant health and development; infant care practices (such as breastfeeding, safe sleep practices); social services received (such as WIC or home visiting); the social context of childbearing (such as intimate partner violence, social support, adverse childhood experiences, and stressful life experiences); and attitudes and feeling about the pregnancy including pregnancy intentions.

At times, jurisdictions may address emerging topics of interest with supplemental modules (pre-grouped questions on a selected topic).

Supplemental modules available for site-specific data collection include disabilities, substance use, COVID-19 experiences, and social experiences. New supplemental modules may be developed to address other emergent issues as they arise. The stillbirth survey may be administered in a smaller number of sites. It includes a single survey instrument.

CDC is seeking approval for an Extension of the PRAMS data collection which currently expires 3/31/2026. OMB approval is requested for three years. The total estimated annual burden is 29,773 hours which is a decrease of 1,495 hours. The change in overall burden results from removal of components already completed: (1) call back surveys (decrease of 1,395 hours), and (2) cognitive and field testing (decrease of 100 hours) since no new questions or supplemental modules are anticipated during the approval window. 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)	Total burden (in hours)
Women who recently delivered a live birth.	PRAMS Phase 9 Questionnaire (core questions plus site selected standard modules).	51,556	1	26/60	22,341
Women who recently delivered a stillbirth.	Supplemental Modules	52,984	1	8/60	7,065
Jurisdictions	PRAMS Stillbirth Questionnaire	160	1	25/60	67
Total	Submission of data file to CDC	50	12	30/60	300
					29,773

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. 2025-20583 Filed 11-20-25; 8:45 am]
BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
[30Day-25-0036]

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 "Division of Vital Statistics Proposal for Access to

Restricted-Use Vital Statistics Data for the National Center for Health Statistics" 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 June 16, 2025, to obtain comments from the public and affected agencies. CDC received two 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

Division of Vital Statistics Proposal for Access to Restricted-Use Vital

Statistics Data for the National Center for Health Statistics—Existing Collection in Use Without an OMB Control Number—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This is a request for OMB Clearance for the Division of Vital Statistics (DVS) Proposal for Access to Restricted-Use Vital Statistics Data for the National Center for Health Statistics. A three-year clearance is requested. The DVS Proposal has been in use since 1998. Recently, as part of the Evidence Act work to develop and implement a federal government-wide Standard Application Process (see 44 U.S.C. 3583) and through OMB consultation, the NCHS DVS became aware that the DVS proposal was, in fact, an information

collection initiative. The NCHS DVS recognizes this and is therefore submitting this OMB clearance package to correct this oversight and obtain OMB approval.

This proposed information collection of information is designed to help facilitate review of the agency's research and will allow the National Center for Health Statistics (NCHS) to determine whether a research study requires access to confidential data. NCHS anticipates it will receive approximately 600 proposals annually. CDC requests OMB approval for an estimated 600 annual burden hours. There is no cost to respondents other than their 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)
Researchers	NCHS Restricted Vital Statistics Data Request Application Form ..	600	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. 2025–20577 Filed 11–20–25; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–25–1260; Docket No. CDC–2025–0684]

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 continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information

collection project titled Maritime Illness Database and Reporting System (MIDRS). This data collection is designed to allow the Vessel Sanitation Program (VSP) to monitor acute gastroenteritis (AGE) illness on cruise ships, conduct sanitation inspections, perform epidemiologic investigations when outbreaks occur, and formulate public health recommendations to prevent future transmission and outbreaks.

DATES: CDC must receive written comments on or before January 20, 2026.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2025–0684 by either of the following methods:

- *Federal eRulemaking Portal:* 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 H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to www.regulations.gov.

Please note: Submit all comments through the Federal eRulemaking portal (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 H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; 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