resources, building capacity to implement and evaluate IPV primary prevention in their state and selected communities, and using evaluation data for quality improvement.

CDC will use the information collected to further understand the facilitators, barriers, and critical factors to implementing specific violence prevention strategies and conducting related program evaluation activities. Data collected will also be used to inform CDC's training and technical assistance, program improvement, progress toward NOFO goals, and the development of future funding opportunities. Program evaluation activities allow CDC to identify and disseminate information about successful prevention strategies implemented by recipients. These functions are central to NCIPC's broad mission of protecting Americans from violence and injury threats. This information collection will allow CDC to monitor the impact of the strategies implemented by the recipients on outcomes related to intimate partner violence prevention. It is also expected to reduce duplication of effort, enhance program impact, and maximize the use of federal funds.

CDC requests OMB approval for an estimated 130 annual burden hours. There are no direct costs 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)
DELTA State Domestic Violence Coalition (SDVC) Project Leads.	Annual Performance Report ..	13	1	10

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. 2026–12279 Filed 6–17–26; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30–Day–26–1393]

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 ''Research Data Center Proposal (RDC) Security Forms for Access to Confidential Data'' 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 April 1, 2026 to obtain comments from the public and affected agencies. CDC received four comments for this 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

Research Data Center (RDC) Data Security Forms for Access to Confidential Data for the National Center for Health Statistics (OMB Control No. 0920–1393)—Reinstatement—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306(b)(4) of the Public Health Service (PHS) Act (42 U.S.C. 242k(b)(4)), as amended, authorizes the Secretary of Health and Human Services (DHHS), acting through NCHS, to receive requests for furnishing statistics to the public. NCHS receives requests for statistics from the public through the Standard Application Process (SAP). The public may apply to access confidential data assets held by a federal statistical agency or unit through the SAP for the purposes of generating statistics and developing evidence. Once an application for confidential data is approved through the SAP, NCHS will collect information to meet its data security requirements using its Data Security Forms. This information collection using the Data Security Forms will occur outside of the SAP. This is a request for approval from OMB to collect information via the Researcher Data Center Data Security Forms over the next three years.

As part of a comprehensive data dissemination program, the Research Data Center (RDC), National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC), requires prospective researchers who need access to confidential data to complete a research application in the SAP. Researchers self-select whether they need access to confidential data to answer their research questions. The RDC requires the researcher to complete

a research application, so NCHS understands the research proposed. The completed application is sent to NCHS through the SAP portal for review and adjudication. If the research application is approved by NCHS, then the researcher must fill out the data security forms. If the researcher will access the data at a Research Data Center, then the ''Data Access Form'' and the ''Designated Agent Form'' would need to be completed and returned to NCHS. If the researcher will access the data through the NCHS Virtual Data Enclave (VDE), then the ''VDE Data Use Agreement Form'', ''Data Access Form'' and the ''Designated Agent Form'' would need to be completed and returned to NCHS.

To capture the information needed to adjudicate a researcher's commitment to protect confidential NCHS data, researchers must complete and sign the data security forms. This request allows for both researcher signature and the time per response for a total estimated annual burden total of 110 hours; 330 hours for a three-year clearance period. There is no cost to a researcher other than their time to complete the forms unless the researcher must pay a nominal notary fee for services incurred. The resulting information in these forms will be used for NCHS internal administrative purposes.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)
Researcher	Research Data Center proposal.	110	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. 2026–12276 Filed 6–17–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-26–0260; Docket No. CDC–2026–1090]

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 Health Hazard Evaluations/Technical Assistance and Emerging Problems. In accordance with its mandates under the Occupational Safety and Health Act of 1970 and the Federal Mine Safety and Health Act of 1977, the National Institute for Occupational Safety and Health (NIOSH) responds to requests for Health Hazard Evaluations (HHEs) to identify chemical, biological or physical hazards in workplaces throughout the United States.

DATES: CDC must receive written comments on or before August 17, 2026.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2026–1090 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 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

Health Hazard Evaluations/Technical Assistance and Emerging Problems (OMB Control No. 0920–0260, Exp. 1/31/2027)—Extension—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).