

CDC requests OMB approval for an estimated 3,126 total annualized burden hours. Participation is voluntary and

there are no costs to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondent	Form	Number of respondents	Number of responses per respondent	Average burden per response (hours)
Persons Screened	Eligibility Screening Form English	5,400	1	5/60
Persons Screened	Eligibility Screening Form Spanish	600	1	5/60
Persons who give permission	Model Project Consent Form English	4,050	1	5/60
Persons who give permission	Model Project Permission Form Spanish	450	1	5/60
Eligible Participants	NEXUS Survey English	4,050	1	30/60
Eligible Participants	NEXUS Survey Spanish	450	1	30/60

Jeffrey M. Zirger,

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Office of Public Health Ethics and
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Disease Control and Prevention.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–25–1360]

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 “CryptoNet Case Report Form” 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 October 21, 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 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

CryptoNet Case Report Form (OMB Control No. 0920–1360, Exp. 1/31/2025)—Revision—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Waterborne Disease Prevention Branch (WDPB) in the Division of Foodborne, Waterborne, and Environmental Diseases (DFWED) works

to prevent domestic and global water-related diseases. The WDPB is comprised of five teams, including the Domestic Waterborne Disease Epidemiology and Response (WDER) Team, which focuses on the prevention and control of waterborne-related diseases and outbreaks in the United States. One of the diseases included in the team’s work is cryptosporidiosis, an acute diarrheal disease caused by infection with *Cryptosporidium* parasites.

The Case Surveillance Program is a subunit within the Domestic WDER Team that focus on the data collection and management activities of five waterborne diseases, including cryptosporidiosis, in the United States. The Case Surveillance Program’s current scope of work includes modernizing data collection and management, enabling data connections, and improving public data access to aid public health action.

CryptoNet is the first molecular tracking system for *Cryptosporidium* in the United States. To meet the needs of the CryptoNet and Case Surveillance Program, and the needs of local officials, the CryptoNet case report form (CRF) was developed. The CRF includes a set of data elements that can be used to identify exposures trends in outbreak- and non-outbreak-associated *Cryptosporidium* cases, to generate hypotheses about the sources of infection in clusters or outbreaks, and to identify strategies to prevent and control *Cryptosporidium* cases, clusters, or outbreaks.

Data from the CRF will be used by federal, state, and local public health officials responsible for conducting interviews with reported cases of cryptosporidiosis in their jurisdiction in order to systemically assess core exposure elements and risk factors among cases of cryptosporidiosis. Collected data will be used by CDC staff to inform cryptosporidiosis sporadic

case, cluster, and outbreak prevention and control strategies. CRF data elements and the CRF were designed for administration via telephone interviews with individuals ill with

cryptosporidiosis, or their designated proxy.

CDC requests OMB approval for an estimated 125 annual burden hours. Providing information is voluntary, and

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)
Individuals ill with cryptosporidiosis, or their designated proxy.	CryptoNet Case Report Form	500	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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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–25–1357; Docket No. CDC–2024–0104]

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 The Greater Access and Impact with NAT (GAIN) Study: Improving HIV Diagnosis, Linkage to Care, and Prevention Services with HIV Point-of-Care Nucleic Acid Tests (NATs). This study will identify facilitators and barriers with implementation of HIV point-of-care (POC) nucleic acid tests (NATs) in clinical settings, estimate the sensitivity and specificity of the HIV POC NAT, and assess the impact of the test in decreased time to receipt of HIV prevention and care.

DATES: CDC must receive written comments on or before March 10, 2025.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0104 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

The Greater Access and Impact with NAT (GAIN) Study: Improving HIV Diagnosis, Linkage to Care, and Prevention Services with HIV Point-of-Care Nucleic Acid Tests (NATs) (OMB Control No. 0920–1357, Exp. 12/31/2024)—Reinstatement—National Center for HIV, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC)

Background and Brief Description

HIV prevention and care services can be improved by the availability of Point-of-care (POC) HIV viral RNA testing in clinical settings. Viral RNA tests are the most sensitive HIV tests for the detection of early infection. The purpose of this data collection is to develop feasible and effective models to integrate HIV POC nucleic acid tests (NATs) in HIV prevention and treatment services. The HIV POC NAT can be used to test persons at high-risk of acquiring HIV infection to reduce the time