

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.
5. Assess information collection costs.

Proposed Project

Medical Monitoring Project (MMP) (OMB Control No. 0920–0740, Exp. 6/30/2021)—Revision—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC), Division of HIV/AIDS Prevention (DHAP) requests a revision of the currently approved Information Collection Request: “Medical Monitoring Project” which expires June 30, 2021. This data collection addresses the need for national estimates of access to, and utilization of HIV-related medical care and services, the quality of HIV-related ambulatory care, and HIV-related behaviors and clinical outcomes. For the proposed project, the same data collection methods will be used as for the currently approved project. Data would be collected from a probability sample of HIV-diagnosed adults in the

U.S. who consent to an interview and abstraction of their medical records. As for the currently approved project, deidentified information would also be extracted from HIV case surveillance records for a dataset (referred to as the minimum dataset), which is used to assess non-response bias, for quality control, to improve the ability of MMP to monitor ongoing care and treatment of HIV-infected persons, and to make inferences from the MMP sample to HIV-diagnosed persons nationally. No other Federal agency collects such nationally representative population-based information from HIV-diagnosed adults. The data are expected to have significant implications for policy, program development, and resource allocation at the state/local and national levels. The changes proposed in this request update the data collection system to meet prevailing information needs and enhance the value of MMP data, while remaining within the scope of the currently approved project purpose. The result is a 10% reduction in burden, or a reduction of 647 total burden hours annually. The reduction in burden was a result of revisions to the interview questionnaire that were made to improve coherence, boost the efficiency of the data collection, and increase the relevance and value of the information, which decreased the time of interview from 45 minutes to 40 minutes.

Changes made, that did not affect the burden, listed below:

- Non-substantive changes have been made to the respondent consent form to decrease the reading comprehension level and make the form more visual.
- Nine data elements were removed from, and three data elements were added to the Minimum Dataset. Because these data elements are extracted from the HIV surveillance system from which they are sampled, these changes do not affect the burden of the project.
- Seven data elements were added to the medical record abstraction data elements to collect information on SARS-CoV–2 (COVID–19) testing. Because the medical records are abstracted by MMP staff, these changes do not affect the burden of the project.

This proposed data collection would supplement the National HIV Surveillance System (NHSS, OMB Control No. 0920–0573, Exp. 11/30/2022) in 23 selected state and local health departments, which collect information on persons diagnosed with, living with, and dying from HIV infection and AIDS. The participation of respondents is voluntary. There is no cost to the respondents other than their time. Through their participation, respondents will help to improve programs to prevent HIV infection as well as services for those who already have HIV. Total estimated annual burden requested is 5,707 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondent | Form name | Number of respondents | Number of responses per respondent | Average hours per response | Total response burden (hours) |
|---|------------------------------------|-----------------------|------------------------------------|----------------------------|-------------------------------|
| Sampled, Eligible HIV-Infected Persons | Interview Questionnaire (Att. 5a). | 7,760 | 1 | 45/60 | 5,173 |
| Facility office staff looking up contact information | Look up contact information. | 1,940 | 1 | 2/60 | 65 |
| Facility office staff approaching sampled persons for enrollment. | Approach persons for enrollment. | 970 | 1 | 5/60 | 81 |
| Facility office staff pulling medical records | Pull medical records | 7,760 | 1 | 3/60 | 388 |
| Total | | | | | 5,707 |

Jeffrey M. Zirger,
Lead, Information Collection Review Office,
Office of Scientific Integrity, 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-20–20KW]
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 “School Health Profiles Test-Retest Reliability Study” 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 March 16, 2020 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

School Health Profiles Test-Retest Reliability Study—New—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The purpose of this request is to obtain OMB approval to conduct the School Health Profiles Test-Retest Reliability Study to establish the reliability of the School Health Profiles ("Profiles"). Profiles is a system of school-based surveys conducted by state and local education and health agencies among school principals and lead health education teachers at the secondary school level to assess school health policies and practices related to health education, physical education and physical activity, tobacco use prevention, nutrition, school-based health services, family and community involvement in school health, and school health coordination. CDC seeks a one-year approval to conduct the School Health Profiles Test-Retest Reliability Study.

Profiles surveys are administered widely. In 2018, 48 states, 21 large

urban school districts, and two territories conducted School Health Profiles. Across all of these jurisdictions, questionnaires were completed by approximately 10,000 principals and by approximately 9,000 lead health education teachers. States and large urban school districts use Profiles as a data source for performance measures for two CDC cooperative agreements: CDC-RFA-PS18-1807, Promoting Adolescent Health Through School-Based HIV Prevention (PS18-1807), and CDC-RFA-DP18-1801 Improving Student Health and Academic Achievement Through Nutrition, Physical Activity and the Management of Chronic Conditions in Schools (DP18-1801). No other surveillance system measures school health policies and programs nationwide.

Between January and June of 2021, approximately 200 principals and 200 lead health education teachers from regular public secondary schools in the United States containing at least one of grades 6 through 12 will complete both a Time 1 and Time 2 Profiles questionnaire. Five questions will be added at the end of both the principal and lead health education teacher questionnaires at the Time 2 administration to gather data on why responses to the same questions may have changed or stayed the same between the two administrations.

There are no costs to respondents except their time. The total estimated annualized burden hours are 686. OMB approval is requested for one year. Participation is voluntary.

ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondents | Form name | Number of respondents | Number of responses per respondent | Average burden per response (in hours) |
|-------------------------------------|---|-----------------------|------------------------------------|--|
| School Principal | School Principal Questionnaire Time 1 | 200 | 1 | 45/60 |
| School Principal | Nonresponse follow-up call | 150 | 1 | 5/60 |
| School Principal | School Principal Questionnaire Time 2 | 200 | 1 | 45/60 |
| School Principal | School Principal Supplemental Questions | 200 | 1 | 5/60 |
| School Principal | Nonresponse follow-up call | 150 | 1 | 5/60 |
| Lead Health Education Teacher | Lead Health Education Teacher Questionnaire Time 1. | 200 | 1 | 45/60 |
| Lead Health Education Teacher | Nonresponse follow-up call | 150 | 1 | 5/60 |
| Lead Health Education Teacher | Lead Health Education Teacher Questionnaire Time 2. | 200 | 1 | 45/60 |
| Lead Health Education Teacher | Lead Health Education Teacher Supplemental Questions. | 200 | 1 | 5/60 |
| Lead Health Education Teacher | Nonresponse follow-up call | 150 | 1 | 5/60 |

Jeffrey M. Zirger,

Lead, Information Collection Review Office,
Office of Scientific Integrity, Office of Science,
Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10371]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare &
Medicaid Services, Health and Human
Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by November 9, 2020.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following

address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number CMS-10371, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of the following:

1. Access CMS' website address at website address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

2. Call the Reports Clearance Office at (410) 786-1326.

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786-4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS-10371 Cooperative Agreements to Support Establishment of State-Operated Health Insurance Exchanges

Under the 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. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request:* Revision of an existing information collection request; *Title of Information Collection:* Cooperative Agreements to Support Establishment of State-Operated Health Insurance Exchanges; *Use:* Section 1311(b) of the Affordable Care Act provides the opportunity for each State to establish

an Exchange (now referred to as an Exchange). Section 1311 of the Affordable Care Act provides for grants to States for the planning and establishment of these Exchanges. Given the innovative nature of Exchanges and the statutorily-prescribed relationship between the Secretary and States in their development and operation, it is critical that the Secretary work closely with States to provide necessary guidance and technical assistance to ensure that States can meet the prescribed timelines, federal requirements, and goals of the statute. Additionally, under 42 CFR 155.1200(b) State Exchanges are required to provide performance monitoring data to CMS. State Exchanges must provide this data at least annually and in the manner, format, and deadlines specified by HHS. The information collection requirements associated with these ICRs will primarily involve programmatic narrative, accompanying budget narrative and appropriate supporting documentation, and provision of performance outcome and operational data by grantees operating their Exchanges. The SBEs are not required to track or submit any personally identifiable data. It is expected that States will create data with readily available word processing and spreadsheet programs relying on source data from information systems developed from grant funding, ACA section 1332 pass-through funding, or state funding sources and submit such information electronically. *Form Number:* CMS-10371 (OMB Control Number: 0938-1119); *Frequency:* Once; *Affected Public:* State Government agencies, non-profit entities; *Number of Respondents:* 17; *Number of Responses:* 37; *Total Annual Hours:* 12,328. For policy questions regarding this collection contact Courtney Williams at (301) 492-5157.

Dated: September 1, 2020.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office
of Strategic Operations and Regulatory
Affairs.

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