

Services Administration will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (240) 276-1243.

Comments are invited on: (a) Whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

**Proposed Project: Projects for Assistance in Transition From Homelessness (PATH) Program Annual Report (OMB No. 0930-0205)—Revision**

The Center for Mental Health Services awards grants each fiscal year to each of the States, the District of Columbia, the Commonwealth of Puerto Rico, the Virgin Islands, Guam, American Samoa, and the Commonwealth of the Northern Mariana Islands from allotments authorized under the PATH program established by Public Law 101-645, 42 U.S.C. 290cc-21 et seq., the Stewart B. McKinney Homeless Assistance Amendments Act of 1990 (section 521 et seq. of the Public Health Service (PHS) Act). Section 522 of the PHS Act requires that the grantee States and Territories must expend their payments

under the Act solely for making grants to political subdivisions of the State, and to non-profit private entities (including community-based veterans' organizations and other community organizations) for the purpose of providing services specified in the Act. Available funding is allotted in accordance with the formula provision of section 524 of the PHS Act.

This submission is for a revision of the current approval of the annual grantee reporting requirements. Section 528 of the PHS Act specifies that not later than January 31 of each fiscal year, a funded entity will prepare and submit a report in such form and containing such information as is determined necessary for securing a record and description of the purposes for which amounts received under section 521 were expended during the preceding fiscal year and of the recipients of such amounts and determining whether such amounts were expended in accordance with statutory provisions.

The proposed changes to the PATH Annual Report Survey are as follows:

**1. Format**

To create a PATH report that is easier to read, the formatting has been modified to be more table driven. In addition, the language has been made more concise. Although the online form and report is close in flow to the previous report, it is necessary to thoroughly read all reporting instructions to insure proper data entry.

**2. Estimated Counts**

The new PATH report does not include entry of estimated counts. Only actual counts should be entered.

**3. Homelessness Management Information System (HMIS) Data Integration**

The Data section of the report is expected to be propagated from the local HMIS when providers use HMIS. This includes client counts, services, referrals, and demographics. This data will be automatically aggregated from client-level data.

**4. Demographic Responses**

In order to facilitate integration of PATH data into HMIS, all data responses have been modified to fully align with valid HMIS responses. For example, the "Hispanic" response has been separated from "Race" and placed in "Ethnicity."

**5. Additional Data Items**

The PATH report now tracks demographic data for persons contacted, as well as those enrolled. For services and referrals, in addition to gathering the number of enrolled persons receiving the service or referral, there is a total count of the number of times that particular service was provided or referral made.

**6. Voluntary Outcome Measures**

The data previously entered as voluntary outcome measures has now been moved to the referral section of the report and are no longer considered "voluntary."

The estimated annual burden for these reporting requirements is summarized in the table below.

Respondents	Number of respondents	Responses/respondent	Burden per response (hrs.)	Total burden
States .....	56	1	19	1,064
Local provider agencies .....	503	1	34	17,102
Total .....	559	.....	.....	18,166

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 8-1099, One Choke Cherry Road, Rockville, MD 20857 or email a copy to [summer.king@samhsa.hhs.gov](mailto:summer.king@samhsa.hhs.gov). Written comments must be received before 60 days after the date of the publication in the **Federal Register**.

**Summer King,**  
*Statistician.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Substance Abuse and Mental Health Services Administration**

**Agency Information Collection Activities: Submission for OMB Review; Comment Request**

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the

Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276-1243.

**Project: Minority AIDS Initiative (MAI) Rapid HIV Testing Clinical Information Form (OMB No. 0930-0295)—Revision**

This request is for a three-year generic clearance to continue rapid HIV testing data collection among 63 TCE-HIV Grantees and their clients and the additional 11 MAI-HIV Grantees and their clients. The primary purpose of the

MAI Rapid HIV Testing Clinical Information Form is to use a standardized data collection instrument to fully capture essential clinical information to enhance preventive services for those who test HIV-negative and refer to quality treatment/medical care those who test HIV-positive.

The aim of the project is to implement and increase rapid HIV testing among racial and ethnic minorities and collect rapid HIV testing data using the MAI Rapid HIV Testing Clinical Information Form. To meet this requirement, all Grantees must offer their clients rapid

HIV preliminary antibody testing during outreach, pretreatment, or program enrollment. In addition, rapid HIV testing may be made available to the sexual and/or injection partners of clients. Grantees must provide onsite rapid HIV testing in accordance with their respective State and local requirements. If a client requests an off-site rapid HIV test, the Grantee must provide a referral to a rapid HIV testing site certified by the local health department.

Grantees are currently using the MAI Rapid HIV Testing Clinical Information

Form in the field to systematically collect information from clients on demographics, previous rapid HIV test results, substance use and sexual risk behaviors, current rapid HIV test results, types of services received, and confirmatory HIV test result. Once a client is offered a rapid HIV test, the Grantee staff completes the MAI Rapid HIV Testing Clinical Information Form with the client present and then enters the data into a secure Web site that allows for real-time data submission.

The estimated annualized burden is summarized below.

Respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total burden hours
MAI Rapid HIV Testing Clinical Information Form (FY 2008 and FY 2009–63 Grantees) .....	10,000	1	0.133	1,330
RHT form for 11 HIV program FY 2011 grantees (public health departments) .....	20,000	1	0.133	2,660
MAI Rapid HIV Testing Clinical Information Form (Re-test) .....	6,000	1	0.133	798
Total .....	30,000	.....	.....	4,788

Written comments and recommendations concerning the proposed information collection should be sent by July 12, 2012 to the SAMHSA Desk Officer at the Office of Information and Regulatory Affairs, Office of Management and Budget (OMB). To ensure timely receipt of comments, and to avoid potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, commenters are encouraged to submit their comments to OMB via email to: *OIRA\_Submission@omb.eop.gov*. Although commenters are encouraged to send their comments via email, commenters may also fax their comments to: 202–395–7285. Commenters may also mail them to: Office of Management and Budget, Office of Information and Regulatory Affairs, New Executive Office Building, Room 10102, Washington, DC 20503.

**Summer King,**  
*Statistician.*

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**Project: National Evaluation of the Comprehensive Community Mental Health Services for Children and Their Families Program: Phase VI (OMB No. 0930–0307)—REVISION**

The Substance Abuse and Mental Health Services Administration (SAMHSA), Center of Mental Health Services is responsible for the national evaluation of the Comprehensive Community Mental Health Services for Children and Their Families Program (Children's Mental Health Initiative—CMHI) that will collect data on child mental health outcomes, family life, and service system development. Data will be collected on 47 service systems, and approximately 6,561 children and families.

Principal changes from the previous Phase VI OMB approval include:

- Addition of nine (9) communities awarded cooperative agreements in FY2010 for data collection.
  - Replacement of intake and follow-up questionnaires for the Child Welfare Sector and Comparison Study with an administrative record review form to lessen burden.
  - Addition of a brief 8-item Education Sector Caregiver Questionnaire to the Education Sector and Comparison Study to capture family involvement in the development and use of Individualized Education Plans (IEPs).
  - Removal of data collection activities for the Alumni Networking Study, the CQI Initiative Evaluation, and the Sustainability Study.
- Data collection for this evaluation will be conducted over a five-year period. Child and family outcomes of interest will be collected at intake and during subsequent follow-up sessions at six-month intervals. The length of time that individual families will participate in the study is up to 24 months. The outcome measures include the following: Child symptomatology and functioning, family functioning, satisfaction, and caregiver strain. The core of service system data will be collected every 18–24 months throughout the 5-year evaluation period. Service utilization and cost data will be tracked and submitted to the national evaluation every six months using two tools: The Flex Fund Tool and the Services and Costs Data Tool to estimate