

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
CRC Survivors	Script for CATI Follow-up	100	1	3/60	5
	Survey of Health Behaviors	1,000	1	40/60	667

Dated: April 1, 2009.
Maryam I. Daneshvar,
Acting Reports Clearance Officer, Centers for Disease Control and Prevention.
 [FR Doc. E9-7789 Filed 4-6-09; 8:45 am]
BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) 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: 2010 National Survey on Drug Use and Health—(OMB No. 0930-0110)—Revision

The National Survey on Drug Use and Health (NSDUH) is a survey of the civilian, non-institutionalized population of the United States 12 years old and older. The data are used to determine the prevalence of use of tobacco products, alcohol, illicit substances, and illicit use of

prescription drugs. The results are used by SAMHSA, ONDCP, Federal government agencies, and other organizations and researchers to establish policy, direct program activities, and better allocate resources.

The 2010 NSDUH will continue conducting a follow-up clinical interview with a subsample of approximately 500 respondents. The design of this study is based on the recommendations from a panel of expert consultants convened by the Center for Mental Health Services (CMHS), SAMHSA, to discuss mental health surveillance data collection strategies. The goal is to create a statistically sound measure that may be used to estimate the prevalence of Serious Mental Illness (SMI) among adults (age 18+).

For the 2010 NSDUH, no questionnaire changes are proposed.

As with all NSDUH/NHSDA surveys conducted since 1999, the sample size of the survey for 2010 will be sufficient to permit prevalence estimates for each of the fifty states and the District of Columbia. The total annual burden estimate is shown below:

	Number of responses	Responses per respondent	Average burden per response (hr.)	Total burden (hrs)
Household Screening	190,800	1	.083	15,836
Interview	67,500	1	1.0	67,500
Clinical Follow-up Interview	500	1	1.0	500
Screening Verification	5,400	1	0.067	362
Interview Verification	10,125	1	0.067	678
	190,800	84,876

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 7-1044, One Choke Cherry Road, Rockville, MD 20857 and e-mail her a copy at summer.king@samhsa.hhs.gov. Written comments should be received within 60 days of this notice.

Dated: April 1, 2009.
Elaine Parry,
Director, Office of Program Services.
 [FR Doc. E9-7788 Filed 4-6-09; 8:45 am]
BILLING CODE 4162-20-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Aging

Agency Information Collection Activities; Proposed Collection; Comment Request; Semi-Annual and Final Reporting Requirements for Older Americans Act Title IV Discretionary Grant Program

AGENCY: Administration on Aging, HHS.

ACTION: Notice.

SUMMARY: The Administration on Aging (AoA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. 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 of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information

collection requirements relating to the continuation of an existing collection for Performance Progress Reports for Older Americans Act Title IV grantees.

DATES: Submit written or electronic comments on the collection of information by May 31, 2009.

ADDRESSES: Submit electronic comments on the collection of information to:

lori.stalbaum@aoa.hhs.gov.

Submit written comments on the collection of information to Lori Stalbaum, Administration on Aging, Washington, DC 20201 or by fax to Lori Stalbaum at 202-357-3469.

FOR FURTHER INFORMATION CONTACT: Lori Stalbaum at 202-357-3452 or *lori.stalbaum@aoa.hhs.gov.*

SUPPLEMENTARY INFORMATION: 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. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request 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 (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The Administration on Aging (AoA) plans to continue an existing approved collection of information for semi-annual and final reports pursuant to requirements in Title IV of the Older Americans Act. Through its Title IV

program, AoA supports projects for the purpose of developing and testing new knowledge and program innovations with the potential for contributing to the well-being of older Americans. Deliverables required by AoA of all Title IV grantees are semi-annual and final reports, as provided for in the Department of Health and Human Services regulations, 45CFR Part 74, Section 74.51. These Title IV grantee performance reporting requirements can be found on AoA's Web site at http://www.aoa.gov/AoARoot/Grants/Reporting_Requirements/docs/FinalReportHandbook.doc. AoA estimates the burden of this collection of information as follows: *Frequency:* Semi-annually with the Final report taking the place of the semi-annual report at the end of the final year of the grant. *Respondents:* States, public agencies, private nonprofit agencies, institutions of higher education, and organizations including tribal organizations. *Estimated Number of Responses:* 600. *Total Estimated Burden Hours:* 12,000.

Dated: April 1, 2009.

Edwin L. Walker,

Acting Assistant Secretary for Aging.

[FR Doc. E9-7847 Filed 4-6-09; 8:45 am]

BILLING CODE 4154-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

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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: National Evaluation of the Comprehensive Community Mental Health Services for Children and Their Families Program: Phase V (OMB No. 0930-0280)—Revision

The Substance Abuse and Mental Health Services Administration (SAMHSA), Center for Mental Health Services (CMHS), is responsible for the National Evaluation of the comprehensive Community Mental Health Services for Children and Their Families Program, which collects data on child mental health outcomes, family life, and service system development and performance. Data will be collected on 30 service systems and roughly 8,810 children and families.

The data collection for this evaluation will be conducted for a three-year period. The core of service system data will be collected twice (every 18 to 24 months) during the three-year evaluation period. A sustainability survey will be conducted in selected years. Service delivery and system variables of interest include the following: maturity of system of care development; adherence to the system of care program model; services received by youth and their families, and the costs of those services; and consumer service experience.

The length of time that individual families will participate in the study ranges from 18 to 36 months depending on when they enter the evaluation. Child and family outcomes of interest will be collected at intake and during subsequent follow-up interviews at six-month intervals. Client service experience information is collected at these follow-up interviews. Measures included in an outcome interview are determined by the type of assessment (intake or follow-up), child's age, and whether the respondent is the caregiver or a youth.

The outcome measures include the following: Child symptomatology and functioning, family functioning, material resources, and caregiver strain. The caregiver interview package includes the Caregiver Information Questionnaire, Child Behavior Checklist, Behavioral and Emotional Rating Scale (BERS), Education Questionnaire, Columbia Impairment Questionnaire, Living Situations Questionnaire, Family Life Questionnaire, and Caregiver Strain Questionnaire at intake, and also