

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE) within the Administration for Children and Families (ACF) is proposing an extension with revisions to the data collection activities conducted as part of the National Survey of Child and Adolescent Well-Being (NSCAW III) (Office of Management and Budget (OMB) #0970–0202). NSCAW is the only source of nationally representative, longitudinal, firsthand information about the functioning and well-being, service needs, and service utilization of children and families who come to the attention of the child welfare system. This request will allow additional time to conduct participant data collections. Minor changes to the instruments are requested to restore an in-person data collection option.

DATES: Comments due within 30 days of publication. OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

ADDRESSES: Written 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. You can also obtain copies of the proposed collection of information by emailing OPREinfocollection@acf.hhs.gov. Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: NSCAW is the only source of nationally representative, longitudinal, firsthand information about the functioning and well-being, service needs, and service utilization of children and families who come to the attention of the child welfare system. The first and second cohorts of NSCAW were initiated in 1999 and 2008, respectively. A major objective for the third cohort of NSCAW [NSCAW III] is to maintain the strengths of previous work, while better positioning the study to address the changing child welfare population. Phase I of NSCAW III, approved November 2016, is complete and included recruitment and sampling process data collection activities. Phase II of NSCAW III, approved July 2017,

includes baseline and follow-up data collection activities, and panel maintenance activities. Phase II follow-up data collection and panel maintenance is still ongoing. Phase III of NSCAW III, approved in September 2020, includes data collection on the child welfare workforce in of participating agencies. Phase III data collection is complete, and analysis of the data is ongoing.

We seek approval for an extension with changes for the currently approved data collection activities, which includes follow-up data collection for Phase II and panel maintenance activities with NSCAW cohort members. As part of this request we are also proposing minor changes to the Phase II information collection. During the COVID–19 pandemic, the in-person option for data collection was removed. We are requesting to restore the previously approved in-person mode as an option for caregiver and child respondents for Phase II data collection.

Respondents: Children and caregivers enrolled in NSCAW III and child welfare agency personnel in participating NSCAW III agencies. Surveys and panel maintenance responses may be obtained by telephone, web, or in person.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Child Follow-up	387	1	.75	290	97
Caregiver Follow-up	409	1	.75	307	102
Caseworker Follow-up	126	3	1.0	379	126
Panel Maintenance with NSCAW Cohort Members	4,723	1	.08	378	126

Estimated Total Annual Burden Hours: 451.

Authority: 42 U.S.C. 628b; Continuing Appropriations Act of 2022.

Mary B. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: Rural Health Network Development Program Performance Improvement Measurement System, OMB No. 0906–0010—Revision

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than September 18, 2023.

ADDRESSES: Submit your comments to paperwork@hrsa.gov or mail the HRSA Information Collection Clearance Officer, Room 14N39, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email paperwork@hrsa.gov or call Samantha Miller, the HRSA Information Collection Clearance Officer, at (301) 443-3983.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the ICR title for reference.

Information Collection Request Title: Rural Health Network Development Program Performance Improvement Measurement System, OMB No. 0906-0010—Revision.

Abstract: The Rural Health Network Development (RHND) program is authorized under section 330A(f) of the Public Health Service Act (42 U.S.C. 254c(f)). The purpose of this program is to support integrated health care networks that collaborate to achieve efficiencies; expand access to, coordinate, and improve the quality of basic health care services and associated health outcomes; and strengthen the rural health care system as a whole. The program supports networks as they address gaps in service, enhance systems of care, and expand capacity of the local health care system.

RHND-funded programs promote population health management and the transition towards value-based care through diverse network participants that includes traditional and non-traditional network partners. Evidence of program impact demonstrated by outcome data and program sustainability are integral components

of the program. This is a 4-year competitive program for networks composed of at least three participants that are existing health care providers. At least 66 percent of network participants must be located in a HRSA-designated rural area.

HRSA currently collects information about RHND awards using an OMB-approved set of performance measures and seeks to revise that approved collection. The proposed revisions are being implemented to better gather award recipient data in response to previously accumulated award recipient feedback, peer-reviewed research, and information gathered from the previously approved RHND measures.

Need and Proposed Use of the Information: This program needs measures that will enable HRSA to provide aggregate program data required by Congress under the Government Performance and Results Act of 1993. These measures cover the principal topic areas of interest to HRSA, including (a) access to care, (b) population demographics, (c) staffing, (d) consortium/network, (e) sustainability, and (f) project specific domains. All measures will evaluate HRSA’s progress toward achieving its goals.

The proposed changes include additional components under questions surrounding the network’s benefits and funding strategies, as well as the types of participant organizations. Questions surrounding Health Information Technology and Telehealth have been modified to reflect an updated telehealth definition based on renewed knowledge on the use of both Health

Information Technology and Telehealth, and to improve understanding of how these important technologies are affecting HRSA award recipients. The Demographics and Services section now includes a question requesting grantees to identify which counties they have served during the project. Finally, revised National Quality Forum and Centers for Medicare & Medicaid Services measures were included to allow uniform collection efforts throughout the HRSA Federal Office of Rural Health Policy. The total number of responses has remained at 44 since the previous ICR. The new RHND grant cycle maintained the same number of award recipients and number of respondents.

Likely Respondents: Respondents will be award recipients of the Rural Health Network Development Program.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Performance Improvement and Measurement System Database	44	1	44	6	264
Total	44	1	44	6	264

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information

technology to minimize the information collection burden.

Maria G. Button,

Director, Executive Secretariat.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Advisory Council on Alzheimer’s Research, Care, and Services; Meeting

AGENCY: Assistant Secretary for Planning and Evaluation, HHS.

ACTION: Notice of meeting.

SUMMARY: This notice announces the public meeting of the Advisory Council on Alzheimer’s Research, Care, and