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Dated: November 30, 2023.

William N. Parham, III,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for (Office of Management and Budget (OMB) Review; Sexual Risk Avoidance Education Program Performance Analysis Study—Extension OMB #0970-0536)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE) and the Family and Youth Services Bureau in the Administration for Children and Families (ACF) request an extension without changes to a currently approved information collection activity as part of the Sexual Risk Avoidance Education (SRAE) Program Performance Analysis Study (PAS). The goal of the study is to collect, analyze, and report on performance measures data for the SRAE program (OMB Control No. 0970-0536; expiration date 12/31/2023). The

purpose of the requested extension is to continue the ongoing data collection and submission of the performance measures by SRAE grantees. Materials under the submission will be updated to reflect only surveys currently in use.

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: The purpose of the SRAE program is to educate youth on how to voluntarily refrain from nonmarital sexual activity and prevent other youth risk behaviors. Data will continue to be used to determine if the SRAE grantees are meeting performance benchmarks related to their program’s mission and priorities.

The SRAE PAS collects performance measures data from SRAE grantees,

program providers, and participants. The data include information on program structure, cost, and support for implementation; program attendance, reach, and dosage; the characteristics of youth involved in programming; youth sexual and other risky behavior prior to program participation; and youth sexual and other risky behavior intentions at program exit. The performance measures help the ACF program office and grantees to monitor and report on progress in implementing SRAE programs and inform technical assistance.

Some of the performance measures data come from youth participants through surveys SRAE grantees administer at program entry and exit. There are separate versions of the entry and exit surveys for middle school youth, which exclude some of the more sensitive items that are included in the versions for high school and older youth. There is also a shorter version of the entry survey for programs conducting impact studies, to reduce the burden on participants in those programs who are likely responding to other surveys as part of their impact study. Although there was a version of the exit survey for programs conducting impact studies in the past, youth in these programs complete the same version of the exit survey as other youth. As the shorter exit surveys are no longer in use, they will be removed through this request.

Respondents: General Departmental (GDSRAE), State (SSRAE), and Competitive (CSRAE) grantees, their subrecipients, and program participants.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
(1) Participant Entry Survey					
GDSRAE participants	378,390	1	0.1333	50,439	16,813
SSRAE participants	952,899	1	0.1333	127,021	42,340
CSRAE participants	60,408	1	0.1333	8,052	2,684
(2) Participant Exit Survey					
GDSRAE participants	302,712	1	0.1667	50,462	16,821
SSRAE participants	762,319	1	0.1667	127,079	42,360
CSRAE participants	48,326	1	0.1667	8,056	2,685
(3) Performance reporting data entry form: grantees					
GDSRAE grantees	119	6	16	11,424	3,808
SSRAE grantees	39	6	16	3,744	1,248
CSRAE grantees	34	6	16	3,264	1,088

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
(4) Performance reporting data entry form: subrecipients					
GDSRAE subrecipients	252	6	13	19,656	6,552
SSRAE subrecipients	426	6	13	33,228	11,076
CSRAE subrecipients	63	6	13	4,914	1,638

Estimated Total Annual Burden Hours: 149,113.

Authority: 42 U.S.C 1310.

Mary B. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0482]

Agency Information Collection Request; 60-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before February 5, 2024.

ADDRESSES: Submit your comments to Sherrette.Funn@hhs.gov or by calling (202) 264–0041 and PRA@HHS.GOV.

FOR FURTHER INFORMATION CONTACT:

When submitting comments or requesting information, please include the document identifier 0990–0482–60D and project title for reference, to Sherrette A. Funn, email: Sherrette.Funn@hhs.gov, PRA@HHS.GOV or call (202) 264–0041 the Reports Clearance Officer.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (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.

Title of the Collection: Continued Evaluation of the National Hypertension Control Initiative.

Type of Collection: Revision.

OMB No.: 0990–0482–OS/Office of Assistant Secretary for Health (OASH)/Office of Minority Health (OMH).

Abstract: As part of the federal response to COVID–19, the U.S. Department of Health and Human Services (HHS)/Office of Secretary (OS)/Office of Assistant Secretary for Health (OASH)/Office of Minority Health (OMH) has funded a new initiative involving two cooperative agreements with the American Heart Association (AHA) to improve COVID–19-related health outcomes by addressing hypertension (high blood pressure) among racial and ethnic minority populations.

The \$32 million project from the HHS Office of Minority Health (OMH) and the Health Resources and Services Administration (HRSA) Bureau of Primary Health Care will support the implementation of the National Hypertension Control Initiative (NHCI), a national initiative to improve blood pressure control among the most at-risk populations, including racial and ethnic minorities.

The NHCI will support 350 participating HRSA-funded health centers by providing patient and provider education and training for effective hypertension control and integration of remote blood pressure monitoring technology into treating hypertension for patients served by participating health centers. The project will also utilize the American Heart Association's targeted media campaigns and existing partnerships with community-based organizations (CBOs) to help reach Black, Latino, and other impacted communities with (i) culturally and linguistically appropriate messages, (ii) access to blood pressure screenings, and (iii) connection to

health centers to encourage proper treatment and management of hypertension of screened individuals. This initiative serves to increase the number of adult patients with controlled hypertension and reduce the potential risk of COVID-related health outcomes.

AHA aims to conduct an evaluation to assess the feasibility of the implementation of each of the three NHCI strategies. The findings of this evaluation will inform the improvement and tailoring of AHA's communication approaches about the importance of and techniques for improving blood pressure control, including the benefits of accurately measuring, rapidly acting, and having a patient-focused approach to blood pressure control.

Methodology

The current proposed evaluation of the NHCI project will use a mixed methods design, integrating both quantitative and qualitative data collection and analyses. Three main goals of data collection will be to: (1) track and monitor Community Health Workers' (CHW) progress on activities related to knowledge and practices for blood pressure control and general health quarterly, (2) assess the reach and success of NHCI project strategies implemented by CHC partners.

Specifically, the AHA will engage in:

Primary Data Collection

CHW Application. Collecting information on participating Community Health Workers (CHWs) at a single point in time to assist with placement in workforce activities related to blood pressure control.

CHW Assessment Form. Monitoring the placement and community-based goals of CHWs participating in the NHCI at a single point in time.

CHW Empowered To Serve (ETS) Program Modules. Administering health lessons and quizzes to Community Health Workers (CHWs) working with Community-based Organizations and Community Health Centers to assess