DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB #: 0970–0536]

Submission for Office of Management and Budget Review; Sexual Risk Avoidance Education Program Performance Analysis Study

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, U.S. 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 (FYSB) in the Administration for Children and Families (ACF) request approval for a temporary extension of currently approved information collection activities and revisions to be implemented in 2026 for the Sexual Risk Avoidance Education (SRAE) Program Performance Analysis Study (PAS)(Office of Management and Budget (OMB) #: 0970–0536; expiration date December 31, 2025). The goal of the study is to collect, analyze and report on performance measures data for the SRAE program. The purpose of the request is to continue the ongoing data collection and submission of the performance measures by SRAE grant recipients. Revisions are proposed to the current performance measures, for implementation in July 2026, to address feedback from grant recipients to simplify and clarify participant surveys and to ensure the measures meet FYSB data needs.

DATES: *Comments due* January 16, 2026. OMB must decide 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: The public may view and comment on this information collection request at: *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202512-0970-004.* 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 grant recipients are meeting their programs' mission and priorities.

The SRAE performance measures are collected from SRAE grant recipients, 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 grant recipients 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 grant recipients 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, it was removed through the previous OMB request, and youth in these programs now complete the same version of the exit survey as other youth.

For continuity, ACF proposes to use the currently approved materials through June 2026, after which revised versions of the performance measures would be implemented. This phased in approach will ease the transition for grant recipients by providing them with ample time for implementation and will allow for data consistency by not changing measures in the middle of a data collection year. The proposed revisions to the current performance measures address feedback from grant recipients to simplify and clarify participant surveys and ensure the measures meet FYSB data needs. The proposed revisions to the participant surveys were cognitively tested with program participants for clarity and to check burden estimates. The changes reduce the burden for completing the participant entry survey from 8 minutes to 5 minutes per response and the participant exit survey from 10 to 7 minutes. Overall, we expect a 61 percent reduction in the annual burden hours under this request compared to the previously approved annual burden.

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

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Annual burden (in hours)
Burden Estimates—Through June 2026				
1. Participant Entry Survey	104,606	1	0.1333	13,944
2 Participant Exit Survey	81,854	1	0.1667	13,645
3. Performance Reporting Data Entry Form	190	1	16	3,040
4. Subrecipient Data Collection and Reporting Form	490	1	13	6,370
Estimated total and annual burden hour estimates through June 2026				36,999

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)
Annual Burden Estimates—July 2026 Forward					
1. Participant Entry Survey	523,030	1	0.0833	43,568	17,427
2. Participant Exit Survey	409,270	1	0.1167	47,762	19,105
3. Performance Reporting Data Entry Form	190	5	16	15,200	6,080
4. Subrecipient Data Collection and Reporting Form	490	5	13	31,850	12,740
Estimated total and annual burden hours July 2026 forward				138,380	55,352
Total Annual Burden Estimates					
Estimated total and annual burden hours				175,379	58,460

Authority: 42 U.S.C. 710(b)(6).

Mary C. 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: Countermeasures Injury Compensation Program—OMB No. 0915–0334—Revision

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

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Before 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 February 17, 2026.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, Room 13N82, 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: Countermeasures Injury Compensation Program—OMB No. 0915–0334—Revision

Abstract: This is a request for continued OMB approval of the information collection requirements for the Countermeasures Injury Compensation Program (CICP or Program). This request includes revisions to improve the efficiency of the information collection process and the quality of the information collected. The revisions streamline questions on the information collection documents and update CICP contact information. HRSA administers CICP in accordance with the Public Readiness and Emergency Preparedness Act (PREP Act) and CICP regulations at 42 CFR part 110. CICP is requesting continued approval for this information collection, which includes documents specified in CICP's regulations (42 CFR part 110).

The PREP Act authorized the establishment of the CICP and provides liability immunity to covered persons for claims of loss caused by, arising out of, relating to, or resulting from the administration or use of covered countermeasures for diseases, threats, and conditions identified in PREP Act declarations. The immunity extended in the PREP Act encourages the development, manufacture, testing, distribution, and administration/use of countermeasures (*e.g.,* vaccine, medication, device) when a disease, health condition, or other threat to health constitutes a public health emergency, or there is a credible risk that it may in the future constitute such an emergency.

Need and Proposed Use of the Information: CICP provides compensation to eligible individuals who suffer serious injuries or death directly caused by a covered countermeasure administered or used pursuant to a PREP Act Declaration or to their estates and/or to certain survivors. An individual who is an injured countermeasure recipient, the estate or survivor(s) of a deceased injured countermeasure recipient, or their representative is responsible for submitting the Request for Benefits (RFB) package, as well as the injured countermeasure recipient's medical records and supporting documentation. Individuals can apply at any time, but eligibility for compensation is subject to meeting applicable filing deadlines and other requirements.

To determine whether a requester is eligible for Program benefits (compensation) for a countermeasure injury, CICP staff must review the RFB package, which includes the following:

(1) RFB Form and Supporting Medical Documentation: Submission of this RFB form and supporting medical documentation initiate the CICP claims review process. CICP assesses the RFB form and supporting medical documentation to gather required information about the requester, document the use or administration of a countermeasure, and obtain medical information about the countermeasure recipient.

(2) Authorization for Use or Disclosure of Health Information Form (Authorization Form): The requester or representative, if applicable, completes the Authorization Form and gives medical providers permission to disclose the countermeasure recipient's health information via medical records to CICP for the purpose of determining eligibility for CICP benefits.