before June 16, 2026. For information on the Commission's privacy policy, including routine uses permitted by the Privacy Act, see *https://www.ftc.gov/site-information/privacy-policy.*

Josephine Liu,

Assistant General Counsel for Legal Counsel.

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BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget #: 0970–0497]

Proposed Information Collection Activity; Personal Responsibility Education Program (PREP) Performance Measures

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 revision to a currently approved information collection activity as part of the Personal Responsibility Education Program (PREP) Performance Measures project (Office of Management and Budget (OMB) #: 0970–0497; expiration date July 31, 2026). The goal of the project is to collect, analyze, and report on performance measures data for the PREP-funded programs. The purpose of the request is to continue the ongoing data collection and submission of the performance measures by PREP grant recipients, with revisions to the current performance measures. We are proposing revisions to the current participant surveys and reporting forms to address feedback from grant recipients to simplify and clarify information collections, and to ensure the measures meet FYSB data needs while reducing burden.

DATES: *Comments due* June 16, 2026.

ADDRESSES: In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above. You can obtain copies of the proposed collection of information and submit comments by emailing *opreinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The purpose of PREP is to provide grants to states, tribes and tribal communities, and community organizations to support evidence-based programs to reduce teen pregnancy and sexually transmitted infections. The programs are required to provide education on both abstinence and contraceptive use. The programs also offer information on adulthood preparation subjects such as healthy relationships, adolescent development, financial literacy, parent–child communication, education and employment skills, and healthy life skills.

The PREP project collects performance measures data from PREP 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 risk behaviors and behaviors related to adulthood preparation prior to program participation; and youth behavior intentions at program exit. The performance measures help the ACF program office and grant recipients to monitor and report on progress in implementing PREP programs and inform technical assistance. In addition, ACF will use the information to continue fulfilling its reporting requirements to Congress and OMB concerning the PREP initiative.

Some of the performance measures data come from youth participants through surveys PREP 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 participants in the Personal Responsibility Education Innovative Strategies (PREIS) and Tribal PREP (TPREP) programs, to reduce the burden on participants in those programs (who are likely responding to other surveys); youth in these programs complete the same version of the exit survey as other youth.

We are proposing revisions to the current performance measures to address feedback from grant recipients and to ensure the measures meet FYSB data needs. Grant recipients have requested various changes to simplify and clarify the measures, including noting particular questions in the participant surveys that youth have difficulty understanding and responding to. In addition, contractor staff have noted which measures most frequently result in help desk contacts or are prone to data quality issues. Finally, FYSB staff identified measures that are no longer needed or could be obtained from other sources. Types of revisions include re-wording to use simpler language, reducing the number of sub-items or response categories, avoiding patterns in which survey respondents need to skip one or more questions based on their response to an earlier question, and removing some measures entirely. The proposed revisions to the participant surveys were cognitively tested with program participants for clarity and to check burden estimates.

Respondents: State PREP (SPREP), TPREP, Competitive PREP (CPREP), and PREIS grant recipients, their subrecipients, and program participants.

Annual Burden Estimates: The changes described above reduced the overall length of the surveys and are expected to reduce the burden for completing the participant entry survey from 8 minutes to 5 minutes per response and the participant exit survey from 7 minutes to 5 minutes per response. Additionally, the estimated number of respondents has been adjusted to reflect the estimated number over the next 3 years, which is reduced compared to previous estimates. Overall, we expect a 40 percent reduction in the annual burden hours under this request compared to the previously approved annual burden.

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)
Instrument 1					
Participant entry survey (all versions)	235,353	1	0.0833	19,605	6,535
Instrument 2					
Participant exit survey (all versions)	195,528	1	0.0833	16,287	5,429
Instrument 3: Performance Reporting System Data Entry Form					
State grant recipients	49	6	18	5,292	1,764
TPREP grant recipients	7	6	18	756	252
CPREP grant recipients	27	6	14	2,268	756
PREIS grant recipients	12	6	14	1,008	336
Instrument 4: Subrecipient Data Collection and Reporting Form					
State subrecipients	269	6	14	22,596	7,532
TPREP subrecipients	25	6	14	2,100	700
CPREP subrecipients	36	6	12	2,592	864
PREIS subrecipients	16	6	12	1,152	384
Estimated Total and Annual Burden Hours				73,656	24,552

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: 42 U.S.C. 1310.

Mary C. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2026–07499 Filed 4–16–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–3099]

Agency Information Collection Activities; Proposed Collection; Comment Request; Radioactive Drug Research Committees

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (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 information collection requirements contained in regulations governing the use of radioactive drugs for basic informational research.

DATES: Either electronic or written comments on the collection of information must be submitted by June 16, 2026.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of June 16, 2026. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for