

DEPARTMENT OF DEFENSE**GENERAL SERVICES
ADMINISTRATION****NATIONAL AERONAUTICS AND
SPACE ADMINISTRATION**

[OMB Control No. 9000–0136; Docket No. 2022–0053; Sequence No. 18]

**Information Collection; Commercial
Acquisitions**

AGENCY: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice and request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, and the Office of Management and Budget (OMB) regulations, DoD, GSA, and NASA invite the public to comment on an extension concerning commercial acquisitions. DoD, GSA, and NASA invite comments on: whether the proposed collection of information is necessary for the proper performance of the functions of Federal Government acquisitions, including whether the information will have practical utility; the accuracy of the estimate of the burden of the proposed information collection; ways to enhance the quality, utility, and clarity of the information to be collected; and ways to minimize the burden of the information collection on respondents, including the use of automated collection techniques or other forms of information technology. OMB has approved this information collection for use through January 31, 2023. DoD, GSA, and NASA propose that OMB extend its approval for use for three additional years beyond the current expiration date.

DATES: DoD, GSA, and NASA will consider all comments received by September 19, 2022.

ADDRESSES: DoD, GSA, and NASA invite interested persons to submit comments on this collection through <https://www.regulations.gov> and follow the instructions on the site. This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. If there are difficulties submitting comments, contact the GSA Regulatory Secretariat Division at 202–501–4755 or GSARegSec@gsa.gov.

Instructions: All items submitted must cite OMB Control No. 9000–0136, Commercial Acquisitions. Comments received generally will be posted without change to <https://www.regulations.gov>, including any

personal and/or business confidential information provided. To confirm receipt of your comment(s), please check www.regulations.gov, approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Zenaida Delgado, Procurement Analyst, at telephone 202–969–7207, or zenaida.delgado@gsa.gov.

SUPPLEMENTARY INFORMATION:**A. OMB Control Number, Title, and any
Associated Form(s)**

9000–0136, Commercial Acquisitions.

B. Need and Uses

This clearance covers the information that offerors must submit to comply with the following Federal Acquisition Regulation (FAR) requirements:

FAR 52.212–3, Offeror Representations and Certifications—Commercial Products and Commercial Services. Paragraph (b)(2) requires offerors to identify the applicable paragraphs at (c) through (v) of this provision that the offeror has completed for the purposes of the relevant solicitation only, if any. The provision stipulates that any changes provided by the offeror under paragraph (b)(2) are applicable to that specific solicitation only, and do not result in an update to the representations and certifications posted electronically in the System for Award Management. The contracting officer will use the information to determine a contractor's eligibility for award, and to incorporate appropriate terms and conditions into the contract award.

C. Annual Burden

Respondents: 140,055.

Total Annual Responses: 414,909.

Total Burden Hours: 207,455.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division by calling 202–501–4755 or emailing GSARegSec@gsa.gov. Please cite OMB Control No. 9000–0136, Commercial Acquisitions.

Janet Fry,

*Director, Federal Acquisition Policy Division,
Office of Governmentwide Acquisition Policy,
Office of Acquisition Policy, Office of
Governmentwide Policy.*

[FR Doc. 2022–15362 Filed 7–18–22; 8:45 am]

BILLING CODE 6820–EP–P

**DEPARTMENT OF HEALTH AND
HUMAN SERVICES****Agency for Healthcare Research and
Quality****Agency Information Collection
Activities: Proposed Collection;
Comment Request**

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) re-approve the proposed information collection project “*Online Submission Form for Supplemental Evidence and Data for Systematic Reviews for the Evidence-based Practice Center Program*.”

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by email at doris.lefkowitz@AHRQ.hhs.gov.

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at doris.lefkowitz@AHRQ.hhs.gov.

SUPPLEMENTARY INFORMATION:**Proposed Project**

“*Online Submission Form for Supplemental Evidence and Data for Systematic Reviews for the Evidence-based Practice Center Program*”

This is an ongoing activity of AHRQ's Evidence-based Practice Center (EPC) Program.

AHRQ's EPC Program develops evidence reports and technology assessments on topics relevant to clinical and other health care organization and delivery issues—specifically those that are common, expensive, and/or significant for the Medicare and Medicaid populations. For example, recent reviews have focused on clinical conditions, such as “Radiation Therapy for Brain Metastases”; health delivery topics, such as “Transitions of Care From Pediatric to Adult Services for Children With Special Healthcare Needs”; and specific technologies, such as “Telehealth for Women's Preventive Services.” These evidence reports include systematic reviews, technical briefs, and rapid reviews; and provide an essential foundation from which to

understand what we know from existing research and what critical research gaps remain. These reports, reviews, and technology assessments are based on rigorous, comprehensive syntheses and analyses of the scientific literature on topics. EPC reports and assessments emphasize explicit and detailed documentation of methods, rationale, and assumptions. EPC reports are conducted in accordance with an established policy on financial and nonfinancial interests. These scientific syntheses may include meta-analyses and cost analyses.

The EPC Program supports AHRQ’s mission by synthesizing and disseminating the available research as a “science partner” with private and public organizations in their efforts to improve the quality, effectiveness, and appropriateness of health care. The EPC Program is a trusted source of rigorous, comprehensive, and unbiased evidence reviews for stakeholders. The resulting evidence reports and technology assessments are used by Federal and State agencies, private-sector professional societies, health delivery systems, providers, payers, and others committed to evidence-based health care. These end-users may use EPC Program evidence reports to inform policy decisions, clinical practice guidelines, and other healthcare decisions.

This research has the following goals:

- Use research methods to gather knowledge on the effectiveness and harms of certain treatments and

healthcare delivery processes and models for medical conditions, both published and unpublished, to evaluate the quality of research studies and the evidence from these studies.

- Promote the use of evidence in healthcare decision making to improve healthcare and health.
- Identify research gaps to inform future research investments.

This study is being conducted by AHRQ pursuant to its statutory authority to conduct and support research on healthcare and on systems for the delivery of such care, including activities with respect to the quality, effectiveness, efficiency, appropriateness and value of healthcare services and with respect to quality measurement and improvement. 42 U.S.C. 299a(a)(1) and (2).

Method of Collection

To achieve the goals of this project the following data collection will be implemented:

- Online Submission Form. This information is collected for the purposes of providing supplemental evidence and data for systematic reviews (SEADS). The online submission form (OSF) collects data from respondents on their name, organization name, description of the submission, medical condition, intervention, and email address. For the purposes of meta-analyses, trial summary data from missing and unidentified studies are sought. For the purposes of constructing evidence tables and quality ratings (e.g., on public

reporting of cost measures or health information exchange), data can vary (e.g., URLs, study designs, and consumer-mediated exchange forms). Information on both completed and ongoing studies is requested. Submitters may alternatively email their submission to the AHRQ EPC mailbox at epc@ahrq.hhs.gov.

The EPC Program currently uses a broad-based announcement via email listserv and a **Federal Register** notice, as needed, to publicize the opportunity to submit scientific information about each topic. AHRQ plans to conduct one SEADS collection per topic. Up to twenty-four topics per year with SEADS portals are anticipated; over the past 5 years the number of SEADS portals has ranged from 11–20, with an average range of 0–5 potential respondents per topic. The EPC Program does not anticipate more than 40 topics per year with SEADS portals.

Estimated Annual Respondent Burden

Exhibit 1 presents estimates of the reporting burden hours for the data collection efforts. Time estimates are based on pilot testing of materials and what can reasonably be requested of respondents. The number of respondents listed in “Number of respondents” of Exhibit 1 reflects a projected upper range response rate per SEADS portal multiplied by the anticipated upper limit of number of SEADS portals per year, based on historical information over the past 3 years.

EXHIBIT 1—ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Hours per response	Total burden hours per SEADS
Online Submission Form (OSF)	200	1	15/60	50
Total	200	1	15/60	50

EXHIBIT 2—ESTIMATED ANNUALIZED COST BURDEN

Form name	Number of respondents	Total burden hours	Average hourly wage rate *	Total cost burden
OSF	200	50	^a \$57.62	\$2,881
Total	200	50	57.62	2,881

* Occupational Employment Statistics, May 2021 National Occupational Employment and Wage Estimates United States, U.S. Department of Labor, Bureau of Labor Statistics. http://www.bls.gov/oes/current/oes_nat.htm#b29-0000.
^aBased on the mean wages for *Public Relations and Fundraising Managers, 11–2030*, the occupational group most likely tasked with completing the OSF.

Request for Comments

In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3520, comments on AHRQ’s information

collection are requested with regard to any of the following: (a) whether the proposed collection of information is necessary for the proper performance of

AHRQ’s health care research and health care information dissemination functions, including whether the information will have practical utility;

(b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) 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 upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Dated: July 14, 2022.

Marquita Cullom,
Associate Director.

[FR Doc. 2022-15373 Filed 7-18-22; 8:45 am]

BILLING CODE 4160-90-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; State Plan for the Temporary Assistance for Needy Families (TANF) (OMB #: 0970-0145)

AGENCY: Office of Family Assistance, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families (ACF) is requesting a 3-year extension of the State Plan for the Temporary Assistance for Needy Families (TANF) (TANF State Plan; OMB #0970-0145, expiration 5/31/2022). There are no changes requested to this information collection.

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 infocollection@acf.hhs.gov. Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The TANF State Plan is a mandatory statement submitted to the Secretary of the Department of Health and Human Services by the state. It consists of an outline specifying how the state's TANF program will be administered and operated and certain required certifications by the state's Chief Executive Officer. It is used to provide the public with information about the program.

Authority to require states to submit a state TANF plan is contained in section 402 of the Social Security Act, as amended by Public Law 104-193, the Personal Responsibility and Work Opportunity Reconciliation Act of 1996. States are required to submit new plans within a 27-month period.

Respondents: The 50 States of the United States, the District of Columbia, Guam, Puerto Rico, and the Virgin Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents per year	Total number of annual responses per respondent	Average burden hours per response	Annual burden hours
Title Amendments	18	1	3	54
State TANF plan	18	1	30	540

Estimated Total Annual Burden Hours: 594.

Authority: Section 402 of the Social Security Act (42 U.S.C. 602), as amended by Pub. L. 104-193, the Personal Responsibility and Work Opportunity Reconciliation Act of 1996.

Mary B. Jones,
ACF/OPRE Certifying Officer.

[FR Doc. 2022-15332 Filed 7-18-22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

[OMB No. 0985-0036]

Agency Information Collection Activities; Proposed Collection; Comment Request; Prevention and Public Health Fund Evidence-Based Chronic Disease Self-Management Education Program Information Collection

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to

comment on the proposed collection of information listed above. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish a 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 proposed extension and solicits comments on the information collection requirements related to ACL's Prevention and Public Health Fund Evidence-Based Chronic Disease Self-Management Education Program Information Collection.

DATES: Comments on the collection of information must be submitted