

## GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0080]  
[Docket No. 2024–0001; Sequence No. 4]

### Submission for OMB Review; General Services Administration Acquisition Regulation; Release of Claims for Construction and Building Service Contracts

**AGENCY:** Office of Acquisition Policy, General Services Administration (GSA).

**ACTION:** Notice.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection regarding release of claims for final payment under construction and building services contracts.

**DATES:** Submit comments on or before: September 3, 2024.

**ADDRESSES:** Written comments and recommendations for this information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under Review—Open for Public Comments” or by using the search function.

**FOR FURTHER INFORMATION CONTACT:** Mr. Bryon Boyer, Procurement Analyst, at [gsarpolicy@gsa.gov](mailto:gsarpolicy@gsa.gov) or 817–850–5580.

#### SUPPLEMENTARY INFORMATION:

##### A. Purpose

The General Services Administration Acquisition Regulation (GSAR) requires construction and building services contractors to submit a release of claims before final payment is made to ensure contractors are paid in accordance with their contract requirements and for work performed. GSA Form 1142, Release of Claims is used to achieve uniformity and consistency in the release of claims process.

##### B. Annual Reporting Burden

*Respondents:* 1,427.  
*Responses per Respondent:* 1.  
*Annual Responses:* 1,427.  
*Hours per Response:* 0.50.  
*Total Burden Hours:* 714.

##### C. Public Comments

A 60-day notice was published in the **Federal Register** at 89 FR 42470 on May 15, 2024. No comments were received.

*Obtaining Copies of Proposals:* Requesters may obtain a copy of the information collection documents from

the Regulatory Secretariat Division (MVCB), at [GSARegSec@gsa.gov](mailto:GSARegSec@gsa.gov). Please cite OMB Control No. 3090–0080; Release of Claims for Construction and Building Service Contracts, in all correspondence.

Jeffrey A. Koses,

*Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.*

[FR Doc. 2024–16981 Filed 7–31–24; 8:45 am]

**BILLING CODE 6820–61–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

#### Supplemental Evidence and Data Request on Blood-Based Tests for Multiple Cancer Screening: A Systematic Review

**AGENCY:** Agency for Healthcare Research and Quality (AHRQ), HHS.

**ACTION:** Request for supplemental evidence and data submission.

**SUMMARY:** The Agency for Healthcare Research and Quality (AHRQ) is seeking scientific information submissions from the public. Scientific information is being solicited to inform our review on *Blood-based Tests for Multiple Cancer Screening: A Systematic Review*, which is currently being conducted by the AHRQ’s Evidence-based Practice Centers (EPC) Program. Access to published and unpublished pertinent scientific information will improve the quality of this review.

**DATES:** *Submission Deadline* on or before September 3, 2024.

#### ADDRESSES:

*Email submissions:* [epc@ahrq.hhs.gov](mailto:epc@ahrq.hhs.gov).

*Print submissions:*

*Mailing Address:* Center for Evidence and Practice Improvement, Agency for Healthcare Research and Quality, ATTN: EPC SEADs Coordinator, 5600 Fishers Lane, Mail Stop 06E53A, Rockville, MD 20857.

*Shipping Address (FedEx, UPS, etc.):* Center for Evidence and Practice Improvement, Agency for Healthcare Research and Quality, ATTN: EPC SEADs Coordinator, 5600 Fishers Lane, Mail Stop 06E77D, Rockville, MD 20857.

#### FOR FURTHER INFORMATION CONTACT:

Kelly Carper, Telephone: 301–427–1656 or Email: [epc@ahrq.hhs.gov](mailto:epc@ahrq.hhs.gov).

**SUPPLEMENTARY INFORMATION:** The Agency for Healthcare Research and Quality has commissioned the

Evidence-based Practice Centers (EPC) Program to complete a review of the evidence for *Blood-based Tests for Multiple Cancer Screening: A Systematic Review*. AHRQ is conducting this review pursuant to Section 902 of the Public Health Service Act, 42 U.S.C. 299a.

The EPC Program is dedicated to identifying as many studies as possible that are relevant to the questions for each of its reviews. In order to do so, we are supplementing the usual manual and electronic database searches of the literature by requesting information from the public (e.g., details of studies conducted). We are looking for studies that report on *Blood-based Tests for Multiple Cancer Screening: A Systematic Review*. The entire research protocol is available online at: <https://effectivehealthcare.ahrq.gov/products/cell-free-dna/protocol>.

This is to notify the public that the EPC Program would find the following information on *Blood-based Tests for Multiple Cancer Screening: A Systematic Review* helpful:

- A list of completed studies that your organization has sponsored for this topic. In the list, please *indicate whether results are available on ClinicalTrials.gov along with the ClinicalTrials.gov trial number.*

- *For completed studies that do not have results on ClinicalTrials.gov, a summary, including the following elements, if relevant: study number, study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, primary and secondary outcomes, baseline characteristics, number of patients screened/eligible/enrolled/lost to follow-up/withdrawn/analyzed, effectiveness/efficacy, and safety results.*

- *A list of ongoing studies that your organization has sponsored for this topic. In the list, please provide the ClinicalTrials.gov trial number or, if the trial is not registered, the protocol for the study including, if relevant, a study number, the study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, and primary and secondary outcomes.*

- *Description of whether the above studies constitute ALL Phase II and above clinical trials sponsored by your organization for this topic and an index outlining the relevant information in each submitted file.*

Your contribution is very beneficial to the Program. Materials submitted must be publicly available or able to be made public. Materials that are considered confidential; marketing materials; study