Filed June 2, 2025 10 a.m. EST Through June 9, 2025 10 a.m. EST

Pursuant to CEQ Guidance on 42 U.S.C. 4332.

Notice: Section 309(a) of the Clean Air Act requires that EPA make public its comments on EISs issued by other Federal agencies. EPA's comment letters on EISs are available at: *https://cdxapps.epa.gov/cdx-enepa-II/public/action/eis/search.*

EIS No. 20250079, Final, BLM, NV, Spring Valley Mine Project, Review Period Ends: 07/14/2025, Contact: Robert Sevon 775–623–1502.

EIS No. 20250080, Draft, USAF, FL, SpaceX Starship-Super Heavy Operations at Cape Canaveral Space Force Station, Comment Period Ends: 07/28/2025, Contact: Ms. Molly Thrash 321–357–7050.

EIS No. 20250081, Draft Supplement, USN, HI, Surveillance Towed Array Sensor System Low Frequency Active Sonar Training and Testing in the Western North Pacific and Indian Oceans, Comment Period Ends: 07/28/2025, Contact: John Burke 808–471–1714.

Dated: June 9, 2025.

Nancy Abrams,

Associate Director, Office of Federal Activities.

[FR Doc. 2025–10807 Filed 6–12–25; 8:45 am]

BILLING CODE 6560–50–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0297; Docket No. 2025–0001; Sequence No. 7]

Submission for OMB Review; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

AGENCY: General Services Administration (GSA).

ACTION: Notice; request for comments.

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 requirement regarding the Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.

DATES: Submit comments on or before July 14, 2025.

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.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Camille Tucker, Office of Governmentwide Policy, GSA, at 202–255–1648, or via email at *customer.experience@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study.

This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management.

Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance.

Such data uses require more rigorous designs that address: the target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential non-response bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior fielding the study.

Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

B. Annual Reporting Burden

Respondents: 1,010,650.
Responses per Respondent: ~1.
Total Annual Responses: 1,010,650.
Hours per Response: ~.063445 hours.
Total Burden Hours: 128,120.

C. Public Comments

A 60-day notice was published in the **Federal Register** at 90 FR 15242 on April 9, 2025. No public comments were received.

Obtaining Copies of Proposals: 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. 3090–0297, Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery, in all correspondence.

William F. Clark,

Director, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2025–10831 Filed 6–12–25; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Patient Safety Organizations: Delistings for the Michigan Surgical Quality Collaborative and Proximie PSO

AGENCY: Agency for Healthcare Research and Quality (AHRQ), Department of Health and Human Services (HHS).

ACTION: Notice of delisting.

SUMMARY: The Patient Safety and Quality Improvement Final Rule (Patient Safety Rule) authorizes AHRQ, on behalf of the Secretary of HHS, to list as a patient safety organization (PSO) an entity that attests that it meets the statutory and regulatory requirements for listing. A PSO can be ''delisted'' by the Secretary if it is found to no longer meet the requirements of the Patient Safety and Quality Improvement Act of 2005 (Patient Safety Act) and Patient Safety Rule, when a PSO chooses to voluntarily relinquish its status as a PSO for any reason, or when a PSO's listing expires. AHRQ accepted a notification of proposed voluntary relinquishment from the Michigan Surgical Quality Collaborative, PSO number P0143, of its status as a PSO,

and has delisted the PSO accordingly. AHRQ delisted the Proximie PSO, PSO number P0244, due to its failure to correct a deficiency.

DATES: The delisting for Proximie PSO was effective at 12:00 Midnight ET (2400) on January 11, 2025. The delisting for Michigan Surgical Quality Collaborative was effective at 12:00 Midnight ET (2400) on April 21, 2025.

ADDRESSES: The directories for both listed and delisted PSOs are ongoing and reviewed weekly by AHRQ. Both directories can be accessed electronically at the following HHS website: *https://www.pso.ahrq.gov/listed.*

FOR FURTHER INFORMATION CONTACT: Cathryn Bach, Center for Quality Improvement and Patient Safety, AHRQ, 5600 Fishers Lane, MS 06N100B, Rockville, MD 20857; Telephone (toll free): (866) 403–3697; Telephone (local): (301) 427–1111; TTY (toll free): (866) 438–7231; TTY (local): (301) 427–1130; Email: *pso@ahrq.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Background

The Patient Safety Act, 42 U.S.C. 299b–21 to 299b–26, and the related Patient Safety Rule, 42 CFR part 3, published in the **Federal Register** on November 21, 2008 (73 FR 70732–70814), establish a framework by which individuals and entities that meet the definition of provider in the Patient Safety Rule may voluntarily report information to PSOs listed by AHRQ, on a privileged and confidential basis, for the aggregation and analysis of patient safety work product.

The Patient Safety Act authorizes the listing of PSOs, which are entities or component organizations whose mission and primary activity are to conduct activities to improve patient safety and the quality of health care delivery.

HHS issued the Patient Safety Rule to implement the Patient Safety Act. AHRQ administers the provisions of the Patient Safety Act and Patient Safety Rule relating to the listing and operation of PSOs. The Patient Safety Rule authorizes AHRQ to list as a PSO an entity that attests that it meets the statutory and regulatory requirements for listing. A PSO can be ''delisted'' if it is found to no longer meet the requirements of the Patient Safety Act and Patient Safety Rule, when a PSO chooses to voluntarily relinquish its status as a PSO for any reason, or when a PSO's listing expires. Section 3.108(d) of the Patient Safety Rule requires AHRQ to provide public notice when it removes an organization from the list of PSOs.

AHRQ has accepted a notification of proposed voluntary relinquishment from the Michigan Surgical Quality Collaborative to voluntarily relinquish its status as a PSO. Accordingly, the Michigan Surgical Quality Collaborative, PSO number P0143, was delisted effective at 12:00 Midnight ET (2400) on April 21, 2025.

Michigan Surgical Quality Collaborative has patient safety work product (PSWP) in its possession. The PSO will meet the requirements of section 3.108(c)(2)(i) of the Patient Safety Rule regarding notification to providers that have reported to the PSO and of section 3.108(c)(2)(ii) regarding disposition of PSWP consistent with section 3.108(b)(3). According to section 3.108(b)(3) of the Patient Safety Rule, the PSO has 90 days from the effective date of delisting and revocation to complete the disposition of PSWP that is currently in the PSO's possession.

Proximie PSO failed to submit a response to the findings of deficiency in the Notice of Proposed Revocation and Delisting by the deadline. As stated in the Patient Safety Rule at 42 CFR 3.108(a)(4)(iii), this failure means that the Notice of Proposed Revocation and Delisting becomes final as a matter of law. It also serves as the basis for AHRQ to revoke acceptance of Proximie PSO certifications for listing, remove Proximie PSO from the list of PSOs, and provide public notice in the **Federal Register** and on the AHRQ PSO website that Proximie PSO has been delisted for cause. Accordingly, the Proximie PSO, P0244, was delisted effective at 12:00 Midnight ET (2400) on January 11, 2025.

More information on PSOs can be obtained through AHRQ's PSO website at *http://www.pso.ahrq.gov.*

Dated: April 28, 2025.

Jeffrey P. Toven,

Executive Officer.

[FR Doc. 2025–10798 Filed 6–12–25; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10398 #37, #87, and #93]

Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit comments regarding our burden estimates or any other aspect of this collection of information, including: the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by June 27, 2025.

ADDRESSES: When commenting, please reference the applicable form number (CMS–10398 #____) and the OMB control number (0938–1148). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection