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 May 31, 2024. 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 April 8, 2024.

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. 9000–0047, Place of Performance. 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)

OMB Control No. 9000–0047, Place of Performance.

B. Need and Uses

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

- FAR 52.214–14, Place of Performance-Sealed Bidding. This FAR provision is prescribed for invitation for bids (*i.e.,* FAR part 14 procurements) where the Government did not specify the place of performance.
- FAR 52.215–6, Place of Performance.

This FAR provision is prescribed for solicitations, when contracting by negotiation (*i.e.,* FAR part 15 procurements), where the Government did not specify the place of performance.

Both provisions ask for identical information from bidders or offerors: whether or not they intend to use one or more plants or facilities located at a different address from the address of the bidder or offeror as indicated in their bid or offer. If the response indicates the intention to use plants or facilities located at a different location than the bidder's or offeror's address, the provisions require that bidders or offerors provide the address(es) of the other place(s) of performance, along with name and address of the owner and operator of such plant or facility (if other than the bidder or offeror).

Contracting officers use the place of performance and the owner of the plant or facility to—

(a) Determine prospective contractor responsibility;

(b) Determine price reasonableness;

(c) Conduct plant or source inspections; and

(d) Determine whether the prospective contractor is a manufacturer or a regular dealer.

C. Annual Burden

Respondents: 6,086.
Total Annual Responses: 964,331.
Total Burden Hours: 43,877.
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. 9000–0047, Place of Performance.

Janet Fry,

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

[FR Doc. 2024–02329 Filed 2–5–24; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Meeting for Software Developers on the Common Formats for Patient Safety Data Collection

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

ACTION: Notice of public meeting.

SUMMARY: AHRQ coordinates the development of sets of standardized definitions and formats (Common Formats) that make it possible to collect, aggregate, and analyze uniformly structured information about health care quality and patient safety for local, regional, and national learning. The Common Formats include technical specifications to facilitate the collection of electronically comparable data by Patient Safety Organizations (PSOs) and other entities. Additional information about the Common Formats can be obtained through AHRQ's PSO website at *https://pso.ahrq.gov/common-formats* and the PSO Privacy Protection Center's website at *https://www.psoppc.org/psoppc_web/publicpages/commonFormatsOverview.*

The purpose of this notice is to announce a meeting to discuss implementation of the Common Formats with software developers and other interested parties. This meeting is designed as an interactive forum where software developers can provide input on use of the formats. AHRQ especially requests participation by and input from those entities which have used AHRQ's technical specifications and implemented, or plan to implement, the Common Formats electronically.

DATES: The meeting will be held from 2:00 to 3:00 p.m. Eastern on Thursday, March 7, 2024.

ADDRESSES: The meeting will be held virtually.

FOR FURTHER INFORMATION CONTACT: Erofile Gripiotis, Program Analyst, Center for Quality Improvement and Patient Safety, AHRQ, 5600 Fishers Lane, 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 and Quality Improvement Act of 2005, 42 U.S.C. 299b–21 to 299b–26 (Patient Safety Act), and the related Patient Safety and Quality Improvement Final Rule, 42 CFR part 3 (Patient Safety Rule), published in the **Federal Register** on November 21, 2008, 73 FR 70731–70814, provide for the Federal listing of Patient Safety Organizations (PSOs), which collect, aggregate, and analyze confidential information (patient safety work product) regarding the quality and safety of health care delivery.

The Patient Safety Act requires PSOs, to the extent practical and appropriate,

to collect patient safety work product from providers in a standardized manner that permits valid comparisons of similar cases among similar providers. (42 U.S.C. 299b–24(b)(1)(F)). The Patient Safety Act also authorizes the development of data standards, known as the Common Formats, to facilitate the aggregation and analysis of non-identifiable patient safety data collected by PSOs and reported to the network of patient safety databases (NPSD). (42 U.S.C. 299b–23(b)). The Patient Safety Act and Patient Safety Rule can be accessed at: *http://www.pso.ahrq.gov/legislation/.*

AHRQ has issued Common Formats for Event Reporting (CFER) for three settings of care—hospitals, nursing homes, and community pharmacies. AHRQ has also issued Common Formats for Event Reporting—Diagnostic Safety (CFER–DS) designed for use in all healthcare settings.

Federally listed PSOs can meet the requirement to collect patient safety work product in a standardized manner to the extent practical and appropriate by using AHRQ's Common Formats. The Common Formats are also available in the public domain to encourage their widespread adoption. An entity does not need to be listed as a PSO or working with one to use the Common Formats. However, the Federal privilege and confidentiality protections only apply to information developed as patient safety work product by providers and PSOs working under the Patient Safety Act.

Agenda, Registration, and Other Information About the Meeting

The Agency for Healthcare Research and Quality (AHRQ) will be hosting this fully virtual meeting to discuss implementation of the Common Formats with members of the public, including software developers and other interested parties. Agenda topics will include recent enhancement to the NPSD dashboards and data submission to the PSO Privacy Protection Center (PSOPPC). Active participation and discussion by meeting participants is encouraged.

AHRQ requests that interested persons send an email to *SDMeetings@infinityconferences.com* for registration information. Before the meeting, an agenda and logistical information will be provided to registrants.

Dated: January 31, 2024.

Marquita Cullom,

Associate Director.

[FR Doc. 2024–02257 Filed 2–5–24; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10552]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

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

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. 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 or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by March 7, 2024.

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.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Implementation of Medicare Programs;—Medicare Promoting Interoperability Program; *Use:* The Centers for Medicare & Medicaid Services (CMS) is requesting approval to collect information from eligible hospitals and critical access hospitals (CAHs). We have finalized changes to this program as discussed in the FY 2024 Inpatient Prospective Payment System (IPPS)/Long-term Care Hospital Prospective Payment System (LTCH PPS) final rule. This is a revision of the information collection request.

The American Recovery and Reinvestment Act of 2009 (Recovery Act) (Pub. L. 111–5) was enacted on February 17, 2009. Title IV of division B of the Recovery Act amended titles XVIII and XIX of the Social Security Act (the Act) by establishing incentive payments to eligible professionals (EPs), eligible hospitals and CAHs, and Medicare Advantage (MA) organizations participating in the Medicare and Medicaid programs that adopt and successfully demonstrate meaningful use of certified EHR technology (CEHRT). These Recovery Act provisions, together with title XIII of division A of the Recovery Act, may be cited as the ''Health Information Technology for Economic and Clinical Health Act'' or the ''HITECH Act.''

The HITECH Act created incentive programs for EPs, eligible hospitals including CAHs, and MA organizations