providing SF 1444, Request for Authorization of Additional Classification and Rate, for the contractor and the Government to enter the recordkeeping and information collection data required by 29 CFR 5.5(a)(1)(ii) prior to transmitting the data to the Department of Labor.

**B. Annual Reporting Burden**

*Number of Respondents:* 3,831.
*Responses per Respondent:* 2.
*Total Annual Responses:* 7,662.
*Review time per response:* 0.5.
*Total Burden Hours:* 3,831.

**C. Public Comments**

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FAR, and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

*Obtaining Copies of Proposals:* Requester may obtain a copy of the justification from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0089, Request for Authorization of Additional Classification and Rate, SF 1444, in all correspondence.

Dated: August 29, 2017.

**Lorin S. Curit,**

*Director, Federal Acquisition Policy Division, Office of Government-wide Acquisition Policy, Office of Acquisition Policy, Office of Government-wide Policy.*

[FR Doc. 2017–18676 Filed 9–1–17; 8:45 am]

**BILLING CODE 6820–EP–P**

## GENERAL SERVICES ADMINISTRATION

**[OMB Control No. 3090–XXXX; Docket No. 2017–0001; Sequence 3]**

## Submission for OMB Review; Ombudsman Inquiry/Request Instrument

**AGENCY:** Office of Acquisition Policy, Office of the Procurement Ombudsman (OPO), General Services Administration (GSA).

**ACTION:** Notice of request for comments regarding a new request for an Office of Management and Budget (OMB) clearance.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the OMB a request to review and approve a new information collection requirement regarding OMB Control No: 3090–XXXX; Ombudsman Inquiry/Request Instrument. A notice was published in the **Federal Register** on May 19, 2017. No comments were received.

**DATES:** Submit comments on or before October 5, 2017.

**ADDRESSES:** Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: Office of Information and Regulatory Affairs of OMB, Attention: Desk Officer for GSA, Room 10236, NEOB, Washington, DC 20503. Additionally submit a copy to GSA by any of the following methods:

- *Regulations.gov: http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching for ''Information Collection 3090–XXXX; Ombudsman Inquiry/Request Instrument.'' Select the link ''Submit a Comment'' that corresponds with ''Information Collection 3090–XXXX; Inquiry/Request Instrument.'' Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–XXXX; Ombudsman Inquiry/Request Instrument'' on your attached document.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Sosa/IC 3090–XXXX; Office of the Ombudsman Inquiry/Request Instrument.

*Instructions:* Please submit comments only and cite Information Collection 3090–XXXX; Inquiry/Request Instrument, in all correspondence related to this collection. Comments received generally will be posted without change to *http://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 (except allow 30 days for posting of comments submitted by mail).

**FOR FURTHER INFORMATION CONTACT:** Ms. Millisa Gary, GSA Procurement/Task & Delivery Order Ombudsman, Office of Acquisition Policy, Office of the Ombudsman, GSA, at telephone 202–501–0699 or via email to *millisa.gary@gsa.gov.*

**SUPPLEMENTARY INFORMATION:**

**A. Purpose**

OPO wants to place an online intake Instrument on the GSA Ombudsman's Web page for receiving inquiries from vendors who are currently doing business with, or interested in doing business with GSA. The inquiries will be collected by the GSA Ombudsman and routed to the appropriate office for resolution and/or implementation in the case of recommendations for process or program improvements. Reporting of the data collected will help highlight thematic issues that vendors encounter with GSA acquisition programs, processes or policies, and identify areas where training is needed. The information collected will also assist in identifying and analyzing patterns and trends to help improve efficiencies and lead to improvements in current practices.

**B. Annual Reporting Burden**

*Maximum Potential Respondents:* 118.

*Responses per Respondent:* 1.

*Total Maximum Potential Annual Responses:* 118.

*Hours per Response:* .25.

*Total Burden Hours:* 29.5.

**C. Public Comments**

Public comments are particularly invited on: Whether this collection of information is necessary, whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

*Obtaining Copies of Proposals:* Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755.

Please cite OMB Control No. 3090–XXXX, Inquiry/Request Instrument, in all correspondence.

**Jeffrey A. Koses,**

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

[FR Doc. 2017–18675 Filed 9–1–17; 8:45 am]

**BILLING CODE 6820–61–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

### Patient Safety Organizations: Voluntary Relinquishment From the Quantros Patient Safety Center

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

**ACTION:** Notice of delisting.

**SUMMARY:** The Patient Safety Rule authorizes AHRQ, on behalf of the Secretary of HHS, to list as a 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 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 has accepted a notification of voluntary relinquishment from the Quantros Patient Safety Center of its status as a PSO, and has delisted the PSO accordingly.

**DATES:** The directories for both listed and delisted PSOs are ongoing and reviewed weekly by AHRQ. The delisting was applicable at 12:00 Midnight ET (2400) on August 15, 2017.

**ADDRESSES:** Both directories can be accessed electronically at the following HHS Web site: *http://www.pso.ahrq.gov/listed.*

**FOR FURTHER INFORMATION CONTACT:** Eileen Hogan, Center for Quality Improvement and Patient Safety, AHRQ, 5600 Fishers Lane, Room 06N94B, 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 b–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 70732–70814, establish a framework by which hospitals, doctors, and other health care providers may voluntarily report information to Patient Safety Organizations (PSOs), on a privileged and confidential basis, for the aggregation and analysis of patient safety events.

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 federally approved PSOs.

AHRQ has accepted a notification from the Quantros Patient Safety Center, a component entity of Quantros Inc., PSO number P0014, to voluntarily relinquish its status as a PSO. Accordingly, the Quantros Patient Safety Center was delisted effective at 12:00 Midnight ET (2400) on August 15, 2017.

The Quantros Patient Safety Center 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. More information on PSOs can be obtained through AHRQ's PSO Web site at *http://www.pso.ahrq.gov.*

**Sharon B. Arnold,**

*Deputy Director.*

[FR Doc. 2017–18707 Filed 9–1–17; 8:45 am]

**BILLING CODE 4160–90–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

**[60Day–17–0765; Docket No. CDC–2017–0062]**

### Proposed Data Collection Submitted for Public Comment and Recommendations

**AGENCY:** Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice with comment period.

**SUMMARY:** The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comments on a request for an extension of an approved information collection entitled, CDC's Fellowship Management System. CDC uses the information collected for processes that aid and enhance the selection of fellowship participants and host sites and to track participant information that helps strengthen the current, emerging, and ever-changing public health workforce.

**DATES:** Written comments must be received on or before November 6, 2017.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC–2017–0062 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

*Instructions:* All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov.*