

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: [psa@ahrq.hhs.gov](mailto:psa@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](http://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](http://Regulations.gov), including any personal information provided. For access to the docket to read background documents or comments received, go to [Regulations.gov](http://Regulations.gov).