

In order to assist AHRQ and CMS to assess the importance, validity, and feasibility of submitted measures, a Subcommittee on Children's Healthcare Quality Measures of the AHRQ National Advisory Council on Healthcare Research and Quality (SNAC) has been established (<http://www.ahrq.gov/chipra/panellist11.htm>). The Subcommittee will consider measures submitted through this public call, and measures submitted by the 7 AHRQ–CMS Centers of Excellence.

CHIPRA asks that measures in the improved core sets be: evidence-based; able to identify disparities by race, ethnicity, socioeconomic status, and special health care need; risk-adjusted as appropriate; and designed to ensure that data are collected and reported in a standard format that permits comparison of quality and data at a State, plan, and provider level.

Dated: February 15, 2012.

**Carolyn M. Clancy,**  
*AHRQ Director.*

[FR Doc. 2012–4267 Filed 2–23–12; 8:45 am]

**BILLING CODE 4160–90–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

#### Patient Safety Organizations: Voluntary Relinquishment From UAB Health System Patient Safety Organization

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

**ACTION:** Notice of Delisting.

**SUMMARY:** AHRQ has accepted a notification of voluntary relinquishment from the UAB Health System Patient Safety Organization of its status as a Patient Safety Organization (PSO). The Patient Safety and Quality Improvement Act of 2005 (Patient Safety Act), Public Law 109–41, 42 U.S.C. 299b–21–b–26, provides for the formation of PSOs, which collect, aggregate, and analyze confidential information regarding the quality and safety of health care delivery. The Patient Safety and Quality Improvement Final Rule (Patient Safety Rule), 42 CFR part 3, 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, including when a PSO chooses to

voluntarily relinquish its status as a PSO for any reason.

**DATES:** The directories for both listed and delisted PSOs are ongoing and reviewed weekly by AHRQ. The delisting was effective at 12:00 Midnight ET (2400) on January 13, 2012.

**ADDRESSES:** Both directories can be accessed electronically at the following HHS Web site: <http://www.pso.AHRQ.gov/index.html>.

#### FOR FURTHER INFORMATION CONTACT:

Susan Grinder, Center for Quality Improvement and Patient Safety, AHRQ, 540 Gaither Road, Rockville, MD 20850; 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 Act authorizes the listing of PSOs, which are entities or component organizations whose mission and primary activity is 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 (PDF file, 450 KB. PDF Help) relating to the listing and operation of PSOs. 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 UAB Health System Patient Safety Organization, PSO number P0042, which is a component entity of the UAB Health System to voluntarily relinquish its status as a PSO. Accordingly, the UAB Health System Patient Safety Organization was delisted effective at 12:00 Midnight ET (2400) on January 13, 2012.

More information on PSOs can be obtained through AHRQ's PSO Web site at <http://www.pso.AHRQ.gov/index.html>.

Dated: February 15, 2012.

**Carolyn M. Clancy,**  
*Director.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

#### Scientific Information Request on Treatment Strategies for Patients With Peripheral Artery Disease (PAD)

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

**ACTION:** Request for scientific information submissions

**SUMMARY:** The Agency for Healthcare Research and Quality (AHRQ) is seeking scientific information submissions from manufacturers of peripheral artery disease treatment medical devices. Scientific information is being solicited to inform our Comparative Effectiveness Review of Treatment Strategies for Patients with Peripheral Artery Disease (PAD), which is currently being conducted by the Evidence-based Practice Centers for the AHRQ Effective Health Care Program. Access to published and unpublished pertinent scientific information on this device will improve the quality of this comparative effectiveness review. AHRQ is requesting this scientific information and conducting this comparative effectiveness review pursuant to Section 1013 of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003, Public Law 108–173.

**DATES:** Submission Deadline on or before March 26, 2012.

#### ADDRESSES:

##### Online Submissions

<http://effectivehealthcare.AHRQ.gov/index.cfm/submitscientific-information-packets/>. Please select the study for which you are submitting information from the list of current studies and complete the form to upload your documents.

*Email submissions:* [ehcsrc@ohsu.edu](mailto:ehcsrc@ohsu.edu) (please do not send zipped files—they are automatically deleted for security reasons).

*Print submissions:* Robin Paynter, Oregon Health and Science University, Oregon Evidence-based Practice Center, 3181 SW. Sam Jackson Park Road, Mail Code: BICC, Portland, OR 97239–3098.

**FOR FURTHER INFORMATION CONTACT:** Robin Paynter, Research Librarian, Telephone: 503–494–0147 or Email: [ehcsrc@ohsu.edu](mailto:ehcsrc@ohsu.edu).

**SUPPLEMENTARY INFORMATION:** In accordance with Section 1013 of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003, Public Law 108–173, the Agency