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#### SUPPLEMENTARY INFORMATION:

##### Background

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 the 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 Close Care Gap, PSO, a component entity of iCareQuality, Inc., PSO number P0145, to voluntarily relinquish its status as a PSO. Accordingly, Close Care Gap, PSO was delisted effective at 12:00 Midnight ET (2400) on July 2, 2015.

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

Sharon B. Arnold,  
Deputy Director.

[FR Doc. 2015-20597 Filed 8-19-15; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Board of Scientific Counselors, National Center for Health Statistics (BSC, NCHS)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC), National Center for Health Statistics (NCHS)

announces the following meeting of the aforementioned committee.

#### TIMES AND DATES:

11:00 a.m.-5:30 p.m., September 24, 2015.

8:30 a.m.-1:00 p.m., September 25, 2015.

**PLACE:** NCHS Headquarters, 3311 Toledo Road, Hyattsville, Maryland 20782.

**STATUS:** This meeting is open to the public; however, visitors must be processed in accordance with established federal policies and procedures. For foreign nationals or non-U.S. citizens, pre-approval is required (please contact Gwen Mustaf, 301-458-4500, [glm4@cdc.gov](mailto:glm4@cdc.gov), or Virginia Cain, [vcain@cdc.gov](mailto:vcain@cdc.gov) at least 10 days in advance for requirements). All visitors are required to present a valid form of picture identification issued by a state, federal or international government. As required by the Federal Property Management Regulations, Title 41, Code of Federal Regulation, Subpart 101-20.301, all persons entering in or on Federal controlled property and their packages, briefcases, and other containers in their immediate possession are subject to being x-rayed and inspected. Federal law prohibits the knowing possession or the causing to be present of firearms, explosives and other dangerous weapons and illegal substances. The meeting room accommodates approximately 100 people.

**PURPOSE:** This committee is charged with providing advice and making recommendations to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NCHS, regarding the scientific and technical program goals and objectives, strategies, and priorities of NCHS.

**MATTERS FOR DISCUSSION:** The agenda will include:

1. Welcome remarks by the Director, NCHS
2. An update on health insurance coverage data
3. A presentation on the National Health and Nutrition Examination Survey (NHANES) and the development of nutritional guidelines
4. A presentation on accessing NCHS data

Requests to make oral presentations should be submitted in writing to the contact person listed below. All requests must contain the name, address, telephone number, and organizational affiliation of the presenter.

Written comments should not exceed five single-spaced typed pages in length

and must be received by September 11, 2015.

The agenda items are subject to change as priorities dictate.

#### CONTACT PERSON FOR MORE INFORMATION:

Virginia S. Cain, Ph.D., Director of Extramural Research, NCHS/CDC, 3311 Toledo Road, Room 7208, Hyattsville, Maryland 20782, telephone (301) 458-4500, fax (301) 458-4024.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Management Analysis and Services Office,  
Centers for Disease Control and Prevention.

[FR Doc. 2015-20553 Filed 8-19-15; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2015-N-0001]

#### Pediatric Advisory Committee; Notice of Meeting; Correction

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; correction.

**SUMMARY:** The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of August 12, 2015 (80 FR 48325). Phenylephrine Hydrochloride was incorrectly linked to DUREZOL (difluprednate ophthalmic emulsion) 0.05% because they were both listed as item number 1 in the numbered list of products to be discussed at the meeting. Phenylephrine Hydrochloride Ophthalmic Solution is a separate stand-alone drug that will be reviewed by the committee and should be listed as item number 2. The other drugs in the numbered list should be renumbered accordingly. This document corrects that error.

**FOR FURTHER INFORMATION CONTACT:** Lisa Granger, Office of Policy, Planning, Legislation, and Analysis, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 3330, Silver Spring, MD 20993-0002, 301-796-9115.

**SUPPLEMENTARY INFORMATION:** In FR Doc. 2015-19729, appearing on page 48325, in the **Federal Register** of Wednesday, August 12, 2015, the following correction is made: