

efficient and effective manner, and perform reviews of quality of care in an area of medical practice where actual performance is measured against objective criteria, which defines acceptable and adequate practice. The selected organization must have a consumer representative on its governing board.

The Omnibus Budget Reconciliation Act of 1987 (Pub. L. 100–203) amended section 1153 of the Act by adding paragraph (i). This provision prohibits CMS from renewing the contract of any QIO that is not an in-State QIO without first publishing in the **Federal Register** a notice announcing when the contract will expire. This notice must be published no later than 6 months before the date the contract expires and must specify the period of time during which an in-State organization may submit a proposal for the QIO contract for that State. If one or more qualified in-State organizations submit a proposal for the QIO contract within the specified period of time, we cannot automatically renew the current contract on a noncompetitive basis, but must instead provide for competition for the contract in the same manner used for a new contract under section 1153(b) of the Act. An in-State QIO is defined under section 1153(i)(3) of the Act as a QIO that has its primary place of business in the State in which review will be conducted (or, be a subsidiary of a parent corporation, whose headquarters is located in that State).

There are currently 4 QIO contracts with entities that do not meet the statutory definition of an in-State QIO. The areas affected for purposes of this notice along with the respective contract expiration dates are as follows:

Vermont—July 31, 2011
 Maine—July 31, 2011
 Idaho—July 31, 2011
 South Carolina—July 31, 2011

II. Provisions of the Notice

This notice announces the scheduled expiration dates of the current contracts between CMS and the out-of-State QIOs responsible for review in the areas mentioned above.

Interested in-State organizations may submit a proposal in competing to become the QIO for these States. In order to be eligible for contract award, the organization must have its primary place of business in the States in which review will be conducted or be a subsidiary of a parent corporation, whose headquarters is located in that State. In order to be eligible for contract award, each interested organization must further demonstrate that it meets the following requirements:

A. Be Either a Physician-Sponsored or a Physician-Access Organization

1. Physician-Sponsored Organization

a. The organization must be composed of a substantial number of the licensed doctors of medicine and osteopathy practicing medicine or surgery in the review area, who are representative of the physicians practicing in the review area.

b. The organization must not be a health care facility, health care facility association, health care facility affiliate, or payor organization. However, statutes and regulations provide that, in the event CMS determines no otherwise qualified non-payor organization is available to undertake a given QIO contract, CMS may select a payor organization which otherwise meets requirements to be eligible to conduct Utilization and Quality Control Peer Review as specified in Part B of Title XI of the Act and its implementing regulations.

c. In order to meet the “substantial number of doctors of medicine and osteopathy” requirements as specified above in paragraph A.1.a, an organization must state and have documentation in its files showing that it is composed of at least 10 percent of the licensed doctors of medicine and osteopathy practicing medicine or surgery in the review area. In order to meet the representation requirements as specified above in paragraph A.1.a, an organization must state and have documentation in its files demonstrating that it is composed of at least 20 percent of the licensed doctors of medicine and osteopathy practicing medicine or surgery in the review area. Alternatively, if the organization does not demonstrate that it is composed of at least 20 percent of the licensed doctors of medicine and osteopathy practicing medicine or surgery in the review area, the organization must demonstrate in its proposal, through letters of support from physicians or physician organizations, or through other means, that it is representative of the area physicians.

2. Physician-Access Organization

a. The organization must have available to it, by arrangement or otherwise, the services of a sufficient number of licensed doctors of medicine or osteopathy practicing medicine or surgery in the review area to ensure adequate peer review of the services furnished by the various medical specialties and subspecialties.

b. The organization must not be a health care facility, health care facility

association, health care facility affiliate, or payor organization.

c. An organization meets the requirements specified above in paragraph A.2.a., if it demonstrates that it has available to it at least one physician in every generally recognized specialty and has an arrangement or arrangements with physicians under which the physicians would conduct review for the organization.

B. Have at Least One Individual Who Is a Representative of Consumers on Its Governing Board

If one or more organizations meet the above requirements in a QIO area and submit proposals for the contracts in accordance with this notice, we will consider those organizations to be potential sources for the 4 contracts upon their expiration. These organizations will be entitled to participate in a full and open competition for the QIO contract to perform the QIO statement of work.

(Catalog of Federal Domestic Assistance Program No. 93.773, Medicare—Hospital Insurance Program; and No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: October 28, 2010.

Donald M. Berwick,

Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. 2010–28817 Filed 11–24–10; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)—Ethics Subcommittee (ES)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the CDC announces the following meeting of the aforementioned subcommittee:

Time and Date: 2 p.m.—3:30 p.m. Eastern Standard Time, January 4, 2011.

Place: Teleconference.

Status: Open to the public, limited only by availability of telephone ports. The public is welcome to participate during the public comment period. A public comment period is tentatively scheduled from 3 p.m.—3:15 p.m. To participate in the teleconference, please dial 1–877–928–1204 and enter conference code 4305992.

Purpose: The ES will provide counsel to the ACD, CDC, regarding a broad range of public health ethics questions and issues arising from programs, scientists and practitioners.

Matter to be Discussed: The ES will discuss a strategy for addressing its charge to provide a preliminary overview to the ACD on ethical issues related to non-communicable disease prevention and control and an ethics framework to guide CDC programs and activities.

The agenda is subject to change as priorities dictate.

Contact Person for More Information: Drue Barrett, Ph.D., Designated Federal Officer, ACD, CDC—ES, CDC, 1600 Clifton Road, NE., M/S D-50, Atlanta, Georgia 30333. Telephone (404) 639-4690. E-mail: d Barrett@cdc.gov.

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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: November 18, 2010.

Elaine L. Baker,

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

[FR Doc. 2010-29774 Filed 11-24-10; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0585]

Neurological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Neurological Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on January 27 and 28, 2011, from 8 a.m. to 6 p.m.

FDA is opening a docket for public comment on this meeting. The docket number is FDA-2010-N-0585. The docket will open for public comment on November 26, 2010. The docket will close on January 25, 2011. Interested persons may submit electronic or written comments regarding this meeting. Submit electronic comments to <http://www.regulations.gov>. Submit

written comments to the Division of Dockets Management, Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit a single copy of electronic comments or a paper copy of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this meeting notice. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Location: Hilton Washington DC North/Gaithersburg, Ballroom, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: James Engles, Food and Drug Administration, Center for Devices and Radiological Health, 10903 New Hampshire Ave., Silver Spring, MD 20993, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512513. Please call the Information Line for up-to-date information on this meeting. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

Agenda: On January 27 and 28, 2011, the committee will discuss and make recommendations regarding the possible reclassification of devices indicated for use in electroconvulsive therapy.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before January 14, 2011. Oral presentations from the public will be scheduled at approximately 10 a.m., immediately following the FDA's presentation, on January 27, 2011. Those individuals interested in making

formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before January 6, 2011. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by January 7, 2011.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets. FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, at 301-796-5966 at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: November 18, 2010.

Thinh Nguyen,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010-29824 Filed 11-24-10; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C.,