

verified from the grantee records to support the information outlined in the FSR.

Failure to submit required reports within the time allowed may result in suspension or termination of an active grant, withholding of additional awards for the project, or other enforcement actions such as withholding of payments or converting to the reimbursement method of payment. Continued failure to submit required reports may result in one or both of the following: (1) The imposition of special award provisions; and (2) the non-funding or non-award of other eligible projects or activities. This applies whether the delinquency is attributable to the failure of the grantee organization or the individual responsible for preparation of the reports.

C. FY 2007 and FY 2008 Single Audit Reports (OMB A-133)

Applicants who have an active SDPI grant are required to be up-to-date in the submission of required audit reports. These are the annual financial audit reports required by OMB A-133, audits of State, local governments, and non-profit organizations that are submitted. Documentation of (or proof of submission) of current FY 2007 and FY 2008 Financial Audit Reports is mandatory. Acceptable forms of documentation include: e-mail confirmation from FAC that audits were submitted; or face sheets from audit reports. Face sheets can be found on the FAC Web site: <http://harvester.census.gov/fac/dissemin/accessoptions.html?submit=Retrieve+Records>.

Telecommunication for the hearing impaired is available at: TTY (301) 443-6394.

## VII. Agency Contacts

- For Grants Budget Management, contact:
  - Denise Clark, Lead Grants Management Specialist, DGO ([denise.clark@ihs.gov](mailto:denise.clark@ihs.gov)), Division of Grants Operations, 801 Thompson Avenue, TMP, Suite 360, Rockville, MD 20852. (301) 443-5204.
  - For Grants.gov electronic application process, contact:
    - Tammy Bagley, Grants Policy, DGP ([tammy.bagley@ihs.gov](mailto:tammy.bagley@ihs.gov)), (301) 443-5204. Grants Policy Web site: [http://www.ihs.gov/NonMedicalPrograms/gogp/index.cfm?module=gogp\\_funding](http://www.ihs.gov/NonMedicalPrograms/gogp/index.cfm?module=gogp_funding).
  - For programmatic questions, contact:
    - Merle Mike, Program Assistant, DDTP ([merle.mike@ihs.gov](mailto:merle.mike@ihs.gov)), (505) 248-4182.

- Lorraine Valdez, Deputy Director, DDTP ([s.lorraine.valdez@ihs.gov](mailto:s.lorraine.valdez@ihs.gov)), (505) 248-4182.

- Area Diabetes Consultants Web site: <http://www.ihs.gov/MedicalPrograms/diabetes/index.cfm?module=peopleADCDirectory>.

Dated: December 22, 2009.

**Yvette Roubideaux,**

Director, Indian Health Service.

[FR Doc. 2010-149 Filed 1-8-10; 8:45 am]

**BILLING CODE 4165-16-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2009-N-0606]

#### General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee; Amendment of Notice

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an amendment to the notice of a meeting of the General and Plastic Surgery Devices Panel of the Medical Devices Advisory Committee. This meeting was announced in the **Federal Register** of November 17, 2009 (74 FR 59194). The amendment is being made to reflect a change in the *Contact Person* and *Procedure* portions of the document, and to provide notice of the availability of a docket for public comment. There are no other changes.

**FOR FURTHER INFORMATION CONTACT:** Margaret McCabe-Janicki, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., White Oak 66, rm. 1535, Silver Spring, MD 20993, 301-796-7029, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512519. Please call the Information Line for up-to-date information on this meeting.

**SUPPLEMENTARY INFORMATION:** In FR Doc. E9-27491, appearing on page 59194, in the **Federal Register** of Tuesday, November 17, 2009, the following corrections are made:

1. On page 59194, in the second column, under *Contact Person*, the first six lines “Peter L. Hudson, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., White Oak 66, rm. 3618, Silver Spring, MD 20993, 301-796-6440 or FDA Advisory” are removed and replaced with “Margaret

McCabe-Janicki, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Avenue., White Oak 66, rm. 1535, Silver Spring, MD 20993, 301-796-7029, or FDA Advisory”.

2. On page 59194, in the third column, a *Comments* portion is added before the *Agenda* portion of the document to read: “*Comments:* FDA is opening a docket for public comment on this document. The docket number is FDA-2009-N-0606. The docket will open for public comment on January 11, 2010. The docket will close on March 19, 2010. Interested persons may submit electronic or written comments regarding this document. 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 two paper copies 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 document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.”

3. On pages 59194 and 59195, beginning on page 59194 in the third column, under *Procedure*, the year “2009” is changed to read “2010” everywhere that it appears.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: January 5, 2010.

**David Dorsey,**

Acting Deputy Commissioner for Policy, Planning and Budget.

[FR Doc. 2010-172 Filed 1-8-10; 8:45 am]

**BILLING CODE 4160-01-S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2009-N-0664]

#### Cardiovascular and Renal Drugs 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:* Cardiovascular and Renal Drugs 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 March 1, 2010, from 8 a.m. to 5 p.m.

*Location:* Hilton Washington DC/ Silver Spring, Maryland Ballroom, 8727 Colesville Road, Silver Spring, MD. The hotel telephone number is 301-589-5200.

*Contact Person:* Elaine Ferguson, Center for Drug Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane, (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301-827-7001, FAX: 301-827-6776, e-mail: [elaine.ferguson@fda.hhs.gov](mailto:elaine.ferguson@fda.hhs.gov), or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 3014512533. 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 March 1, 2010, the committee will discuss biologics license application (BLA) 125288, for belatacept injectable, by Bristol Myers Squibb, to be used in patients with kidney transplants to prevent rejection of the transplanted kidney.

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 February 12, 2010. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those desiring to make

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 February 4, 2010. 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 February 5, 2010.

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 Elaine Ferguson 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: January 5, 2010.

**David Dorsey,**  
*Acting Deputy Commissioner for Policy,  
Planning and Budget.*

[FR Doc. 2010-173 Filed 1-8-10; 8:45 am]

**BILLING CODE 4160-01-S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2009-N-0664]

#### The General Hospital and Personal Use 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:* General Hospital and Personal Use 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 March 5, 2010, from 8 a.m. to 5 p.m.

*Location:* Hilton Washington DC North/Gaithersburg, Salons C, D and E, 620 Perry Pkwy., Gaithersburg, MD.

*Contact Person:* Tracy Phillips, Center for Devices and Radiological Health, Food and Drug Administration, 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 3014512520. 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 March 5, 2010, the committee will discuss and make recommendations regarding clinical risks and benefits of post-market actions in response to insulin pump failures. Insulin pumps are intended for continuous delivery of insulin at set and variable rates and as an aid in the management of diabetes mellitus in persons requiring insulin.

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 February 22, 2010. Oral presentations from the public will be scheduled immediately following