clearly address the following sections of the CFRs: § 418.52, § 418.54(a)(2), § 418.56(a), § 418.56(d), § 418.56(e), § 418.58, § 418.58(a)(2), § 418.58(c)(2), § 418.58(d)(1), § 418.60(b)(2)(ii), § 418.62(b), § 418.62(c), § 418.64(a)(1–3), § 418.64(b)(1), § 418.64(d)(3)(iv), § 418.72, § 418.76(a)(1), § 418.76(b)(3)(i), § 418.76(c), § 418.76(e), § 418.76(h)(1), § 418.76(j)(2), § 418.76(k), § 418.76(k)(2), § 418.100(b), § 418.100(c)(2), § 418.100(f)(1)(i), § 418.100(g)(3), § 418.104(d), § 418.104(f), § 418.106(b)(1), § 418.106(c)(1), § 418.106(e)(1), § 418.108(c)(3), § 418.110(a), § 418.110(c)(1)(i), § 418.110(c)(1)(ii), § 418.110(e), § 418.110(e)(2), § 418.110(f)(1), § 418.110(f)(3)(iv), § 418.110(f)(3)(vi), § 418.112(f), and § 418.116(b)(2).

B. Term of Approval

Based on the review and observations described in section III of this final notice, we have determined that CHAP’s accreditation program for hospices meet or exceed our requirements. Therefore, we approve CHAP as a national accreditation organization for hospices that request participation in the Medicare program, effective November 20, 2012 through November 20, 2018.

V. Collection of Information Requirements

This document does not impose information collection and recordkeeping requirements. Consequently, it need not be reviewed by the Office of Management and Budget under the authority of the Paperwork Reduction Act of 1995 (44 U.S.C. 35).

VI. Regulatory Impact Statement

In accordance with the provisions of Executive Order 12866, this proposed notice was not reviewed by the Office of Management and Budget.

(Docket No. FDA–2012–N–0001)

Endocrinologic and Metabolic Drugs Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Endocrinologic and Metabolic Drugs Advisory Committee. This meeting was announced in the Federal Register of October 10, 2012 (77 FR 61609). The amendment is being made to reflect a change in the Location and Procedure portions of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Paul Tran, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001; FAX: 301–847–8533; email: EMDAC@fda.hhs.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), and follow the prompts to the desired center or product area. Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the Federal Register of October 10, 2012, FDA announced that a meeting of the Endocrinologic and Metabolic Drugs Advisory Committee would be held on November 8, 2012. On page 61609, in the second column, the Location portion of the document is changed to read as follows:

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Building 31 Conference Center, the Great Room, (Rm. 1503), Silver Spring, MD 20993–0002. Information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: http://www.fda.gov/AdvisoryCommittees/default.htm; under the heading “Resources for You,” click on “Public Meetings at the FDA White Oak Campus.” Please note that visitors to the White Oak Campus must enter through Building 1.

On page 61609, in the third column, the Procedure portion of the document is changed to read as follows:

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 November 2, 2012.

Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. 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 October 25, 2012. 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 October 26, 2012.

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.


Jill Hartzler Warner,
Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2012–25741 Filed 10–18–12; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0001]

Nonprescription Drugs Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Nonprescription Drugs Advisory Committee. This meeting was announced in the Federal Register of August 30, 2012 (77 FR 52743). The amendment is being made to reflect a change in the Location and Contact Person portions of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Minh Doan, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Building 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001,
DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Notice of Diabetes Mellitus Interagency Coordinating Committee Meeting

SUMMARY: The Diabetes Mellitus Interagency Coordinating Committee (DMICC) will hold a meeting on November 15, 2012, from 9:00 to 11:30 a.m. at the Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817. The meeting is open to the public.

DATES: The meeting will be held on November 15, 2012, from 9:00 to 11:30 a.m. Individuals wanting to present oral comments must notify the contact person at least 10 days before the meeting date.

ADDITIONAL INFORMATION:
- The meeting will be held at Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817.
- For further information contact: Further information concerning this meeting, see the DMICC Web site, www.diabetescommittee.gov, or contact B. Tibor Roberts, Executive Secretary of the Diabetes Mellitus Interagency Coordinating Committee, National Institute of Diabetes and Digestive and Kidney Diseases, 31 Center Drive, Building 31A, Room 9A19, MSC 2560, Bethesda, MD 20892–2560, telephone: 301–496–6623; FAX: 301–480–6741; email: dmicc@mail.nih.gov.

SUPPLEMENTARY INFORMATION: The DMICC, chaired by the National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK) comprising members of the Department of Health and Human Services and other federal agencies that support diabetes-related activities, facilitates cooperation, communication, and collaboration on diabetes among government entities. DMICC meetings, held several times a year, provide an opportunity for Committee members to learn about and discuss current and future diabetes programs in DMICC member organizations and to identify opportunities for collaboration. The November 15, 2012, DMICC meeting will focus on “Federal Initiatives To Address Gestational Diabetes Mellitus.”

Jill Hartzler Warner, Acting Associate Commissioner for Special Medical Programs.

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