Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to http://www.regulations.gov and insert the docket number, found in brackets in the heading of this document, into the “Search” box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

Submit written requests for single copies of the guidance to the Policy and Regulations Staff (HFV–6), Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your requests. See the SUPPLEMENTARY INFORMATION section for electronic access to the guidance document.

FOR FURTHER INFORMATION CONTACT:
Dragan Momcilovic, Center for Veterinary Medicine (HFV–226), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–5944, email: dragan.momcilovic@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background
Following publication of the proposed rule to update FDA’s veterinary feed directive (VFD) regulation in December 2013 (78 FR 75515), in the Federal Register of December 1, 2013 (80 FR 75119), FDA published the notice of availability for a draft guidance entitled “Veterinary Feed Directive Common Format Questions and Answers” giving interested persons until February 1, 2016, to comment on the draft guidance. FDA received several comments on the draft guidance and those comments were considered as the guidance was finalized. The guidance announced in this notice finalizes the draft guidance dated December 2015.

II. Significance of Guidance
This level 1 guidance is being issued consistent with FDA’s good guidance practices regulation (21 CFR 10.115). The guidance represents the current thinking of FDA on Veterinary Feed Directive Common Format Questions and Answers. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

III. Paperwork Reduction Act of 1995
This guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR 514.1 have been approved under OMB Control No. 0910–0032. The collections of information in 21 CFR 558.6 have been approved under OMB Control No. 0910–0363.

IV. Electronic Access
Persons with access to the Internet may obtain the guidance at either http://www.fda.gov/AnimalVeterinary/GuidanceComplianceEnforcement/GuidanceforIndustry/default.htm or http://www.regulations.gov.

Dated: September 16, 2016.

Leslie Kux,
Associate Commissioner for Policy.

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Renewal of Charter for the Advisory Committee on Organ Transplantation

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HRSA is giving notice that the Advisory Committee on Organ Transplantation (ACOT) has been rechartered. The effective date of the renewed charter is September 1, 2016.

FOR FURTHER INFORMATION CONTACT:
Robert Walsh, Executive Secretary, Advisory Committee on Organ Transplantation, HRSA, Room 08W60, 5600 Fishers Lane, Rockville, MD 20857. Phone: (301) 443–6839; fax: (301) 594–6095; email: rwalsh@hrsa.gov.

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. Section 227a, Section 222 of the Public Health Service Act, as amended, 42 CFR 121.12 (2000), and in accordance with the Federal Advisory Committee Act (FACA), Public Law 92–463, ACOT was initially chartered on September 1, 2000, and was renewed at appropriate intervals.

ACOT provides advice to the Secretary of HHS (the Secretary) on all aspects of organ donation, procurement, allocation, and transplantation, and on such other matters that the Secretary determines. The recommendations of ACOT facilitate Department efforts to oversee the Organ Procurement and Transplantation Network, as set forth in the National Organ Transplant Act of 1984, as amended.

On August 31, 2016, the Secretary approved the ACOT charter to be renewed. The filing date of the renewed charter was September 1, 2016. Renewal of the ACOT charter gives authorization for the Committee to operate until September 1, 2018.

A copy of the ACOT charter is available on the ACOT Web site at http://www.organdonor.gov/legislation/advisory.html. A copy of the charter also can be obtained by accessing the FACA database that is maintained by the Committee Management Secretariat under the General Services Administration. The Web site address for the FACA database is http://www.facadatabase.gov/.

Jason E. Bennett,
Director, Division of the Executive Secretariat.

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting Announcement for the Technical Advisory Panel on Medicare Trustee Reports

ACTION: Notice of public meeting.

SUMMARY: This notice announces the meeting date for the second Technical Advisory Panel on Medicare Trustee Reports on Friday, September 30, 2016 in Washington, DC.

DATES: The meeting will be held on Friday, September 30, 2016 from 9:00 a.m. to 5:00 p.m., Eastern Daylight Time (EDT) and it is open to the public.

ADDRESS: This will be a virtual meeting held via WebEx.

FOR FURTHER INFORMATION CONTACT: Dr. Donald Oellerich, Designated Federal Officer, at the Office of Human Services Policy, Assistant Secretary for Planning and Evaluation, U.S. Department of Health and Human Services, 200 Independence Ave. SW., Washington, DC 20201. (202) 690–8410.

SUPPLEMENTARY INFORMATION:

I. Purpose: The Panel will discuss the long-term rate of change in health spending and may make recommendations to the Secretary on how the Medicare Trustees might more accurately estimate health spending in the short and long run. The Panel’s discussion is expected to be very technical in nature and will focus on the actuarial and economic assumptions and methods by which Trustees might more accurately measure health spending. This Committee is governed...
by the provisions of the Federal Advisory Committee Act, as amended (5 U.S.C. App. 2, section 10(a)(1) and (a)(2)). The Committee is composed of 9 members appointed by the Assistant Secretary for Planning and Evaluation.

II. Agenda. The Panel will likely hear presentations from the HHS Office of the Actuary on issues they wish the panel to address. This may be followed by HHS staff presentations regarding the methods and assumptions for the short range (10 year) Part A, Part B and Part D. After any presentations, the Panel will deliberate openly on the topics. Interested persons may observe the deliberations, but the Panel will not hear public comments during this time. The Panel will also allow an open public session for any attendee to address issues specific to the topic.

III. Meeting Attendance. The Friday, September 30, 2016 meeting is open to the public through WebEx; however, WebEx attendance is limited to space available.

Meeting Registration
The public may join the meeting through WebEx. Space is limited and registration is required in order to attend. Registration may be completed by emailing or faxing all the following information to Dr. Donald Oellerich at don.oellerich@hhs.gov or fax 202–690–6562:
Name.
Company name.
Postal address.
Email address.

A confirmation email with the WebEx information will be sent to the registrants shortly after completing the registration process. If language interpretation or other reasonable accommodation for a disability is needed, please contact Dr. Oellerich, no later than Sept 24, 2016 by sending an email message to don.oellerich@hhs.gov or calling 202–690–8410.

IV. Special Accommodations.
Individuals requiring special accommodations must include the request for these services during registration.

V. Copies of the Charter.
The Secretary’s Charter for the Technical Advisory Panel on Medicare Trustee Reports is available upon request from Dr. Donald Oellerich at don.oellerich@hhs.gov or by calling 202–690–8410.

Dated: September 16, 2016.

Kathryn E. Martin,
Acting Assistant Secretary for Planning and Evaluation.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

National Institute of General Medical Sciences; 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., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of General Medical Sciences Special Emphasis Panel: Review of MIRA Applications.
Date: October 27–28, 2016.
Time: 8:00 a.m. to 3:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Hilton Garden Inn Bethesda, 7301 Waverly Street, Bethesda, MD 20814.

Contact Person: Brian K. Pike, Ph.D., Scientific Review Officer, Office of Scientific Review, National Institute of General Medical Sciences, National Institutes of Health, 45 Center Drive, Room 3AN18, Bethesda, MD 20892, 301–504–3907, pikber@mail.nih.gov.

Name of Committee: National Institute of General Medical Sciences Special Emphasis Panel: To Review COBRE Grant Applications.
Date: November 2, 2016.
Time: 8:00 a.m. to 4:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Hilton Garden Inn Bethesda, 7301 Waverly Street, Bethesda, MD 20814.

Contact Person: Nina Sidorova, Ph.D., Scientific Review Officer, Office of Scientific Review, National Institute of General Medical Sciences, National Institutes of Health, 45 Center Drive, Room 3AN22, Bethesda, MD 20892–6200, 301–504–3663, sidorova@ngms.nih.gov.

[Catalogue of Federal Domestic Assistance Program Nos. 93.375, Minority Biomedical Research Support; 93.821, Cell Biology and Biophysics Research; 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.862, Genetics and Developmental Biology Research; 93.88, Minority Access to Research Careers; 93.96, Special Minority Initiatives; 93.859, Biomedical Research and Research Training, National Institutes of Health, HHS]
Dated: September 16, 2016.

Melanie J. Gray,
Program Analyst, Office of Federal Advisory Committee Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

National Institute of Neurological Disorders and Stroke; 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., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Neurological Sciences Training Initial Review Group: NST–1 Subcommittee.
Date: October 10–11, 2016.
Time: 8:00 a.m. to 6:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Embassy Suites Alexandria Old Town, 1900 Diagonal Road, Alexandria, VA 22314.

Contact Person: William Benzing, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Research, NINDS/NIH/DHHS, Neuroscience Center, 6001 Executive Blvd., Suite 3204, MSC 9529, Bethesda, MD 20892–9529, 301–496–0660, Benzing@mail.nih.gov.

Name of Committee: National Institute of Neurological Disorder and Stroke Initial Review Group; Neurological Sciences and Disorders A.
Date: October 24–25, 2016.
Time: 8:00 a.m. to 6:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Melrose Hotel, 2430 Pennsylvania Ave. NW., Washington, DC 20037.

Contact Person: Natalia Strunnikova, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Research, NINDS/NIH/DHHS, Neuroscience Center, 6001 Executive Blvd., Suite 3204, MSC 9529, Bethesda, MD 20892–9529, 301–946–0286, natalia.strunnikova@nih.gov.

Name of Committee: Neurological Sciences Training Initial Review Group: NST–2 Subcommittee.
Date: October 24, 2016.
Time: 8:00 a.m. to 5:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Embassy Suites Alexandria Old Town, 1900 Diagonal Road, Alexandria, VA 22314.

Contact Person: Elizabeth Webber, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Research,