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75515), in the **Federal Register** of December 1, 2015 (80 FR 75119), FDA published the notice of availability for a draft guidance entitled ''Veterinary Feed Directive Common Format Questions and Answers'' giving interested persons until February 1, 2016, to comment on the draft guidance. FDA received several comments on the draft guidance and those comments were considered as the guidance was finalized. The guidance announced in this notice finalizes the draft guidance dated December 2015.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.

[FR Doc. 2016–22775 Filed 9–21–16; 8:45 am]

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 217a, 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.

[FR Doc. 2016–22858 Filed 9–21–16; 8:45 am]

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.

ADDRESSES: 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