

appropriate advisory committee meeting 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 October 14, 2016. Oral presentations from the public will be scheduled between approximately 10 a.m. to 10:30 a.m. and 3 p.m. to 3:30 p.m. on October 25, 2016, and between approximately 10:15 a.m. to 10:45 a.m. and 2:30 p.m. to 3 p.m. on October 26, 2016. 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 6, 2016. 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 7, 2016.

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 disabilities. If you require accommodations due to a disability, please contact AnnMarie Williams at AnnMarie.Williams@fda.hhs.gov or 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: September 16, 2016.

Janice M. Soreth,

Acting Associate Commissioner, Special Medical Programs.

[FR Doc. 2016-22808 Filed 9-21-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0155]

Veterinary Feed Directive Common Format Questions and Answers; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a guidance for industry #233 entitled "Veterinary Feed Directive Common Format Questions and Answers." FDA had received comments requesting that we require a uniform Veterinary Feed Directive (VFD) form. We declined this request because we think that requiring a specific VFD form would be too prescriptive. However, we acknowledge that a common VFD format would help veterinarians, their clients (*i.e.*, animal producers), and distributors (including feed mills) quickly identify relevant information on the VFD. We are issuing this guidance to recommend a common VFD format. We expect this guidance will reduce potential errors on VFDs.

DATES: Submit either electronic or written comments on Agency guidances at any time.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <http://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the

manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2010-N-0155 for "Veterinary Feed Directive Common Format Questions and Answers." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <http://www.regulations.gov> or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- **Confidential Submissions—**To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <http://www.regulations.gov>. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <http://www.fda.gov/regulatoryinformation/dockets/default.htm>.

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]

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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]

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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