Estimated Number of Respondents per Level:

- LOA 1 Account: 114,256.
- LOA 2 Account: 17,848.
  - In Person ID Proofing (subset of LOA 2): 5,354.
  - Online\Remote ID Proofing (subset of LOA 2): 12,494.

Estimated Total Number of Respondents: 132,104.

Estimated Number of Responses per Respondent: 1.

Estimated Total Annual Burden on Respondents: 26,239.8 hours.

Comments are invited on (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) the accuracy of the agency's estimate of burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of the information on those who respond, through the use of appropriate automated, electronic, mechanical, technological or other forms of information technology collection methods. Copies of the information collection may be obtained from Mr. Zeimet by calling or emailing your request to the contact information above in the **FOR FURTHER INFORMATION** section. All responses to this notice will be summarized and included in the request for OMB approval. All comments will become a matter of public record.

Dated: July 11, 2016.

Jonathan Alboum,

Chief Information Officer, Office of the Chief Information Officer.

[FR Doc. 2016–16817 Filed 7–21–16; 8:45 am]

BILLING CODE 3410–KR–P

DEPARTMENT OF AGRICULTURE

Food Safety and Inspection Service

[Docket No. FSIS–2016–0024]

Codex Alimentarius Commission: Meeting of the Codex Committee on Residues of Veterinary Drugs in Food

AGENCY: Office of the Deputy Under Secretary for Food Safety, USDA.

ACTION: Notice of public meeting and request for comments.

SUMMARY: The Office of the Deputy Under Secretary for Food Safety, U.S. Department of Agriculture (USDA), and the Food and Drug Administration (FDA), are sponsoring a public meeting on September 22, 2016. The objective of the public meeting is to provide information and receive public comments on agenda items and draft United States (U.S.) positions to be discussed at the 23rd Session of the Codex Committee on Residues of Veterinary Drugs in Foods (CCRVDF) of the Codex Alimentarius Commission (Codex), taking place in Houston, Texas, October 17–21, 2016. The Deputy Under Secretary for Food Safety and the FDA recognize the importance of providing interested parties the opportunity to obtain background information on the 23rd Session of the CCRVDF and to address items on the agenda.

DATES: The public meeting is scheduled for Thursday, September 22, 2016, from 1:00 p.m.–4:00 p.m.

ADDRESSES: The public meeting will take place at the USDA, Jamie L. Whitten Building, 1400 Independence Avenue SW., Room 107–A, Washington, DC 20250.

Documents related to the 23rd Session of the CCRVDF will be accessible via the Internet at the following address: *http://www.codexalimentarius.org/meetings-reports/en/.*

Brandi Robinson, U.S. Delegate to the 23rd Session of the CCRVDF, invites U.S. interested parties to submit their comments electronically to the following email address: *Brandi.Robinson@fda.hhs.gov.*

Call-In-Number

If you wish to participate in the public meeting for the 23rd Session of the CCRVDF by conference call, please use the following call-in-number:

Call-in-Number: 1–888–844–9904.

The participant code will be posted on the following Web page: *http://www.fsis.usda.gov/wps/portal/fsis/topics/international-affairs/us-codex-alimentarius/public-meetings.*

Registration

Attendees may register to attend the public meeting by emailing *uscodex@fsis.usda.gov* by September 16, 2016. Early registration is encouraged as it will expedite entry into the building. The meeting will be held in a Federal building. Attendees should bring photo identification and plan for adequate time to pass through the security screening systems. Attendees who are not able to attend the meeting in person, but who wish to participate, may do so by phone.

FOR FURTHER INFORMATION ABOUT THE 23RD SESSION OF THE CCRVDF CONTACT: Brandi Robinson, ONADE International Coordinator, Center for Veterinary Medicine, U.S. Food and Drug Administration, 7500 Standish Place, HFV–100, Rockville, MD 20855. Telephone: (240) 402–0645, Email: *Brandi.Robinson@fda.hhs.gov.*

For Further Information About the Public Meeting Contact

Kenneth Lowery, U.S. Codex Office, 1400 Independence Avenue SW., South Agriculture Building, Room 4861, Washington, DC 20250. Telephone: (202) 690–4042, Fax: (202) 720–3157, Email: *Kenneth.Lowery@fsis.usda.gov*

SUPPLEMENTARY INFORMATION:

Background

Codex was established in 1963 by two United Nations organizations, the Food and Agriculture Organization and the World Health Organization (FAO/WHO). Through adoption of food standards, codes of practice, and other guidelines developed by its committees, and by promoting their adoption and implementation by governments, Codex seeks to protect the health of consumers and ensure fair practices in the food trade.

The CCRVDF is responsible for determining priorities for the consideration of residues of veterinary drugs in foods, recommending maximum levels of such substances, developing codes of practice as may be required, and considering methods of sampling and analysis for the determination of veterinary drug residues in foods.

The Committee is hosted by the United States.

Issues To Be Discussed at the Public Meeting

The following items on the Agenda for the 23rd Session of the CCRVDF will be discussed during the public meeting:

- Matters referred to the Committee by Codex or its subsidiary bodies;
- Matters of interest arising from the FAO/WHO and from the 81st Meeting of the Joint FAO/WHO Expert Committee on Food Additives (JECFA);
- Report of the World Organisation for Animal Health activities, including the International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products;
- Proposed draft Risk Management Recommendations (RMR) for gentian violet at Step 3;
- Proposed draft Maximum Residue Limits (MRLs) for ivermectin (cattle muscle) and lasalocid sodium (chicken, turkey, quail, and pheasant kidney, liver, muscle, skin and fat) at Step 4;
- Proposed draft MRLs for ivermectin (cattle fat, kidney, muscle),

teflubenzuron (salmon fillet, muscle) and zilpaterol hydrochloride (cattle fat, kidney, liver, muscle) at Step 3;

- Discussion paper on the unintended presence of residues of veterinary drugs in food commodities resulting from the carry-over of drug residues into feed;
- Discussion paper on the establishment of a rating system to establish priority for the CCRVDF work;
- Global survey to provide information to the CCRVDF to move compounds from the database on countries' needs for MRLs to the JECFA Priority List (Report of Environmental Working Group) and Database on countries' needs for MRLs;
- Draft priority list of veterinary drugs requiring evaluation or re-evaluation by JECFA; and
- Other Business & Future Work.

Each issue listed will be fully described in documents distributed, or to be distributed, by the Secretariat before the Meeting. Members of the public may access or request copies of these documents (see **ADDRESSES**).

Public Meeting

At the September 22, 2016, public meeting, draft U.S. positions on the agenda items will be described and discussed, and attendees will have the opportunity to pose questions and offer comments. Written comments may be offered at the meeting or sent to the U.S. Delegate for the 23rd Session of the CCRVDF, Brandi Robinson (see **ADDRESSES**). Written comments should state that they relate to the activities of the 23rd Session of the CCRVDF.

Additional Public Notification

Public awareness of all segments of rulemaking and policy development is important. Consequently, FSIS will announce this **Federal Register** publication on-line through the FSIS Web page located at: *http://www.fsis.usda.gov/federal-register.*

FSIS also will make copies of this publication available through the FSIS Constituent Update, which is used to provide information regarding FSIS policies, procedures, regulations, **Federal Register** notices, FSIS public meetings, and other types of information that could affect or would be of interest to our constituents and stakeholders. The Update is available on the FSIS Web page. Through the Web page, FSIS is able to provide information to a much broader, more diverse audience. In addition, FSIS offers an email subscription service which provides automatic and customized access to selected food safety news and information. This service is available at: *http://www.fsis.usda.gov/subscribe.* Options range from recalls to export information, regulations, directives, and notices. Customers can add or delete subscriptions themselves, and have the option to password protect their accounts.

USDA Non-Discrimination Statement

No agency, officer, or employee of the USDA shall, on the grounds of race, color, national origin, religion, sex, gender identity, sexual orientation, disability, age, marital status, family/parental status, income derived from a public assistance program, or political beliefs, exclude from participation in, deny the benefits of, or subject to discrimination any person in the United States under any program or activity conducted by the USDA.

How To File a Complaint of Discrimination

To file a complaint of discrimination, complete the USDA Program Discrimination Complaint Form, which may be accessed online at *http://www.ocio.usda.gov/sites/default/files/docs/2012/Complain_combined_6_8_12.pdf,* or write a letter signed by you or your authorized representative.

Send your completed complaint form or letter to USDA by mail, fax, or email:

Mail: U.S. Department of Agriculture, Director, Office of Adjudication, 1400 Independence Avenue SW., Washington, DC 20250–9410.

Fax: (202) 690–7442.

Email: program.intake@usda.gov.

Persons with disabilities who require alternative means for communication (Braille, large print, audiotape, etc.), should contact USDA's TARGET Center at (202) 720–2600 (voice and TDD).

Done at Washington, DC, on July 19, 2016.

Paulo Almeida,

U.S. Manager for Codex Alimentarius.

[FR Doc. 2016–17377 Filed 7–21–16; 8:45 am]

BILLING CODE 3410–DM–P

DEPARTMENT OF AGRICULTURE

Forest Service

Missoula Resource Advisory Committee Meeting

AGENCY: Forest Service, USDA.

ACTION: Notice of meeting.

SUMMARY: The Missoula Resource Advisory Committee (RAC) will meet in Frenchtown, Montana. The committee is authorized under the Secure Rural Schools and Community Self-Determination Act (the Act) and operates in compliance with the Federal Advisory Committee Act. The purpose of the committee is to improve collaborative relationships and to provide advice and recommendations to the Forest Service concerning projects and funding consistent with Title II of the Act. RAC information can be found at the following Web site: *http://www.fs.usda.gov/main/lolo/workingtogether/advisorycommittees.*

DATES: The meeting will be held on Wednesday, August 3, 2016, at 6 p.m.

All RAC meetings are subject to cancellation. For status of meeting prior to attendance, please contact the person listed under **FOR FURTHER INFORMATION CONTACT**.

ADDRESSES: The meeting will be held at Frenchtown Rural Fire Station 1, 16875 Marion Street, Frenchtown, Montana.

Written comments may be submitted as described under **SUPPLEMENTARY INFORMATION**. All comments, including names and addresses when provided, are placed in the record and are available for public inspection and copying. The public may inspect comments received at Ninemile Ranger District.

FOR FURTHER INFORMATION CONTACT: Sari Lehl, RAC Coordinator, by phone at 406–626–5201 or via email at *slehl@fs.fed.us.*

Individuals who use telecommunication devices for the deaf (TDD) may call the Federal Information Relay Service (FIRS) at 1–800–877–8339 between 8:00 a.m. and 8:00 p.m., Eastern Standard Time, Monday through Friday.

SUPPLEMENTARY INFORMATION: The purpose of the meeting is for RAC project proposal presentations.

The meeting is open to the public. The agenda will include time for people to make oral statements of three minutes or less. Individuals wishing to make an oral statement should request in writing by August 1, 2016, to be scheduled on the agenda. Anyone who would like to bring related matters to the attention of the committee may file written statements with the committee staff before or after the meeting. Written comments and requests for time to make oral comments must be sent to Sari Lehl, RAC Coordinator, Ninemile Ranger District, 20325 Remount Road, Huson, Montana 59846; by email to *slehl@fs.fed.us,* or via facsimile to 406–626–5201.

Meeting Accommodations: If you are a person requiring reasonable accommodation, please make requests in advance for sign language interpreting, assistive listening devices, or other reasonable accommodation. For access to the facility or proceedings, please contact the person listed in the