○ Are there uniform and recognized differences in the lengths of rental and leasing agreements? Do these lead to special funding considerations for the government?

○ Are there specific differences in other terms and conditions, such as maintenance, warranty, insurance, taxes, storage, and transportation? What are the specific distinctions?

• FAR 7.402(a) and (b) provide circumstances indicating when the purchase or lease method is appropriate. Should rent be added as a third method? If so, when is renting a more appropriate method than purchasing or leasing and in what way would it impact the determination of most effective procurement approach (*e.g.,* cost savings and efficiencies)?

• Does short-term rental offer cost savings and efficiencies unavailable through leasing?

• What additional guidance might be provided at FAR subpart 7.4 to clarify when and how to perform the required analysis?

Dated: July 9, 2013.

William Clark,

Acting Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2013–16920 Filed 7–15–13; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting; Subcommittee for Dose Reconstruction Reviews, Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board), National Institute for Occupational Safety and Health (NIOSH)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC), announces the following meeting for the aforementioned subcommittee:

Time and Date: 10:00 a.m.–5:00 p.m., Eastern Time, August 7, 2013.

Place: Audio Conference Call via FTS Conferencing. The USA toll-free, dial-in number is 1–866–659–0537, and the pass code is 9933701.

Status: Open to the public, but without an oral public comment period.

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000, to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines that have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction, which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort.

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2013.

Purpose: The Advisory Board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee for Dose Reconstruction Reviews was established to aid the Advisory Board in carrying out its duty to advise the Secretary, HHS, on dose reconstruction.

Matters To Be Discussed: The agenda for the Subcommittee meeting includes: dose reconstruction program quality management and assurance activities, including current findings from NIOSH internal dose reconstruction blind reviews; and discussion of dose reconstruction cases under review (sets 8–9, and Savannah River Site, Rocky Flats Plant, and Los Alamos National Laboratory cases from sets 10–13).

The agenda is subject to change as priorities dictate.

In the event an individual cannot attend, written comments may be submitted. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

Contact Person for More Information: Theodore Katz, Designated Federal Official, NIOSH, CDC, 1600 Clifton Road, Mailstop E–20, Atlanta, Georgia 30333, Telephone: (513) 533–6800, Toll Free 1 (800) CDC–INFO, Email: *ocas@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dana Redford,

Acting Director, Management Analysis and Services Office Centers for Disease Control and Prevention.

[FR Doc. 2013–16964 Filed 7–15–13; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting; State, Tribal, Local and Territorial (STLT) Subcommittee, Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting of the aforementioned subcommittee:

Time and Date: 8:30 a.m.–4:00 p.m., EDT, August 9, 2013.

Place: CDC, Building 19, Rooms 254 and 255, 1600 Clifton Road NE., Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available. The meeting rooms accommodate approximately 20 people. The public is welcome to participate during the public comment period, which is tentatively scheduled from 3:15 p.m. to 3:35 p.m. This meeting will also be available by teleconference. Please dial (888) 233–0592 and enter code 33288611.

Purpose: The Subcommittee will provide advice to the CDC Director through the ACD on strategies and future needs and challenges faced by State, Tribal, Local and Territorial health agencies, and will provide guidance on opportunities for CDC.

Matters To Be Discussed: The STLT Subcommittee members will discuss implementation of ACD-adopted recommendations related to the health department of the future and how CDC can best support STLT health departments.

The agenda is subject to change as priorities dictate.

Contact Person for More Information: Judy Monroe, M.D., Designated Federal Officer, STLT Subcommittee—ACD, CDC, 1600 Clifton Road, NE., M/S E–70, Atlanta, Georgia 30333, Telephone: (404) 498–0300, Email: *OSTLTSDirector@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

Dana Redford,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2013–16965 Filed 7–15–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–D–0254]

Salmonella Contamination of Dry Dog Food; Withdrawal of Compliance Policy Guide

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; Withdrawal.

SUMMARY: The Food and Drug Administration (FDA) is announcing the withdrawal of the compliance policy guide (CPG) entitled ''Sec. 690.700 *Salmonella* Contamination of Dry Dog Food.'' This CPG is obsolete.

DATES: The withdrawal is effective July 16, 2013.

FOR FURTHER INFORMATION CONTACT: Diane D. Jeang, Office of Regulatory Affairs, Food and Drug Administration, 12420 Parklawn Dr., Rockville, MD 20857, 301–796–3890.

SUPPLEMENTARY INFORMATION: FDA issued the CGP entitled ''Sec. 690.700 *Salmonella* Contamination of Dry Dog Food (CPG 690.700)'' on October 1, 1980. CPG 690.700 was issued as a result of a human case of salmonellosis traced to dry dog food; a subsequent FDA-conducted survey of dry dog food; a risk analysis; and the development of an appropriate sampling technique to test dry dog food for *salmonella* organisms.

Elsewhere in this issue of the **Federal Register**, FDA is announcing the availability of a new CPG to address all food for animals that may contain *salmonella* organisms, including dry dog food. This new CPG, entitled ''Compliance Policy Guide Sec. 690.800 *Salmonella* in Food for Animals,'' supersedes CPG 690.700 and makes CPG 690.700 obsolete. The notice of availability for CPG ''Sec. 690.800 *Salmonella* in Food for Animals'' is published elsewhere in this issue of the **Federal Register**.

FDA is withdrawing CPG 690.700, in its entirety, to eliminate obsolete compliance policy.

Dated: July 10, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–16973 Filed 7–15–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–D–0378]

Compliance Policy Guide Sec. 690.800 *Salmonella* in Food for Animals; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing the availability of guidance for our staff entitled ''Compliance Policy Guide Sec. 690.800 *Salmonella* in Food for Animals'' (the CPG). The CPG provides guidance to FDA staff on *Salmonella*-contaminated food for animals.

DATES: Submit either electronic or written comments on the CPG at any time.

ADDRESSES: Submit written requests for single copies of the CPG to the Food and Feed Policy Staff, Office of Policy and Risk Management, Office of Regulatory Affairs, Food and Drug Administration, 12420 Parklawn Dr., Rockville, MD 20850. Send one self-addressed adhesive label to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the CPG.

Submit electronic comments on the CPG to *http://www.regulations.gov.* Submit written comments on the CPG to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Kim Young, Center for Veterinary Medicine (HFV–230), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9207, *kim.young@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

We are announcing the availability of a guidance document entitled ''Compliance Policy Guide Sec. 690.800 *Salmonella* in Food for Animals'' (the CPG). The CPG provides guidance to FDA staff on *Salmonella*-contaminated food for animals. The CPG is being issued consistent with our good guidance practices regulation (21 CFR 10.115). The CPG represents FDA's current thinking on *Salmonella*-contaminated food for animals. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statutes and regulations.

In the **Federal Register** of August 2, 2010 (75 FR 45130), we announced the availability of a draft CPG entitled ''Compliance Policy Guide Sec. 690.800 *Salmonella* in Animal Feed,'' and gave interested persons an opportunity to submit comments by November 1, 2010, for us to consider before beginning our work on the final version of the CPG. In the **Federal Register** of October 29, 2010 (75 FR 66769), we published a notice extending the comment period until December 31, 2010. We received numerous comments on the draft CPG and have modified the final CPG where appropriate. The CPG announced in this notice finalizes the draft CPG announced on August 2, 2010.

Changes to the CPG include:

- The title of the CPG is changed from ''*Salmonella* in Animal Feed'' to ''*Salmonella* in Food for Animals.'' FDA made this change to clarify that the CPG covers all animal food. The term ''food for animals'' here includes pet food and animal feed.
- The term ''Direct Human Contact Animal Feed'' has been removed from the CPG, because commenters found the term to be confusing. The term pet food is now used instead. It is defined to mean food for pets and includes treats and chews for pets.

The CPG explains criteria that FDA personnel should consider in recommending enforcement action against food for animals that is adulterated due to the presence of *Salmonella.* In particular, the CPG provides regulatory action guidance relating to pet food or pet food ingredients that are contaminated with *Salmonella.* In addition, the CPG provides regulatory action guidance relating to animal feed and animal feed ingredients that are contaminated with certain *Salmonella* serotypes that are pathogenic to the particular species of animal for which the animal feed or animal feed ingredients are intended. The CPG also contains information that may be useful to regulated industry and the public.

This notice is related to two notices published elsewhere in this issue of the **Federal Register**, in which FDA is announcing: (1) The removal of 21 CFR 500.35 ''Animal feeds contaminated with *Salmonella* microorganisms,'' and