Prevention and the Agency for Toxic Substances and Disease Registry.

Catherine Ramadei,

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

[FR Doc. 2016–05265 Filed 3–8–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

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

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: 11:30 a.m.–1:00 p.m. EDT, April 08, 2016.

Place: This meeting will be held by teleconference.

Status: This meeting is open to the public, limited only by the availability of telephone ports. The public is welcome to participate during the public comment, which is tentatively scheduled from 12:40 p.m. to 12:45 p.m. To participate on the 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 for Discussion: The STLT Subcommittee members will discuss progress on implementation of ACD-adopted recommendations related to the health departments of the future, additional developments that may expand these recommendations, and how CDC can best support STLT health departments.

The agenda is subject to change as priorities dictate.

Contact Person for More Information: John Auerbach, MBA, Designated Federal Officer, STLT Subcommittee, ACD, CDC, 4770 Buford Highway, MS E70, Atlanta, Georgia 30341, Telephone (404) 498–0300, Email: *OSTLTSDirector@cdc.gov.* Please submit comments to *OSTLTSDirector@cdc.gov* by April 1, 2016.

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.

Claudette Grant,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Evaluation of Domestic Human Trafficking Demonstration Projects.

OMB No.: New Collection.

Description: The Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS) is proposing a 2-year data collection as part of the ''Evaluation of Domestic Human Trafficking Demonstration Projects'' study. This notice addresses the cross-site process evaluation to be conducted with the FY 2015 domestic human trafficking demonstration sites funded by the Family and Youth Services Bureau (FYSB).

The objective of the process evaluation is to describe program operations and implementation experience, such as start-up efforts, service provision to a wide array of trafficking victims, collaboration development, training, and sustainability actions. Information from the evaluation will assist federal, state, and community policymakers and funders in laying the groundwork for the refinement of program models to serve domestic victims of human trafficking, as well as evaluation strategies for future programs targeting trafficking victims.

The evaluation of domestic human trafficking demonstration projects will document and describe each site's community and organizational capacity; partnership composition and functioning; comprehensive, victim-centered services; and survivor characteristics, experiences, and outcomes. Primary data for the evaluation will be collected via qualitative interviews, including key informant interviews, case narrative interviews, client interviews, bimonthly telephone interviews, and cost modules (*i.e.,* structured interviews with project directors to collect information on costs). Data will be collected via two site visits per year, during 2016 and 2017. Case narrative interviews will include follow up interviews. Interviews from multiple perspectives will enhance the government's understanding of strategies by which grantees can identify, engage and serve diverse populations of victims of severe forms of human trafficking.

Respondents: Project directors and case managers at the three FY 2015 FYSB funded demonstration projects; staff (*e.g.,* program managers and directors) from partner organizations that are working with the three FY 2015 FYSB-funded demonstration projects; and clients who have received services from the three FY 2015 FYSB-funded demonstration projects.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Project Director Interview	6	3	1	2	6
Case Manger Interview	30	15	1	1.25	19
Partner Interviews	30	15	1	1.25	19
Case Narrative Interview	30	15	1	1	15
Client Interview	30	15	1	1	15
Human Trafficking Evaluation Cost Module/Human Trafficking Evaluation Labor Module	6	3	1	1	3
Bi-monthly Project Director Calls	6	3	1	6	18

Estimated total annual burden hours: 95.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

ACF Certifying Officer.

[FR Doc. 2016–05240 Filed 3–8–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–D–1180]

Ensuring Safety of Animal Feed Maintained and Fed On-Farm; 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 (GFI) # 203 entitled ''Ensuring Safety of Animal Feed Maintained and Fed On-Farm.'' This guidance is intended to help animal producers (persons who feed animals) develop and implement on-farm practices to ensure the safety of animal feed maintained and fed to animals on the farm.

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–2014–D–1180 for ''Ensuring Safety of Animal Feed Maintained and Fed On-Farm.'' 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: Phares Okelo, Center for Veterinary Medicine (HFV–226), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–5921, email: *phares.okelo@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of March 20, 2015 (80 FR 15014), FDA published the notice of availability for a draft GFI #203 entitled ''Ensuring Safety of Animal Feed Maintained and Fed On-Farm'' giving interested persons until June 3, 2015, to comment on the draft guidance. FDA received several comments on the