

- The nominee's contact information and current occupation or position;
- The nominee's resume or curriculum vitae, including prior or current membership on other National Institute for Occupational Safety and Health (NIOSH), CDC, or HHS advisory committees or other relevant organizations, associations, and committees;
- The category of membership (epidemiologist, environmental medicine or environmental health specialist, occupational physician with experience treating WTC rescue and recovery workers, occupational physician, representative of WTC responders, or toxicologist) that the candidate is qualified to represent;
- A summary of the background, experience, and qualifications that demonstrates the nominee's suitability for the nominated membership category;
- Articles or other documents the nominee has authored that indicate the nominee's knowledge and experience in relevant subject categories; and
- A statement that the nominee is aware of the nomination, is willing to regularly attend and participate in STAC meetings, and has no known conflicts of interest that would preclude membership on the Committee.

STAC members will be selected upon the basis of their relevant experience and competence in their respective categorical fields. The information received through this nomination process, in addition to other relevant sources of information, will assist the WTC Program Administrator in appointing members to serve on the STAC. In selecting members, the WTC Program Administrator will consider individuals nominated in response to this **Federal Register** notice as well as other qualified individuals.

The CDC is committed to bringing greater diversity of thought, perspective, and experience to its advisory committees. Nominees from all races, genders, ages, and persons living with disabilities are encouraged to apply. Nominees must be U.S. citizens.

Candidates invited to serve will be asked to submit the "Confidential Financial Disclosure Report," OGE Form 450. This form is used by CDC to determine whether there is a financial conflict between that person's private interests and activities and their public responsibilities as a Special Government Employee as well as any appearance of a loss of impartiality, as defined by Federal regulation. The form may be viewed and downloaded at <http://www.oge.gov/Forms-Library/OGE-Form-450-Confidential-Financial-Disclosure->

Report/. This form should not be submitted as part of a nomination.

DATES: Nominations must be submitted (postmarked or electronically received) by March 31, 2015.

Submissions must be electronic or by mail. Submissions should reference docket 229-C. Electronic submissions: You may electronically submit nominations, including attachments, to nioshdocket@cdc.gov. Attachments in Microsoft Word are preferred. Regular, Express, or Overnight Mail: Written nominations may be submitted (one original and two copies) to the following address only: NIOSH Docket 229-C, c/o Mia Wallace, Committee Management Specialist, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1600 Clifton Rd. NE., MS: E-20, Atlanta, Georgia 30333. Telephone and facsimile submissions cannot be accepted. For further information contact: Paul Middendorf, Senior Health Scientist, 1600 Clifton Rd. NE., MS: E-20, Atlanta, GA 30333; telephone (404) 498-2500 (this is not a toll-free number); email pmiddendorf@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.

Elaine L. Baker,

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

[FR Doc. 2015-03682 Filed 2-23-15; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10407]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register**

concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by March 26, 2015.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395-5806 OR Email: OIRA_submission@omb.eop.gov.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at <http://www.cms.hhs.gov/PaperworkReductionActof1995>.
2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.
3. Call the Reports Clearance Office at (410) 786-1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786-1326.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies

to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Summary of Benefits and Coverage and Uniform Glossary; *Use:* Section 2715 of the PHS Act directs the Department of Health and Human Services (HHS), the Department of Labor (DOL), and the Department of the Treasury (collectively, the Departments), in consultation with the National Association of Insurance Commissioners (NAIC) and a working group comprised of stakeholders, to “develop standards for use by a group health plan and a health insurance issuer in compiling and providing to applicants, enrollees, and policyholders and certificate holders a summary of benefits and coverage explanation that accurately describes the benefits and coverage under the applicable plan or coverage.” To implement these disclosure requirements, collection of information requests relate to the provision of the following: Summary of benefits and coverage, which includes coverage examples; a uniform glossary of health coverage and medical terms; and a notice of modifications. *Form Number:* CMS–10407 (OMB control number 0938–1146); *Frequency:* Annual; *Affected Public:* Private Sector—Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 126,500; *Number of Responses:* 41,153,858; *Total Annual Hours:* 322,411. (For policy questions regarding this collection, contact Heather Raeburn at 301–492–4224.)

Dated: February 18, 2015.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2015–03650 Filed 2–23–15; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No.: 0970–0139]

Uniform Project Description (UPD) Project Narrative Format for Discretionary Grant Application Forms; Correction

AGENCY: Administration for Children and Families, HHS.

ACTION: Notice; correction.

SUMMARY: The Administration for Children and Families published a document in the **Federal Register** of February 17, 2015, concerning a request for comments on a proposed information collection. The document contained an incorrect citation.

FOR FURTHER INFORMATION CONTACT: Christopher Beach, Senior Grants Policy Specialist, Division of Grants Policy, Office of Administration, Administration for Children and Families, telephone (202) 401–1539.

Correction: In the **Federal Register** of February 17, 2015, in FR. Doc. 2015–03144, on page 8324, in the third column, correct the last sentence of the “Description” caption to read: “Guidance for the content of information requested in the Uniform Project Description is based in 45 CFR 75.203 and Appendix I to 45 CFR part 75.”

Christopher Beach,

Senior Grants Policy Specialist, Office of Administration.

[FR Doc. 2015–03627 Filed 2–23–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–D–0235]

Evaluating the Effectiveness of New Animal Drugs for the Reduction of Pathogenic Shiga Toxin-Producing *Escherichia coli* in Cattle; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry (GFI #229) entitled “Evaluating the Effectiveness of New Animal Drugs for the Reduction of Pathogenic Shiga

Toxin-Producing *E. coli* in Cattle.” The purpose of this document is to provide recommendations to industry relating to study design and describe criteria the Center for Veterinary Medicine (CVM) thinks are the most appropriate for the evaluation of the effectiveness of new animal drugs that are intended to reduce pathogenic Shiga toxin-producing *E. coli* (STEC) in cattle.

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by April 27, 2015.

ADDRESSES: Submit written requests for single copies of the guidance to the Communications Staff (HFV–12), 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 draft guidance document.

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

FOR FURTHER INFORMATION CONTACT: Joshua R. Hayes, Center for Veterinary Medicine (HFV–133), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–402–0651, Joshua.hayes@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance for industry (GFI #229) entitled “Evaluating the Effectiveness of New Animal Drugs for the Reduction of Pathogenic Shiga Toxin-Producing *E. coli* in Cattle.” This draft guidance provides recommendations to industry relating to study design and describes criteria CVM thinks are the most appropriate for the evaluation of the effectiveness of new animal drugs that are intended to reduce pathogenic STEC in cattle. It discusses general considerations regarding the development of protocols, study conduct, animal welfare, substantial evidence of effectiveness, experimental parameters, nutritional content of experimental diets, and the assessment of drug concentrations in experimental diets. It also discusses the studies and analyses CVM recommends for sponsors