entering or exiting elevators. GSA requires users to operate their Segway at a speed no greater than a walking pace of three (3) miles per hour. Security personnel, as well as GSA personnel and other agency personnel in delegated buildings, shall monitor, to the extent practical, the safe and responsible operation of Segway devices by their users. Should security or building management personnel observe the unsafe operation of a Segway device, these individuals shall remind the user of their responsibility for the safe operation of the Segway and the user's accountability for not operating the device at a speed that exceeds three (3) miles per hour.

Security (where access is controlled by security personnel and screening devices)

Magnetometer and X-ray security screening devices are ineffective for the evaluation of Segways. Segway representatives have reported that magnetometer devices affect the operation and software programs of the transporter device when coming in contact with or close proximity to magnetic fields. Accordingly, Segway recommends that the transporter device not be operated within five (5) feet of any magnetometer or X-ray security screening device. If within five (5) feet of a magnetometer or X-ray security screening device, the user of the transporter device should place the unit in standby mode or powered off mode to prevent damage to the transporter device.

GSA and associated building security personnel reserve the right to inspect Segway devices upon entrance to a Federal building. Upon entering a Federal building under the jurisdiction, custody or control of GSA, security personnel shall notify the individual that only those persons with a mobility impairment are authorized to operate a Segway within the building. The security personnel may not ask a person using the device questions about the nature and extent of the person's disability. Security personnel shall inspect Segways in the same manner as other motorized devices that enter the building, including electric wheelchairs and scooters. Security personnel may request that the Segway user demonstrate that the device is operational thereby ensuring that the device contains its propulsion and battery equipment. The Segway user will be subject to the appropriate screening protocols established by security personnel for the protection of occupants, visitors and facilities, while maintaining the dignity of all persons who enter the building.

Vertical Transportation

Segway devices are permitted on elevators, but are not permitted on escalators, as per manufacturer guidance.

[FR Doc. E7–25592 Filed 1–4–08; 8:45 am]

BILLING CODE 6820–RH–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: Grants.gov–4040–0002]

Agency Information Collection Request. 60-Day Public Comment Request

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed information collection request for public comment. Interested persons are invited to send comments regarding this 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.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *Sherette.funncoleman@hhs.gov*, or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be received with 60 days.

Proposed Project: SF–424 Mandatory—OMB No. 4040–0002-Revision—Grants.gov.

Abstract: This collection is the government-wide form used for mandatory grant programs. Proposed revision to the form includes the addition of a data block that will collect the ''Descriptive Title of Applicant's Project.'' The data field labeled ''County'' will be revised to read ''County/Parish.'' The instructions are also being revised to incorporate the new descriptive title block and also, revisions to the instructions for areas affected by funding and the congressional district. Changes to the instructions will increase data quality and clarity for the collection.

Adding an additional data block is necessary to comply with the requirements of the Federal Funding Accountability and Transparency Act (FFATA). FFATA was signed into law on September 26, 2006 (Pub. L. 109–282). The legislation requires the Office of Management and Budget (OMB) to establish a publicly available, online database containing information about entities that are awarded federal grants, loans, and contracts. The revised form will assist agencies in collecting the required data elements for the database through the SF–424 applications. This form will be utilized on occasion by up to 26 Federal grant making agencies with mandatory grant programs. We are requesting a 2-year clearance of this form. The affected public includes, Federal, State, local or tribal governments, business or other for profit, and not for profit institutions.

ESTIMATED ANNUALIZED BURDEN TABLE

Agency	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
DOL	110	2.6	1	286
DOT	50	1.1	1	55
DoED	114	1	1	114
NEA	65	1	32/60	35
USDA	317	1	1	317

ESTIMATED ANNUALIZED BURDEN TABLE—Continued

Agency	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Total	.....	.....	.....	807

Dated: December 20, 2007.

Terry Nicolosi,

Office of the Secretary, Director, Office of Resources Management.

[FR Doc. E7–25431 Filed 1–4–08; 8:45 am]

BILLING CODE 4151–AE–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0487]

Meeting Being Planned to Obtain Public Input for Ensuring the Safety of Pet Food

AGENCY: Food and Drug Administration, HHS

ACTION: Notice of intent to schedule and hold public meeting.

SUMMARY: The Food and Drug Administration (FDA) is announcing its intention to schedule and hold a public meeting early in 2008 to obtain input from stakeholder groups, including, but not limited to, the Association of American Feed Control Officials (AAFCO), veterinary medical associations, animal health organizations, and pet food manufacturers for the development of ingredient, processing, and labeling standards to ensure the safety of pet food. These standards were mandated by the FDA Amendments Act of 2007 (FDAAA).

Date, Time, and Location: The date, time, and location for the 2008 public meeting will be announced in a subsequent notice that will be published in the **Federal Register** a later date.

Addresses: A docket has been opened at FDA to receive any comments in advance of the public meeting. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to either *http://www.fda.gov/dockets/ecomments* or *http://www.regulations.gov*.

FOR FURTHER INFORMATION CONTACT: Walter Osborne, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9024, FAX: 240–276–9101, or e-mail: *walter.osborne@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

FDAAA was signed into law by the President on September 27, 2007 (Public Law 110–085). Title X of the FDAAA includes several provisions pertaining to food safety, including the safety of pet food. Sec. 1002(a)of the new law directs that, within 2 years, FDA is to issue new regulations to establish ingredient standards and definitions, processing standards, and updated standards for labeling to include nutritional and ingredient information. This same provision of the law also directs that, in developing these new regulations, FDA obtain input from its stakeholders, including AAFCO, veterinary medical associations, animal health organizations, and pet food manufacturers. In order to obtain such input, FDA intends to hold a public meeting to hear directly from interested stakeholders.

II. Public Meeting Details

Because FDA is mandated by Congress to establish the new pet food requirements within 2 years of enactment of the FDAAA, it is imperative that the agency begin the rulemaking process as soon as possible. To that end, FDA intends to hold a public meeting in the greater Rockville, MD area sometime within the first 3 months of 2008. After the meeting, FDA will review all of the comments submitted to the docket prior to initiating the regulation drafting process.

III. Comments

FDA will publish a subsequent notice in the **Federal Register** announcing the details of the 2008 public meeting. However, anyone wishing to submit general comments about the new law as it relates to pet food safety or the planned public meeting may do so to the Division of Dockets Management (see *Addresses*). Submit a single copy of electronic comments or two paper copies of any written comments, except that individuals may submit one paper copy. Comments should be identified with the full title and the docket number found in brackets in the heading of this document. Received comments will be available for public examination in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. You may also view received comments on the FDA's Internet site at: *http://www.fda.gov/ohrms/dockets*.

Dated: December 27, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–25599 Filed 1–4–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Child Health and Human Development; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of a meeting of the National Advisory Child Health and Human Development Council.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and/or contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications and/or contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Advisory Child Health and Human Development Council.

Date: January 24, 2008.

Open: 8 a.m. to 12:15 p.m.