awards for the three MIPPA priority areas. Funding will be distributed through a formula as identified in statute. The amounts allocated are based upon factors defined in statute and will be distributed to each priority area based on the formula. ACL will fund total project periods of up to three (3) years contingent upon availability of federal funds.

Priority Area 1—SHIP: $7.5 million in FY 14 for state agencies that administer the SHIP Program.

Priority Area 2—AAA: $7.5 million in FY 14 for SUAs for Area Agencies on Aging and for Native American programs. Funding for Native American Programs ($264,000) is deducted from Priority 2 and is being allocated through a separate process.

Priority Area 3—ADRC: $5 million in FY 2014 for state agencies that administer ADRC programs that were established prior to March 2014.

III. Eligibility Criteria and Other Requirements

1. Eligible Applicants

MIPPA Priority Areas 1, 2 and 3: Awards made under this announcement, by statute, will be made only to agencies of State Governments.

Priority Area 1: Only existing SHIP grant recipients are eligible to apply.

Priority Area 2: Only State Units on Aging are eligible to apply.

Priority Area 3: Only State Agencies that received an ACL and CMS Aging and Disability Resource Center (ADRC) grant where the ADRC was established by March, 2014 are eligible in FY 2014. Eligibility may change if future funding is available.

2. Cost Sharing or Matching

Cost Sharing does not apply.

3. DUNS Number

All grant applicants must obtain and keep current a D–U–N–S number from Dun and Bradstreet. It is a nine-digit identification number, which provides unique identifiers of single business entities. The D–U–N–S number can be obtained from: *https://iupdate.dnb.com/iUpdate/viewiUpdateHome.htm*.

4. Intergovernmental Review

Executive Order 12372, Intergovernmental Review of Federal Programs, is not applicable to these grant applications.

IV. Submission Information

1. Application Kits

Application kits/Program Instructions are available at *www.grantsolutions.gov.* Instructions for completing the application kit will be available on the site.

2. Submission Dates and Times

To receive consideration, applications must be submitted by 11:59 p.m. Eastern time on August 4, 2014.

VII. Agency Contacts

Direct inquiries regarding programmatic issues to U.S. Department of Health and Human Services, Administration on Aging, Office of Supportive and Caregiver Services, Washington, DC 20201, attention: Katherine Glendening or by calling 202–357–3859, or by email *Katherine.Glendening@acl.hhs.gov*.

Dated: June 24, 2014.

Kathy Greenlee,

Administrator and Administration on Aging.

[FR Doc. 2014–15149 Filed 6–26–14; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1155]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Food Labeling Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Food Labeling Regulations'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: On December 31, 2013, the Agency submitted a proposed collection of information entitled ''Food Labeling Regulations'' to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0381. The approval expires on June 30, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: June 23, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–15036 Filed 6–26–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0179]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: On December 13, 2013, the Agency submitted a proposed collection of information entitled ''Prior Notice of Imported Food Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002'' to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0520. The approval expires on June 30, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: June 23, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–15034 Filed 6–26–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–D–1009]

Draft Guidance for Industry on Use of Nanomaterials in Food for Animals; 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 #220) entitled ''Use of Nanomaterials in Food for Animals.'' The draft guidance describes FDA's current thinking regarding the use of nanomaterials or the application of nanotechnology in food for animals. It is intended to assist industry and other stakeholders in identifying potential issues related to safety or regulatory status of food for animals containing nanomaterials or otherwise involving the application of nanotechnology.

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 September 10, 2014.

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: Dragan Momcilovic, Center for Veterinary Medicine (HFV–226), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–453–6856, *dragan.momcilovic@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance for industry (GFI #220) entitled ''Use of Nanomaterials in Food for Animals.'' This draft guidance applies to food ingredients that are intended for use in food for animals and either: (1) Consist entirely of nanomaterials, (2) contain nanomaterials as a component, or (3) otherwise involve the application of nanotechnology.

This guidance is not applicable to other products regulated by FDA, including food substances intended for use in food for humans. This guidance also does not apply to food contact substances or color additives intended for use in food for animals or food for humans.

Medicated feed contains new animal drugs approved for use in or on animal food. This guidance does not apply to a nanomaterial form of a new animal drug or drug component (e.g., drug carrier) in medicated feed. However, it does apply to nanomaterial animal food ingredients in medicated feed.

This guidance is not intended to bring into question the regulatory status of animal food ingredients that naturally exist in the nanoscale range or that contain incidental amounts of particles in the nanoscale range, and that have already been determined to be generally recognized as safe or approved in response to a food additive petition.

A notice announcing the availability of another draft guidance (GFI #221) entitled ''Recommendations for Preparation and Submission of Animal Food Additive Petitions'' was published in the **Federal Register** on September 11, 2013 (78 FR 55727). GFI #221, when finalized, would provide information regarding the submission of food additive petitions (FAPs) for animal food additives. This draft guidance (GFI #220) would provide additional information that would be useful when submitting FAPs for nanomaterial animal food additives and would supplement the information provided in GFI #221.

II. Significance of Guidance

This level 1 draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the Agency's current thinking on this topic. 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.

III. Paperwork Reduction Act of 1995

This draft guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR 571.1 and 571.6 have been approved under OMB control number 0910–0546; the collections of information in 21 CFR 70.25, 71.1, 170.35, 171.1, 21 CFR parts 172, 173, 179, and 180, and in Form FDA 3503, have been approved under OMB control number 0910–0016.

IV. Comments

Interested persons may submit either electronic comments regarding this document to *http://www.regulations.gov* or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at *http://www.regulations.gov.*

V. Electronic Access

Persons with access to the Internet may obtain the draft guidance at either *http://www.fda.gov/AnimalVeterinary/GuidanceComplianceEnforcement/GuidanceforIndustry/default.htm or http://www.regulations.gov.*

Dated: June 23, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–15030 Filed 6–26–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–D–0636]

Global Unique Device Identification Database; Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of the guidance entitled