ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
CDC Partners, Public Health Professionals, Health Care Professionals, General Public	25,000	1	27/60	11,250
Total	25,000			11,250

Dated: August 6, 2010.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010–19911 Filed 8–11–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Aging

Agency Information Collection Activities; Submission for OMB Review; Comment Request; National Survey of Older Americans Act Title III Service Recipients

AGENCY: Administration on Aging, HHS.

ACTION: Notice.

SUMMARY: The Administration on Aging (AoA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by September 13, 2010.

ADDRESSES: Submit written comments on the collection of information by fax 202.395.6974 to the OMB Desk Officer for AoA, Office of Information and Regulatory Affairs, OMB.

FOR FURTHER INFORMATION CONTACT: Valerie Cook 202–357–3583.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, AoA has submitted the following proposed collection of information to OMB for review and clearance.

The National Survey of Older Americans Act Title III Service Recipients information collection, which builds on earlier national pilot studies and surveys, as well as performance measurement tools developed by AoA grantees in the Performance Outcomes Measures Project (POMP), will include consumer assessment surveys for the Congregate and Home-delivered meal nutrition programs; Case Management, Homemaker, and Transportation Services; and the National Family Caregiver Support Program. This information will be used by AoA to track performance outcome measures; support budget requests; comply with Government Performance and Results Act (GPRA) reporting requirements; provide national benchmark information for POMP grantees; and inform program development and management initiatives. Descriptions of previous National Surveys of Older Americans Act Participants can be found under the section on Performance Outcomes on AoA's Web site at: *http://www.aoa.gov/AoARoot/Program_Results/OAA_Performance.aspx.* Copies of the survey instruments and data from previous National Surveys of OAA Participants can be found and queried using the AGing Integrated Database (AGID) at *http://www/agidnet.org/.*

AoA estimates the burden of this collection of information as follows: *Respondents:* Individuals; *Number of Respondents:* 6,250; *Number of Responses per Respondent:* one; *Average Burden per Response:* 6,000 at 30 minutes, 250 at 4 hours: Total Burden: 6,250 hours.

Dated: August 9, 2010.

Kathy Greenlee,

Assistant Secretary for Aging.

[FR Doc. 2010–19957 Filed 8–11–10; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, Coordinating Center for Infectious Diseases: Notice of Charter Amendment

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Board of Scientific Counselors, Coordinating Center for Infectious Diseases, Department of Health and Human Services, has amended their charter to reflect the change in the name of the board to the Board of Scientific Counselors, Office of Infectious Diseases.

For information, contact Robin Mosely, M.A., Designated Federal Officer, Board of Scientific Counselors, Office of Infectious Diseases, CDC, 1600 Clifton Road, NE., Mailstop D10, Atlanta, Georgia 30333, telephone 404/639–4461 or fax 404/639–1255.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: August 4, 2010

Elaine L. Baker,

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

[FR Doc. 2010–19908 Filed 8–11–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–D–0313]

Draft Guidance for Industry: Prevention of *Salmonella* Enteritidis in Shell Eggs During Production, Storage, and Transportation; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance entitled "Prevention of *Salmonella* Enteritidis in Shell Eggs During Production, Storage, and Transportation" (the draft guidance). The draft guidance, when finalized, will provide guidance to egg producers on how to comply with certain provisions contained in FDA's final rule "Prevention of *Salmonella* Enteritidis in Shell Eggs During Production, Storage, and Transportation" (the final rule), including how to implement *Salmonella*

Enteritidis (SE) prevention measures, how to sample for SE, and how to maintain records documenting compliance with the final rule.

DATES: Although you can comment on any guidance at any time (see § 10.115(g)(5) (21 CFR 10.115(g)(5))), to ensure that the agency considers your comments on the draft guidance before it begins work on the final version of the guidance, submit either electronic or written comments on the draft guidance by October 12, 2010.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Division of Plant and Dairy Food Safety/Office of Food Safety, Center for Food Safety and Applied Nutrition (HFS–315), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, or fax your request to 301–436–1070. Send two self-addressed adhesive labels to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance. 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: Nancy Bufano, Center for Food Safety and Applied Nutrition (HFS–316), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–1493.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of July 9, 2009 (74 FR 33030), FDA issued the final rule requiring shell egg producers to implement measures to prevent SE from contaminating eggs on the farm and from further growth during storage and transportation, and requiring these producers to maintain records concerning their compliance with the final rule and to register with FDA. The final rule became effective September 8, 2009.

FDA is issuing the draft guidance as a level 1 draft guidance consistent with FDA's good guidance practices regulation (§ 10.115). The draft guidance, when finalized, will represent the agency's current thinking on how to comply with certain measures designed to prevent SE from contaminating eggs on the farm, as well as how to sample for SE and maintain records documenting compliance with the final rule. 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.

II. 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 118.5, 118.6, 118.10, and 118.11 have been approved under OMB control number 0910–0660.

III. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) either electronic or written comments regarding this document. It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed 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.

IV. Electronic Access

Persons with access to the Internet may obtain the draft guidance at *http://www.fda.gov/FoodGuidances* or *http://www.regulations.gov*.

Dated: August 9, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010–19905 Filed 8–11–10; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Intent To Award Patient Protection and Affordable Care Act Funding to Approved But Unfunded Applications (ABU) Formerly Received in Response to the American Recovery and Reinvestment Act of 2009 (ARRA) Centers for Disease Control and Prevention Funding Opportunity DP09–912ARRA09, "Communities Putting Prevention to Work (CPPW)"

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: This notice provides notice of CDC's intent to fund additional Approved but Unfunded (ABU) cooperative agreement applications previously received and competed in response to CDC Funding Opportunity, CDC–RFA–DP09–912ARRA09, "Communities Putting Prevention to Work" (CPPW). It is the intent of CDC to fund additional previously received applications with Patient Protection Affordable Care Act (PPACA), Section 4002, appropriations. To this end, CDC will remove the following ARRA–Specific Requirements published in the aforementioned funding opportunity announcement:

—Catalogue of Domestic Assistance Number 93.724

—*Recovery Act-Specific Reporting Requirements*

Recipients of Federal awards from funds authorized under Division A of the Recovery Act must comply with all requirements specified in Division A of the Recovery Act (Pub. L. 111–5), including reporting requirements outlined in Section 1512 of the Act and designated Recovery Act outcome and output measures as detailed at the end of this section. For purposes of reporting, Recovery Act recipients must report on Recovery Act sub-recipient (sub-grantee and sub-contractor) activities as specified below.

Not later than 10 days after the end of each calendar quarter, starting with the quarter ending _____; and reporting by _____, the recipient must submit quarterly reports to HHS that will posted to Recovery.gov, containing the following information:

a. The total amount of Recovery Act funds under this award;

b. The amount of Recovery Act funds received under this award that were obligated and expended to projects or activities;

c. The amount of unobligated award balances;

d. A detailed list of all projects or activities for which Recovery Act funds under this award were obligated and expended, including

- The name of the project or activity;
- A description of the project or activity;
- An evaluation of the completion status of the project or activity;
- An estimate of the number of jobs created and the number of jobs retained by the project or activity (see OMB Guidance M–09–21, June 22, 2009) and;
- For infrastructure investments made by State and local governments, the purpose, total cost, and rationale of the agency for funding the infrastructure investment with funds made available under this Act, and the name of the person to contact at the agency if there