

## 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.

**BILLING CODE 4163–18–P**

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Disease Control and Prevention

**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 Disease Control and Prevention.


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

**BILLING CODE 4154–01–P**

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### Table: Estimated Annualized Burden Hours

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>CDC Partners, Public Health Professionals, Health Care Professionals, General Public</td>
<td>25,000</td>
<td>1</td>
<td>27/60</td>
<td>11,250</td>
</tr>
<tr>
<td>Total</td>
<td>25,000</td>
<td></td>
<td></td>
<td>11,250</td>
</tr>
</tbody>
</table>
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.


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.

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 under this Act, and the name of the person to contact at the agency if there