Additionally, the evaluation will capture information to describe the quality of the implementation of the various mentor-coaching approaches including consistency of the mentor-coach implementation with the planned approach, the frequency and content of interactions between the mentor-coaches and the teaching staff, and apparent changes in teaching staff behavior, including their own professional development. The evaluation will also capture information about the characteristics of those who provided coaching, the characteristics of teaching staff that were mentored, as well as the characteristics of the settings and the systems in which the mentor-coaching was embedded. Lastly, the evaluation will document the factors that appear to be most critical to successful implementation and implementation challenges.

The data collection will include a census survey of all grantees; a census survey of all mentor-coaches; telephone interviews with a sub-sample of administrators, mentor-coaches, and teaching staff; and a mentor-coach activity snapshot.

Respondents: Grantee and center administrative staff, mentor-coaches, teaching staff.

### ANNUAL BURDEN ESTIMATES

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Annual number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total annual burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grantee Census Survey</td>
<td>131</td>
<td>1</td>
<td>0.5</td>
<td>66</td>
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<tr>
<td>Mentor-Coach Census Survey</td>
<td>400</td>
<td>1</td>
<td>0.5</td>
<td>200</td>
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<tr>
<td>Administrator Telephone Interview</td>
<td>85</td>
<td>1</td>
<td>1.0</td>
<td>85</td>
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<tr>
<td>Mentor-Coach Telephone Interview</td>
<td>65</td>
<td>1</td>
<td>1.0</td>
<td>65</td>
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<tr>
<td>Teaching Staff Telephone Interview</td>
<td>130</td>
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<td>1.0</td>
<td>130</td>
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<tr>
<td>Mentor-Coach Activity Snapshot</td>
<td>65</td>
<td>2</td>
<td>0.25</td>
<td>33</td>
</tr>
</tbody>
</table>

Estimated Total Annual Burden Hours: 579.

Additional Information: In compliance with the requirements of Section 3506(C)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. E-mail address: OPREinfocollection@acf.hhs.gov.

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the Federal Register. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: May 17, 2011.

Steven M. Hamner, OPRE Reports Clearance Officer.

[FR Doc. 2011–12787 Filed 5–25–11; 8:45 am]

BILLING CODE 4184–22–M

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Food and Drug Administration**

[Docket No. FDA–2010–D–0153]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Draft Guidance for Industry and Food and Drug Administration Staff: Food and Drug Administration and Industry Procedures for Section 513(g) Requests for Information Under the Federal Food, Drug, and Cosmetic Act

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Fax written comments on the collection of information by June 27, 2011.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or e-mailed to oira_submission@omb.eop.gov. All comments should be identified with the OMB control number 0910–NEW and title “Draft Guidance for Industry and FDA Staff: FDA and Industry Procedures for Section 513(g) Requests for Information Under the Federal Food, Drug, and Cosmetic Act.” Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, Daniel.Gittleson@fda.hhs.gov.

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

Draft Guidance for Industry and FDA Staff: FDA and Industry Procedures for Section 513(g) Requests for Information under the Federal Food, Drug, and Cosmetic Act—(OMB Control Number 0910–NEW)

Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from OMB for each collection of information they conduct or sponsor. “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
Respondents to this collection of information are mostly device manufacturers; however, anyone may submit a 513(g) request for information. The total number of annual responses is based on the average number of 513(g) requests received each year by the Agency. FDA based its estimates on the number of 513(g) requests for information received by both CDRH and CBER from 2007 to 2009.

Dated: May 20, 2011.

Leslie Kux,
Acting Assistant Commissioner for Policy.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration
[Docket No. FDA–2011–N–0320]

Agency Information Collection Activities; Proposed Collection; Comment Request; Experimental Study on Consumer Responses to Whole Grain Labeling Statements on Food Packages

AGENCY: Food and Drug Administration, HHHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the Federal Register concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on a study entitled: “Experimental Study on Consumer Responses to Whole Grain Labeling Statements on Food Packages.”

DATES: Submit either electronic or written comments on the collection of information by July 25, 2011.

ADDRESSES: Submit electronic comments on the collection of information to http://www.regulations.gov. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the