

1996), as well as the William Wilberforce Trafficking Victims Protection Reauthorization Act of 2008 (Pub. L. 110–457), which authorizes post release services under certain conditions to eligible children. All programs must comply with the Flores Settlement Agreement, Case No. CV85–4544–RJK (C.D. Cal. 1996), pertinent regulations and ORR policies and procedures.

**Christopher Beach,**

Office of Administration, Office of Financial Services, Division of Grants Policy.

[FR Doc. 2016–31014 Filed 12–22–16; 8:45 am]

**BILLING CODE 4184–45–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Proposed Information Collection Activity; Comment Request**

*Proposed Projects: Childcare.gov.*

*Title: CCDF Grantee Consumer Education Database Linkages with Childcare.gov.*

*OMB No.: New.*

*Description:* The Child Care and Development Block Grant (CCDBG) Act of 2014 requires HHS to create a national Web site for consumer education. The National Web site will be hosted at *childcare.gov*. CCDBG grantees are also required to stand up child care consumer education Web sites that meet the requirements of the law. The CCDBG Final Rule aligns the National and State Web sites by requiring Lead Agencies to provide HHS

with linkages to their databases that store consumer education information. The *Childcare.gov* Web site, maintained by Office of Child care will collect child care specific information from State and Territory databases and make that information available for parents using the *childcare.gov* Web site to search for child care that meets their needs. *Childcare.gov* will provide consumers, directly or through linkages to State and Territory data sources, with the following minimum information and services:

(1) A localized list of all eligible child care providers, differentiating between licensed and license-exempt providers;

(2) Child care provider-specific information from a quality rating and improvement system or information about other quality indicators, to the extent that such information is publicly available and practicable.

*Respondents:* CCDBG grantees in States and Territories.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
<i>Childcare.gov</i> data collection: Establish and maintain Web-based data connection in subsequent years .....	56	260	.57	8,299

**Estimated Total Annual Burden Hours: 8,299 hours.**

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chap 35), 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 of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov*. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on

respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

**Robert Sargis,**

Reports Clearance Officer.

[FR Doc. 2016–30982 Filed 12–22–16; 8:45 am]

**BILLING CODE 4184–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2012–N–1021]

**Medical Device User Fee and Modernization Act; Notice to Public of Web Site Location of Fiscal Year 2017 Proposed Guidance Development**

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or the Agency) is announcing the Web site location where the Agency will post two lists of guidance documents that the Center for Devices and Radiological Health (CDRH

or the Center) intends to publish in fiscal year (FY) 2017. In addition, FDA has established a docket where interested persons may comment on the priority of topics for guidance, provide comments and/or propose draft language for those topics, suggest topics for new or different guidance documents, comment on the applicability of guidance documents that have issued previously, and provide any other comments that could benefit the CDRH guidance program and its engagement with stakeholders. This feedback is critical to the CDRH guidance program to ensure that we meet stakeholder needs.

**DATES:** Submit either electronic or written comments by February 21, 2017.  
**ADDRESSES:** You may submit comments as follows:

*Electronic Submissions*

Submit electronic comments in the following way:

- Federal eRulemaking Portal: *https://www.regulations.gov*. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are