**Department of Health and Human Services**

**Administration for Children and Families**

**Proposed Information Collection Activity; Comment Request**

**Proposed Projects:**

**Title:** DRA TANF Final Rule.

**OMB No.:** 0970–0338.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**ANNUAL BURDEN ESTIMATES**

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Preparation and Submission of Data Verification Procedures §§ 261.60–261.63</td>
<td>54</td>
<td>1</td>
<td>640</td>
<td>34,560</td>
</tr>
<tr>
<td>Caseload Reduction Documentation Process, ACF–202 §§ 261.41 &amp; 261.44</td>
<td>54</td>
<td>1</td>
<td>120</td>
<td>6,480</td>
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<tr>
<td>Reasonable Cause/Corrective Compliance Documentation Process §§ 262.4, 262.6, &amp; 262.7; § 261.51</td>
<td>54</td>
<td>2</td>
<td>240</td>
<td>25,920</td>
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<tr>
<td>TANF Data Report Part 265</td>
<td>54</td>
<td>4</td>
<td>2,201</td>
<td>475,416</td>
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<tr>
<td>SSP–MOE Data Report Part 265</td>
<td>29</td>
<td>4</td>
<td>714</td>
<td>82,824</td>
</tr>
</tbody>
</table>

**Estimated Total Annual Burden Hours:** 625,200.

In compliance with the requirements of Section 506(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 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.

**BILLING CODE 4184–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA–2016–N–1380]

**The Role of Hospitals in Modernizing Evidence Generation for Device Evaluation: Harnessing the Digital Revolution for Surveillance; Public Workshop; Request for Comments**

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

***ACTION***: Notice of public workshop; request for comments.

**SUMMARY**: The Food and Drug Administration (FDA) is announcing the following public workshop entitled “The Role of Hospitals in Modernizing Evidence Generation for Device Evaluation: Harnessing the Digital Revolution for Surveillance.” Hospitals play a critical role in the development...