DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control; Special Emphasis Panel (SEP): Ability of Individual and Integrated Tick Management (ITM) Technologies To Reduce the Entomological Risk of Lyme Disease, Funding Opportunity Announcement (FOA) CK11–005, Initial Review

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the aforementioned meeting:

**Time and Date:** 12 p.m.–2 p.m., May 10, 2011 (Closed).

**Place:** Teleconference.

**Status:** The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of 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 Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**Dated:** February 4, 2011.

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

**Contact Person for More Information:**
Amy Yang, PhD, Scientific Review Officer, CDC, 1600 Clifton Road, NE., Mailstop E60, Atlanta, Georgia 30333, Telephone: (404) 498–2733.

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 Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

**BILLING CODE 4163–18–P**

<table>
<thead>
<tr>
<th>Instrument or requirement</th>
<th>Number of respondents</th>
<th>Yearly submittals</th>
<th>Average burden hours per response</th>
<th>Final rule total annual burden hours</th>
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<tr>
<td>Preparation and Submission of Data Verification Procedures—§§ 261.60–261.63</td>
<td>54</td>
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<td>640</td>
<td>34,560</td>
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<td>Caseload Reduction Documentation Process, ACF–202—§§ 261.41 &amp; 261.44</td>
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<td>Reasonable Cause/Corrective Compliance Documentation Process—§§ 262.4, 262.6, &amp; 262.7; § 261.51</td>
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<td>TANF Data Report—Part 265</td>
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<td>SSP–MOE Data Report—Part 265</td>
<td>29</td>
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**Total Burden Hours:** 625,200.

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 of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L’Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer, E-mail address: infocollect@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


Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance on Consultation Procedures: Foods Derived From New Plant Varieties

**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