performance of functions of the FAR, and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

DATES: Submit comments on or before August 26, 2011.

ADDRESSES: Submit comments identified by Information Collection 9000–0164, Contractor Business Ethics Compliance Program and Disclosure Requirements, by any of the following methods:

- Regulations.gov: http://www.regulations.gov. Submit comments via the Federal eRulemaking portal by inputting “Information Collection 9000–0164, Contractor Business Ethics Compliance Program and Disclosure Requirements”, under the heading “Enter Keyword or ID” and selecting “Search”. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0164, Contractor Business Ethics Compliance Program and Disclosure Requirements”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0164, Contractor Business Ethics Compliance Program and Disclosure Requirements”, on your attached document.
- Instructions: Please submit comments only and cite Information Collection 9000–0164, Contractor Business Ethics Compliance Program and Disclosure Requirements, in all correspondence related to this collection. All comments received will be posted without change to http://www.regulations.gov, including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT: Mr. Anthony Robinson, Procurement Analyst, Acquisition Policy Division, GSA (202) 501–2658 or e-mail Anthony.Robinson@gsa.gov.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Secretary’s Advisory Committee on Human Research Protections

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of the Assistant Secretary for Health.

ACTION: Notice.

SUMMARY: Pursuant to Section 10(a) of the Federal Advisory Committee Act, U.S.C. Appendix 2, notice is hereby given that the Secretary’s Advisory Committee on Human Research Protections (SACHRP) will hold its twenty-fifth meeting. The meeting will be open to the public. Information about SACHRP and the meeting agenda will be posted on the SACHRP Web site at: http://www.dhhs.gov/ohrp/sachrp/mtngs/index.html.

DATES: The meeting will be held on Tuesday, July 19, 2011 from 8:30 a.m. until 5 p.m. and Wednesday, July 20, 2011 from 8:30 a.m. until 5 p.m.


FOR FURTHER INFORMATION CONTACT: Jerry Menikoff, M.D., J.D., Director, Office for Human Research Protections (OHRP), or Julia Gorey, J.D., Executive Director, SACHRP; U.S. Department of Health and Human Services, 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852; 240–453–8141; fax: 240–453–6909; e-mail address: Julia.Gorey@hhs.gov.

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, SACHRP was established to provide expert advice and recommendations to the Secretary of Health and Human Services and the Assistant Secretary for Health on issues and topics pertaining to or associated with the protection of human research subjects.

On July 19, 2011, SACHRP will hear a presentation by the Executive Director of the Presidential Commission for the Study of Bioethical Issues focusing on the work of the Commission; this will be followed by SACHRP discussion. After lunch, SACHRP will hear the report of the Subpart A Subcommittee (SAS). SAS is charged with developing recommendations for consideration by SACHRP regarding the application of subpart A of 45 CFR part 46 in the current research environment; this Subcommittee was established by SACHRP in October 2006.

Recommendations to be discussed focus on the return of research results to subject, internet-based research, and improvements to the informed consent process.

On July 20, 2011, the morning will open with a report from the Subcommittee on Harmonization (SOH). The SOH was established by SACHRP at its July 2009 meeting, and is charged with identifying and prioritizing areas in which regulations and/or guidelines for human subjects research adopted by various agencies or offices within HHS would benefit from harmonization, consistency, clarity, simplification and/or coordination. In the afternoon, SACHRP will hear a panel of speakers discussing consequences and processes surrounding scientific misconduct and fraud.

Public Comment will be heard on both days.

Public attendance at the meeting is limited to space available. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated persons. Members of the public will have the opportunity to provide comments on
both days of the meeting. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed materials distributed to SACHRP members for this scheduled meeting should submit materials to the Executive Director, SACHRP, prior to the close of business July 15, 2011.

Dated: June 21, 2011.
Jerry Menikoff,
Director, Office for Human Research Protections, Executive Secretary, Secretary’s Advisory Committee on Human Research Protections.

[FR Doc. 2011–16051 Filed 6–24–11; 8:45 am]
BILLING CODE 4150–36–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Administration for Children and Families

**Submission for OMB Review; Comment Request**

**Title:** Head Start Program Information Report.

**OMB No.:** 0980–0017.

**Description:** The Office of Head Start within the Administration for Children and Families, United States Department of Health and Human Services, is proposing to renew authority to collect information using the Head Start Program Information Report (PIR). The PIR provides information about Head Start and Early Head Start services received by the children and families enrolled in Head Start programs. The information collected in the PIR is used to inform the public about these programs and to make periodic reports to Congress about the status of children in Head Start programs as required by the Head Start Act.

**Respondents:** Head Start and Early Head Start Program Grant Recipients.

**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>
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</thead>
<tbody>
<tr>
<td>Annual PIR</td>
<td>2,690</td>
<td>1</td>
<td>4 hours</td>
<td>10,760</td>
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<tr>
<td>Monthly Enrollment (Grantees only)</td>
<td>1,600</td>
<td>12</td>
<td>3 minutes</td>
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**Estimated Total Annual Burden**

Hours: 12,392.

**Additional Information:** Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. E-mail address: infocollection@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 making a decision concerning the collection of information between 30 and 60 days after publication of this document in the Federal Register. 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: 202–395–7285, E-mail: OIRA_SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families. Jerry Menikoff, Director, Office for Human Research Protections, Executive Secretary, Secretary’s Advisory Committee on Human Research Protections.

[FR Doc. 2011–16014 Filed 6–24–11; 8:45 am]
BILLING CODE 4184–01–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Administration for Children and Families

**Submission for OMB Review; Comment Request**

**Title:** ACF–535 LIHEAP Quarterly Allocation Estimates.

**OMB No.:** 0970–0037.

**Description:** The LIHEAP Quarterly Allocation Estimates, ACF Form-535 is a one-page form that is sent to 50 State grantees and to the District of Columbia. It is also sent to Tribal Government grantees that receive over $1 million annually for the Low Income Home Energy Assistance Program (LIHEAP). Grantees are asked to complete and submit the form in the 4th quarter of each year. The data collected on the form are grantees estimates of obligations they expect to make each quarter for the upcoming fiscal year for the LIHEAP program. This is the only method used to request anticipated distributions of the grantees LIHEAP funds. The information is used to develop apportionment requests to OMB and to make grant awards based on grantees anticipated needs. Information collected on this form is not available through any other Federal source.

**Submission of the form is voluntary.**

**Respondents:** State Governments.

**Annual Burden Estimates:**

<table>
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<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>
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<td>0.25</td>
<td>13.75</td>
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</table>

**Estimated Total Annual Burden**

Hours: 13.75.

**Additional Information:** Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. E-mail address: infocollection@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.