without change to http://www.regulations.gov, including any personal information provided. For additional information on submitting comments, see the “Request for Comments” heading of the SUPPLEMENTARY INFORMATION section.

Docket: For access to the docket to read background documents or comments received, go to http://www.regulations.gov and insert the docket number, found in brackets in the heading of this document, into the “Search” box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT:
Mary Weick-Brady, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5426, 301–796–6089, FAX: 301–847–8510, email: Mary.Brady@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:
I. Background

In the Federal Register of January 7, 2013 (78 FR 951), FDA published a notice of public workshop with a 90-day comment period to request comments on all aspects of the public workshop, including topics outlined in section II of that document (78 FR 951 at 952).

The agency has received a request for an extension of the comment period until May 30, 2013. The request conveyed concern that the current comment period does not allow sufficient time to develop a meaningful or thoughtful response that allows for consideration of presentations by FDA and other stakeholders at the public workshop on April 29 and 30, 2013. FDA has considered the request and is extending the comment period for the notice of public workshop until May 30, 2013. The request is providing FDA with a sufficient time to develop a meaningful response that allows for consideration of presentations by FDA and other stakeholders at the public workshop on April 29 and 30, 2013.

The agency has received a request for an extension of the comment period until May 30, 2013. The request conveyed concern that the current comment period does not allow sufficient time to develop a meaningful or thoughtful response that allows for consideration of presentations by FDA and other stakeholders at the public workshop on April 29 and 30, 2013. FDA has considered the request and is extending the comment period for the notice of public workshop until May 30, 2013. The agency believes that the extension allows adequate time for interested persons to submit comments without significantly delaying consideration of these important issues.

II. Request for Comments

Regardless of attendance at the public workshop, interested persons may submit either electronic comments regarding this document to http://www.regulations.gov or written comments to the Division of Dockets Management (see ADDRESSES). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. In addition, when responding to specific questions as outlined in section II of the notice of public workshop (78 FR 951 at 952), please identify the question you are addressing. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at http://www.regulations.gov.

Dated: March 20, 2013.

Leslie Kux,
Assistant Commissioner for Policy.
[FR Doc. 2013-06725 Filed 3–22–13; 8:45 am]

SUPPLEMENTARY INFORMATION:

II. Request for Comments

Regardless of attendance at the public workshop, interested persons may submit either electronic comments regarding this document to http://www.regulations.gov or written comments to the Division of Dockets Management (see ADDRESSES). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. In addition, when responding to specific questions as outlined in section II of the notice of public workshop (78 FR 951 at 952), please identify the question you are addressing. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at http://www.regulations.gov.

Dated: March 20, 2013.

Leslie Kux,
Assistant Commissioner for Policy.
[FR Doc. 2013-06725 Filed 3–22–13; 8:45 am]

Department of Health and Human Services

National Institutes of Health

Proposed Collection; Comment Request: NIH Office of Intramural Training & Education Application

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the Office of Intramural Training & Education/OIR/OD, the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection

Title: NIH Office of Intramural Training & Education Application. Type of Information Collection Request: Revision. Form Number: 0925–0299. Expiration Date: March 31, 2014. Need and Use of Information Collection: The Office of Intramural Training & Education (OITE) administers a variety of programs and initiatives to recruit pre-college through post-doctoral educational level individuals into the National Institutes of Health Intramural Research Program (NIH–IRP) to facilitate development into future biomedical scientists. The proposed information collection is necessary in order to determine the eligibility and quality of potential awardees for traineeships in these programs. The applications for admission consideration include key areas such as: personal information, eligibility criteria, contact information, student identification number, training program selection, scientific discipline interests, educational history, standardized examination scores, reference information, resume components, employment history, employment interests, dissertation research details, letters of recommendation, financial aid history, sensitive data, future networking contact, travel information, as well as feedback questions about interviews and application submission experiences. Sensitive data collected on the applicants, race, gender, ethnicity, disability, and recruitment method, are made available only to OITE staff members or in aggregate form to select NIH offices and are not used by the admission committee for admission consideration; optional to submit.

Frequency of Response: On occasion. Affected Public: Individuals seeking intramural training opportunities and references for these individuals. Type of Respondents: students, post-baccalaureates, technicians, graduate students, post-doctorates, references, and alumni. There are no capital costs, operating costs, and/or maintenance costs to report.

The annual reporting burden is displayed in the following table:

<table>
<thead>
<tr>
<th>Type of respondent</th>
<th>Estimated No. of respondents</th>
<th>Estimated total annual burden hours</th>
<th>Estimated total annual burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Summer Internship Program in Biomedical Research (SIP)</td>
<td>6,820.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>Biomedical Engineering Summer Internship Program (BESIP)</td>
<td>80.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>Post-baccalaureate Training Program (PBT)</td>
<td>1,885.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>Community College Summer Enrichment Program (CCSEP)</td>
<td>100.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>Technical Training Program (PTT)</td>
<td>115.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>Graduate Partnerships Program (GPP)—Application (Select Institutional Partnerships)</td>
<td>250.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>Graduate Partnerships Program (GPP)—Registration (Select Institutional Partnerships + Individual Partnership)</td>
<td>140.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>National Graduate Student Research Conference (NGSRC)</td>
<td>800.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
</tbody>
</table>
**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Center for Scientific Review; Notice of Closed Meeting**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

**Name of Committee:** Center for Scientific Review Special Emphasis Panel; Member Conflict: Neuroscience

**Date:** April 8, 2013.

**Time:** 2:00 p.m. to 4:00 p.m.

**Agenda:** To review and evaluate grant applications.

**Place:** National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892.

(Telephone Conference Call)

**Contact Person:** Richard D Crosland, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4158, MSC 7850, Bethesda, MD 20892, 301–435–1220, rc218u@nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.


Dated: March 18, 2013.

Melanie J. Gray,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–06685 Filed 3–22–13; 8:45 am]

BILLING CODE 4140–01–P

---

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Cancer Institute Amended; Notice of Meeting**

Notice is hereby given of a change in the meeting of the National Cancer Institute Special Emphasis Panel, April 24, 2013, 8:30 a.m.–4:30 p.m., 9000 Rockville Pike, Building 45, Conference Room D, Bethesda, MD 20892 which was published in the Federal Register on March 8, 2013, 78FR15021.

This notice is being amended to change the meeting format from a face to face meeting to a teleconference on Wednesday April 24, 2013, 8:30 a.m.–4:30 p.m. The meeting is closed to the public.

Dated: March 18, 2013.

Melanie J. Gray,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–06686 Filed 3–22–13; 8:45 am]

BILLING CODE 4140–01–P

---

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Center for Scientific Review; Closed Meetings**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the