This proposed information collection is being submitted to the Office of Management and Budget, Office of Information and Regulatory Affairs, for review and clearance in accordance with the provisions of the Paperwork Reduction Act of 1995, unless it displays a currently valid OMB control number. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and you are not required to respond to, an information collection unless it displays a currently valid OMB control number.

**SUMMARY:** Under the provisions of Section 3507(a)(1) of the Paperwork Reduction Act of 1995, the National Institutes of Health, National Cancer Institute (NCI), the National Human Genome Research Institute, and the National Institute on Aging are soliciting comments on this information collection request.

**Department of Health and Human Services**

**National Institutes of Health**

**DHHS**

**NIH**

**NCI**

**NHGRI**

**Director, DCCPS, NCI/NIH, 6130 Executive Blvd, Bethesda, MD 20892; telephone 301–594–6652 or e-mail your comments to willis@mail.nih.gov.**

**Comment Request; Questionnaire Cognitive Interview and Pretesting**

This proposed information collection is being submitted to the Office of Management and Budget, Office of Information and Regulatory Affairs, for review and clearance in accordance with the provisions of the Paperwork Reduction Act of 1995, unless it displays a currently valid OMB control number. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and you are not required to respond to, an information collection unless it displays a currently valid OMB control number.

**SUMMARY:** Under the provisions of Section 3507(a)(1) of the Paperwork Reduction Act of 1995, the National Institutes of Health, National Cancer Institute (NCI), the National Human Genome Research Institute, and the National Institute on Aging are soliciting comments on this information collection request.

<table>
<thead>
<tr>
<th>Type of Respondents</th>
<th>Number of Respondents</th>
<th>Frequency of Response</th>
<th>Burden Hours Total</th>
<th>Per response Burden Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>General Public</td>
<td>1,200</td>
<td>1</td>
<td>3,600.0</td>
<td>(1.25)</td>
</tr>
<tr>
<td>Experts in their Field</td>
<td>600</td>
<td>1</td>
<td>600.0</td>
<td>(1.25)</td>
</tr>
<tr>
<td>Physicians, Scientists, and similar Respondents</td>
<td>1,200</td>
<td>1</td>
<td>1,200.0</td>
<td>(1.25)</td>
</tr>
</tbody>
</table>

**Comment Request; Questionnaire Cognitive Interview and Pretesting**

This proposed information collection is being submitted to the Office of Management and Budget, Office of Information and Regulatory Affairs, for review and clearance in accordance with the provisions of the Paperwork Reduction Act of 1995, unless it displays a currently valid OMB control number. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and you are not required to respond to, an information collection unless it displays a currently valid OMB control number.

**SUMMARY:** Under the provisions of Section 3507(a)(1) of the Paperwork Reduction Act of 1995, the National Institutes of Health, National Cancer Institute (NCI), the National Human Genome Research Institute, and the National Institute on Aging are soliciting comments on this information collection request.

<table>
<thead>
<tr>
<th>Type of Respondents</th>
<th>Number of Respondents</th>
<th>Frequency of Response</th>
<th>Burden Hours Total</th>
<th>Per response Burden Hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>General Public</td>
<td>1,200</td>
<td>1</td>
<td>3,600.0</td>
<td>(1.25)</td>
</tr>
<tr>
<td>Experts in their Field</td>
<td>600</td>
<td>1</td>
<td>600.0</td>
<td>(1.25)</td>
</tr>
<tr>
<td>Physicians, Scientists, and similar Respondents</td>
<td>1,200</td>
<td>1</td>
<td>1,200.0</td>
<td>(1.25)</td>
</tr>
</tbody>
</table>
DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of 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 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: Experimental Oncology.

Date: March 17, 2011.

Time: 1 p.m. to 3 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: Syed M Quadri, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6210, MSC 7804, Bethesda, MD 20892, 301–435–1211, quadris@csr.nih.gov.

Comment: Applications: NeuroAIDS Applications.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Project: NeuroAIDS Applications.

Date: March 30–31, 2011.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Eduardo A Montalvo, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5108, MSC 7852, Bethesda, MD 20892, (301) 435–1168, montalve@csr.nih.gov.

Comment: Applications: Visceral Pain Relief.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Visceral Pain Relief.

Date: March 30, 2011.

Time: 1 p.m. to 2 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: M Catherine Bennett, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5182, MSC 7846, Bethesda, MD 20892, 301–435–1766, bennettec3@csr.nih.gov.

Comment: Applications: NeuroAIDS Applications.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Project: Cell Biology.

Date: March 29–30, 2011.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: David Balasundaram, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5189, MSC 7840, Bethesda, MD 20892, 301–435–1022, balasundaram@csr.nih.gov.

Comment: Applications: NeuroAIDS Applications.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Project: NeuroAIDS Applications.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Comments are invited on: (a) Whether the proposed collections of information are 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) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on