2,500 hours). Thus, the total estimated burden is 3,906 hours. This estimate is 1,352 hours lower than the 5,258 hours published in the 60-day notice and reflects 20 fewer hours for the pretest invitation, 12 fewer hours for the pretest, and 1,320 fewer hours for the survey invitation. Recent evidence available to the Agency suggests the study will not need to send as many pretest or survey invitations as originally estimated to achieve its target sample sizes in the pretest and survey. The number of pretests was changed from 200 to 150 to correct an error that was made in the 60-day notice.

FDA estimates the burden of this collection of information as follows:

<table>
<thead>
<tr>
<th>Activity</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Total annual responses</th>
<th>Average burden per response</th>
<th>Total hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cognitive Interview Screener</td>
<td>72</td>
<td>1</td>
<td>72</td>
<td>0.083 (5 min.)</td>
<td>6</td>
</tr>
<tr>
<td>Cognitive Interview</td>
<td>9</td>
<td>1</td>
<td>9</td>
<td>1</td>
<td>9</td>
</tr>
<tr>
<td>Pretest</td>
<td>1,000</td>
<td>1</td>
<td>1,000</td>
<td>0.032 (2 min.)</td>
<td>33</td>
</tr>
<tr>
<td>Pretest Invitation</td>
<td>150</td>
<td>1</td>
<td>150</td>
<td>0.25 (15 min.)</td>
<td>38</td>
</tr>
<tr>
<td>Survey Invitation</td>
<td>40,000</td>
<td>1</td>
<td>40,000</td>
<td>0.033 (2 min.)</td>
<td>1,320</td>
</tr>
<tr>
<td>Survey</td>
<td>10,000</td>
<td>1</td>
<td>10,000</td>
<td>0.25 (15 min.)</td>
<td>2,500</td>
</tr>
</tbody>
</table>

Total                                                                                                           3,906

*There are no capital costs or operating and maintenance costs associated with this collection of information.*

**II. References**

The following references have been placed on display in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday. (FDA has verified the Web site addresses, but we are not responsible for any subsequent changes to the Web sites after this document is published in the *Federal Register.*)


Dated: December 22, 2011.

Leslie Kux,
Acting Assistant Commissioner for Policy.

[PR Doc. 2011–33303 Filed 12–28–11; 8:45 am]

**BILLING CODE 4160–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Office of Inspector General

[Docket ID OIG 910–N]

Privacy Act; System of Records

**AGENCY:** Office of Inspector General, HHS.

**ACTION:** Notice of amendment to system of existing records.

**SUMMARY:** In accordance with the Privacy Act of 1974, as amended, the Office of Inspector General gives notice of a proposed amendment to its Privacy Act system of records entitled “Consolidated Data Repository” (09–90–1000). This system of records is being amended to include records regarding Federal and State benefit programs and service providers in Federal health care programs.
DATES: Effective Date: This system of records will become effective without further notice on February 27, 2012, unless comments received on or before that date result in a contrary determination.

Comment Date: Comments on this amendment to the system of records will be considered if we receive them at the addresses provided below no later than 5 p.m. on January 30, 2012.

ADDRESSES: You may submit your written comments, identified by OIG–910–N, by any of the following methods:

• Federal Rulemaking Portal: http://www.regulations.gov. Follow the online instructions for submitting comments.


Instructions: We do not accept comments by facsimile (FAX) transmission. All submissions received must include the agency name and docket number for this Federal Register document. The general policy for comment submissions from members of the public is to make these available for public viewing on http://www.regulations.gov after receipt. All comments, including attachments and other supporting materials, received are subject to public disclosure.

FOR FURTHER INFORMATION CONTACT: Patrice Drew, OIG Regulatory Officer, External Affairs, (202) 619–1368.

SUPPLEMENTARY INFORMATION: In accordance with the Privacy Act of 1974 (5 U.S.C 552a), an agency is to publish a notice in the Federal Register when there is a revision, change, or addition to its system of records. OIG is proposing to amend its system of records entitled “Consolidated Data Repository” (SORN 09–90–1000). OIG is adding record sources to the system. This system fulfills our responsibilities under the Inspector General Act of 1978 (5 U.S.C. App.) “to conduct and supervise audits and investigations relating to the administration and operations” of the Department of Health and Human Services (HHS). This amendment will assist OIG in performing timely and independent audits, evaluations and inspections, and investigations of the Medicare and Medicaid programs.

SYSTEM NAME: Consolidated Data Repository–HHS–OIG (SORN 09–90–1000).

RECORD SOURCE CATEGORIES:

Description of the Change: Remove the current entry and in its place add the following: "Sources of information in this records system include: Federal, State, and local government records regarding Medicare, Medicaid, and other benefit programs; Department documents and records; materials regarding service providers in Federal health care programs furnished by nongovernmental sources; and public source materials."

Dated: December 22, 2011.

Daniel R. Levinson, Inspector General.

[FR Doc. 2011–33346 Filed 12–28–11; 8:45 am]

BILLING CODE 4150–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; 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 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 Skeletal Healing, Regeneration and Repair.

Date: January 18, 2012.

Time: 11 a.m. to 1 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: Priscilla B. Chen, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4104, MSC 7814, Bethesda, MD 20892. (301) 435–1787, chenpl@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel PAR Panel: NHLBI Systems Biology.

Date: January 19–20, 2012.

Time: 8 a.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

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

(Virtual Meeting).

Contact Person: Ai-Ping Zou, MD, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4118, MSC 7814, Bethesda, MD 20892. (301) 435–1777, zouai@csr.nih.gov.