pursuant to Exemptions (8), and (9)(A)(ii) and (9)(B).

2. Consideration of supervisory activities. Closed pursuant to Exemptions (9)(A)(ii) and (9)(B).

FOR FURTHER INFORMATION CONTACT: Mary Rupp, Secretary of the Board, Telephone: 703–518–6304.

Mary Rupp,

Board Secretary.

[FR Doc. E8–24560 Filed 10–10–08; 11:15 am]

BILLING CODE 7535–01–P

THE NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

Meetings of Humanities Panel

AGENCY: The National Endowment for the Humanities.

ACTION: Notice of meetings.

SUMMARY: Pursuant to the provisions of the Federal Advisory Committee Act (Pub. L. 92–463, as amended), notice is hereby given that the following meetings of Humanities Panels will be held at the Old Post Office, 1100 Pennsylvania Avenue, NW., Washington, DC 20506.

FOR FURTHER INFORMATION CONTACT: Michael P. McDonald, Advisory Committee Management Officer, National Endowment for the Humanities, Washington, DC 20506; telephone (202) 606–8322. Hearing-impaired individuals are advised that information on this matter may be obtained by contacting the Endowment's TDD terminal on (202) 606–8282.

SUPPLEMENTARY INFORMATION: The proposed meetings are for the purpose of panel review, discussion, evaluation and recommendation on applications for financial assistance under the National Foundation on the Arts and the Humanities Act of 1965, as amended, including discussion of information given in confidence to the agency by the grant applicants. Because the proposed meetings will consider information that is likely to disclose trade secrets and commercial or financial information obtained from a person and privileged or confidential and/or information of a personal nature the disclosure of which would constitute a clearly unwarranted invasion of personal privacy, pursuant to authority granted me by the Chairman's Delegation of Authority to Close Advisory Committee meetings, dated July 19, 1993, I have determined that these meetings will be closed to the public pursuant to subsections (c)(4), and (6) of section 552b of Title 5, United States Code.

1. *Date:* November 3, 2008.
Time: 9 a.m. to 5 p.m.
Room: 421.
Program: This meeting will review applications for Interpreting America's Historic Places Grants Program, submitted to the Division of Public Programs, at the August 27, 2008 deadline.

2. *Date:* November 6, 2008.
Time: 9 a.m. to 5 p.m.
Room: 421.
Program: This meeting will review applications for America's Media Makers Grants Program, submitted to the Division of Public Programs, at the August 27, 2008 deadline.

3. *Date:* November 6, 2008.
Time: 9 a.m. to 5 p.m.
Room: 415.
Program: This meeting will review applications for World Studies I in Preservation and Access Humanities Collections and Resources, submitted to the Division of Preservation and Access, at the July 31, 2008 deadline.

4. *Date:* November 18, 2008.
Time: 9 a.m. to 5 p.m.
Room: 415.
Program: This meeting will review applications for Music and Performing Arts in Preservation and Access Humanities Collections and Resources, submitted to the Division of Preservation and Access, at the July 31, 2008 deadline.

5. *Date:* November 20, 2008.
Time: 9 a.m. to 5 p.m.
Room: 415.
Program: This meeting will review applications for U.S. History and Culture III in Preservation and Access Humanities Collections and Resources, submitted to the Division of Preservation and Access, at the July 31, 2008 deadline.

Michael P. McDonald,

Advisory Committee Management Officer.

[FR Doc. E8–24352 Filed 10–14–08; 8:45 am]

BILLING CODE 7536–01–P

NUCLEAR REGULATORY COMMISSION

[Docket No. NRC–2008–0368]

Agency Information Collection Activities: Submission for the Office of Management and Budget (OMB) Review; Comment Request

AGENCY: U.S. Nuclear Regulatory Commission (NRC).

ACTION: Notice of the OMB review of information collection and solicitation of public comment.

SUMMARY: The NRC has recently submitted to OMB for review the following proposal for the collection of information under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). The NRC hereby informs potential respondents that an agency may not conduct or sponsor, and that a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The NRC published a **Federal Register** Notice with a 60-day comment period on this information collection on July 9, 2008.

1. *Type of submission, new, revision, or extension:* Extension.

2. *The title of the information collection:* Registration Certificate In-Vitro Testing with Byproduct Material under General License.

3. *Current OMB approval number:* 3150–0038.

4. *The form number if applicable:* NRC Form 483.

5. *How often the collection is required:* There is a one-time submittal of information to receive a validated copy of NRC Form 483 with an assigned registration number. In addition, any changes in the information reported on NRC Form 483 must be reported in writing to the Commission within 30 days after the effective date of such change.

6. *Who will be required or asked to report:* Any physician, veterinarian in the practice of veterinary medicine, clinical laboratory or hospital which desires a general license to receive, acquire, possess, transfer, or use specified units of byproduct material in certain *in vitro* clinical or laboratory tests.

7. *An estimate of the number of annual responses:* 85 (15 NRC Licensees and 70 Agreement State Licensees).

8. *The estimated number of annual respondents:* 85 (15 NRC Licensees and 70 Agreement State Licensees).

9. *An estimate of the total number of hours needed annually to complete the requirement or request:* 12.4 hours (Record keeping: 1.13 hours + Reporting: 2 hours NRC licensees and 9.3 hours Agreement State licensees).

10. *Abstract:* Section 31.11 of 10 CFR establishes a general license authorizing any physician, clinical laboratory, veterinarian in the practice of veterinary medicine, or hospital to possess certain small quantities of byproduct material for *in vitro* clinical or laboratory tests not involving the internal or external administration of the byproduct material or the radiation there from to human beings or animals. Possession of byproduct material under 10 CFR 31.11 is not authorized until the physician,