Corporate Credit Unions, Technical Corrections.
4. NCUA’s 2011 Operating Budget.
5. NCUA’s Overhead Transfer Rate.
6. NCUA’s Operating Fee Scale.

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

Mary Rupp, Board Secretary.

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NATIONAL TRANSPORTATION SAFETY BOARD

SES Performance Review Board

AGENCY: National Transportation Safety Board.

ACTION: Notice.

SUMMARY: Notice is hereby given of the appointment of members of the National Transportation Safety Board Performance Review Board (PRB).

FOR FURTHER INFORMATION CONTACT:

SUPPLEMENTARY INFORMATION: Section 4314(c)(1) through (5) of Title 5, United States Code requires each agency to establish, in accordance with regulations prescribed by the Office of Personnel Management, one or more SES Performance Review Boards. The board reviews and evaluates the initial appraisal of a senior executive’s performance by the supervisor and considers recommendations to the appointing authority regarding the performance of the senior executive.

The following have been designated as members of the Performance Review Board of the National Transportation Safety Board:
The Honorable Christopher A. Hart, Vice Chairman, National Transportation Safety Board; PRB Chair.
The Honorable Mark R. Rosekind, Member, National Transportation Safety Board.

Steven Goldberg, Chief Financial Officer, National Transportation Safety Board.

Dr. John Cavolowsky, Director, Airspace Systems Program Office, Aeronautics Research Mission Directorate, National Aeronautics and Space Administration.

Jerold Gidner, Special Counselor to the Assistant Secretary–Indian Affairs, Department of the Interior.

David L. Mayer, Managing Director, National Transportation Safety Board.

The Honorable Robert L. Sumwalt, III, Member, National Transportation Safety Board. (Alternate).

Florence Carr, Deputy Managing Director, Federal Maritime Commission. (Alternate).

Christopher W. Warner, General Counsel, U.S. Chemical Safety and Hazard Investigation Board. (Alternate).


Candi Bing, Federal Register Coordinator.

[FR Doc. 2010–28652 Filed 11–12–10; 8:45 am]

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NUCLEAR REGULATORY COMMISSION

[Docket No. NRC–2010–0276]

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

AGENCY: 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 August 12, 2010.

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

2. The title of the information collection: 10 CFR Part 35 “Medical Use of Byproduct Material”

3. Current OMB approval number: 3150–0010

4. The form number if applicable: N/A

5. How often the collection is required: Reports of medical events, doses to an embryo/fetus or nursing child, or leaking sources are reportable on occurrence. A certifying entity desiring to be recognized by the NRC must submit any one-time report for medical use of byproduct material or radiation therefrom to humans for medical use.

6. Who will be required or asked to report: Physicians and medical institutions holding an NRC license authorizing the administration of byproduct material or radiation therefrom to humans for medical use.


8. The estimated number of annual respondents: 8,610 (1,148 for NRC Licenses and 7,462 for Agreement States).

9. An estimate of the total number of hours needed annually to complete the requirement or request: 1,173,785 hours (156,538 for NRC Licenses and 1,017,247 for Agreement States).

10. Abstract: 10 CFR part 35, “Medical Use of Byproduct Material,” contains NRC’s requirements and provisions for the medical use of byproduct material and for issuance of specific licenses authorizing the medical use of this material. These regulations and provisions provide for the radiation safety of workers, the general public, patients, and human research subjects. The 10 CFR part 35 contains mandatory requirements that apply to NRC licensees authorized to administer byproduct material or radiation therefrom to humans for medical use.

The information in the required reports and records is used by the NRC to ensure that public health and safety is protected, and that the possession and use of byproduct material is in compliance with the license and regulatory requirements.

A copy of the final supporting statement may be viewed free of charge at the NRC Public Document Room, One White Flint North, 11555 Rockville Pike, Room O–1 F21, Rockville, Maryland 20852. OMB clearance requests are available at the NRC worldwide Web site: http://www.nrc.gov/public-involve/doc-comment/omb/index.html. The document will be available on the NRC home page site for 60 days after the signature date of this notice.

Comments and questions should be directed to the OMB reviewer listed below by December 15, 2010. Comments received after this date will be considered if it is practical to do so, but assurance of consideration cannot be given to comments received after this date.

Christine J. Kymn, Desk Officer, Office of Information and Regulatory Affairs (3150–0010), NMB–10202, Office of Management and Budget, Washington, DC 20503.