NUCLEAR REGULATORY COMMISSION

Documents Containing Reporting or Recordkeeping Requirements: Office of Management and Budget (OMB) Review

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).

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

2. The title of the information collection:

—Final rule, 10 CFR part 35, Medical Use of Byproduct Material

—NRC Form 313, Application for Material License, and Supplemental Forms

NRC Form 313A, Training and Experience, and

NRC Form 313B, Preceptor Statement

3. The form number if applicable: NRC Form 313, 313A and 313B

4. 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 request recognition.

5. 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.

6. An estimate of the number of responses: 214,402 (61,182 NRC licensees, 153,220 Agreement State licensees). In addition, 23 organizations are expected to prepare requests for recognition.

NRC Form 313: 7 (2 NRC licensees, 5 Agreement State licensees) applications for new modalities.

7. The estimated number of annual respondents: 5793 (1,655 NRC licensees and 4,138 Agreement State licensees).

8. An estimate of the total number of hours needed annually to complete the requirement or request: Part 35: 889,754 hours (254,059 hours for NRC licensees and 635,695 hours for Agreement State licensees) (an average of 154 hours per licensee). In addition, there is a one-time burden of 368 hours on certifying boards involved in their preparing requests for recognition. NRC Form 313: 673 hours (193 hours for NRC licensees and 480 hours for Agreement State licensees).

9. An indication of whether Section 3507(d), Pub. L. 104–13 applies: Applicable

10. Abstract: 10 CFR Part 35, ''Medical Use of Byproduct Material'', is being restructured into a more risk-informed, more performance-based regulation. The final rule 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 supporting statement may be viewed free of charge at the NRC Public Document Room, One White Flint North, 11555 Rockville Pike, Room O–1 F23, Rockville, MD 20852. OMB clearance packages are available at the NRC worldwide web site: http://www.nrc.gov/NRC/PUBLIC/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 by April 16, 2001:

Amy Farrell, Office of Information and Regulatory Affairs (3150–0010, and –0120), NEOB–10202, Office of Management and Budget, Washington DC 20503.

Comments can also be submitted by telephone at (202) 395–7318.

The NRC Clearance Officer is Brenda Jo. Shelton, 301–415–7233.

Dated at Rockville, Maryland, this 9th day of March 2001.

For the Nuclear Regulatory Commission.

Brenda Jo. Shelton,

NRC Clearance Officer, Office of the Chief Information Officer.

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NUCLEAR REGULATORY COMMISSION

Advisory Committee on the Medical Uses of Isotopes: Meeting Notice

AGENCY: U.S. Nuclear Regulatory Commission.

ACTION: Notice of meeting.

SUMMARY: The U.S. Nuclear Regulatory Commission will convene a meeting of the Advisory Committee on the Medical Uses of Isotopes (ACMUI) on April 18, 2001. The meeting will take place at the address provided below. The entire meeting will be open to the public. Topics of discussion will include: (1) status of issuance of the new 10 CFR part 35, Medical Use of Byproduct Material; (2) transition and implementation issues for the new 10 CFR part 35; (3) recognition of certification boards for training and experience qualifications; and (4) licensing issues for brachytherapy.

DATES: The meeting will be held on April 18, 2001, from 8:00 a.m. to 5:00 p.m.

ADDRESSES: U.S. Nuclear Regulatory Commission, Two White Flint North Building, Conference Room T2B3, 11545 Rockville Pike, Rockville, MD 20852–2738.

FOR FURTHER INFORMATION CONTACT: Angela R. Williamson, telephone (301) 415–5030, e-mail arw@nrc.gov, of the Office of Nuclear Material Safety and Safeguards, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001.

Conduct of the Meeting

Manuel D. Cerqueira, M.D., will chair the meeting. Dr. Cerqueira will conduct the meeting in a manner that will facilitate the orderly conduct of business. The following procedures apply to public participation in the meeting:

1. Persons who wish to provide a written statement should submit reproducible copy to Angela Williamson (address previously listed) by April 11, 2001. Statements must pertain to the topics on the agenda for the meeting.

2. Questions from members of the public will be permitted during the meeting, at the discretion of the Chairman.

3. The transcript and written comments will be available for inspection and copying for a fee, at the NRC Public Document Room, 11555 Rockville Pike, Rockville, MD 20852–2738, telephone (800) 397–4209, on or about May 20, 2001. Minutes of the meeting will be available on or about June 8, 2001.

4. Seating for the public will be on a first-come, first served basis.

This meeting will be held in accordance with the Atomic Energy Act of 1954, as amended (primarily Section 161a); the Federal Advisory Committee Act (5 U.S.C. App); and the Commission's regulations in Title 10, *U.S. Code of Federal Regulations,* Part 7.