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.

[FR Doc. 01–6617 Filed 3–15–01; 8:45 am]

BILLING CODE 7590–01–P

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.

Dated: March 12, 2001.

Andrew L. Bates,

Advisory Committee Management Officer.

[FR Doc. 01–6615 Filed 3–15–01; 8:45 am]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

Advisory Committee on Reactor Safeguards, Subcommittee Meeting on Planning and Procedures; Notice of Meeting

The ACRS Subcommittee on Planning and Procedures will hold a meeting on April 4, 2001, Room T–2B1, 11545 Rockville Pike, Rockville, Maryland.

The entire meeting will be open to public attendance, with the exception of a portion that may be closed pursuant to 5 U.S.C. 552b(c)(2) and (6) to discuss organizational and personnel matters that relate solely to internal personnel rules and practices of ACRS, and information the release of which would constitute a clearly unwarranted invasion of personal privacy.

The agenda for the subject meeting shall be as follows:

Wednesday, April 4, 2001—2:30 p.m. until the conclusion of business

The Subcommittee will discuss proposed ACRS activities and related matters. The purpose of this meeting is to gather information, analyze relevant issues and facts, and to formulate proposed positions and actions, as appropriate, for deliberation by the full Committee.

Oral statements may be presented by members of the public with the concurrence of the Subcommittee Chairman; written statements will be accepted and made available to the Committee. Electronic recordings will be permitted only during those portions of the meeting that are open to the public, and questions may be asked only by members of the Subcommittee, its consultants, and staff. Persons desiring to make oral statements should notify the cognizant ACRS staff person named below five days prior to the meeting, if possible, so that appropriate arrangements can be made.

Further information regarding topics to be discussed, the scheduling of sessions open to the public, whether the meeting has been canceled or rescheduled, the Chairman's ruling on requests for the opportunity to present oral statements, and the time allotted therefor can be obtained by contacting the cognizant ACRS staff person, Dr. John T. Larkins (telephone: 301/415–7360) between 7:30 a.m. and 4:15 p.m. (EST). Persons planning to attend this meeting are urged to contact the above named individual one or two working days prior to the meeting to be advised of any changes in schedule, etc., that may have occurred.

Dated: March 9, 2001.

James E. Lyons,

Associate Director for Technical Support, ACRS/ACNW.

[FR Doc. 01–6614 Filed 3–15–01; 8:45 am]

BILLING CODE 7590–01–P

NUCLEAR REGULATORY COMMISSION

[Docket No. 50–247]

License No. DPR–26; Consolidated Edison Company of New York, Inc.; Receipt of Petition for Director's Decision Under 10 CFR 2.206

Notice is hereby given that by Petition dated December 4, 2000, Deborah Katz, Marilyn Elie, Tim Judson, Kyle Rabin, Mark Jacobs, Paul Gunter, and Jim Riccio (petitioners) have requested that the Nuclear Regulatory Commission (NRC) take the following six actions with regard to Indian Point Nuclear Generating Unit No. 2 (IP2): (1) Suspend the license for the IP2 reactor because of the licensee's ''persistent and pervasive, negligent management of the reactor,'' (2) investigate whether the potential misrepresentation of material fact by the utility regarding ''significantly insufficient'' engineering calculations was due to a lack of rigor and thoroughness or was deliberate, (3) revoke the IP2 operating license if it is found that the licensee deliberately provided insufficient and false information, (4) if the license is not revoked, maintain IP2 on the ''list of agency's focus reactors'' until management demonstrates it can fulfill its regulatory requirements and commitments, (5) not approve the transfer of the IP2 license until management can demonstrate that the Updated Final Safety Analysis Report (UFSAR), the condition report backlog, and the maintenance requirements are up-to-date and workers have been retrained, and (6) not allow the IP2 reactor to restart until the fundamental breakdown in management is analyzed and corrected.

As a basis for this request, the petitioners state that the NRC inspections and other plant performance measurement processes have uncovered serious weaknesses and inaccuracies in the UFSAR, the Technical Specifications, the design and licensing bases, communications, maintenance, procedures, and worker training which, in the aggregate, point to a systemic mismanagement problem. The petitioners further state that without solid evidence that the licensee has addressed the root causes of systemic mismanagement, brought the reactor within compliance with its licensing and design bases, and established that the material condition of safety-significant reactor components is within safe limits, the licensee is no more prepared to operate IP2 than it was before the two recent operating events.

The Petition has been accepted for review pursuant to 10 CFR 2.206 of the Commission's regulations, and has been referred to the Director of the Office of Nuclear Reactor Regulation (NRR). In accordance with Section 2.206, appropriate action will be taken on this Petition within a reasonable time. The NRR Petition Review Board (PRB) met on December 20, 2000, to consider Requested Action 6, that the NRC prevent the IP2 reactor from restarting. The PRB recommended that the request be denied, and the Director denied it. The Director denied Requested Action 6 because the Petitioners' bases for prohibiting IP2's restart had been previously evaluated individually and in aggregate by the NRC for regulatory and safety significance. The Director found that the issues did not warrant prohibiting the restart of IP2. The petitioners Deborah Katz, Tim Judson, Kyle Rabin, Mark Jacobs, Paul Gunter, and Jim Riccio met with the NRR PRB on January 24, 2001, to discuss the Petition. The results of that discussion were considered in the board's determination regarding the schedule for the review of the Petition. The Petition and the NRC's acknowledgment letter are available in ADAMS for inspection at the Commission's Public Document Room, located at One White Flint North, 11555 Rockville Pike (first floor), Rockville, Maryland, and from the ADAMS Public Library component on the NRC's Web site, *http://www.nrc.gov* (the Public Electronic Reading Room) at accession nos. ML010580302 and ML010510218, respectively. Information regarding this Petition can also be found on the Indian Point Unit 2 Event page on the NRC's Web site, *http://www.nrc.gov/NRC/REACTOR/IP/index.html*

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

For the Nuclear Regulatory Commission.

Samuel J. Collins,

Director, Office of Nuclear Reactor Regulation.

[FR Doc. 01–6619 Filed 3–15–01; 8:45 am]

BILLING CODE 7590–01–P