impacts on organizations that are members of ERCs. Results will be used for continuous program performance improvement and external reporting, e.g., for the Government Performance and Results Act.

Estimate of Burden: Public reporting burden for this collection of information is estimated to average 30 minutes per response.

Respondents: Individuals.

Estimated Number of Responses per Form: 400.

Estimated Total Annual Burden on Respondents: 200 hours.

Frequency of Responses: One time.

Dated: November 13, 2001.

Suzanne H. Plimpton,

Reports Clearance Officer.

[FR Doc. 01–28766 Filed 11–16–01; 8:45 am]

BILLING CODE 7555–01–M

NUCLEAR REGULATORY COMMISSION

[Docket Nos. 50–254 and 50–265]

Exelon Generation Company, LLC; Notice of Consideration of Issuance of Amendment to Facility Operating License and Opportunity for a Hearing

The U.S. Nuclear Regulatory Commission (the Commission) is considering issuance of an amendment to Facility Operating License Nos. DPR–29 and DPR–30, issued to Exelon Generation Company, LLC (Exelon, the licensee), for operation of the Quad Cities Nuclear Power Station, Units 1 and 2, located in Rock County, Illinois.

The proposed amendment would allow an increase in the licensed power level from 2511 megawatts thermal (MWt) to 2957 MWt. This change represents an increase of approximately 17.8 percent above the current licensed thermal power at Quad Cities Nuclear Power Station, Units 1 and 2, and is considered an extended power uprate. The proposed amendment would also change the operating licenses and the technical specifications appended to the operating licenses to provide for implementing uprated power operation.

The original amendment request, dated December 27, 2000, was submitted by Commonwealth Edison Company (ComEd). ComEd was subsequently merged into Exelon Generation Company, LLC. By letter dated February 7, 2001, Exelon informed the NRC that it assumed responsibility for all pending NRC actions that were requested by ComEd. The original application was supplemented by letters dated February 12, April 6 and 13, May 3, 18, and 29, June 5, 7, and 15, July 6 and 23, August 7, 8, 9, 13 (two letters), 14 (two letters), 29, and 31 (two letters), September 5 (two letters), 14, 19, 25, 26, and 27 (two letters), and November 2, 2001 (two letters).

Before issuance of the proposed license amendment, the Commission will have made findings required by the Atomic Energy Act of 1954, as amended (the Act), and the Commission's regulations.

By December 19, 2001, the licensee may file a request for a hearing with respect to issuance of the amendment to the subject facility operating license, and any person whose interest may be affected by this proceeding and who wishes to participate as a party in the proceeding must file a written request for a hearing and a petition for leave to intervene. Requests for a hearing and petitions for leave to intervene shall be filed in accordance with the Commission's ''Rules of Practice for Domestic Licensing Proceedings'' in 10 CFR part 2. Interested persons should consult a current copy of 10 CFR 2.714, which is available at the Commission's Public Document Room, located at One White Flint North, 11555 Rockville Pike (first floor), Rockville, Maryland or electronically on the Internet at the NRC Web site *http://www.nrc.gov/NRC/CFR/index.html.* If there are problems in accessing the document, contact the Public Document Room Reference staff at 1–800–397–4209, 301–415–4737, or by e-mail to *pdr@nrc.gov.* If a request for a hearing or petition for leave to intervene is filed by the above date, the Commission or an Atomic Safety and Licensing Board, designated by the Commission or by the Chairman of the Atomic Safety and Licensing Board Panel, will rule on the request and/or petition; and the Secretary or the designated Atomic Safety and Licensing Board will issue a notice of hearing or an appropriate order.

As required by 10 CFR 2.714, a petition for leave to intervene shall set forth with particularity the interest of the petitioner in the proceeding, and how that interest may be affected by the results of the proceeding. The petition must specifically explain the reasons why intervention should be permitted with particular reference to the following factors: (1) The nature of the petitioner's right under the Act to be made a party to the proceeding; (2) the nature and extent of the petitioner's property, financial, or other interest in the proceedings; and (3) the possible effect of any order that may be entered in proceeding on the petitioner's interest. The petition must also identify the specific aspect(s) of the subject matter of the proceeding as to which petitioner wishes to intervene. Any person who has filed a petition for leave to intervene or who has been admitted as a party may amend the petition without requesting leave of the Board up to 15 days prior to the first prehearing conference scheduled in the proceeding, but such an amended petition must satisfy the specifically requirements described above.

Not later than 15 days prior to the first prehearing conference scheduled in the proceeding, a petitioner shall file a supplement to the petition to intervene that must include a list of the contentions that the petitioner seeks to have litigated in the hearing. Each contention must consist of a specific statement of the issue of law or fact to be raised or controverted. In addition, the petitioner shall provide a brief explanation of the bases of each contention and a concise statement of the alleged facts or expert opinion that support the contention and on which the petitioner intends to rely in providing the contention at the hearing. The petitioner must also provide references to those specific sources and documents of which the petitioner is aware and on which the petitioner intends to rely to establish those facts or expert opinion. The petitioner must provide sufficient information to show that a genuine dispute exists with the applicant on a material issue of law or fact. Contentions shall be limited to matters within the scope of the amendment under consideration. The contention must be one that, if proven, would entitle the petitioner to relief. A petitioner who fails to file such a supplement that satisfies these requirements with respect to at least one contention will not be permitted to participate as a party.

Those permitted to intervene become parties to the proceeding, subject to any limitations in the order granting leave to intervene, and have the opportunity to participate fully in the conduct of the hearing, including the opportunity to present evidence and cross-examine witnesses.

A request for a hearing and petition for leave to intervene must be filed with the Secretary of the Commission, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001, Attention: Rulemakings and Adjudications Staff, or may be delivered to the Commission's Public Document Room, located at One White Flint North, 11555 Rockville Pike (first floor), Rockville, Maryland, by the above date. A copy of the request for a hearing and the petition should also be sent to the Office of the General Counsel, U.S. Nuclear Regulatory

Commission, Washington, DC 20555–0001, and to Mr. Edward J. Cullen, Jr., Vice President and General Counsel, Exelon Generation Company, LLC, 300 Exelon Way, Kennett Square, PA 19348, attorney for the licensee.

Nontimely filings of petitions for leave to intervene, amended petitions, supplemental petitions and/or requests for a hearing will not be entertained absent a determination by the Commission, the presiding officer, or the Atomic Safety and Licensing Board that the petition and/or request should be granted based upon a balancing of the factors specified in 10 CFR 2.714(a)(1)(i)–(v) and 2.714(d).

If a request for a hearing is received, the Commission's staff may issue the amendment after it completes its technical review and prior to the completion of any required hearing if it publishes a further notice for public comment of its proposed finding of no significant hazards consideration in accordance with 10 CFR 50.91 and 50.92.

For further details with respect to this action, see the application for amendment dated December 27, 2000, as supplemented by letters dated February 12, April 6 and 13, May 3, 18, and 29, June 5, 7, and 15, July 6 and 23, August 7, 8, 9, 13 (two letters), 14 (two letters), 29, and 31 (two letters), September 5 (two letters), 14, 19, 25, 26, and 27 (two letters), and November 2, 2001 (two letters), which are available for public inspection at the Commission's Public Document Room, located at One White Flint North, 11555 Rockville Pike (first floor), Rockville, Maryland, Publicly available records will be accessible electronically from the Agencywide Documents Access and Management Systems (ADAMS) Public Electronic Reading Room on the Internet at the NRC Web site, *http://www.nrc.gov/NRC/ADAMS/index.html.* Persons who do not have access to ADAMS or who encounter problems in accessing the documents located in ADAMS, should contact the NRC Public Document Room Reference staff by telephone at 1–800–397–4209, 301–415–4737 or by e-mail to pdr@nrc.gov.

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

For the Nuclear Regulatory Commission.

Lawrence W. Rossbach,

Project Manager, Section 2 Project Directorate III, Division of Licensing Project Management, Office of Nuclear Reactor Regulation.

[FR Doc. 01–28645 Filed 11–16–01; 8:45 am]

BILLING CODE 7590–01–M

NUCLEAR REGULATORY COMMISSION

Advisory Committee on the Medical Uses of Isotopes: Call for Nominations

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

ACTION: Call for nominations.

SUMMARY: The NRC is advertising for nominations for the position Interventional Cardiology Physician on the Advisory Committee on the Medical Uses of Isotopes (ACMUI).

DATES: Nominations are due on or before January 18, 2002.

ADDRESSES: Submit four copies of the nominee's resume to the Office of Human Resources, Attn: Ms. Joyce Riner, Mail Stop T2D32, U.S. Nuclear Regulatory Commission, Washington, DC 20555.

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

SUPPLEMENTARY INFORMATION: The ACMUI advises NRC on policy and technical issues related to the regulation of the medical use of byproduct material. Responsibilities include providing comments on changes to NRC rules, regulations, and guidance documents; evaluating certain non-routine uses of byproduct material; providing technical assistance in licensing, inspection, and enforcement cases; and bringing key issues to the attention of NRC for appropriate action.

ACMUI members possess the medical and technical skills needed to address evolving issues. The current membership is comprised of the following professionals: (a) Nuclear medicine physician; (b) nuclear cardiologist; (c) medical physicist in nuclear medicine; unsealed byproduct material; (d) therapy physicist; (e) radiation safety officer; (f) nuclear pharmacist; (g) two radiation oncologists; (h) patients' rights advocate; (i) Food and Drug Administration representative; (j) State representative; and (k) health care administrator.

NRC is inviting nominations for an interventional cardiologist physician appointment to the ACMUI. This is a new position. Nominees should be interventional cardiologist physicians with experience in intravascular brachytherapy use of radiation sources. Committee members serve a 3-year term, with possible reappointment to an additional 3-year term.

Nominees must be U.S. citizens and be able to devote approximately 80 hours per year to committee business. Members who are not Federal employees are compensated for their service. In addition, members are reimbursed travel expenses (including per-diem, in lieu of subsistence); and are also reimbursed secretarial and correspondence expenses. Members who are full-time Federal employees are reimbursed travel expenses only. Nominees will undergo a security background check and will be required to complete financial disclosure statements to avoid conflict-of-interest issues.

Dated at Rockville, Maryland, this 13th day of November, 2001.

For the Nuclear Regulatory Commission.

Andrew L. Bates,

Advisory Committee Management Officer.

[FR Doc. 01–28817 Filed 11–16–01; 8:45 am]

BILLING CODE 7590–01–P

OFFICE OF PERSONNEL MANAGEMENT

[RI 20–64 and RI 20–64A]

Submission for OMB Review Comment Request for Review of an Information Collection

AGENCY: Office of Personnel Management.

ACTION: Notice.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995 (Public Law 104–13, May 22, 1995), this notice announces that the Office of Personnel Management (OPM) has submitted to the Office of Management and Budget a request for review of an information collection. RI 20–64, Former Spouse Survivor Annuity Election, is used by the Civil Service Retirement System to provide information about the amount of annuity payable after a survivor reduction and to obtain a survivor benefits election from annuitants who are eligible to elect to provide survivor benefits for a former spouse. RI 20–64A, Information on Electing a Survivor Annuity for Your Former Spouse, is a pamphlet that provides important information to retirees under the Civil Service Retirement System who want to provide a survivor annuity for a former spouse.

Approximately 30 RI 20–64 forms are completed annually. The form takes approximately 45 minutes to complete. The annual burden is 23 hours.

For copies of this proposal, contact Mary Beth Smith-Toomey on (202) 606–