centralized review under Executive Order 12866.

This action does not preempt or modify any provision of state law; nor does it impose enforcement responsibilities on any state; nor does it diminish the power of any state to enforce its own laws. Accordingly, this action does not have federalism implications warranting the application of Executive Order 13132.

The Deputy Administrator hereby certifies that this action will have no significant impact upon small entities whose interests must be considered under the Regulatory Flexibility Act, 5 U.S.C. 601 *et seq.* The establishment of aggregate production quotas for schedules I and II controlled substances is mandated by law and by international treaty obligations. The quotas are necessary to provide for the estimated medical, scientific, research and industrial needs of the United States, for export requirements and the establishment and maintenance of reserve stocks. While aggregate production quotas are of primary importance to large manufacturers, their impact upon small entities is neither negative nor beneficial. Accordingly, the Deputy Administrator has determined that this action does not require a regulatory flexibility analysis.

This action meets the applicable standards set forth in Sections 3(a) and 3(b)(2) of Executive Order 12988 Civil Justice Reform.

This action will not result in the expenditure by state, local, and tribal governments, in the aggregate, or by the private sector, of $126,400,000 or more in any one year, and will not significantly or uniquely affect small governments. Therefore, no actions were deemed necessary under the provisions of the Unfunded Mandates Reform Act of 1995.

This action is not a major rule as defined by Section 804 of the Small Business Regulatory Enforcement Fairness Act of 1996. This action will not result in an annual effect on the economy of $100,000,000 or more; a major increase in costs or prices; or significant adverse effects on competition, employment, investment, productivity, innovation, or on the ability of United States-based companies to compete with foreign-based companies in domestic and export markets.

Dated: September 2, 2010.

Michele M. Leonhart,

Deputy Administrator.

[FR Doc. 2010–22785 Filed 9–13–10; 8:45 am]

BILLING CODE 4410–09–P

OFFICE OF NATIONAL DRUG CONTROL POLICY

Appointment of Members of Senior Executive Services Performance Review Board.

AGENCY: Office of National Drug Control Policy [ONDCP].

ACTION: Notice of Appointments.

Heading: Appointment of Members of Senior Executive Services Performance Review Board.

SUMMARY: The following persons have been appointed to the ONDCP Senior Executive Service Performance Review Board: Dr. Terry Zobeck, Ms. Martha Gagne, Ms. Christine Leonard, and Mr. Patrick Ward.

FOR FURTHER INFORMATION CONTACT: Please direct any questions to Linda V. Priebe, Deputy General Counsel (202) 395–6622, Office of National Drug Control Policy, Executive Office of the President, Washington, DC 20503.

Linda V. Priebe,

Deputy General Counsel.

[FR Doc. 2010–22794 Filed 9–13–10; 8:45 am]

BILLING CODE 3180–02–P

NUCLEAR REGULATORY COMMISSION

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

AGENCY: U.S. Nuclear Regulatory Commission.

ACTION: Call for Nominations.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) is advertising for nominations for the patients' rights advocate position on the Advisory Committee on the Medical Uses of Isotopes (ACMUI). Nominees should have professional or personal experience with or knowledge about patient advocacy. Also, involvement or leadership with patient advocacy organizations is preferred.

DATES: Nominations are due on or before November 15, 2010.

NOMINATION PROCESS: Submit an electronic copy of resume or curriculum vitae, along with a cover letter, to Ms. Ashley Cockerham, *ashley.cockerham@nrc.gov.* The cover letter should describe the nominee's current involvement with patients' rights advocacy and express the nominee's interest in the position. Please ensure that resume or curriculum vitae includes the following information, if applicable: education; certification; professional association membership and committee membership activities; and number of years, recentness, and type of setting for patient advocacy.

FOR FURTHER INFORMATION CONTACT: Ms. Ashley Cockerham, U.S. Nuclear Regulatory Commission, Office of Federal and State Materials and Environmental Management Programs; (240) 888–7129; *ashley.cockerham@nrc.gov.*

SUPPLEMENTARY INFORMATION: The patients' rights advocate provides advice to NRC staff on patients' issues associated with the regulation of medical applications of byproduct material. This advice includes ensuring patients' rights are represented during the development and implementation of NRC medical-use policy. This individual is appointed based on his or her professional and personal experience with and/or knowledge about patient advocacy, involvement and/or leadership with patient advocacy organizations, and other information obtained in letters or during the selection process. Nominees should have the demonstrated ability to establish effective work relationships with peers and implement successful approaches to problem solving and conflict resolution. ACMUI members currently serve a four-year term and may be considered for reappointment to an additional term. The current membership is comprised of the following professionals: (a) Nuclear medicine physician; (b) nuclear cardiologist; (c) nuclear medicine physicist; (d) therapy medical physicist; (e) radiation safety officer; (f) nuclear pharmacist; (g) two radiation oncologists; (h) patients' rights advocate; (i) Food and Drug Administration representative; (j) Agreement State representative; (k) health care administrator; and (l) diagnostic radiologist. For additional information about membership on the ACMUI, visit the ACMUI Membership Web page, *http://www.nrc.gov/about-nrc/regulatory/advisory/acmui/membership.html.*

Nominees must be U.S. citizens and be able to devote approximately 160 hours per year to Committee business. Members are expected to attend semi-annual meetings in Rockville, Maryland and to participate in teleconferences, as needed. Members who are not Federal employees are compensated for their service. In addition, these members are reimbursed for travel and correspondence expenses. Full-time Federal employees are reimbursed for travel expenses only.

Security Background Check: The selected nominee will undergo a