a check in that amount to the Consent Decree Library at the stated address.

Maureen Katz,

Assistant Chief, Environmental Enforcement Section, Environment and Natural Resources Division.

[FR Doc. 2010–28599 Filed 11–12–10; 8:45 am] **BILLING CODE 4410–15–P**

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to Title 21 Code of Federal Regulations 1301.34(a), this is notice that on May 14, 2010, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the basic classes of controlled substances listed in schedule II:

Drug	Schedule
Methamphetamine (1105)	II II

The company plans to import the listed controlled substances to manufacture bulk controlled substances for sale to its customers.

No comments, objections, or requests for any hearings will be accepted on any application for registration or reregistration to import crude opium, poppy straw, concentrate of poppy straw, and coca leaves. As explained in the Correction to Notice of Application pertaining to Rhodes Technologies, 72 FR 3417 (2007), comments and requests for hearings on applications to import narcotic raw material are not appropriate.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule I or II, which fall under the authority of section 1002(a)(2)(B) of the Act [(21 U.S.C. 952(a)(2)(B)] may, in the circumstances set forth in 21 U.S.C. 958(i), file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement

Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than December 15, 2010.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the Federal Register on September 23, 1975, (40 FR 43745–46), all applicants for registration to import a basic class of any controlled substance in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: November 1, 2010.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Marine Well Containment Venture

Notice is hereby given that, on September 29, 2010, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 et seq. ("the Act"), Marine Well Containment Venture ("MWCV") has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, new entities are now participating in the MWCV. Pursuant to Section 6(b) of the Act, the identities of the new entities participating in the venture are: Chevron Gulf of Mexico Response Co. LLC, Houston, TX; ConocoPhillips Marine Containment Holdings Co. LLC, Houston, TX; ExxonMobil Offshore Well Containment LLC, Houston, TX; and Shell Offshore Response Co. LLC, Houston, TX.

No other changes have been made in either the membership or planned

activity of the venture. The composition of members in this venture may change, and MWCV intends to file additional written notifications disclosing all changes in membership.

On August 18, 2010, MWCV filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on October 12, 2010 (75 FR 62570).

Patricia A. Brink,

Deputy Director of Operations Antitrust Division.

[FR Doc. 2010–28558 Filed 11–12–10; 8:45 am]

BILLING CODE 4410-11-M

NATIONAL CREDIT UNION ADMINISTRATION

Sunshine Act Notice

TIME AND DATE: 10 a.m., Wednesday, November 17, 2010.

PLACE: Board Room, 7th Floor, Room 7047, 1775 Duke Street, Alexandria, VA 22314–3428.

STATUS: Closed.

Matters To Be Considered

- 1. Pilot Programs (3). Closed pursuant to some or all of the following: Exemptions (4) and (8).
- 2. Insurance Appeals (3). Closed pursuant to some or all of the following: Exemptions (4) and (6).
- 3. Personnel (2). Closed pursuant to some or all of the following: Exemption (2).
- 4. Consideration of Supervisory Activities (4). Closed pursuant to some or all of the following: Exemptions (8), (9)(A)(ii) and 9(B).

FOR FURTHER INFORMATION CONTACT:

Mary Rupp, Secretary of the Board, Telephone: 703–518–6304.

Mary Rupp,

 $Board\ Secretary.$

[FR Doc. 2010–28795 Filed 11–10–10; 4:15 pm] **BILLING CODE P**

NATIONAL CREDIT UNION ADMINISTRATION

Sunshine Act; Notice of Agency Meeting

TIME AND DATE: 9 a.m., Thursday, November 18, 2010.

PLACE: Board Room, 7th Floor, Room 7047, 1775 Duke Street, Alexandria, VA 22314–3428.

STATUS: Open.

MATTERS TO BE CONSIDERED:

1. Interim Final Rule—Part 704 of NCUA's Rules and Regulations,

Corporate Credit Unions, Technical Corrections.

- 2. Proposed Rule—Part 704 of NCUA's Rules and Regulations, Corporate Credit Unions.
- 3. Insurance Fund Report and Premium/Assessment Ranges.
 - 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.

[FR Doc. 2010–28808 Filed 11–10–10; 4:15 pm]

BILLING CODE P

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:

Emily Carroll, Chief, Human Resources Division, Office of Administration, National Transportation Safety Board, 490 L'Enfant Plaza, SW., Washington, DC 20594–0001, (202) 314–6233.

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).

Dated: November 8, 2010.

Candi Bing,

Federal Register Coordinator.

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

BILLING CODE P

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. Ĉurrent 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 a one-time request for recognition and revise the information on occurrence.

- 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.
- 7. An estimate of the number of annual responses: 246,581 ((NRC: 31,732 + 1,148 recordkeepers = 32,880) + (Agreement States: 206,239 + 7,462 recordkeepers = 213,701)).
- 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 requirements 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/doccomment/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), NEOB–10202, Office of Management and Budget, Washington, DC 20503.