Dated: March 8, 2006.

Brenda E. Dyer,

Department Clearance Officer, Department of Justice.

[FR Doc. E6–3512 Filed 3–10–06; 8:45 am]

BILLING CODE 4810–FY–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under 21 U.S.C. 952(a)(2)(B) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on August 10, 2005, ISP Freetown Fine Chemicals, 238 South Main Street, Assonet, Massachusetts 02702, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The company plans to import Phenylacetone to manufacture amphetamine.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may 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 written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than (30 days from publication).

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 listed in Schedule I or II are, and will continue to be required 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: March 6, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. 06–2363 Filed 3–10–06; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 11, 2005, and published in the **Federal Register** on August 19, 2005, (70 FR 48779), Abbott Laboratories, DBA Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey 07981, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedules I and II:

Drug	Schedule
Dihydromorphine (9145)	I
Hydromorphone (9150)	II

The company plans to manufacture bulk product and dosage units for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Abbott Laboratories, DBA Knoll Pharmaceutical Company to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Abbott Laboratories, DBA Knoll Pharmaceutical Company to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: March 6, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. 06–2341 Filed 3–10–06; 8:45 am]

BILLING CODE 4410–09–P

NATIONAL CREDIT UNION ADMINISTRATION

Notice of Meeting

Time and Date: 10 a.m., Thursday, March 16, 2006.

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

Status: Open.

Matters To Be Considered:

1. Requests form two (2) Federal Credit Unions to Convert to Community Charters.
2. NCUA's Annual Performance Budget 2006.
3. NCUA's Strategic Plan 2006–2011.
4. Interim Final Rule and Request for Comments: Part 745 of NCUA's Rules and Regulations, Share Insurance Coverage.

For Further Information Contact: Mary Rupp, Secretary of the Board, Telephone: 703–518–6304.

Paul M. Peterson

Acting Secretary of the Board.

[FR Doc. 06–2456 Filed 3-9–06; 3:42 pm]

BILLING CODE 7535–01–M

NATIONAL SCIENCE FOUNDATION

Comment Request: Biological Sciences Proposal Classification Form

AGENCY: National Science Foundation.

ACTION: Notice.

SUMMARY: The National Science Foundation (NSF) is announcing plans to establish clearance of this collection. In accordance with the requirement of section 3506(c)(2)(A) of the Paper Reduction Act of 1995, we are providing opportunity for public comment on this action. After obtaining and considering public comment, NSF will prepare the submission requesting OMB clearance of this collection for no longer than 3 years.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance