Alexandria, Virginia 22301; and must be filed no later than (60 days from publication).

Dated: January 11, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–720 Filed 1–20–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 31, 2005, Clariant LSM (Missouri), Inc., 2460 W. Bennett Street (or P.O. Box 1246, Zip: 65801), Springfield, Missouri 65807–1229, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Phenylacetone (8501), and Methadone Intermediate (9254), a basic class of controlled substances listed in Schedule II.

The company plans to manufacture in bulk, for sale to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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 March 24, 2006.

Dated: January 11, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–719 Filed 1–20–06; 8:45 am]

BILLING CODE 4410–09–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, Clinical Trial Services (US), 2661 Audubon Road, Audubon, Pennsylvania 19403, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Fentanyl (9801) and Oxycodone (9143), basic classes of controlled substance listed in Schedule II.

The company plans to import small quantities of the listed controlled substance in dosage form to conduct clinical trials.

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 February 22, 2006.

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 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: January 11, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–717 Filed 1–20–06; 8:45 am]

BILLING CODE 4410–09–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 July 27, 2005, Cody Laboratories Inc., 601 Yellowstone Avenue, Cody, Wyoming 82414–9321, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in Schedule II:

Drug	Schedule
Raw Opium (9600)	II
Poppy Straw (9650)	II
Concentrate of Poppy Straw (9670).	II

The company plans to import narcotic raw materials for manufacturing and further distribution to its customers.

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 February 22, 2006.

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 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: January 11, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–718 Filed 1–20–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 12, 2005, National Center for Natural Products Research—NIDA MProject, University of Mississippi, 135 Coy Waller Lab Complex, University, Mississippi 38577, 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:

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I

The company plans to cultivate marihuana for the National Institute of Drug Abuse for research approved by the Department of Health and Human Services.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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 March 24, 2006.

Dated: January 11, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–716 Filed 1–20–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 18, 2005, and published in the **Federal Register** on August 19, 2005, (70 FR 48780), Sigma Aldrich Research Biochemicals, Inc., 1–3 Strathmore Road, Natick, Massachusetts 01760, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of N-Benzylpiperazine (7493), a basic class of controlled substance listed in Schedules I.

The company plans to manufacture the listed controlled substances in bulk 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 Sigma Aldrich Research, Biochemicals, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Sigma Aldrich Research, Biochemicals, Inc. 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 class of controlled substance listed.

Dated: January 11, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–715 Filed 1–20–06; 8:45 am]

BILLING CODE 4410–09–P

NATIONAL SCIENCE FOUNDATION

National Science Board; Workshop on Hurricane Science and Engineering; Toward A National Agenda for Hurricane Science and Engineering: Perspectives From Federal Agencies

Date and Time: January 24, 2005, 8 a.m.–5:15 p.m. (ET).

Place: National Science Foundation, 4201 Wilson Boulevard, Room 1235, Arlington, VA 22230.

Public Meeting Attendance: All visitors must report to the NSF's visitor's desk at the 9th and N. Stuart Streets entrance to receive a visitor's badge.

Contact Information: Please refer to the National Science Board Web site (*http://www.nsf.gov/nsb*) for updated Agenda. NSB Office: (703) 292–7000.

Status: This Workshop will be open to the public.

Provisional Workshop Agenda

8–8:05 a.m. Welcoming Remarks.
8:05–8:20 a.m Motivation, Purpose and Goals.
8:20–8:30 a.m. Process and Logistics for NSB Workshops.
8:30–9:30 a.m. Panel Session I: Physical, Biological and Ecological Sciences.
9:30–10 a.m. Roundtable Discussion.
10–10:15 a.m. Break.
10:15–11:15 a.m. Panel Session II: Social, Behavioral, and Economic Sciences.
11:15–11:45 a.m. Roundtable Discussion.
1–2 p.m. Panel Session III: Engineering and Infrastructure.
2–2:30 p.m. Roundtable Discussion.
2:30–4 p.m. Break-Out Groups (3 concurrent sessions).
4–5 p.m. Break-Out Group Reports and Discussion.
5–5:15 p.m. Summary and Next Steps.
5:15 p.m. Adjourn.

Michael P. Crosby,

Executive Officer and NSB Office Director.

[FR Doc. E6–711 Filed 1–20–06; 8:45 am]

BILLING CODE 7555–01–P