shall be the Commission investigative attorneys, party to this investigation; and

(3) For the investigation so instituted, the Honorable Paul J. Luckern is designated as the presiding administrative law judge.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with § 210.13 of the Commission's rules of practice and procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(d) and 210.13(a), such responses will be considered by the Commission if received no later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and to authorize the administrative law judge and the Commission, without further notice to that respondent, to find the facts to be as alleged in the complaint and this notice and to enter both an initial determination and a final determination containing such findings, and may result in the issuance of a limited exclusion order or a cease and desist order or both directed against that respondent.

Issued: January 23, 2002.

By order of the Commission.

Marilyn R. Abbott,

Acting Secretary.

[FR Doc. 02–2140 Filed 1–28–02; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated August 30, 2001, and published in the **Federal Register** on September 10, 2001, (66 FR 47039), Applied Science Labs, Inc., A Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Heroin (9200)	I
Cocaine (9041)	II
Codeine (9050)	II
Meperidine (9230)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to import these controlled substances for the manufacture of references standards.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Applied Science Labs, Inc. to import the listed controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Applied Science Labs, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, section 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: January 16, 2002.

Laura M. Nagel,

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

[FR Doc. 02–2080 Filed 1–28–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By notice dated July 13, 2001, and published in the **Federal Register** on July 23, 2001, (66 FR 38321), Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of 4-Bromo-2, 5-dimethoxphenethylamine (7392), a basic class of controlled substance listed in Schedule I.

The firms plans to manufacture small quantities of the listed controlled substance for reference standards.

No comments or objections were received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Applied Science Labs to manufacture the listed controlled substance is consistent with the public interest at this time. DEA has investigated Applied Science Labs on a regular basis to ensure that the company's continued registration is consistent with the public interest. This investigation has included inspection and testing of the company's physical security systems, audits of the company's records, 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: January 16, 2002.

Laura M. Nagel,

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

[FR Doc. 02–2081 Filed 1–28–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By notice dated June 19, 2001, and published in the **Federal Register** on July 3, 2001 (66 FR 35269), the National Center for Natural Products Research-NIDA MProject, University of Mississippi, 135 Coy Walker Complex, University, Mississippi 38677, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the controlled substance listed below:

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

The firm will cultivate marihuana for the National Institute of Drug Abuse for research approved by the Department of Health and Human Services.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of National Center for Natural Products Research-NIDA MProject to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated National Center for Natural Products Research-NIDA MProject to ensure that the company's registration is consistent with the public interest. This investigation 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: January 16, 2002.

Laura M. Nagel,

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

[FR Doc. 02–2082 Filed 1–28–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated August 2, 2001, and published in the **Federal Register** on August 10, 2001 (66 FR 42240), Sigma Chemical Company, Subsidiary of Sigma-Aldrich Company, which has changed its name to Sigma-Aldrich Company, 3500 Dekalb Street, St. Louis, Missouri 63118, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Cathinone (1235)	I
Methcathinone (1237)	I
Aminorex (1585)	I
Gamma hydroxbutyric acid (2010)	I
Methaqualone (2565)	I
Ibogaine (7260)	I
Lysergic acid diethylamide (7315)	I
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
4-Bromo-2, 5-dimethoxyamphetamine (7391).	I
4-Bromo-2, 5-dimethoxyphenethylamine (7392).	I
2, 5-Dimethoxyamphetamine (7396).	I
3, 4-Methylenedioxyamphetamine (7400).	I
N-Hydroxy-3, 4-methylenedioxyamphetamine (7402).	I
3, 4-Methylenedioxy-N-ethylamphetamine (7404).	I
3, 4-Methylenedioxymethamphetamine (7405).	I
4-Methoxyamphetamine (7411)	I
Bufotenine (7433)	I
Psilocyn (7438)	I
Heroin (9200)	I
Normorphine (9313)	I
Etonitazene (9624)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Diprenorphine (9058)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylecgonine (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Opium powdered (9639)	II
Oxymorphone (9653)	II
Fentanyl (9801)	II

The firm plans to repackage and offer as pure standards controlled substances in small milligram quantities for drug testing and analysis.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Sigma-Aldrich Company is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Sigma-Aldrich Company on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: January 16, 2002.

Laura M. Nagel,

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

[FR Doc. 02–2079 Filed 1–28–02; 8:45 am]

BILLING CODE 4410–09–M

MISSISSIPPI RIVER COMMISSION

Sunshine Act Meetings

Agency Holding the Meetings: Mississippi River Commission.

Time and Date: 8:30 a.m., March 4, 2002.

Place: On board MISSISSIPPI V at City Front, Cairo, IL.

Status: Open to the public.

Matters to be Considered: (1) State of the Valley Report by President of the Commission on general conditions of the Mississippi River and Tributaries project and regional and national issues affecting the Corps of Engineers programs and projects; (2) District Commander's report on the Mississippi River and Tributaries project within Memphis District area; and (3) Presentations by public participants on Corps of Engineers issues.

Time and Date: 9 a.m., March 5, 2002.

Place: On board MISSISSIPPI V at Mud Island River Park Landing, Memphis, TN.

Status: Open to the public.

Matters to be Considered: (1) State of the Valley Report by President of the Commission on general conditions of the Mississippi River and Tributaries project and regional and national issues affecting the Corps of Engineers programs and projects; (2) District Commander's report on the Mississippi River and Tributaries project within Memphis District area; and (3) Presentations by public participants on Corps of Engineers issues.

Time and Date: 3:00 p.m., March 6, 2002.

Place: On board MISSISSIPPI V at City Front, Vicksburg, MS.

Status: Open to the public.

Matters to be Considered: (1) State of the Valley Report by President of the Commission on general conditions of the Mississippi River and Tributaries