complete a Record of Decision. It is expected that the Board of Directors of the Agency will adopt a Notice of Determination pursuant to CEQA in December 1998.

ADDRESSES: Copies of the FEIS/FEIR may be requested from Mr. Tad Berkebile, Sacramento County Water Agency, 827 Seventh Street, Room 301, Sacramento CA 95814; telephone: (916) 874–6851. See Supplementary Information section for locations where the FEIS/FEIR is available for public inspection and review.

FOR FURTHER INFORMATION CONTACT: Mr. Tad Berkebile at (916) 874–6851 or Mr. Cecil Lesley, Bureau of Reclamation, 7794 Folsom Dam Road, Folsom CA 95630, telephone: (916) 989–7221 or TDD (916) 989–7285.

SUPPLEMENTARY INFORMATION: Public Law 101–514, Section 206, authorizes and directs the Secretary of the Interior to enter into long-term municipal and industrial water supply contracts to meet the immediate water needs of Sacramento County. Contracts under Public Law 101–514, Section 206, are not subject to the prohibition on new Reclamation contracts of Public Law 102–575, Title XXXIV. However, any contracts under Public Law 101–514 are required to have terms and conditions that allow the Secretary to amend the contracts as necessary to meet the obligations of applicable State and Federal laws. The law specifically directs the Secretary to enter into contracts up to 22,000 acre-feet per year with Sacramento County and up to 13,000 acre-feet per year with the San Juan Water District (serving a part of northeastern Sacramento County). Water delivered annually under these contracts is at the discretion of the Secretary, who will make a determination of the amount to be made available ''based upon the quantity of water actually needed after considering reasonable efforts to (i) promote full utilization of existing water entitlement within Sacramento County, (ii) implement water conservation and metering programs within areas served by the contract, and (iii) implement programs to maximize to the extent feasible conjunctive use of surface water and groundwater'' (Public Law 101–514, Section 206 [b][(1)]. Of its annual allocation of 22,000 acre-feet per year, the Agency intends to provide up to 7,000 acre-feet per year to the City of Folsom through a subcontract.

No potentially affected Indian Trust Assets (ITA's) have been identified by Reclamation for the proposed project or alternatives.

Copies of the FEIS/FEIR are available for public inspection and review at the following locations:

- Bureau of Reclamation, Central California Area Office at 7794 Folsom Dam Road in Folsom, CA.
- Bureau of Reclamation at 2800 Cottage Way, Room E–1704 in Sacramento, CA.
- Sacramento County Water Agency at 827 Seventh Street, Room 301 in Sacramento, CA.
- Sacramento County Clerk-Recorder's Office at 600 Eighth Street in Sacramento, CA.
- San Juan Water District at 9935 Auburn-Folsom Road in Granite Bay, CA.
- Sacramento Central Library at 828 I Street in Sacramento, CA.
- Folsom Library at 300 Purcifer Street in Folsom, CA.
- City of Folsom, Public Works Department at 50 Natoma Street in Folsom, CA.

Dated: December 14, 1998.

Kirk C. Rodgers,

Deputy Regional Director.

[FR Doc. 98–34025 Filed 12–22–98; 8:45 am]

BILLING CODE 4310–94–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on October 29, 1998, B.I. Chemical, Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of amphetamine (1100), a basic class of controlled substance listed in Schedule II.

The firm plans to bulk manufacture amphetamine for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 22, 1999.

Dated: December 14, 1998.

John H. King,

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

[FR Doc. 98–33878 Filed 12–22–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Registration

By Notice dated October 1, 1998, and published in the **Federal Register** on October 9, 1998, (63 FR 544512), Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121–4340, 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
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
Phencyclidine (7471)	II
Phenylaceton (8501)	II
Cocaine (9041)	II

The firm plans to import small quantities of the listed controlled substances to make reagents for distribution to the biomedical research community.

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 Calbiochem-Novabiochem Corporation 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 the firm 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: December 14, 1998.

John H. King,

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

[FR Doc. 98–33885 Filed 12–22–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (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 Section 1002(a) 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 Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on October 26, 1998, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Methamphetamine (1105)	II
Phenylacetone (8501)	II

The firm plans to import the phenylacetone to manufacture methamphetamine and to import racemic methamphetamine for resolution into the d- and 1-stereoisomers.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

This procedure is to be conducted simulateneously with and independent of the procedures described in 21 CFR 1301.34 (b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), 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 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: December 14, 1998.

John H. King,

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

[FR Doc. 98–33879 Filed 12–22–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on October 26, 1998, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of methamphetamine (1105), a basic class of controlled substance listed in Schedule II.

The firm plans to bulk manufacture methamphetamine to produce products for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 22, 1999.

Dated: December 14, 1998.

John H. King,

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

[FR Doc. 98–33880 Filed 12–22–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated September 2, 1998, and published in the **Federal Register** on September 10, 1998, (63 FR 48522), Dupont Pharmaceuticals, The Dupont Merck Pharmaceutical Co., 1000 Stewart Avenue, Garden City, New York 11530, 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 below:

Drug	Schedule
Oxycodone (9143)	II
Hydrocodone (9193)	II
Oxymorphone (9652)	II

The firm plans to manufacture the listed controlled substances to make finished products.

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 Dupont Pharmaceuticals to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Dupont Pharmaceuticals 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 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.