direct sale offer will qualify the purchaser to make application for conveyance of those mineral interests. Purchasers must submit a non refundable $50.00 filing fee for the conveyance of the mineral estate upon request by the Bureau of Land Management.

Detailed information concerning the sale, including the reservations, sale procedures and conditions, and planning and environmental documents, is available at the Coos Bay District Office, 1300 Airport Lane, North Bend, OR 97459.

For a period of 45 days from the date of publication of this notice in the **Federal Register**, interested parties may submit comments to the District Manager, Bureau of Land Management, at the above address. Objections will be reviewed by the State Director who may sustain, vacate, or modify this realty action. In absence of any objections, this realty action will become the final determination of the Department of the Interior.

FOR FURTHER INFORMATION CONTACT: Linda Petterson, Realty Specialist, Umpqua Field Office, at 1300 Airport Lane, North Bend, Oregon 97459, (Telephone 541 756–0100).

Dated: August 12, 1999.

Neal Middlebrook,

Associate District Manager.

[FR Doc. 99–21679 Filed 8–19–99; 8:45 am]

BILLING CODE 4710–33–M

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[OR–125–08–1430–00; GP9.–0284; OR 53839]

Coos Bay District; Notice of Realty Action: Direct Sale of Public Land in Coos County, OR

AGENCY: Bureau of Land Management.

SUMMARY: The following land is suitable for direct sales under Section 203 of the Federal Land Policy and Management Act of 1976, 43 U.S.C. 1713, at no less than the appraised fair market value. The land will not be offered for sale until at least 60 days after publication of this notice:

Willamette Meridian, Oregon

T. 27 S., R. 11 W.,
Sec. 5 Lot 6 containing 1.82 acres.

The above described land is hereby segregated from appropriation under the public land laws, including the mining laws, but not from sale under the above cited statute, for 270 days or until title transfer is completed or the segregation is terminated by publication in the **Federal Register**, whichever occurs first.

This land is difficult and uneconomic to manage as part of the public lands and is not suitable for management by another Federal agency. No significant resource values will be affected by this disposal. The sale is consistent with BLM's planning for the land involved and the public interest will be served by the sale.

Purchasers must be U.S. citizens, 18 years of age or older, a state or a state instrumentally authorized to Purchasers must be U.S. citizens, 18 years of age or older, a state or a state instrumentally authorized to hold property, or a corporation authorized to own real estate in the state in which the land is located.

The land is being offered in Coos County, Oregon using the direct sale procedures authorized under 43 CFR 2713.3–3. The parcel will be offered to Leslie N. Crum, who holds a homesite lease on the subject parcel.

The terms, conditions and reservations applicable to the sale are as follows;

1. A right-of-way for ditches and canals will be reserved to the United States under 43 U.S.C. 945.

2. Patents will be issued subject to all valid existing rights and reservations of record.

3. The mineral interest being offered for conveyance have no known mineral values and may be conveyed simultaneously, in accordance with Section 209 of the Federal Land Policy and Management Act. Acceptance of the direct sale offer will qualify the purchaser to make application for conveyance of those mineral interests. Purchasers must submit a non refundable $50.00 filing fee for the conveyance of the mineral estate upon request by the Bureau of Land Management.

Detailed information concerning the sale, including the reservations, sale procedures and conditions, and planning and environmental documents, is available at the Coos Bay District Office, 1300 Airport Lane, North Bend, OR 97459.

For a period of 45 days from the date of publication of this notice in the **Federal Register**, interested parties may submit comments to the District Manager, Bureau of Land Management, at the above address. Objections will be reviewed by the State Director who may sustain, vacate, or modify this realty action. In absence of any objections, this realty action will become the final determination of the Department of the Interior.

FOR FURTHER INFORMATION CONTACT: Linda Petterson, Realty Specialist, Umpqua Field Office, at 1300 Airport Lane, North Bend, Oregon 97459, (Telephone 541 756–0100).

Dated: August 12, 1999.

Neal Middlebrook,

Associate District Manager.

[FR Doc. 99–21680 Filed 8–19–99; 8:45 am]

BILLING CODE 4310–33–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 June 4, 1999, Guilford Pharmaceuticals, Inc., 6611 Tributary Street, Baltimore, Maryland 21224, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of cocaine (9014), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture methyl-3-beta-(4-trimethylstannylphenyl)-tropane-2-carboxylate as a final intermediate for the production of dopascan injection.

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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than October 19, 1999.

Dated: August 6, 1999.

John H. King,

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

[FR Doc. 99–21586 Filed 8–19–99; 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 June 17,

1999, ISP Freetown Acquisition, Corp., 238 South Main Street, Freetown, Massachusetts 02702, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of 2,5-Dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The firm plans to manufacture bulk 2,5-Dimethoxyamphetamine for conversion into a noncontrolled substance.

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 Assistance Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than October 19, 1999.

Dated: August 5, 1999.

John H. King,

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

[FR Doc. 99–21587 Filed 8–19–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated February 5, 1999, and published in the **Federal Register** on February 26, 1999, (64 FR 9541), Medeva Pharmaceuticals CA, Inc., 3501 West Garry Avenue, Santa Ana, California 92704, 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
Methylphenidate (1724)	II
Diphenoxylate (9170)	II

The firm plans to manufacture the listed controlled substances to make finished dosage forms for distribution to its customers.

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 Medeva Pharmaceuticals CA, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Medeva Pharmaceuticals CA, 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 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: August 5, 1999.

John H. King,

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

[FR Doc. 99–21584 Filed 8–19–99; 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 April 5, 1999, Morton Grove Pharmaceuticals, Inc., 6451 W. Main Street, Morton Grove, Illinois 60053, made application to the Drug Enforcement Administration to be registered as an importer of codeine (9050), a basic class of controlled substance listed in Schedule II.

The firm plans to import the codeine to produce controlled substances in Schedule III through V.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than September 20, 1999.

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 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: August 5, 1999.

John H. King,

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

[FR Doc. 99–21588 Filed 8–19–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Registration

By Notice dated May 14, 1999, and published in the **Federal Register** on May 25, 1999 (64 FR 28214), Research Biochemicals, Limited Partnership, 1–3 Strathmore Road, Natick, Massachusetts 01760, 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
Methaqualone (2565)	I
Alpha-Ethyltryptamine (7249)	I
Ibogaine (7260)	I