

9, 1998, Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey 07981, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

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

The firm plans to produce bulk product and finished dosage units 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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than March 5, 1999.

Dated: December 23, 1998.

John H. King,

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

[FR Doc. 98-34812 Filed 12-31-98; 8:45 am]

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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 November 13, 1998, Wildlife Laboratories, Inc., 1401 Duff Drive, Suite 600, Fort Collins, Colorado 80524, 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
Etorphine Hydrochloride (9059) ..	II
Carfentanil (9743)	II

The firm plans to import the listed controlled substances to produce finished products for distribution to its customers.

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, DC 20537, Attention: DEA Federal Register Representative (CCR), 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 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 23, 1998.

John H. King,

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

[FR Doc. 98-34813 Filed 12-31-98; 8:45 am]

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DEPARTMENT OF JUSTICE

Foreign Claims Settlement Commission

Sunshine Act Meeting

[F.C.S.C. Meeting Notice No. 1-99]

The Foreign Claims Settlement Commission, pursuant to its regulations

(45 CFR Part 504) and the Government in the Sunshine Act (5 U.S.C. 552b), hereby gives notice in regard to the scheduling of meetings and oral hearings for the transaction of Commission business and other matters specified, as follows:

Date and Time: Monday, January 11, 1999, 10:00 a.m.

Subject Matter: Hearings on the Record on Objections to Proposed Decisions on claims against Albania, as follows:

Claim No.

ALB-072 Thomas Toma
ALB-092 Thanos A. Laske
ALB-173 Marigo Tellios, et al.
ALB-220 Gjergji Gjeli
ALB-315 Afroditi Botsis

Status: Open.

All meetings are held at the Foreign Claims Settlement Commission, 600 E Street, N.W., Washington, DC. Requests for information, or advance notices of intention to observe an open meeting, may be directed to: Administrative Officer, Foreign Claims Settlement Commission, 600 E Street, N.W., Room 6002, Washington, DC 20579. Telephone: (202) 616-6988.

Dated at Washington, DC, December 30, 1998.

David E. Bradley,

Chief Counsel.

[FR Doc. 98-34828 Filed 12-30-98; 1:26 pm]

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DEPARTMENT OF LABOR

Employment Standards Administration, Wage and Hour Division

Minimum Wages for Federal and Federally Assisted Construction; General Wage Determination Decisions

General wage determination decisions of the Secretary of Labor are issued in accordance with applicable law and are based on the information obtained by the Department of Labor from its study of local wage conditions and data made available from other sources. They specify the basic hourly wage rates and fringe benefits which are determined to be prevailing for the described classes of laborers and mechanics employed on construction projects of a similar character and in the localities specified therein.

The determinations in these decisions of prevailing rates and fringe benefits have been made in accordance with 29 CFR Part 1, by authority of the Secretary of Labor pursuant to the provisions of the Davis-Bacon Act of March 3, 1931, as amended (46 Stat. 1494, as amended, 40 U.S.C. 276a) and of other Federal