The Commission will transmit its determination in this investigation to the Secretary of Commerce on August 15, 2004. The views of the Commission are contained in USITC Publication 3792 (August 2005), entitled Metal Calendar Slides from Japan: Investigation No. 731–TA–1094 (Preliminary).

By order of the Commission.
Issued: August 15, 2005.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 05–16425 Filed 8–18–05; 8:45 am]

BILLING CODE 7020–02–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 March 16, 2005, Abbott Laboratories, DBA Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey 07981, 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 and II:

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

The company plans to manufacture bulk product and dosage units for distribution 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 October 18, 2005.

Dated: August 11, 2005.

William J. Walker,

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

[FR Doc. 05–16468 Filed 8–18–05; 8:45 am]

BILLING CODE 4410–09–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 March 7, 2005, Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, 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 and II:

Drug	Schedule
Gamma hydroxybutyric acid (2010).	I
Amphetamine (1100)	II
Methylphenidate (1724)	II

The company plans to manufacture bulk products for finished dosage units and distribution 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 October 18, 2005.

Dated: August 11, 2005.

William J. Walker,

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

[FR Doc. 05–16469 Filed 8–18–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 29, 2005, and published in the **Federal Register** on April 6, 2005, (70 FR 17473), Polaroid Corporation, 1265 Main Street, Building W6, Waltham, Massachusetts 02454, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of 2,5-Dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The company plans to manufacture the listed controlled substance in bulk for conversion into non-controlled substances.

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 Polaroid Corporation to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Polaroid Corporation 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: August 12, 2005.

William J. Walker,

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

[FR Doc. 05–16466 Filed 8–18–05; 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