Authority: This review is being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.62 of the Commission's rules.

Issued: August 24, 1999.

By order of the Commission.

Donna R. Koehnke,

Secretary.

[FR Doc. 99–22346 Filed 8–26–99; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigations Nos. 731–TA–846 through 850 (Preliminary)]

Certain Seamless Carbon and Alloy Steel Standard, Line, and Pressure Pipe from the Czech Republic, Japan, Mexico, Romania, and South Africa

Determinations

On the basis of the record [1] developed in the subject investigations, the United States International Trade Commission determines, pursuant to section 733(a) of the Tariff Act of 1930 (19 U.S.C. 1673b(a)), that there is a reasonable indication that an industry in the United States is materially injured by reason of imports from the Czech Republic, Japan, Romania, and South Africa of small diameter (less than or equal to 4.5 inches in outside diameter) seamless carbon and alloy (other than stainless) steel standard, line, and pressure pipe (including redraw hollows), provided for in subheadings 7304.10.10, 7304.10.50, 7304.31.30, 7304.31.60, 7304.39.00, 7304.51.50, 7304.59.60, and 7304.59.80 of the Harmonized Tariff Schedule of the United States (HTS), that are alleged to be sold in the United States at less than fair value (LTFV). The Commission also determines that there is a reasonable indication that an industry in the United States is materially injured by reason of imports from Japan and Mexico of large diameter (greater than 4.5 inches up to and including 16 inches in outside diameter) seamless carbon and alloy (other than stainless) steel standard, line, and pressure pipe, provided for in subheadings 7304.10.10, 7304.10.50, 7304.39.00, and 7304.59.80 of the HTS, that are alleged to be sold in the United States at LTFV.

Commencement of Final Phase Investigations

Pursuant to § 207.18 of the Commission's rules, the Commission also gives notice of the commencement of the final phase of its investigations. The Commission will issue a final phase notice of scheduling that will be published in the **Federal Register** as provided in § 207.21 of the Commission's rules upon notice from the Department of Commerce (Commerce) of affirmative preliminary determinations in the investigations under section 733(b) of the Act, or, if the preliminary determinations are negative, upon notice of affirmative final determinations in those investigations under section 735(a) of the Act. Parties that filed entries of appearance in the preliminary phase of these investigations need not enter a separate appearance for the final phase of the investigations. Industrial users and, if the merchandise under investigation is sold at the retail level, representative consumer organizations have the right to appear as parties in Commission antidumping and countervailing duty investigations. The Secretary will prepare a public service list containing the names and addresses of all persons, or their representatives, who are parties to the investigations.

Background

On June 30, 1999, petitions were filed with the Commission and the Department of Commerce by Koppel Steel Corp., Beaver Falls, PA; Sharon Tube Co., Sharon, PA; U.S. Steel Group, Fairfield, AL; USS/Kobe Steel Co., Lorain, OH; and Vision Metals' Gulf States Tube Div., Rosenberg, TX; alleging that an industry in the United States is materially injured or threatened with material injury by reason of LTFV imports of small diameter seamless carbon and alloy steel standard, line, and pressure pipe from the Czech Republic, Japan, Romania, and South Africa; and by reason of LTFV imports of large diameter seamless carbon and alloy steel standard, line, and pressure pipe from Japan and Mexico. [2] Accordingly, effective June 30, 1999, the Commission instituted antidumping investigations Nos. 731–TA–846 through 850 (Preliminary).

Notice of the institution of the Commission's investigations and of a public conference to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** of July 8, 1999 (64 FR 36920). The conference was held in Washington, DC, on July 21, 1999, and all persons who requested the opportunity were permitted to appear in person or by counsel.

The Commission transmitted its determinations in these investigations to the Secretary of Commerce on August 23, 1999. The views of the Commission are contained in USITC Publication 3221 (August 1999), entitled Certain Seamless Carbon and Alloy Steel Standard, Line, and Pressure Pipe from the Czech Republic, Japan, Mexico, Romania, and South Africa: Investigations Nos. 731–TA–846 through 850 (Preliminary).

By order of the Commission.

Issued: August 23, 1999.

Donna R. Koehnke,

Secretary.

[FR Doc. 99–22341 Filed 8–26–99; 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 April 20, 1999, B.I. Chemical, Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, 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
Amphetamine (1100)	II
Methadone (9250)	II
Levo-alphacetylmethadol (LAAM) (9648).	II

The firm plans to bulk manufacture the listed controlled substances for formulation into finished pharmaceuticals.

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 October 26, 1999.

[1] The record is defined in § 207.2(f) of the Commission's rules of practice and procedure (19 CFR 207.2(f)).

[2] Koppel, Sharon, and Vision are not petitioners in the investigations regarding large diameter subject products.

Dated: August 17, 1999.

John H. King,

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

[FR Doc. 99–22309 Filed 8–26–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances Notice of Registration

By Notice dated March 1, 1999, and published in the **Federal Register** on April 9, 1999, (64 FR 17416), Lipomed, Inc., One Broadway, Cambridge, Massachusetts 02142, made application 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
Methaqualone (2565)	I
Lysergic acid diethylamide (7315).	I
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
3, 4, 5–Trimethoxyamphetamine (7390).	I
4–Bromo–2, 5–dimethoxyamphetamine (7391).	I
4–Methyl–2, 5–dimethoxyamphetamine (7395).	I
2, 5–Dimethoxyamphetamine (7396).	I
2, 5–Dimethoxy–4–ethylamphetamine (7399).	I
3, 4–Methylenedioxyamphetamine (7400).	I
3, 4–Methylenedioxy–N–ethylamphetamine (7404).	I
3, 4–Methylenedioxymethamphetamine (7405).	I
Psilocybin (7437)	I
Psilocyn (7438)	I
Acetyldihydrocodeine (9051)	I
Dihydromorphine (9145)	I
Heroin (9200)	I
Tilidine (9750)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Amobarbital (2125)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylecgonine (9180)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Methadone (9250)	II
Dextroproproxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Alfentanil (9737)	II
Fentanyl (9801)	II

The firm plans to import small reference standard quantities of finished commercial product from its sister company in Switzerland for sale to its customers for drug testing and pharmaceutical research and development.

No comments or objections have been received. DEA has considered the factors of Title 21, United States Code, Section 823(a) and determined that the registration of Lipomed, 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 Lipomed, 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 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: August 17, 1999.

John H. King,

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

[FR Doc 99–22307 Filed 8–26–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated April 16, 1999, and published in the **Federal Register** on April 29, 1999, (64 FR 23114), Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, 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
Tetrahydrocannabinols (7370)	I.
Amphetamine (1100)	II.
Methylphenidate (1724)	II.
Cocaine (9041)	II.
Codeine (9050)	II.
Diprenorphine (9058)	II.
Etorphine Hydrochloride (9059)	II.
Dihydrocodeine (9120)	II.
Oxycodone (9143)	II.
Hydromorphone (9150)	II.
Diphenoxylate (9170)	II.
Hydrocodone (9193)	II.
Levorphanol (9220)	II.
Meperidine (9230)	II.
Methadone (9250)	II.
Methadone-intermediate (9254)	II.
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II.
Morphine (9300)	II.
Thebaine (9333)	II.
Opium extracts (9610)	II.
Opium fluid extract (9620)	II.
Opium tincture (9630)	II.
Opium powdered (9639)	II.
Opium granulated (9640)	II.
Levo-alphacetylmethadol (9648)	II.
Oxymorphone (9652)	II.
Noroxymorphone (9668)	II.
Alfentanil (9637)	II.
Sufentanil (9740)	II.
Fentanyl (1980)	II.

The firm plans to manufacture the controlled substances for distribution as bulk products to its customers.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Mallinckrodt Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Mallinckrodt Chemicals, 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.