INTERNATIONAL TRADE COMMISSION

[Inv. No. 337–TA–496]

In the Matter of: Certain Home Vacuum Packaging Products; Notice of Commission Decision Not To Review an Initial Determination Granting a Motion To Withdraw Two Patent Claims From the Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined not to review the presiding administrative law judge's (''ALJ's'') initial determination (''ID'') granting a motion to withdraw claims 24 and 25 of U.S. Patent No. 4,941,310, from the above-captioned investigation.

FOR FURTHER INFORMATION CONTACT: Timothy P. Monaghan, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone (202) 205–3152. Copies of the Commission order, and all other nonconfidential documents filed in connection with this investigation are or will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone (202) 205–2000. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810. General information concerning the Commission may also be obtained by accessing its Internet server (*http://www.usitc.gov*). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov*.

SUPPLEMENTARY INFORMATION: On August 18, 2003, the Commission instituted this investigation based upon a complaint filed by Tilia, Inc. and Tilia International (collectively, ''Tilia''). 68 FR 49521. At the same time, the Commission provisionally accepted a motion for temporary relief filed by Tilia. In its complaint, Tilia alleges that the accused imported products infringe claims 3, 4, 6, 24–25, and 34 of U.S. Patent No. 4,941,310 (''the '310 patent''). The notice of investigation named Applica, Inc., Applica Consumer Products, Inc.; ZeroPack Co., Ltd.; The Holmes Group, Inc.; and The Rival Company as respondents. On January 15, 2004, the Commission determined not to review an ID denying Tilia's motion for temporary relief. On February 3, 2004, Tilia moved pursuant to rules 210.21(a)(1) to withdraw claims 24 and 25 of the '310 patent from the investigation. Respondents and the Commission investigative attorney did not oppose the motion. On February 18, 2004, the ALJ issued an ID granting Tilia's motion to withdraw claims 24 and 25 of the '310 patent from the investigation. No petitions of the ID were filed.

This action is taken under the authority of section 337 of the Tariff Act of 1930, 19 U.S.C. 1337, and § 210.42 of Commission's Rules of Practice and Procedure, 19 CFR 210.42.

By order of the Commission.

Issued: March 10, 2004.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 04–5803 Filed 3–12–04; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on February 18, 2004, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Methamphetamine (1105), and Hydromorphone (9150), basic class of controlled substances listed in Schedule II. The firm had inadvertently dropped the two basic classes from its renewal application submitted on August 25, 2003, and published in the **Federal Register** on February 18, 2004 (69 FR 7656).

The firm plans to manufacture the listed controlled substances in bulk to supply 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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than May 14, 2004.

Dated: March 5, 2004.

William J. Walker,

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

[FR Doc. 04–5774 Filed 3–12–04; 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(1)), 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 registration 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 January 30, 2004, Johnson Matthey Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey 08066, 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
Phenylacetone (8501)	II
Raw Opium (9600)	II
Concentrate of Poppy Straw (9670).	II

The firm plans to import the listed controlled substances as raw materials for use in the manufacture of bulk controlled substances for distribution to its customers.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of the basic classes of controlled substances listed 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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than April 14, 2004. 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 1975 notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import 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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: March 5, 2004.

William J. Walker,

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

[FR Doc. 04–5779 Filed 3–12–04; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 20, 2004, Lin Zhi International, Inc., 687 North Pastoria Avenue, Sunnyvale, California 94085, 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
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxymethamphetamine (7405).	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Benzoylecgonine (9180)	II
Methadone (9250)	II
Dextropropoxyphene (9273)	II
Morphine (9300)	II

The firm plans to manufacture small quantities of controlled substances to make drug testing reagents and controls.

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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than May 14, 2004.

Dated: March 5, 2004.

William J. Walker,

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

[FR Doc. 04–5778 Filed 3–12–04; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 5, 2004, Mallinckrodt Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, 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
Tetrahydrocannabinols (7370)	I
Codeine-N-oxide (9053)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Heroin (9200)	I
Morphine-N-oxide (9307)	I
Nicomorphine (9312)	I
Normorphine (9313)	I
Norlevorphanol (9634)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Diprenorphine (9058)	II
Etorphine HCL (9059)	II
Dihydrocodeine (9120)	II
Hydromorphone (9150)	II
Oxycodone (9143)	II
Diphenoxylate (9170)	II
Benzoylecgonine (9180)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone Intermediate (9254)	II
Metopon (9260)	II
Dextropropoxyphene (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
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances for internal use and for sale to other companies.

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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than May 14, 2004.

Dated: March 5, 2004.

William J. Walker,

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

[FR Doc. 04–5773 Filed 3–12–04; 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(1)), 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 registration 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 January 6, 2004, Mallinckrodt Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below: