Dated: February 21, 1996.

Paul Andrews,

Acting District Manager.

[FR Doc. 96–4868 Filed 3–1–96; 8:45 am]

BILLING CODE 4310–DQ–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated September 29, 1995, and published in the Federal Register on October 11, 1995, (60 FR 52923), Ciba-Geigy Corporation, Pharmaceuticals Division Regulatory Compliance, 556 Morris Avenue, Summit, New Jersey 07901, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of methylphenidate (1724), a basic class of controlled substance listed in Schedule II.

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 Ciba-Geigy Corporation to manufacture the listed controlled substance is consistent with the public interest at this time. Therefore, pursuant to Section 303 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 and Title 21, Code of Federal Regulations, Section 1301.54(e), 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 class of controlled substance listed above is granted.

Gene R. Haislip,

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

[FR Doc. 96–4946 Filed 3–1–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 19, 1995, and published in the Federal Register on October 25, 1995, (60 FR 54707), Eli Lilly Industries, Inc., Chemical Plant, Kilometer 146 7, State Road 2, Mayaguez, Puerto Rico 00680, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of dextropropoxyphene, bulk (non-dosage forms) (9273), a basic class of controlled substance listed in Schedule II.

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 Eli Lilly Industries, Inc. to manufacture the listed controlled substance is consistent with the public interest at this time. Therefore, pursuant to Section 303 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 and Title 21, Code of Federal Regulations, Section 1301.54(e), 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 class of controlled substance listed above is granted.

Dated: February 26, 1996.

Gene R. Haislip,

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

[FR Doc. 96–4947 Filed 3–1–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on November 13, 1995, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396)	I
Difenoxin (9168)	I
Methylphenidate (1724)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Meperidine intermediate-A (9232)	II
Meperidine intermediate-B (9233)	II
Meperidine intermediate-C (9234)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Oxymorphone (9652)	II
Morphine (9300)	II
Oxymorphone (9652)	II
Sufentanil (9740)	II
Carfentanil (9743)	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances in bulk to supply final dosage form manufacturers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application.

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 May 3, 1996.

Dated: February 26, 1996.

Gene R. Haislip,

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

[FR Doc. 96–4944 Filed 3–1–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 19, 1995, MD Pharmaceutical, Inc., 3501 West Garry Avenue, Santa Ana, California 92704, made application to the Drug Enforcement Administration (DEA) for registration 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.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objects to the issuance of the above application.

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 May 3, 1996.

Dated: February 26, 1996.

Gene R. Haislip,

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

[FR Doc. 96–4945 Filed 3–1–96; 8:45 am]

BILLING CODE 4410–09–M

Importer of Controlled Substances; Notice of Registration

By Notice dated December 15, 1995, and published in the Federal Register on December 28, 1995, (60 FR 67141), North Pacific Trading Company, 1505 SE Gideon Street, Portland, Oregon 97202, made application to the Drug Enforcement Administration to be registered as an importer of Marihuana (7360), a basic class of controlled substance listed in Schedule I.

No comment 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 North Pacific Trading Company to import the listed controlled substance is consistent with the public interest at this time. 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 class of controlled substance listed above.

Dated: February 26, 1996.

Gene R. Haislip,

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

[FR Doc. 96–4948 Filed 3–1–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 29, 1995, and published in the Federal Register on July 6, 1995, (60 FR 35225), Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Pholcodine (9314)	I
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Benzoylecgonine (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	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
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

A registered manufacturer filed a comment requesting that Penick's application for registration be denied for considerations of the public interest. The commenter also questioned whether Penick has the manufacturing and processing capabilities to manufacture the listed controlled substances. DEA has conducted inspection of Penick and determined that Penick has complied with the factors in Title 21, United States Code, Section 823(a). Penick's current application was filed to renew a manufacturer registration which the firm has maintained for several years and under which the firm manufactured controlled substances in the past in conformance with the Controlled Substances Act and its implementing regulations. Therefore, pursuant to section 303 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 and Title 21, Code of Federal Regulations, Section 1301.54(e), 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: February 26, 1996.

Gene R. Haislip,

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

[FR Doc. 96–4949 Filed 3–1–96; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Mine Safety and Health Administration

Petitions for Modification

The following parties have filed petitions to modify the application of mandatory safety standards under section 101(c) of the Federal Mine Safety and Health Act of 1977.

1. Mackie J. Coal Company, Inc.

[Docket No. M–95–169–C]

Mackie J. Coal Company, Inc., Route 2, Box 530, Grundy, Virginia 24614 has filed a petition to modify the application of 30 CFR 75.1710–1 (canopies or cabs; self-propelled electric face equipment; installation requirements) to its Mine No. 4 (I.D. No. 44–06051) located in Buchanan County, Virginia. The petitioner proposes to use self-propelled electric face equipment without cabs or canopies in mining heights of 48 inches or less. The petitioner states that application of the standard would result in a diminution of safety to the equipment operator.

2. Marfork Coal Company, Inc.

[Docket No. M–95–170–C]

Marfork Coal Company, Inc., P.O. Box 457, Whitesville, West Virginia 25209 has filed a petition to modify the application of 30 CFR 75.333(d)(1) (ventilation controls) to its Outpost East Mine (I.D. No. 46–08296); its Outpost West Mine (I.D. No. 46–08295); its White Queen Mine (I.D. No. 46–08297); its Brushy Eagle Mine (I.D. No. 46–08315); its Low Gap Mine (I.D. No. 46–08442); and its Birch Fork Mine (I.D. No. 46–08493) all located in Raleigh County, West Virginia. The petitioner proposes to use electronically operated Roll-Down Doors constructed of rubber material similar to those used in conveyor belts to control ventilation within the air course in the main entries instead of using heavy Metal Doors. The petitioner asserts that the proposed alternative method would provide at least the same measure of protection as would the mandatory standard.

3. Leeco, Inc.

[Docket No. M–95–171–C]

Leeco, Inc., 100 Coal Drive, London, Kentucky 40741 has filed a petition to modify the application of 30 CFR 75.388(a)(1) (boreholes in advance of mining) to its Mine No. 63 (I.D. No. 15–16413); its Mine No. 68 (I.D. No. 15–17497) located in Perry County, Kentucky; its Mine No. 60 (I.D. No. 15–12941); and its Mine No. 66 (I.D. No. 15–17172) located in Leslie County, Kentucky. Instead of drilling boreholes,