

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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on February 12, 1996, Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application to the Drug Enforcement Administration to be registered as an importer of phenylacetone (8501) a basic class of controlled substance listed in Schedule II.

The firm plans to import the phenylacetone to manufacture dextroamphetamine sulfate.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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 1301.54 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than May 6, 1996.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42 (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), (b), (c), (d), (e), and (f) are satisfied.

Dated: March 27, 1996.

Gene R. Haislip,

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

[FR Doc. 96-8306 Filed 4-3-96; 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 January 31, 1996, Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, 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
Tetrahydrocannabinols (7370) ....	I
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-alphaacetylmethadol (9648)	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The firm plans to produce bulk finished products 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 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 3, 1996.

Dated: March 27, 1996.

Gene R. Haislip,

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

[FR Doc. 96-8307 Filed 4-3-96; 8:45 am]

BILLING CODE 4410-09-M

### 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 of 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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on January 31, 1996, Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Coca Leaves (9040) .....	II
Opium, raw (9600) .....	II
Opium poppy (9650) .....	II
Poppy Straw Concentrate (9670)	II

The firm plans to import the listed controlled substances to manufacture bulk, finished product.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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 1301.54 in such form as prescribed by 21 CFR 1316.47.

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 May 6, 1996.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42(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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: March 27, 1996.

Gene R. Haislip,

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

[FR Doc. 96-8308 Filed 4-3-96; 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 January 31, 1996, Radian Corporation, P.O. Box 201088, 8501 Mopac Blvd., Austin, Texas 78720, has made written request to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Cathinone (1235) .....	I
Methcathinone (1237) .....	I
N-Ethylamphetamine (1475) .....	I
N,N-Dimethylamphetamine (1480) .....	I
Aminorex (1585) .....	I
4-Methylaminorex (cis isomer) (1590) .....	I
Methaqualone (2565) .....	I
Lysergic acid diethylamide (7315) .....	I
Tetrahydrocannabinols (7370) ....	I
Mescaline (7381) .....	I
3,4-Methylenedioxymphetamine (7400) .....	I
3,4-Methylenedioxy-N-ethylamphetamine (7404) .....	I
3,4-Methylenedioxymethamphetamine (7405) .....	I
4-Methoxyamphetamine (7411) ..	I
Psilocybin (7437) .....	I
Psilocyn (7438) .....	I
Dihydromorphine (9145) .....	I
Normorphine (9313) .....	I
Acetylmethadol (9601) .....	I
Alphacetylmethadol except Levo-Alphacetylmethadol (9603) .....	I
Normethadone (9635) .....	I
3-Methylfentanyl (9813) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Methylphenidate (1724) .....	II
Amobarbital (2125) .....	II
Pentobarbital (2270) .....	II
Secobarbital (2315) .....	II
Phencyclidine (7471) .....	II
1-Piperidinocyclohexanecarbonitrile (8603) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II

Drug	Schedule
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II
Benzoylcegonine (9180) .....	II
Ethylmorphine (9190) .....	II
Hydrocodone (9193) .....	II
Isomethadone (9226) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Methadone-intermediate (9254) ..	II
Morphine (9300) .....	II
Levo-alphaacetylmethadol (9648) ..	II
Oxymorphone (9652) .....	II
Alfentanil (9737) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture small quantities of the listed controlled substances to make drug reference standards.

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 triplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. Federal Register Representative (CCR), and must be filed no later than June 3, 1996.

Dated: March 27, 1996.

Gene R. Haislip,

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

[FR Doc. 96-8309 Filed 4-3-96; 8:45 am]

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### 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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on January 31, 1996, Radian Corporation, 8501 Mopac Blvd., P.O. Box 201088, Austin, Texas 78720, made written request to the Drug Enforcement Administration to be registered as an

importer of the basic classes of controlled substances listed below:

Drug	Schedule
Ibogaine (7260) .....	I
Etorphine (except HCl) (9056) ...	I
Heroin (9200) .....	I
Cocaine (9041) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Dextropropoxyphene, bulk (non-dosage forms) (9273) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Oxymorphone (9652) .....	II

The firm plans to import small quantities of the listed controlled substances to make drug reference standards.

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 1301.54 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed 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 May 6, 1996.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42-(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 basic classes of any controlled substances 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 27, 1996.

Gene R. Haislip,

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

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