

(2004); Anne Lazar Thorn, M.D., 62 FR 847 (1997).

Here, it is clear Dr. Edwin is not currently licensed to handle controlled substances in Illinois, where he is registered with DEA. Therefore, he is not entitled to maintain that registration. Because Dr. Edwin is not entitled to a DEA registration in Illinois due to lack of state authorization to handle controlled substances, the Deputy Administrator concludes it is unnecessary to address whether Dr. Edwin's registration should be revoked based upon the remaining public interest grounds asserted in the Order to Show Cause/Immediate Suspension of Registration. See Fereida Walker-Graham, M.D., 68 FR 24761 (2003); Nathaniel-Aikens-Afful, M.D., 69 FR 16871 (1997); Sam F. Moore, D.V.M., 58 FR 14428 (1993).

Accordingly, the Deputy Administrator of the Drug Enforcement Administration, pursuant to the authority vested in her by 21 U.S.C. 823 and 824 and 28 CFR 0.100(b) and 0.104, hereby orders that DEA Certificate of Registration, AE7528295, issued to Paramaballoth Edwin, M.D., be, and it hereby is, revoked. The Deputy Administrator further orders that any pending applications for renewal or modification of such registration be, and they hereby are, denied. This order is effective November 1, 2004.

Dated: September 13, 2004.

**Michele M. Leonhart,**

*Deputy Administrator.*

[FR Doc. 04-21967 Filed 9-29-04; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to 21 CFR 1301.33(a), this is a notice that on June 10, 2004 Aldrich Chemical Company Inc., DBA Isotec, 3858 Benner Road, Miamisburg, Ohio 45342-4304, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed:

Drug	Schedule
Cathinone (1235) .....	I
Methcathinone (1237) .....	I
N-Ethylamphetamine (1475) .....	I
N,N-Dimethylamphetamine (1480) .....	I
Aminorex (1585) .....	I
Gamma hydroxybutyric acid (2010) .....	I
Methaqualone (2565) .....	I

Drug	Schedule
Lysergic acid dethylamide (7315) .....	I
Tetrahydrocannabinols (7370) .....	I
Mescaline (7381) .....	I
2,5-Dimethoxyamphetamine (7396) .....	I
3,4-Methylenedioxyamphetamine (7400) .....	I
3,4-Methylenedioxy-N-ethylamphetamine (7404) .....	I
3,4-Methylenedioxy-methamphetamine (7405) .....	I
4-Methoxyamphetamine (7411) .....	I
Psilocybin (7437) .....	I
Psilocyn (7438) .....	I
N-Ethyl-1-phenylcyclohexylamine (7455) .....	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
1-Phenylcyclohexylamine (7460) .....	II
Phencyclidine (7471) .....	II
Phenylacetone (8501) .....	II
1-Piperidinocyclohexane-carbonitrile (8603) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Benzoyllecgonine (9180) .....	II
Ethylmorphine (9190) .....	II
Hydrocodone (9193) .....	II
Isomethadone (9226) .....	II
Meperidine (9230) .....	II
Meperidine intermediate-A (9232) .....	II
Meperidine intermediate-B (9233) .....	II
Methadone (9250) .....	II
Methadone intermediate (9254) .....	II
Dextropropoxyphene, bulk, (non-dosage forms) (9273) .....	II
Levo-alphacetylmethadol (9648) .....	II
Oxymorphone (9652) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture small quantities of the listed controlled substances to produce isotope labeled standards for drug analysis.

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 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 (CCD)

and must be filed no later than (60 days from publication).

Dated: September 16, 2004.

**William J. Walker,**

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

[FR Doc. 04-21944 Filed 9-29-04; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Sheldon Kantor, D.P.M.; Revocation of Registration

On March 28, 2003, the then-Acting Administrator of the Drug Enforcement Administration (DEA) issued an Order to Show Cause/Immediate Suspension of Registration to Sheldon Kantor, D.P.M. (Dr. Kantor) of Hollywood, Florida. Dr. Kantor was notified of an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration, AK4080545, as a practitioner, and deny any pending applications for renewal or modification of such registration pursuant to 21 U.S.C. 823(f) and 824(a) for reason that his continued registration would be inconsistent with the public interest. Dr. Kantor was further notified that his DEA registration was immediately suspended as an imminent danger to the public health and safety pursuant to 21 U.S.C. 824(d).

The Order to Show Cause/Immediate Suspension alleged in relevant part, that during the week of February 24, 2003, and again on March 3, 2003, DEA received information from a registered distributor of controlled substances that Dr. Kantor had ordered and received large quantities of Schedule III and IV controlled substances. In response, DEA investigators presented Dr. Kantor with a Notice of Inspection, however, he refused to consent to the inspection. While speaking with investigators, Dr. Kantor admitted he had not maintained a log of controlled substances dispensed. When an investigator inquired as to the location of previously received controlled substances, Dr. Kantor stated that they were in the trunk of his car. He then refused to disclose the whereabouts of that vehicle. The Order to Show Cause alleged that Dr. Kantor also refused to consent to a subsequent inspection of his registered location.

The Order to Show Cause alleged that Dr. Kantor had been convicted in federal court and sentenced to 21 months imprisonment, commencing March 23, 2003, for his involvement in a scheme

to defraud the Medicare program of over one million dollars. The Order to Show Cause also outlined how one of Dr. Kantor's patient files revealed his distribution of large quantities of Schedule III and IV controlled substances to a single patient over a seven month period, including 19,200 dosage units of hydrocodone and four prescriptions for OxyContin, a Schedule II controlled substance, to that same patient. When DEA investigators executed a search warrant of the patient's residence on March 12, 2003, the search revealed the patient had several bottles of 100-count hydrocodone tablets. However, the vast majority of hydrocodone tablets, as well as other controlled substances, could not be accounted for.

According to the investigative file, the Order to Show Cause/Immediate Suspension of Registration was personally delivered to Dr. Kantor on March 31, 2003, at a federal detention center in Miami, Florida. More than thirty days have passed since the Order to Show Cause/Immediate Suspension of Registration was served on Dr. Kantor and DEA has not received a request for hearing or any other reply from Dr. Kantor or anyone purporting to represent him in this matter.

Therefore, the Deputy Administrator of DEA, finding that (1) thirty days having passed since the delivery of the Order to Show Cause/Immediate Suspension of Registration to Dr. Kantor, and (2) no request for hearing having been received, concludes that Dr. Kantor is deemed to have waived his hearing right. See David W. Linder, 67 FR 12579 (2002). After considering material from the investigative file in this matter, the Deputy Administrator now enters her final order without a hearing pursuant to 21 CFR 1301.43(d) and (e) and 1301.46.

The Deputy Administrator finds Dr. Kantor is currently registered with DEA as a practitioner under DEA Registration, AK4080545, in Schedules II through V. That registration expires on December 31, 2004. At the time DEA initiated action to revoke his DEA registration, Dr. Kantor was licensed under Florida law as a podiatrist. However, following the issuance and delivery of the Order to Show Cause/Immediate Suspension of Registration, the Deputy Administrator obtained a copy of a Final Order adopted by the State of Florida, Board of Podiatric Medicine (Board) on November 20, 2003. A review of the Final Order reveals that on the above date, the Board permanently revoked Dr. Kantor's license of podiatry.

The investigative file contains no evidence that the Board's Final Order revoking Dr. Kantor's podiatry license has been lifted, stayed or that his license has been reinstated. Therefore, the Deputy Administrator finds that Dr. Kantor is not currently authorized to practice medicine in the State of Florida and as a result, it is reasonable to infer that he is also without authorization to handle controlled substances in that state.

DEA does not have statutory authority under the Controlled Substances Act to issue or maintain a registration if the applicant or registrant is without state authority to handle controlled substances in the state in which he conducts business. See 21 U.S.C. 802(21), 823(f) and 824(a)(3). This prerequisite has been consistently upheld. See Stephen J. Graham, M.D., 69 FR 11661 (2004); Dominick A. Ricci, M.D., 58 FR 51104 (1993); Bobby Watts, M.D., 53 FR 11919 (1988).

Here, it is clear Dr. Kantor's podiatry license has been revoked and therefore, he is not currently licensed to handle controlled substances in Florida, the state where he maintains a DEA controlled substance registration. Therefore, Dr. Kantor is not entitled to a DEA registration in that state. Because Dr. Kantor is not entitled to a DEA registration in Florida due to his lack of state authorization to handle controlled substances, the Deputy Administrator concludes it is unnecessary to address whether his registration should be revoked based upon the public interest grounds asserted in the Order to Show Cause/Immediate Suspension of Registration. See Fereida Walker-Graham, M.D., 68 FR 24761 (2003); Nathaniel-Aikens-Afful, M.D., 62 FR 16871 (1997); Sam F. Moore, D.V.M., 58 FR 14428 (1993).

Accordingly, the Deputy Administrator of the Drug Enforcement Administration, pursuant to the authority vested in her by 21 U.S.C. 823 and 824 and 28 CFR 0.100(b) and 0.104, hereby orders that DEA Certificate of Registration, AK 4080545, issued to Sheldon Kantor, D.P.M. be, and it hereby is, revoked. The Deputy Administrator further orders that any pending applications for renewal or modification of such registration be, and they hereby are, denied. This order is effective November 1, 2004.

Dated: September 13, 2004.

**Michele M. Leonhart,**

*Deputy Administrator.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Registration

By Notice dated May 18, 2004, and published in the **Federal Register** on June 3, 2004, (69 FR 31412-31413), 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:

Drug	Schedule
Dihydromorphine (9145) .....	I
Hydromorphine (9150) .....	II

The company plans to manufacture bulk product and finished dosage units for distribution to its customers.

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 Abbott Laboratories to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Abbott Laboratories 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 classes of controlled substances listed.

**William J. Walker,**

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

[FR Doc. 04-21952 Filed 9-29-04; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to 21 CFR 1301.33(a), this is notice that on July 22, 2004, Lifepoint, Inc., 10400 Trademark Street, Rancho Cucamonga, California 91730, made application by renewal to the Drug