

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 is granted.

Dated: November 19, 2003.

Laura M. Nagel,

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

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 3-4]

Anthony D. Dinozzi, D.D.S., Revocation of Registration

On September 25, 2002, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause to Anthony David Dinozzi, D.D.S. (Respondent) notifying him of an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration, BD4361692 under 21 U.S.C. 824(a)(2), (a)(3), and (a)(4). The Order to Show Cause further sought to deny any pending applications for renewal or modification of the Respondent's registration for reasons that he was convicted of a felony offense related to controlled substances, is not authorized to handle controlled substances, and his continued registration would be inconsistent with the public interest.

Specifically, the Order to Show Cause alleged that the Respondent is not authorized under state law to handle controlled substances based upon the March 31, 2001, expiration of his Pennsylvania state license to practice dentistry. The Order to Show Cause further alleged that the Respondent was convicted in Clermont County, Ohio on charges of Tampering with Evidence (a third degree felony) and Aggravated Trafficking in Drugs under Bulk (a fourth degree felony).

By letter dated October 11, 2002, the Respondent, acting *pro se*, timely requested a hearing in this matter. On October 25, 2002, the presiding Administrative Law Judge Mary Ellen Bittner (Judge Bittner) issued to the Government as well as the Respondent an Order for Prehearing Statements.

In view of filing a prehearing statement, the Government filed Government's Request for Stay of

Proceedings and Motion for Summary Judgment. The Government asserted that the Respondent is without authorization to handle controlled substances in the State of Pennsylvania, and as a result, further proceedings in the matter were not required. On November 6, 2002, Judge Bittner issued a Memorandum to Counsel staying the Order for Filing Prehearing Statements, and afforded the Respondent until November 25, 2002, to respond to the Government's Motion. The Respondent did not file a response.

Accordingly, on January 13, 2003, Judge Bittner issued her Opinion and Recommended Decision of the Administrative Law Judge (Opinion and Recommended Decision). As part of her recommended ruling, Judge Bittner granted the Government's Motion for Summary Disposition and found that the Respondent lacked authorization to handle controlled substances in Pennsylvania, the jurisdiction in which the is registered with DEA. In granting the Government's motion, Judge Bittner also recommended that the Respondent's DEA registration be revoked and any pending applications for modification or renewal be denied. No exceptions were filed by either party to Judge Bittner's Opinion and Recommended Decision, and on February 19, 2003, the record of these proceedings was transmitted to the Office of the DEA Deputy Administrator.

The Acting Deputy Administrator has considered the record in its entirety and pursuant to 21 CFR 1316.67, hereby issues her final order based upon findings of fact and conclusions of law as hereinafter set forth. The Acting Deputy Administrator adopts, in full, the Opinion and Recommended Decision of the Administrative Law Judge.

The Acting Deputy Administrator finds that the Respondent currently possesses DEA Certificate of Registration BD4361692, and is registered to handle controlled substances in Pennsylvania. The Acting Deputy Administrator further finds that on March 31, 2001, the Respondent license to practice dentistry expired. There is no evidence before the Acting Deputy Administrator that the Respondent has applied for, and been granted renewal of his Pennsylvania dental license. Therefore, the Acting Deputy Administrator finds that the Respondent is currently not licensed to practice dentistry in Pennsylvania 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 *Kanwaljit S. Serai, M.D.*, 68 FR 48943 (2003); *Dominick A. Ricci, M.D.*, 58 FR 51104 (1993); *Bobby Watts, M.D.*, 53 FR 11919 (1988). The agency has further held that a person may not hold a DEA registration even if the loss of state authority is due to the expiration of state licensure without further action by the state. *William D. Levitt, D.O.*, 64 FR 49,822 (1999).

Here, it is clear that the Respondent is not currently licensed to handle controlled substances in Pennsylvania, where he is registered with DEA. Therefore, he is not entitled to maintain that registration. Because the Respondent is not entitled to a DEA registration in Pennsylvania due to his lack of state authorization to handle controlled substances, the Acting Deputy Administrator concludes that it is unnecessary to address whether the Respondent's registration should be revoked based upon the other grounds asserted in the Order to Show Cause. See *Cordell Clark, M.D.*, 68 FR 48942 (2003); *Nathaniel-Aikens-Afful, M.D.*, 62 FR 16871 (1997); *Sam F. Moore, D.V.M.*, 58 FR 14428 (1993).

Accordingly, the Acting 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, BD4361692, issued to Anthony David Dinozzi, D.D.S., be, and it hereby is, revoked. The Acting Deputy Administrator further orders that any pending applications for renewal of such registration be, and they hereby are, denied. This order is effective January 2, 2004.

Dated: November 13, 2003.

Michele M. Leonhart,

Acting Deputy Administrator.

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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 25,

2003, Lifepoint, Inc., 10400 Trademark Street, Rancho Cucamonga, California 91730, 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-Methylenedioxyamphetamine (7400).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxy-N-methylamphetamine (7405).	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
phencyclidine (7471)	II
Benzoylcegonine (9180)	II
Morphine (9300)	II

The firm plans to produce small quantities of controlled substances for use in drug test kits.

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 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 February 2, 2004.

Dated: November 14, 2003.

Laura M. Nagel,

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

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Drug	Schedule
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxy-N-methylamphetamine (7405).	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Cocaine (9041)	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.

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 LinZhi International, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Lin-Zhi International, Inc. to ensure that the company's registration is consistent with the public interest. This 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 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 bulk manufacturer of the basic classes of controlled substances listed is granted.

Dated: November 19, 2003.

Laura M. Nagel,

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

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BILLING CODE 4410-09-M

Order to Show Cause alleged that the Respondent's DEA Certificate of Registration should be revoked because the Respondent was without authorization to handle controlled substances. The Order to Show Cause further sought denial of any pending applications for registration based on allegations that the Respondent's continued registration would be inconsistent with the public interest. Specifically, the Order to Show Cause alleged that effective March 15, 2002, the California Medical Board (Medical Board) ordered that Respondent be prohibited from handling controlled substances based upon acts of negligence in both his care of patients and billing practices. The Order to Show Cause further alleged that a DEA investigation revealed the Respondent's failure to adhere to various DEA-recordkeeping requirements.

By letter dated September 30, 2002, the Respondent, acting *pro se*, timely requested a hearing in this matter. On October 15, 2002, the presiding Administrative Law Judge Mary Ellen Bittner (Judge Bittner) issued to the Government as well as the Respondent an Order for Prehearing Statements.

In lieu of filing a prehearing statement, the Government filed Government's Request for Stay of Proceedings and Motion for Summary Judgment. The Government argued that the Respondent is without authorization to handle controlled substances in the State of California, and as a result, further proceedings in the matter were not required. Attached to the Government's motion was a copy of a declaration from the Medical Board's Chief of Enforcement who averred among other things, that on March 15, 2002, the Medical Board issued an Interim Order of Suspension summarily suspending the Respondent's medical license. The Medical Board representative further stated that as of October 25, 2002, the Medical Board's Interim Order of Suspension remained in effect. On November 7, 2002, Judge Bittner issued a Memorandum to Counsel staying the filing or prehearing statements and afforded the Respondent until November 26, 2002, to respond to the Government's Motion.

On or around October 30, 2002, the Respondent filed a prehearing statement where he disputed allegations that he maintained inadequate records of his handling of controlled substances. The Respondent maintained that his procedures for handling controlled substances were proper, and that prosecution witnesses offered biased testimony in the previous Board proceeding involving the Respondent's

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 19, 2003, and published in the **Federal Register** on September 2, 2003, (68 FR 52225), LinZhi International, Inc., 687 North Pastoria Avenue, Sunnyvale, California 94085, made application to the Drug Enforcement Administration for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 03-2]

Jules M. Lusman, M.D., Revocation of Registration

On September 6, 2002, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause to Jules Lusman, M.D. (Respondent) notifying him of an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration, BL2210300 under 21 U.S.C. 824(a)(3) and (a)(4). The