

A person who is licensed as a physician, however, is “registered” and exempt from the statute’s registration fee and application requirements. Ga. Code Ann. § 16–13–35(g)(2) (Westlaw, current through acts passed during the 2019 Session of the General Assembly). The Georgia Code defines “physician” as “a person licensed to practice medicine.” Ga. Code Ann. § 43–34–21(2) (Westlaw, current through acts passed during the 2019 Session of the General Assembly). Under Georgia law, “to practice medicine” includes “attaching the title ‘M.D.’ . . . to one’s name, indicating that such person is engaged in the treatment or diagnosis of disease, defects, or injuries to human beings.” Ga. Code Ann. § 43–34–21(3) (Westlaw, current through acts passed during the 2019 Session of the General Assembly).

Here, the undisputed evidence in the record, including Respondent’s admission, is that Respondent currently lacks authority to practice medicine in Georgia. Government Motion, Exhs. 2–4; RMRC, at 2. As already discussed, a “physician,” under Georgia law, is a person licensed to practice medicine. Further, under Georgia law, a “physician” is registered to dispense controlled substances. Because Respondent lacks authority to practice medicine in Georgia, he is not registered to handle controlled substances in Georgia according to Georgia law. Accordingly, Respondent is not eligible to maintain a DEA registration and I will order that Respondent’s DEA registration be revoked.⁶

Order

Pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 824(a), I hereby revoke DEA Certificate of Registration No. BS4014332 issued to John Yolman Salinas, M.D. Pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 823(f), I hereby deny any pending application of John Yolman Salinas to renew or to modify this registration, and any pending application of John Yolman Salinas to be registered in the state of Georgia. This Order is effective October 21, 2019.

⁶ Given my findings that Respondent is registered with the DEA in Georgia, that his Georgia medical license is revoked, and that he lacks authority in Georgia to dispense controlled substances, I find that both of Respondent’s CAPs—changing his registered address to another state or a Territory of the United States, and “procuring a position as an undercover physician to infiltrate pill mills and help in the war against drugs”—provide no basis for me to discontinue or defer this proceeding. 21 U.S.C. 824(c)(3).

Dated: September 9, 2019.

Uttam Dhillon,

Acting Administrator.

[FR Doc. 2019–20420 Filed 9–19–19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

**Importer of Controlled Substances
Application: Fisher Clinical Services,
Inc.**

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 21, 2019. Such persons may also file a written request for a hearing on the application on or before October 21, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 19, 2019, Fisher Clinical Services, Inc., 700A–C Nestle Way, Breinigsville, Pennsylvania 18031–1522 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Psilocybin	7437	I
Methylphenidate	1724	II
Levorphanol	9220	II
Noroxymorphone	9668	II
Tapentadol	9780	II

The company plans to import the listed controlled substances for clinical trials.

Dated: August 19, 2019.

Neil D. Doherty,

Acting Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

**Importer of Controlled Substances
Application: Galephar Pharmaceutical
Research Inc.**

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 21, 2019. Such persons may also file a written request for a hearing on the application on or before October 21, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on June 10, 2019, Galephar Pharmaceutical Research Inc., #100 Carr 198 Industrial Park, Juncos, Puerto Rico, 00777 applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Hydromorphone	9150	II

The company plans to import the listed controlled substance in finished dosage form for analytical purpose only.

Dated: August 22, 2019.

Neil D. Doherty,

Acting Assistant Administrator.

[FR Doc. 2019–20412 Filed 9–19–19; 8:45 am]

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