

State law to engage in the . . . dispensing of controlled substances.” With respect to a practitioner, DEA has also long held that the possession of authority to dispense controlled substances under the laws of the state in which a practitioner engages in professional practice is a fundamental condition for obtaining and maintaining a practitioner’s registration. *See, e.g., James L. Hooper, D.O.*, 76 FR 71371, 71372 (2011), *pet. for rev. denied*, 481 F. App’x 826 (4th Cir. 2012); *Frederick Marsh Blanton, D.O.*, 43 FR 27616, 27617 (1978).³

According to Kansas statute, “dispense” means “to deliver a controlled substance to an ultimate user or research subject by or pursuant to the lawful order of a practitioner, including the packaging, labeling or compounding necessary to prepare the substance for that delivery” Kan. Stat. Ann. sec. 65–4101(l) (2024). Further, a “practitioner” includes, among others, “a person licensed to practice medicine and surgery” *Id.* sec. 65–4101(ll).

Here, the undisputed evidence in the record is that Registrant lacks authority to practice medicine in Kansas. As discussed above, an individual must be a licensed practitioner to dispense a controlled substance in Kansas. Thus, because Registrant lacks authority to practice medicine in Kansas and, therefore, is not authorized to handle controlled substances in Kansas, Registrant is not eligible to maintain a DEA registration. Accordingly, the Agency will order that Registrant’s DEA registration be revoked.

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

³ This rule derives from the text of two provisions of the Controlled Substances Act (CSA). First, Congress defined the term “practitioner” to mean “a physician . . . or other person licensed, registered, or otherwise permitted, by . . . the jurisdiction in which he practices . . . , to distribute, dispense, . . . [or] administer . . . a controlled substance in the course of professional practice.” 21 U.S.C. 802(21). Second, in setting the requirements for obtaining a practitioner’s registration, Congress directed that “[t]he Attorney General shall register practitioners . . . if the applicant is authorized to dispense . . . controlled substances under the laws of the State in which he practices.” 21 U.S.C. 823(g)(1). Because Congress has clearly mandated that a practitioner possess state authority in order to be deemed a practitioner under the CSA, DEA has held repeatedly that revocation of a practitioner’s registration is the appropriate sanction whenever he is no longer authorized to dispense controlled substances under the laws of the state in which he practices. *See, e.g., James L. Hooper*, 76 FR at 71371–72; *Sheran Arden Yeates, D.O.*, 71 FR 39130, 39131 (2006); *Dominick A. Ricci, D.O.*, 58 FR 51104, 51105 (1993); *Bobby Watts, D.O.*, 53 FR 11919, 11920 (1988); *Frederick Marsh Blanton*, 43 FR 27617.

of Registration No. BM8877473 issued to William J. Mack, M.D. Further, pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 823(g)(1), I hereby deny any pending applications of William J. Mack, M.D., to renew or modify this registration, as well as any other pending application of William J. Mack, M.D., for additional registration in Kansas. This Order is effective November 12, 2024.

Signing Authority

This document of the Drug Enforcement Administration was signed on October 2, 2024, by Administrator Anne Milgram. That document with the original signature and date is maintained by DEA. For administrative purposes only, and in compliance with requirements of the Office of the Federal Register, the undersigned DEA Federal Register Liaison Officer has been authorized to sign and submit the document in electronic format for publication, as an official document of DEA. This administrative process in no way alters the legal effect of this document upon publication in the **Federal Register**.

Heather Achbach,

Federal Register Liaison Officer, Drug Enforcement Administration.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Robert P. Hansen, M.D.; Decision and Order

On November 29, 2023, the Drug Enforcement Administration (DEA or Government) issued an Order to Show Cause (OSC) to Robert P. Hansen, M.D., of Fresno, California (Registrant). Request for Final Agency Action (RFAA), Exhibit (RFAAX) 2, at 1, 4. The OSC proposed the revocation of Registrant’s Certificate of Registration No. BH4921727, alleging that Registrant’s registration should be revoked because Registrant is “currently without authority to prescribe, administer, dispense, or otherwise handle controlled substances in the State of California, the state in which [he is] registered with DEA.” *Id.* at 2 (citing 21 U.S.C. 824(a)(3)).

The OSC notified Registrant of his right to file with DEA a written request for hearing, and that if he failed to file such a request, he would be deemed to have waived his right to a hearing and be in default. *Id.* at 2–3 (citing 21 CFR 1301.43). Here, Registrant did not

request a hearing. RFAA, at 3.¹ “A default, unless excused, shall be deemed to constitute a waiver of the registrant’s/applicant’s right to a hearing and an admission of the factual allegations of the [OSC].” 21 CFR 1301.43(e).

Further, “[i]n the event that a registrant . . . is deemed to be in default . . . DEA may then file a request for final agency action with the Administrator, along with a record to support its request. In such circumstances, the Administrator may enter a default final order pursuant to [21 CFR] § 1316.67.” *Id.* § 1301.43(f)(1). Here, the Government has requested final agency action based on Registrant’s default pursuant to 21 CFR 1301.43(c), (f), 1301.46. RFAA, at 1; *see also* 21 CFR 1316.67.

Findings of Fact

The Agency finds that, in light of Registrant’s default, the factual allegations in the OSC are admitted. According to the OSC, effective August 18, 2023, Registrant surrendered his California physician and surgeon license. RFAAX 2, at 2. According to California online records, of which the Agency takes official notice, Registrant’s California physician and surgeon license remains surrendered.² California DCA License Search, <https://search.dca.ca.gov> (last visited date of signature of this Order). Accordingly, the Agency finds that Registrant is not licensed to practice as a physician in

¹ Based on the Government’s submissions in its RFAA dated February 26, 2024, the Agency finds that service of the OSC on Registrant was adequate. Specifically, the included Declaration from a DEA Diversion Investigator (DI) indicates that on January 10, 2024, Registrant was successfully served a copy of the OSC via email to his registered email address, as the email was not returned as undeliverable. RFAAX 1, at 1, 3; *Mohammed S. Aljanaby, M.D.*, 82 FR 34552, 34552 (2017) (finding that service by email satisfies due process where the email is not returned as undeliverable and other methods have been unsuccessful). The DI notes in the Declaration that Registrant was mailed a copy of the OSC on or about December 8, 2023, but on January 10, 2024, the mailing was returned as undeliverable. RFAAX 1, at 1.

² Under the Administrative Procedure Act, an agency “may take official notice of facts at any stage in a proceeding—even in the final decision.” United States Department of Justice, Attorney General’s Manual on the Administrative Procedure Act 80 (1947) (Wm. W. Gaunt & Sons, Inc., Reprint 1979). Pursuant to 5 U.S.C. 556(e), “[w]hen an agency decision rests on official notice of a material fact not appearing in the evidence in the record, a party is entitled, on timely request, to an opportunity to show the contrary.” Accordingly, Registrant may dispute the Agency’s finding by filing a properly supported motion for reconsideration of findings of fact within fifteen calendar days of the date of this Order. Any such motion and response shall be filed and served by email to the other party and to Office of the Administrator, Drug Enforcement Administration at dea.addo.attorneys@dea.gov.

California, the state in which he is registered with DEA.

Discussion

Pursuant to 21 U.S.C. 824(a)(3), the Attorney General is authorized to suspend or revoke a registration issued under 21 U.S.C. 823 “upon a finding that the registrant . . . has had his State license or registration suspended . . . [or] revoked . . . by competent State authority and is no longer authorized by State law to engage in the . . . dispensing of controlled substances.” With respect to a practitioner, DEA has also long held that the possession of authority to dispense controlled substances under the laws of the state in which a practitioner engages in professional practice is a fundamental condition for obtaining and maintaining a practitioner’s registration. *See, e.g., James L. Hooper, M.D.*, 76 FR 71371, 71372 (2011), *pet. for rev. denied*, 481 F. App’x 826 (4th Cir. 2012); *Frederick Marsh Blanton, M.D.*, 43 FR 27616, 27617 (1978).³

According to California statute, “dispense” means “to deliver a controlled substance to an ultimate user or research subject by or pursuant to the lawful order of a practitioner, including the prescribing, furnishing, packaging, labeling, or compounding necessary to prepare the substance for that delivery.” Cal. Health & Safety Code sec. 11010 (West 2024). Further, a “practitioner” means a person “licensed, registered, or otherwise permitted, to distribute, dispense, conduct research with respect to, or administer, a controlled substance in the course of professional practice or research in [the] state.” *Id.* sec. 11026(c).

Here, the undisputed evidence in the record is that Registrant currently lacks authority to practice as a physician in

California. As discussed above, an individual must be a licensed practitioner to dispense a controlled substance in California. Thus, because Registrant currently lacks authority to practice as a physician in California and, therefore, is not currently authorized to handle controlled substances in California, Registrant is not eligible to maintain a DEA registration. Accordingly, the Agency will order that Registrant’s DEA registration be revoked.

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. BH4921727 issued to Robert P. Hansen, M.D. Further, pursuant to 28 CFR 0.100(b) and the authority vested in me by 21 U.S.C. 823(g)(1), I hereby deny any pending applications of Robert P. Hansen, M.D., to renew or modify this registration, as well as any other pending application of Robert P. Hansen, M.D., for additional registration in California. This Order is effective November 12, 2024.

Signing Authority

This document of the Drug Enforcement Administration was signed on October 2, 2024, by Administrator Anne Milgram. That document with the original signature and date is maintained by DEA. For administrative purposes only, and in compliance with requirements of the Office of the Federal Register, the undersigned DEA Federal Register Liaison Officer has been authorized to sign and submit the document in electronic format for publication, as an official document of DEA. This administrative process in no way alters the legal effect of this document upon publication in the **Federal Register**.

Heather Achbach,

Federal Register Liaison Officer, Drug Enforcement Administration.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Janet S. Pettyjohn, D.O.; Decision and Order

On June 21, 2023, the Drug Enforcement Administration (DEA or Government) issued an Order to Show Cause and Immediate Suspension of Registration (OSC/ISO) to Janet S. Pettyjohn, D.O., of Tampa, Florida (Registrant). Request for Final Agency

Action (RFAA), Exhibit (RFAAX) 2, at 1. The OSC/ISO informed Registrant of the immediate suspension of her DEA registration, Control No. AP6641713,¹ pursuant to 21 U.S.C. 824(d), alleging that Registrant’s continued registration constitutes “‘an imminent danger to the public health or safety.’” *Id.* (quoting 21 U.S.C. 824(d)). The OSC/ISO also proposed the revocation of Registrant’s registration, alleging that Registrant’s continued registration is inconsistent with the public interest. *Id.* (citing 21 U.S.C. 823(g)(1), 824(a)(4)).

The OSC/ISO notified Registrant of her right to file with DEA a written request for hearing within 30 days after the date of receipt of the OSC/ISO. RFAAX 2, at 6. The OSC/ISO also notified Registrant that if she failed to file such a request, she would be deemed to have waived her right to a hearing and be in default. *Id.* (citing 21 CFR 1301.43). Here, Registrant did not request a hearing. RFAA, at 1–2.² “A default, unless excused, shall be deemed to constitute a waiver of the [registrant’s] right to a hearing and an admission of the factual allegations of the [OSC/ISO].” 21 CFR 1301.43(e).

Further, “[i]n the event that a registrant . . . is deemed to be in default . . . DEA may then file a request for final agency action with the Administrator, along with a record to support its request. In such circumstances, the Administrator may enter a default final order pursuant to [21 CFR] § 1316.67.” *Id.* § 1301.43(f)(1). Here, the Government has requested final agency action based on Registrant’s default pursuant to 21 CFR 1301.43(c), (f), because Registrant has not timely requested a hearing nor filed an Answer to the June 21, 2023 OSC/ISO. *See also id.* § 1316.67.

I. Findings of Fact

The Agency finds that, in light of Registrant’s default, the factual allegations in the OSC/ISO are admitted. 21 CFR 1301.43(e). Accordingly, Registrant admits that between August 2021 and February 2023, she issued 60 prescriptions for controlled substances to six individuals without conducting

¹ This registration expired on March 31, 2024. RFAAX 1. The fact that a registrant allows her registration to expire during the pendency of an administrative enforcement proceeding does not impact the Agency’s jurisdiction or prerogative to adjudicate the OSC to finality. *Jeffrey D. Olsen, M.D.*, 84 FR 68474, 68479 (2019).

² Based on the Government’s submissions in its RFAA dated December 18, 2023, the Agency finds that service of the OSC/ISO on Registrant was adequate. Attached to the Government’s RFAA is the Declaration of a DEA Diversion Investigator asserting that on June 21, 2023, the OSC/ISO was served on Registrant’s counsel, who confirmed receipt. RFAAX 3, appendix A, at 1.

³ This rule derives from the text of two provisions of the Controlled Substances Act (CSA). First, Congress defined the term “practitioner” to mean “a physician . . . or other person licensed, registered, or otherwise permitted, by . . . the jurisdiction in which he practices . . . , to distribute, dispense, . . . [or] administer . . . a controlled substance in the course of professional practice.” 21 U.S.C. 802(21). Second, in setting the requirements for obtaining a practitioner’s registration, Congress directed that “[t]he Attorney General shall register practitioners . . . if the applicant is authorized to dispense . . . controlled substances under the laws of the State in which he practices.” 21 U.S.C. 823(g)(1). Because Congress has clearly mandated that a practitioner possess state authority in order to be deemed a practitioner under the CSA, DEA has held repeatedly that revocation of a practitioner’s registration is the appropriate sanction whenever he is no longer authorized to dispense controlled substances under the laws of the state in which he practices. *See, e.g., James L. Hooper*, 76 FR 71371–72; *Sheran Arden Yeates, D.O.*, 71 FR 39130, 39131 (2006); *Dominick A. Ricci, D.O.*, 58 FR 51104, 51105 (1993); *Bobby Watts, D.O.*, 53 FR 11919, 11120 (1988); *Frederick Marsh Blanton*, 43 FR 27617.