purposes of this Investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel,² solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.³

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission. Issued: April 1, 2024.

Lisa Barton,

Secretary to the Commission. [FR Doc. 2024–07215 Filed 4–4–24; 8:45 am] BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 23-64]

Traesa A. Brown, M.D.; Decision and Order

On August 31, 2023, the Drug Enforcement Administration (DEA or Government) issued an Order to Show Cause and Immediate Suspension of Registration (OSC/ISO) to Traesa A. Brown, M.D. (Respondent) of Florence, South Carolina. OSC, at 1, 5. The OSC/ ISO informed Respondent of the immediate suspension of her DEA Certificate of Registration (registration or COR), Control No. BB9937624, pursuant to 21 U.S.C. 824(d), alleging that Respondent's continued registration constitutes "' an imminent danger to the public health or safety.' '' Id. at 1 (quoting 21 U.S.C. 824(d)). The OSC/ ISO also proposed the revocation of Respondent's registration, alleging that Respondent's continued registration is inconsistent with the public interest and alleging that Respondent has no state authority to handle controlled substances. Id. (citing 21 U.S.C. 823(g)(1), 824(a)(3), 824(a)(4)).

On September 20, 2023, Respondent requested a hearing. On October 13, 2023, the Government filed a Motion for Summary Disposition only pertaining to the allegation that Respondent lacks state authority to handle controlled substances.¹ See Government's Notice of Filing of Evidence and Motion for Summary Disposition (Motion for Summary Disposition), dated October 13, 2023.² Respondent did not respond to the Government's Motion for Summary Disposition. On October 23, 2023, Administrative Law Judge Paul E. Soeffing (the ALJ) granted the Government's Motion for Summary Disposition and recommended the revocation of Respondent's registration, finding that because Respondent lacks state authority to handle controlled substances in South Carolina, the state in which she is registered with DEA, "there is no other fact of consequence for this tribunal to decide in order to determine whether or not she is entitled to hold a COR." Order Granting the Government's Motion for Summary Disposition, and Recommended Rulings, Findings of Fact, Conclusions of Law, and Decision of the Administrative Law Judge (RD), at 6. Respondent did not file exceptions to the RD.

Having reviewed the entire record, the Agency adopts and hereby incorporates by reference the entirety of the ALJ's rulings, findings of fact, conclusions of law, and recommended sanction as found in the RD and summarizes and expands upon portions thereof herein.

Findings of Fact

The Government asserts that on October 1, 2022, Respondent's South Carolina controlled substance registration expired. RD, at 3–4.³ Further, the Government asserts that on June 30, 2023, Respondent's South

² The Government originally filed a Motion for Summary Disposition on October 12, 2023, and therein asserted that Respondent had failed to timely file an Answer to the allegations in the OSC/ ISO. RD, at 2 n.4; Motion for Summary Disposition, dated October 12, 2023, at 3–4. Later on October 12, 2023, the Government was informed that Respondent had filed an Answer on October 10, 2023, and was provided with a copy of Respondent's Answer. RD, at 2 n.4. On October 13, 2023, the Government filed its amended Motion for Summary Disposition, referenced in this Decision, with revisions based on its receipt of the copy of Respondent's Answer. *Id.; see also* Motion for Summary Disposition, dated October 13, 2023.

³ See also Motion for Summary Disposition, dated October 13, 2023, Exhibit (GX) 1; Motion for Summary Disposition, dated October 13, 2023, at 4– 5. Carolina medical license expired. RD, at $4.^4$

According to South Carolina online records, of which the Agency takes official notice, Respondent's South Carolina controlled substance registration is expired.⁵ SC DHEC Bureau of Drug Control, Controlled Substances Registration Verification, https://.dhec.sc.gov//Licensing/Home/ Verify (last visited date of signature of this Order). Further, Respondent's South Carolina medical license is listed as "lapsed." South Carolina Board of Medical Examiners, Licensee Lookup, https://verify.llronline.com/LicLookup/ Med/Med.aspx (last visited date of signature of this Order).

Accordingly, the Agency finds that Respondent is not currently licensed to engage in the practice of medicine nor to handle controlled substances in South Carolina, the state in which she is registered with the DEA.

Discussion

Pursuant to 21 U.S.C. 824(a)(3), the Attorney General is authorized to suspend or revoke a registration issued under section 823 of the Controlled Substances Act (CSA) "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, the 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

⁵ 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, Respondent 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.

² All contract personnel will sign appropriate nondisclosure agreements.

³Electronic Document Information System (EDIS): https://edis.usitc.gov.

¹ This suggests that the Government has dropped the public interest allegation included in the OSC/ ISO; as such, the Agency will only consider the lack of state authority allegation from the OSC/ISO.

⁴ See also Motion for Summary Disposition, dated October 13, 2023, at 4. As noted by the ALJ, the Government did not submit documentary evidence regarding the status of Respondent's South Carolina medical license as they had for Respondent's South Carolina controlled substance registration, *see supra* n.3. RD, at 4 n.8.

F. App'x 826 (4th Cir. 2012); *Frederick Marsh Blanton, M.D.,* 43 FR 27616, 27617 (1978).⁶

According to South Carolina statute, "[e]very person who manufactures, distributes, or dispenses any controlled substance or who proposes to engage in the manufacture, distribution, or dispensing of any controlled substance, shall obtain a registration issued by the [Department of Health and Environmental Control] in accordance with its rules and regulations." S.C. Code section 44-53-290(a) (2024). Further, "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, administering, packaging, labeling, or compounding necessary to prepare the substance for the delivery." Id. section 44-53-110(15).

Here, the undisputed evidence in the record is that Respondent currently lacks authority to dispense controlled substances in South Carolina because her South Carolina controlled substance registration is expired. As discussed above, an individual must hold a controlled substance registration to dispense a controlled substance in South Carolina. Thus, because Respondent lacks authority to handle controlled substances in South Carolina, Respondent is not eligible to maintain a DEA registration. RD, at 6. Accordingly, the Agency will order that Respondent'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. BB9937624 issued to Traesa A. Brown, 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 Traesa A. Brown, M.D., to renew or modify this registration, as well as any other pending application of Traesa A. Brown, M.D., for additional registration in South Carolina. This Order is effective May 6, 2024.

Signing Authority

This document of the Drug Enforcement Administration was signed on April 1, 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. [FR Doc. 2024–07237 Filed 4–4–24; 8:45 am] BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 23-63]

Ralph Reach, M.D.; Decision And Order

On August 30, 2023, the Drug Enforcement Administration (DEA or Government) issued an Order to Show Cause (OSC) to Ralph Reach, M.D. (Respondent). OSC, at 1, 4. The OSC proposed the revocation of Respondent's DEA Certificates of Registration Nos. FR0673548 and FR0004589 at the registered addresses of 142 Mall Church Road, Cedar Bluff, Virginia 24609 and 102 North Broadway Street, Johnson City, Tennessee 37601, respectively. Id. at 1. The OSC alleged that Respondent's DEA registrations should be revoked because Respondent is "without authority to prescribe, administer, dispense, or otherwise handle controlled substances in the State of Tennessee and the Commonwealth of Virginia, the jurisdictions in which [he is] registered with DEA." Id. at 2 (citing 21 U.S.C. 824(a)(3)).

On September 14, 2023, Respondent requested a hearing. On September 27, 2023, the Government filed a Motion for Summary Disposition, which Respondent opposed. On November 7, 2023, Administrative Law Judge Teresa A. Wallbaum (the ALJ) granted the Government's Motion for Summary Disposition and recommended the revocation of Respondent's registration, finding that because Respondent lacks state authority to handle controlled substances in Tennessee and Virginia, the states in which he is registered with DEA, "[t]here is no genuine issue of material fact in this case." Order Granting the Government's Motion for Summary Disposition, and Recommended Rulings, Findings of Fact, Conclusions of Law, and Decision of the Administrative Law Judge (RD), at 7. On November 9, 2023, Respondent filed a document titled "Notice of Appeal"¹ in response to the RD.

Having reviewed the entire record, the Agency adopts and hereby incorporates by reference the entirety of the ALJ's rulings, findings of fact, conclusions of law, and recommended sanction as found in the RD and summarizes and expands upon portions thereof herein.

Findings of Fact

Effective June 30, 2023, the Tennessee Department of Health revoked Respondent's Tennessee medical license. RD, at 5.² Further, effective July 6, 2023, the Virginia Department of Health Professions suspended Respondent's Virginia medical license. *Id*.³

According to Tennessee and Virginia online records, of which the Agency takes official notice, Respondent's Tennessee medical license remains revoked and Respondent's Virginia medical license remains suspended.⁴

² See also Government's Notice of Filing of Evidence and Motion for Summary Disposition, Exhibit (GX) 3, at 1.

⁴ 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, Respondent 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

⁶ This rule derives from the text of two provisions of the 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) (this section, formerly § 823(f), was redesignated as part of the Medical Marijuana and Cannabidiol Research Expansion Act, Pub. L. 117-215, 136 Stat. 2257 (2022)). Because Congress has clearly mandated that a practitioner possess state authority in order to be deemed a practitioner under the CSA, the 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, M.D., 71 FR 39130, 39131 (2006); Dominick A. Ricci, M.D., 58 FR 51104, 51105 (1993); Bobby Watts, M.D., 53 FR 11919, 11920 (1988); Frederick Marsh Blanton, 43 FR at 27,617.

¹The document blankly asserts that that Respondent appeals the RD without explaining the basis therefor or otherwise identifying his exceptions to the RD pursuant to 21 CFR 1316.66. *See* Respondent's Notice of Appeal.

³ See also GX 1, at 1–2.