Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is a notice that on July 7, 2026, Prof Compounding CTS of Ameri, 9901 South Wilcrest Drive, Houston, Texas 77099–5132, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Morphine	9300	II

The company plans to import Morphine Sulfate USP as a narcotic raw material for domestic bulk manufacturing and supply under the ASPR Strategic Active Pharmaceutical Ingredients Reserve (SAPIR) Program. The imported material will be used to support the strategic domestic supply of essential controlled substance Active Pharmaceutical Ingredients for national preparedness and resilience, including conversion into finished pharmaceutical products by qualified U.S. manufacturers as needed. No other activity for this drug code is authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Justin Wood,

Acting Deputy Assistant Administrator.

[FR Doc. 2026–17285 Filed 8–24–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1758]

Importer of Controlled Substances Application: Galephar Pharmaceutical Research Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Galephar Pharmaceutical Research Inc. has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before September 24, 2026. Such persons may also file a written request for a hearing on the application on or before September 24, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 20, 2026, Galephar Pharmaceutical Research Inc., #100 Carr 198, Industrial Park, Juncos, Puerto Rico 00777–3873, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Hydromorphone	9150	II
Morphine	9300	II

The company plans to import the listed controlled substances for analytical purposes only. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Justin Wood,

Acting Deputy Assistant Administrator.

[FR Doc. 2026–17284 Filed 8–24–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Catina Allen, N.P.; Decision and Order

On October 22, 2025, the Drug Enforcement Administration (DEA or Government) issued an Order to Show Cause (OSC) to Catina Allen, N.P., of Muncie, Indiana (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. MA8433233, alleging that Registrant is ''currently without authority to prescribe, administer, dispense, or otherwise handle controlled substances in the State of Indiana, the state in which [she is] registered with DEA.'' *Id.* at 2 (citing 21 U.S.C. 824(a)(3)).[1]

The OSC notified Registrant of her right to file a written request for hearing, and 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.* at 2–3 (citing 21 CFR 1301.43). Here, Registrant did not request a hearing, and the Agency finds her to be in default. RFAA, at 2–3.[2] ''A

[1] According to Agency records, Registrant's DEA registration expired on June 30, 2026. The fact that a registrant allows her registration to expire during the pendency of an OSC does not impact the Agency's jurisdiction or prerogative under the Controlled Substances Act (CSA) to adjudicate the OSC to finality. *Jeffrey D. Olsen, M.D.,* 84 FR 68474, 68476–68479 (2019).

[2] Based on the Government's submissions in its RFAA dated March 25, 2026, the Agency finds that

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