Controlled substance	Drug code	Schedule
Noroxymorphone	9668	II
Phenazocine	9715	II
Thiafentanil	9729	II
Piminodine	9730	II
Racemethorphan	9732	II
Racemorphan	9733	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II
Carfentanil	9743	II
Tapentadol	9780	II
Bezitramide	9800	II
Fentanyl	9801	II
Moramide-intermediate	9802	II

The company plans to import the listed controlled substances for distribution for analytical testing purposes. 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.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–07279 Filed 4–25–25; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1512]

Bulk Manufacturer of Controlled Substances Application: Pharmaron Manufacturing Services (US), LLC; Correction

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice; correction.

SUMMARY: The Drug Enforcement Administration (DEA) published a document in the **Federal Register** on March 26, 2025, concerning a notice of application for bulk manufacturer of Controlled Substances. As that document indicated the registrant's incorrect plans for the listed controlled substances.

SUPPLEMENTARY INFORMATION:

Correction

In the **Federal Register** (FR) of March 26, 2025, in FR Doc. 2025–05055 (90 FR 13782), on page 13782, Column 3, under **SUPPLEMENTARY INFORMATION** correction to the paragraph underneath controlled substance table should read as follows: The company plans to bulk manufacture Noroxymorphone (9668) as an intermediate product to be sold to its customers under the Contract Manufacture Organization (CMO) which the material will be shipped to, and converted to a non-controlled substance. Oxymorphone (9652) will be used as a starting material to be converted to Noroxymorphone. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–07281 Filed 4–25–25; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1518]

Bulk Manufacturer of Controlled Substances Application: Siemens Healthcare Diagnostics Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Siemens Healthcare Diagnostics, Inc. has applied to be registered as a bulk manufacturer 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 June 27, 2025. Such persons may also file a written request for a hearing on the application on or before June 27, 2025.

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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on January 15, 2025, Siemens Healthcare Diagnostics, Inc., 100 GBC Drive, Mailstop 108, Newark, Delaware 19702–2461, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Ecgonine	9180	II

The company plans to bulk manufacture the listed controlled substance in bulk to be used in the manufacture of the DEA exempt products. No other activity for this drug code is authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–07282 Filed 4–25–25; 8:45 am]

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