Background

On February 14, 2024, Cornerstone Chemical Company, Waggaman, Louisiana, filed petitions with the Commission and Commerce, alleging that an industry in the United States is materially injured or threatened with material injury by reason of subsidized imports of melamine from Germany, India, Qatar, and Trinidad and Tobago and LTFV imports of melamine from Germany, India, Japan, Netherlands, Qatar, and Trinidad and Tobago. Accordingly, effective February 14, 2024, the Commission instituted countervailing duty investigation Nos. 701–TA–706–709 and antidumping duty investigation Nos. 731–TA–1667–1672 (Preliminary).

Notice of the institution of the Commission's investigations and of a public conference to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** of February 21, 2024 (89 FR 13090). The Commission conducted its conference on March 6, 2024. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to §§ 703(a) and 733(a) of the Act (19 U.S.C. 1671b(a) and 1673b(a)). It completed and filed its determinations in these investigations on April 1, 2024. The views of the Commission are contained in USITC Publication 5503 (April 2024), entitled *Melamine from Germany, India, Japan, Netherlands, Qatar, and Trinidad and Tobago: Investigation Nos. 701 TA–706–709 and 731–TA–1667–1672 (Preliminary).*

By order of the Commission.

Issued: April 1, 2024.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2024–07181 Filed 4–3–24; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1350]

Bulk Manufacturer of Controlled Substances Application: Sterling Wisconsin, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Sterling Wisconsin, LLC 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 3, 2024. Such persons may also file a written request for a hearing on the application on or before June 3, 2024.

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 February 27, 2024, Sterling Wisconsin, LLC, W130N10497 Washington Drive, Germantown, Wisconsin 53022–4448, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Lysergic Acid Diethylamide.	7315	I
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Mescaline	7381	I
5-Methoxy-N-N-Dimethyltryptamine.	7431	I
Psilocybin	7437	I
Oliceridine	9245	II
Thebaine	9333	II
Alfentanil	9737	II

The company plans to bulk manufacture the listed controlled substances for commercial sale to its customers. In reference to drug codes 7350 (Marihuana Extract), 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

Marsha Ikner,

Acting Deputy Assistant Administrator.

[FR Doc. 2024–07110 Filed 4–3–24; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1348]

Bulk Manufacturer of Controlled Substances Application: Patheon Pharmaceuticals Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Patheon Pharmaceuticals 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 3, 2024. Such persons may also file a written request for a hearing on the application on or before June 3, 2024.

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 February 28, 2024, Patheon Pharmaceuticals Inc., 2110 East Galbraith Road, Cincinnati, Ohio 45237–1625, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):