in the United States at less than fair value (''LTFV'').[2] [3]

Background

The Commission instituted these investigations effective December 18, 2024, following receipt of petitions filed with the Commission and Commerce by the American Active Anode Material Producers, the members of which are Anovion Technologies, Sanborn, New York; Syrah Technologies LLC, Vidalia, Louisiana; NOVONIX Anode Materials LLC, Chattanooga, Tennessee; Epsilon Advanced Materials, Leland, North Carolina; and SKI US, Inc., Marietta, Georgia. The final phase of the investigations was scheduled by the Commission following notification of preliminary determinations by Commerce that imports of active anode material from China were subsidized within the meaning of section 703(b) of the Act (19 U.S.C. 1671b(b)) and sold at LTFV within the meaning of 733(b) of the Act (19 U.S.C. 1673b(b)). Notice of the scheduling of the final phase of the Commission's investigations and of a public hearing 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** on August 13, 2025 (90 FR 38993).[4] The Commission conducted its hearing on February 12, 2026. All persons who requested the opportunity were permitted to participate.

The Commission made these determinations pursuant to §§ 705(b) and 735(b) of the Act (19 U.S.C. 1671d(b) and 19 U.S.C. 1673d(b)). It completed and filed its determinations in these investigations on March 31, 2026. The views of the Commission are contained in USITC Publication 5719 (March 2026), entitled *Active Anode Material from China: Investigation Nos. 701–TA–752 and 731–TA–1730 (Final).*

By order of the Commission.

Issued: March 31, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026–06488 Filed 4–2–26; 8:45 am]

BILLING CODE 7020–02–P

[2] 90 FR 22465, May 28, 2025, and 90 FR 34423, July 22, 2025.

[3] Commissioner Jason E. Kearns dissenting.

[4] Due to the lapse in appropriations and ensuing cessation of Commission operations, the Commission tolled its schedule for this proceeding. The schedule was revised in a subsequent notice published in the **Federal Register** on December 11, 2025 (90 FR 57484).

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1699]

Importer of Controlled Substances Application: Blue Rabbit Veterinary LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Blue Rabbit Veterinary LLC 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 May 4, 2026. Such persons may also file a written request for a hearing on the application on or before May 4, 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 February 23, 2026, Blue Rabbit Veterinary LLC, 405 Heron Drive, Suite 300, Swedesboro, New Jersey 08085–1749, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Etorphine HCL	9059	II
Thiafentanil	9729	II

The company plans to import the listed controlled substances for sale to their customers (veterinarians). 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.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2026–06520 Filed 4–2–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1700]

Importer of Controlled Substances Application: Lipomed/LGC Standards

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Lipomed/LGC Standards 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 May 4, 2026. Such persons may also file a written request for a hearing on the application on or before May 4, 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