INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–737 and 731–TA–1712 (Final)]

Hexamine (Hexamethylenetetramine) From China

Determinations

On the basis of the record [1] developed in the subject investigations, the United States International Trade Commission (''Commission'') determines, pursuant to the Tariff Act of 1930 (''the Act''), that an industry in the United States is materially injured by reason of imports of hexamine from China, provided for in subheading 2933.69.50 of the Harmonized Tariff Schedule of the United States, that have been found by the U.S. Department of Commerce (''Commerce'') to be sold in the United States at less than fair value (''LTFV'') and subsidized by the government of China.[2] [3]

Background

The Commission instituted these investigations effective September 30, 2024, following receipt of petitions filed with the Commission and Commerce by Bakelite Synthetics (Atlanta, Georgia). The Commission scheduled the final phase of the investigations following notification of preliminary determinations by Commerce that imports of hexamine 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 May 22, 2025 (90 FR 21948 and as revised in 90 FR 31241, July 14, 2025). The Commission conducted its hearing on July 18, 2025. 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 September 3, 2025. The views of the Commission are contained in USITC Publication 5660 (September 2025), entitled *Hexamine (Hexamethylenetetramine) from China: Investigation Nos. 701–TA–737 and 731–TA–1712 (Final).*

By order of the Commission.

Issued: September 3, 2025.

Lisa Barton,

Secretary to the Commission.

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[1] The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

[2] 90 FR 33922 and 33923 (July 18, 2025).

[3] Commissioner David S. Johanson determines that that an industry in the United States is threatened with material injury by reason of imports of hexamine from China that have been found by Commerce to be sold in the United States at LTFV and subsidized by the government of China.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1561]

Importer of Controlled Substances Application: VA Cooperative Studies Program

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: VA Cooperative Studies Program 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 October 8, 2025. Such persons may also file a written request for a hearing on the application on or before October 8, 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. 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 May 9, 2025, VA Cooperative Studies Program, 2401 Centre Avenue South East, Albuquerque, New Mexico 87106, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Tetrahydrocannabinols ..	7370	I

The company plans to import finished dosage unit products containing the above listed controlled substances for research and clinical trial studies 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. 2025–17210 Filed 9–5–25; 8:45 am]

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NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

National Endowment for the Arts

Subject 30-Day Notice for the ''2025 Final Descriptive Report Update'' Proposed Collection; Comment Request

AGENCY: National Endowment for the Arts.

ACTION: Notice.

SUMMARY: The National Endowment for the Arts (NEA) has submitted the following public information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995: