provide the public and other Federal agencies with an opportunity to comment on new, proposed, revised, and continuing collections of information. This helps us assess the impact of our information collection requirements and minimize the public's reporting burden. It also helps the public understand our information collection requirements and provide the requested data in the desired format.

A **Federal Register** notice with a 60-day public comment period soliciting comments on this collection of information was published on March 13, 2026 (91 FR 12441). We did not receive any comments in response to that Notice.

As part of our continuing effort to reduce paperwork and respondent burdens, we invite the public and other Federal agencies to comment on new, proposed, revised, and continuing collections of information. This helps us assess the impact of our information collection requirements and minimize the public's reporting burden. It also helps the public understand our information collection requirements and provide the requested data in the desired format.

We are especially interested in public comments addressing the following:

(1) Whether or not the collection of information is necessary for the proper performance of the functions of the agency, including whether or not the information will have practical utility.

(2) The accuracy of our estimate of the burden for this collection of information, including the validity of the methodology and assumptions used.

(3) Ways to enhance the quality, utility, and clarity of the information to be collected.

(4) How might the agency minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of response.

Comments that you submit in response to this notice are a matter of public record. We will include or summarize each comment in our request to the Office of Management and Budget (OMB) to approve this Information Collection Request (ICR). Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

Abstract: The NPS is authorized by section 4 of the National Trails System Act (16 U.S.C. 1243) and Secretarial Order No. 3319 to administer the National Recreation Trail (NRT) program, which establishes National Water Trails (NWT) as a class of National Recreation Trails and directs that such trails collectively be considered in a National Water Trails System.

The NPS uses forms 10–1002: *Application for Designation as National Water Trail* and 10–1003: *Application for Designation as National Recreation Trail* to collect information NPS requires when submitting suitable trails or trail systems and water trails to the Secretary of the Interior for designation. The applications are evaluated for adherence to NRT requirements and criteria. NPS evaluation of an application is based on (1) the sufficiency of information provided on the application form and in supporting documentation, such as photographs, maps, and written landowner consents that accompany the form, and (2) successfully meeting the NRT requirements and criteria. Successful applications are forwarded to the Secretary of the Interior for approval.

Title of Collection: Application for Designation as National Recreation Trail or National Water Trail.

OMB Control Number: 1024–0283.

Form Number: NPS 10–1002: *Application for Designation as National Water Trail* and NPS 10–1003: *Application for Designation as National Recreation Trail.*

Type of Review: Reinstatement of a previously approved collection.

Description of Respondents: Private individuals; businesses; educational institutions; nonprofit organizations; state, tribal, and local governments; and Federal agency land units.

Total Estimated Number of Annual Respondents: 30.

Estimated Completion Time per Response: Varies by activity, 30 minutes to 8 hours.

Total Estimated Number of Annual Burden Hours: 130 hours.

Respondent's Obligation: Required to obtain or retain a benefit.

Frequency of Collection: On occasion.

Total Estimated Annual Non hour Burden Cost: None.

An agency may not conduct, or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

The authority for this action is the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*).

Phadrea Ponds,

Information Collection Clearance Officer, National Park Service.

[FR Doc. 2026–17379 Filed 8–25–26; 8:45 am]

BILLING CODE 4312–52–P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–802–804 and 731–TA–1799–1803 (Preliminary)]

Linear Hydraulic Cylinders From Canada, China, India, Mexico, and South Korea; Revised Schedule for the Subject Investigations

AGENCY: United States International Trade Commission.

ACTION: Notice.

DATES: August 21, 2026.

FOR FURTHER INFORMATION CONTACT: Stamen Borisson ((202) 205–3125), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its internet server (*https://www.usitc.gov*). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

SUPPLEMENTARY INFORMATION: On July 29, 2026, the Commission established a schedule for the conduct of the final phase of the subject investigations (91 FR 49442, August 4, 2026). Subsequently, the Department of Commerce (''Commerce'') extended the deadline for its initiation determination to September 8, 2026 (91 FR 53848, August 20, 2026). The Commission, therefore, is revising its schedule to conform with Commerce's new schedule.

The Commission must reach preliminary determinations within 25 days after the date on which the Commission receives notice from Commerce of initiation of the investigations, and the Commission's views must be transmitted to Commerce within five business days thereafter.

For further information concerning this proceeding, see the Commission's

notice cited above and the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A and C (19 CFR part 207).

Authority: These investigations are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 207.21 of the Commission's rules.

By order of the Commission.

Issued: August 21, 2026.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2026–17367 Filed 8–25–26; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1760]

Importer of Controlled Substances Application: Curia New York Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Curia New York 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 25, 2026. Such persons may also file a written request for a hearing on the application on or before September 25, 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 10, 2026, Curia New York Inc., 33 Riverside Avenue, Rensselaer, New York 12144, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
4-Anilino-N-phenethyl-4-piperidine (ANPP).	8333	II
Poppy Straw Concentrate.	9670	II

The company plans to import the listed controlled substances for bulk manufacturing into other controlled substances to be distributed to their customers. 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. 2026–17388 Filed 8–25–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1744]

Bulk Manufacturer of Controlled Substances Application: Chemtos, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Chemtos, 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 October 26, 2026. Such persons may also file a written request for a hearing on the application on or before October 26, 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 7, 2026, Chemtos, LLC, 16713 Picadilly Court, Round Rock, Texas 78664–8544, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Amineptine	1219	I
Mesocarb	1227	I
3-Fluoro-N-methylcathinone (3-FMC)	1233	I
Cathinone	1235	I
Methcathinone	1237	I
4-Fluoro-N-methylcathinone (4-FMC)	1238	I