

DEPARTMENT OF JUSTICE**Antitrust Division****Notice Pursuant to the National Cooperative Research and Production Act of 1993—Global Synchronizer Foundation**

Notice is hereby given that, on February 25, 2025, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), Global Synchronizer Foundation (“GSF”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Hong Kong Monetary Authority, Central Hong Kong, PEOPLE’S REPUBLIC OF CHINA; Copper Markets (Switzerland) AG, New York, NY; Goldman Sachs & Co. LLC, Jersey City, NJ; and Bank of New York Mellon Corporation, New York, NY, have been added as parties to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and GSF intends to file additional written notifications disclosing all changes in membership.

On September 18, 2024, GSF filed its original notification pursuant to section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to section 6(b) of the Act on October 11, 2024 (89 FR 82632).

Suzanne Morris,
Deputy Director Civil Enforcement Operations, Antitrust Division.

[FR Doc. 2025–03661 Filed 3–6–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE**Antitrust Division****Notice Pursuant to the National Cooperative Research and Production Act of 1993—Consortium for Rare Earth Technologies**

Notice is hereby given that, on February 12, 2025, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), Consortium for Rare Earth Technologies (“CREaTe”) has filed written notifications simultaneously with the

Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, AsterTech LLC, Dayton, OH, has been added as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and CREaTe intends to file additional written notifications disclosing all changes in membership.

On April 22, 2022, CREaTe filed its original notification pursuant to section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to section 6(b) of the Act on May 13, 2022 (87 FR 29384).

The last notification was filed with the Department on October 11, 2024. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on January 24, 2025 (90 FR 8153).

Suzanne Morris,
Deputy Director Civil Enforcement Operations, Antitrust Division.

[FR Doc. 2025–03653 Filed 3–6–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE**Drug Enforcement Administration**

[Docket No. DEA–1505]

Importer of Controlled Substances Application: LTS Therapy Systems, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: LTS Therapy Systems, 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 April 7, 2025. Such persons may also file a written request for a hearing on the application on or before April 7, 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 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on November 20, 2024, LTS Therapy Systems, LLC, 1685 Marthaler Lane, West Saint Paul, Minnesota 55118–3517, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Dimethyltryptamine	7435	I

The company plans to import the above controlled substance as bulk API for internal research, development, and analytical purposes only. No other activity for this drug code is 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. 2025–03648 Filed 3–6–25; 8:45 am]

BILLING CODE P