

(3) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

(a) The complainant is:

Vicor Corporation, 25 Frontage Road,
Andover, MA 01810

(b) The respondents are the following entities alleged to be in violation of section 337, and are the parties upon which the complaint is to be served:

Delta Electronics, Inc., 186, Ruey Kuang Road, Neihu Dist., Taipei 114501, Taiwan

Delta Electronics (Americas) Ltd., 46101 Fremont Blvd., Fremont, CA 94538

Delta Electronics (USA) Inc., 2925 E. Plano Pkwy., Plano, TX 75074.

Cyntec Co., Ltd., 2 R&D 2nd Rd., Science-Based Industry Park, Hsinchu 30076, Taiwan

Quanta Computer Inc., No. 211, Wenhua 2nd Rd., Guishan Dist., Taoyuan City 333, Taiwan

Quanta Cloud Technology Inc., 1F, No. 211 Wenhua 2nd Rd., Guishan Dist., Taoyuan City 33377, Taiwan

Quanta Cloud Technology USA LLC, 1010 Rincon Circle, San Jose, CA 95131

Quanta Computer USA Inc., 45630 Northport Loop East, Fremont, CA 94538

Hon Hai Precision Industry Co. Ltd. (d/b/a, Foxconn Technology Group), No. 2, Zihyou St., Tucheng Dist. New Taipei City 236, Taiwan

Foxconn Industrial internet Co. Ltd., 2F C1 Foxconn Technology Park, 2 Donghuan 2 Road Longhua, Shenzhen, 518109 China

FII USA Inc. (a/k/a Foxconn Industrial, Internet USA Inc.), 611 East Wisconsin Ave., Milwaukee, WI 53202,

Ingrasys Technology Inc., 5F., No. 1188, Nanqing Rd., Luzhu Dist., Taoyuan City, Taiwan

Ingrasys Technology USA Inc., 2025 Gateway Place, Ste. 190, San Jose, CA 95110

(c) The Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street SW, Suite 401, Washington, DC 20436; and

(4) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with section 210.13 of the Commission's Rules of Practice and

Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), as amended in 85 FR 15798 (March 19, 2020), such responses will be considered by the Commission if received not later than 20 days after the date of service by the complainant of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: August 14, 2023.

Lisa Barton,

Secretary to the Commission.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1234]

Importer of Controlled Substances Application: Curium US LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Curium US 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 September 18, 2023. Such persons may also file a written request for a hearing on the application on or before September 18, 2023.

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 July 12, 2023, Curium US LLC, 2703 Wagner Place, Maryland Heights, Missouri 63043-3421, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Ecgonine	9180	II

The company plans to import small quantities of a derivative form of the listed controlled substance to be used for manufacturing purposes. 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.

Claude Redd,

Acting Deputy Assistant Administrator.

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