

granted a registration as an importer for schedule II controlled substances to the above listed companies.

Dated: August 16, 2019.

Neil D. Doherty,

Acting Assistant Administrator.

[FR Doc. 2019-18686 Filed 8-28-19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: The registrants listed below have applied for and been granted registration by the Drug Enforcement

Administration (DEA) as importers of schedule I and II controlled substances.

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as an importers of various basic classes of schedule I and II controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for a hearing were submitted for these notices.

Companies	FR docket	Published
Unither Manufacturing, LLC	84 FR 13961	April 8, 2019.
Shertech Laboratories, LLC	84 FR 26446	June 6, 2019.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable various basic classes of schedule I and II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance

with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule II controlled substances to the above listed companies.

Dated: August 16, 2019.

Neil D. Doherty,

Acting Assistant Administrator.

[FR Doc. 2019-18688 Filed 8-28-19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: The registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of schedule I and II controlled substances.

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as an importers of various basic classes of schedule I and II controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for a hearing were submitted for these notices.

Companies	FR docket	Published
United States Pharmacopeial Convention	84 FR 23582	May 22, 2019.
Bellwyck Clinical Services	84 FR 31622	July 2, 2019.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable various basic classes of schedule I and II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has

granted a registration as an importer for schedule I and II controlled substances to the above listed companies.

Dated: August 16, 2019.

Neil D. Doherty,

Acting Assistant Administrator.

[FR Doc. 2019-18685 Filed 8-28-19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: Cerilliant Corporation

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 28, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 15, 2019, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665-2402 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances: