DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

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

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

Company	FR docket	Published
Noramco Inc	83 FR 53107	October 19, 2018.
Catalent CTS, LLC	83 FR 54613	October 30, 2018.
United States Pharmacopeial Convention	83 FR 54611	October 30, 2018.
Fisher Clinical Services, Inc	83 FR 54612	October 30, 2018.
Cambrex High Point, Inc	83 FR 54610	October 30, 2018.
Sharp (Bethlehem), LLC	83 FR 54612	October 30, 2018.

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 basic classes of schedule I or 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 company's maintenance of effective controls against diversion by inspecting and testing each of the 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 or II controlled substances to the above listed companies.

Dated: December 21, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01519 Filed 2–6–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration

ACTION: Notice of registration.

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

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as bulk manufacturers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted for these notices.

Company	FR Docket	Published
AMPAC Fine Chemicals Virginia, LLC	83 FR 48334	September 24, 2018.
AMPAC Fine Chemicals, LLC	83 FR 49578	October 2, 2018.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of this registrant to manufacture the applicable basic classes of 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 the company's maintenance of effective controls against diversion by inspecting and testing their physical security systems, verifying their compliance with state and local laws, and reviewing each of the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed companies.

Dated: December 21, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01525 Filed 2–6–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 2018–43]

Kurt L. Pflieger, M.D.; Order Dismissing Order To Show Cause

On July 12, 2018, the Assistant Administrator, Diversion Control Division, Drug Enforcement Administration (hereinafter, DEA or Government), issued an Order to Show Cause to Kurt L. Pflieger, M.D. (hereinafter, Respondent), of Rockwall, Texas. Order to Show Cause (hereinafter, OSC), at 1. The OSC proposes the revocation of Respondent's Certificate of Registration on the ground that he does not have authority to handle controlled substances in the State of Texas, the State in which he is registered with the DEA. *Id.*

After the Administrative Law Judge (hereinafter, ALJ) certified and transmitted the record to me along with his Recommended Decision, the Government submitted a ''Motion to Dismiss Order to Show Cause'' (hereinafter, Motion). According to the Motion, the Texas Medical Board held