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

a hearing after the ALJ granted the Government's Motion for Summary Disposition and recommended revocation of Respondent's registration. Motion, at 1. The Motion also states that, as a result of the hearing, the Texas Medical Board suspended Respondent's medical license, ordered that the suspension be stayed, and placed Respondent on five years' probation under various terms and conditions. *Id.*

The Motion concludes by stating that, ''Since . . . [Respondent] now has state authority to handle controlled substances, the DEA respectfully request[s] the Administrator to dismiss the pending Order to Show Cause and recommended ruling from the Administrative Law Judge.'' *Id.* at 2. Accordingly, I shall dismiss the Order to Show Cause.

Order

Pursuant to 28 CFR 0.100(b) and the authority thus vested in me by 21 U.S.C. 823(f) and 824(a), I order that the Order to Show Cause issued to Kurt L. Pflieger, M.D. be, and it hereby is, dismissed. This Order is effective March 11, 2019.

Dated: January 17, 2019.

Uttam Dhillon,

Acting Administrator.

[FR Doc. 2019–01528 Filed 2–6–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Chattem Chemicals

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 March 11, 2019. Such persons may also file a written request for a hearing on the application on or before March 11, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also 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.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (''Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on October 31, 2018, Chattem Chemicals Inc., 3801 Saint Elmo Avenue, Chattanooga, Tennessee 37409–1237 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Methamphetamine	1105	II
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Phenylacetone	8501	II
Opium, raw	9600	II
Poppy Straw Concentrate	9670	II
Tapentadol	9780	II

The company plans to import the listed controlled substances to manufacture bulk controlled substances for sale to its customers.

The company plans to import an intermediate of tapentadol (9780), to bulk manufacture tapentadol for distribution to its customers.

Dated: December 21, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01512 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: Registrants listed below have applied for and been granted a registration by the Drug Enforcement Administration (DEA) as bulk manufacturers 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 are listed in the table below. No comments or objections were submitted for these notices.

Company	FR Docket	Published
National Center for Natural Products Research NIDA MPROJECT	83 FR 48334	August 31, 2018.
Halo Pharmaceutical, Inc	83 FR 48334	September 24, 2018.
Nanosyn, Inc	83 FR 48867	September 27, 2018.