Dated: September 23, 2019.

Thomas W. Prevoznik,

Acting Assistant Administrator, Deputy Assistant Administrator.

[FR Doc. 2019–21320 Filed 9–30–19; 8:45 am]

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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 have applied for and been granted a registration by the Drug Enforcement Administration (DEA) as a bulk manufacturer of a various classes of schedule I and II controlled substances.

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as a bulk manufacturer of various classes of scheduled I and II controlled substances. Information on a previously published notices is listed below. No comments or objections were submitted for these notices.

Company	FR docket	Published
Absolute Standards, Inc	84 FR 31620	July 2, 2019.
Pisgah Laboratories, Inc	84 FR 31622	July 2, 2019.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants 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 each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing 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: September 24, 2019.

Thomas W. Prevoznik,

Acting Assistant Administrator, Deputy Assistant Administrator.

[FR Doc. 2019–21312 Filed 9–30–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration; Siemens Healthcare Diagnostics, Inc.

ACTION: Notice of registration.

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

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as a bulk manufacturer of a basic class of schedule I and II controlled substances. Information on previously published notices is listed below. No comments or objections were submitted for the notice.

Company	FR Docket	Published
Siegfried USA, LLC	84 FR 7129	March 1, 2019.
Patheon Pharmaceuticals, Inc	84 FR 8114	March 6, 2019.
S & B Pharma Inc	84 FR 8116	March 6, 2019.
Siemens Healthcare Diagnostics, Inc	84 FR 10534	March 21, 2019.
Synthcon, LLC	84 FR 13962	April 8, 2019.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants 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 each of the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each of the company's compliance with state and local laws, and reviewing each 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: September 23, 2019.

Thomas W. Prevoznik,

Acting Assistant Administrator, Deputy Assistant Administrator.

[FR Doc. 2019–21313 Filed 9–30–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Noramco, Inc.

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 31, 2019. Such persons may also file a written request for a hearing on the application on or before October 31, 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 a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a 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. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 18, 2019, Noramco Inc., 1550 Olympic Drive, Athens, Georgia 30601 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Nabilone	7379	II
Phenylacetone	8501	II
Thebaine	9333	II
Poppy Straw Concentrate	9670	II
Tapentadol	9780	II

The company plans to import phenylacetone (8501), and poppy straw concentrate (9670) to bulk manufacture other controlled substances for distribution to its customers. The company plans to import an intermediate form of tapentadol (9780) to bulk manufacture tapentadol (9780) for distribution to its customers.

The company plans to import impurities of buprenorphine that have been determined by DEA to be captured under drug code (9333) thebaine.

In reference to drug codes 7360 and 7370, the company plans to import a synthetic cannabidiol and a synthetic tetrahydrocannabinol. No other activity for these drug codes is authorized for this registration. Placement of these drug codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances. 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 FDA approved or non-approved finished dosage forms for commercial sale.

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: September 23, 2019

Thomas W. Prevoznik,

Acting Assistant Administrator, Deputy Assistant Administrator.

[FR Doc. 2019–21321 Filed 9–30–19; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: CreaGen Inc.

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 December 2, 2019.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on August 14, 2019, CreaGen Inc., 299 Washington Street, Unit A, Woburn, Massachusetts 01801–2795 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Dated: September 23, 2019.

Thomas W. Prevoznik,

Acting Assistant Administrator Deputy Assistant Administrator.

[FR Doc. 2019–21319 Filed 9–30–19; 8:45 am]

BILLING CODE 4410–09–P

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
Restek Corporation	84 FR 35691	July 24, 2019.
AMRI Rensselaer, Inc	84 FR 35692	July 24, 2019.
Alcami Carolinas Corporation	84 FR 36941	July 30, 2019.
Cambrex Charles City	84 FR 36945	July 30, 2019.
Chattem Chemicals, Inc	83 FR 39129	August 8, 2019.