the public interest in this investigation, as appropriate, and provide the Commission with findings of fact and a recommended determination on this issue, which shall be limited to the statutory public interest factors set forth in 19 U.S.C. 1337(d)(1), (f)(1), (g)(1);

(4) 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:

INVT SPE LLC, One Market Plaza, Spear Tower, 42nd Floor, San Francisco, CA 94105.

(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:

Apple Inc., 1 Infinite Loop, Cupertino, CA 95014.

HTC Corporation, 23 Xinghua Road, Taoyuan City, Taoyuan County 330, Taiwan.

HTC America, Inc., 308 Occidental Ave. S, Suite 300, Seattle, WA 98104.

ZTE Corporation, ZTE Plaza, Keji Road South, Hi-Tech Industrial Park, Nanshan District, Guangdong Province, 518057, China.

ZTE (USA) Inc., 2425 N Central Expressway, Suite 800, Richardson, TX 75080.

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

(5) 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), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission 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: October 16, 2018.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2018–22869 Filed 10–18–18; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Consortium for Execution of Rendezvous and Servicing Operations

Notice is hereby given that, on September 10, 2018, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), Consortium for Execution of Rendezvous and Servicing Operations (''CONFERS'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the identities of the parties to the venture and (2) the nature and objectives of the venture. The notifications were filed for the purpose of invoking the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the identities of the parties to the venture are: Altius Space Machines, Inc., Broomfield, CO; Analytical Graphics, Inc., Exton, PA; Astroscale PTE. LTD., Tokyo, JAPAN; Ball Aerospace and Technology Corp., Boulder, CO; Chandah Space Technologies, Houston, TX; Hoffer Flow Controls, Inc. Elizabeth City, NC; iBOSS gmbH, Aachen, GERMANY; MacDonald, Dettwiler and Associates, Inc., Brampton, Ontario, CANADA; Thales Alenia Space, Courbevioe, FRANCE; The Aerospace Corporation, El Segundo, CA; and XL Catlin, LLC, New York, NY.

The general area of CONFERS' planned activity is to establish an independent, self-sustaining industry forum to advocate and promote on-orbit satellite maintenance, servicing, and rendezvous operations by collaborating to research, develop, and publish voluntary, consensus technical and safety standards, and engaging with governments on policy and oversight of satellite servicing activities. To fulfill its mission, CONFERS will recruit a broad array of members from satellite operators, service providers, insurers and underwriters, and engage other stakeholders from industry, academia, and governments. The process will be fully collaborative and will include dedicated outreach activities to engage the global commercial satellite community. The members of the CONFERS believe that future standards should be based on a set of guiding principles that will help establish responsible norms of behavior.

Suzanne Morris,

Chief, Premerger and Division Statistics Unit, Antitrust Division.

[FR Doc. 2018–22826 Filed 10–18–18; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration

ACTION: Notice of registration.

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

SUPPLEMENTARY INFORMATION: The company listed below applied to be registered as a bulk manufacturer of various basic classes of controlled substances. Information on a previously published notice is listed in the table below. No comments or objections were submitted for this notice.

Company	FR Docket	Published
Siegfried USA, LLC	83 FR 32905 ..	July 16, 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 the company's physical security systems, verifying the 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 company.

Dated: October 10, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–22829 Filed 10–18–18; 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: 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 are listed in the table below. No comments or objections were submitted and no requests for hearings were submitted for these notices.

Company	FR Docket	Published
Chattem Chemicals, Inc	83 FR 39129 ..	August 8, 2018.
Myoderm	83 FR 39130 ..	August 8, 2018.
Mylan Pharmaceuticals, Inc	83 FR 22517 ..	May 15, 2018.
Anderson Brecon, Inc	83 FR 39128 ..	August 8, 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 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: October 10, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–22830 Filed 10–18–18; 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 November 19, 2018. Such persons may also file a written request for a hearing on the application on or before November 19, 2018.

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 hearings should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 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: 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 July 4, 2018, 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