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 June 5, 2017, Fisher Clinical Services, Inc., 700 A–C Nestle Way, Breinigsville, Pennsylvania 18031–1522 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Methylphenidate	1724	II
Levorphanol	9220	II
Noroxymorphone	9668	II
Tapentadol	9780	II

The company plans to import the listed controlled substances in finished dosage form for testing, and clinical trials purposes only. This authorization does not extend to the import of a finished Food and Drug Administration (FDA) approved or non-approved dosage form for commercial distribution in the United States.

Dated: March 15, 2018.

Susan A. Gibson,

Deputy Assistant Administrator.

[FR Doc. 2018–06321 Filed 3–28–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Lannett Company, 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 April 30, 2018. Such persons may also file a written request for a hearing on the application on or before April 30, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 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/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 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 February 24, 2014, Lannett Company, Inc., 9001 Torresdale Avenue, Philadelphia, Pennsylvania 19136 applied to be registered as an importer of tetrahydrocannabinols (7370), a basic class of controlled substance listed in schedule I.

The company plans to import the finished dosage forms to support their abbreviated new drug application (ANDA) submission to the U.S. Food and Drug Administration (FDA). No other activity for this drug code is authorized for this registration.

Dated: March 15, 2018.

Susan A. Gibson,

Deputy Assistant Administrator.

[FR Doc. 2018–06313 Filed 3–28–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Novitium Pharma, LLC

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 April 30, 2018. Such persons may also file a written request for a hearing on the application on or before April 30, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 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 request for hearing 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/DRW, 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 January 8, 2018, Novitium Pharma, LLC., 70 Lake Drive, East Windsor, NJ 08520 applied to be registered as an importer of the Schedule II controlled substance Levorphanol (9220).

The company plans to import the controlled substance to develop the manufacturing process for a drug product that will in turn be used to produce a tablet equivalent to the current brand product.

Dated: March 15, 2018.

Susan A. Gibson,

Deputy Assistant Administrator.

[FR Doc. 2018–06318 Filed 3–28–18; 8:45 am]

BILLING CODE 4410–09–P