

interest, and bonding filed in response to the Commission Notice, the Commission has determined that the appropriate form of relief in this investigation is: (1) An LEO prohibiting the unlicensed entry of infringing road-milling machines and components thereof covered by one or more of claim 29 of the '309 patent or claims 2, 5, 16, or 23 of the '530 patent that are manufactured abroad for or on behalf of, or imported by or on behalf of, any of the Respondents or any of their affiliated companies, parents, subsidiaries, or other related business entities, or their successors or assigns; and (2) a CDO directed against Caterpillar Paving Products, Inc. and Caterpillar Inc., and their affiliated companies, parents, subsidiaries, or other related business entities, or their successors or assigns.

The Commission has further determined that the public interest factors enumerated in subsections (d)(1) and (f)(1) (19 U.S.C. 1337(d)(1), (f)(1)) do not preclude issuance of the above-referenced remedial orders. Additionally, the Commission has determined to impose a bond of fourteen (14) percent of entered value of the covered products during the period of Presidential review (19 U.S.C. 1337(j)). The investigation is terminated.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, and in Part 210 of the Commission's Rules of Practice and Procedure, 19 CFR part 210.

By order of the Commission.

Issued: July 18, 2019.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2019-15689 Filed 7-23-19; 8:45 am]

BILLING CODE 7020-02-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Cardinal Health

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 August 23, 2019. Such persons may also file a written request for a hearing on the application on or before August 23, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 31, 2019, Cardinal Health, 15 Ingram Boulevard, LaVergne, Tennessee applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Secobarbital	2315	II

The company plans to only distribute to licensed registrants for the purpose of medical use.

Dated: July 15, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019-15728 Filed 7-23-19; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Restek Corporation

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 August 23, 2019. Such persons may also file a written request for a hearing on the application on or before August 23, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.34(a), this is notice that on April 12, 2019, Restek Corporation, 110 Benner Circle, Bellefonte, Pennsylvania 16823-8433 applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols	7370	I

The company plans to import the listed controlled substance in bulk for manufacture of analytical reference material which, in its final form, is an exempted product.

Dated: July 15, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019-15738 Filed 7-23-19; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: AMRI Rensselaer, 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 August 23, 2019. Such persons may also file a written request for a hearing on the application on or before August 23, 2019.

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

22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.34(a), this is notice that on May 7, 2019, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144 applied to be registered as an importer of the following basic class of controlled substance:

Table with 3 columns: Controlled substance, Drug code, Schedule. Row: Poppy Straw Concentrate, 9670, II

The company plans to import the listed controlled substance to manufacture a bulk controlled substance for distribution to its customers.

Dated: July 15, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019-15737 Filed 7-23-19; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

[Docket No. DEA-392]

Drug Enforcement Administration Bulk Manufacturer of Controlled Substances Application: IsoSciences, 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 September 23, 2019.

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

SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.33(a), this is notice that on May 21, 2019, IsoSciences, LLC, 340 Mathers Road, Ambler, Pennsylvania 19002-3420 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Table with 3 columns: Controlled substance, Drug code, Schedule. Lists various substances like Cathinone, Methcathinone, Lysergic acid diethylamide, etc.