DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Research Triangle Institute

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 30, 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 May 23, 2019, Research Triangle Institute, 3040 East Cornwallis Road, Hermann Building, Room 106, Research Triangle Park, North Carolina 27709, applied to be registered as a bulk manufacturer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols	7370	I

The company will manufacture via synthesis, Tetrahydrocannabinols (7370) for use by customers as analytical reference standards.

Dated: July 16, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019–16175 Filed 7–29–19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: AMPAC Fine Chemicals 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 30, 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 March 18, 2019, AMPAC Fine Chemicals LLC, Highway 50 and Hazel Avenue, Building 05001, Rancho Cordova, California 95670 applied to be registered as bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Methylphenidate	1724	II
Levomethorphan	9210	II
Levorphanol	9220	II
Thebaine	9333	II
Tapentadol	9780	II
Remifentanil	9739	II

The company plans to manufacture the listed controlled substances for distribution to its customers.

Dated: July 16, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019–16177 Filed 7–29–19; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Southern Ohio Correctional Facility

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration or the proposed authorization to import on or before August 29, 2019. Such persons may also file a written request for a hearing on the application for registration and for authorization to import on or before August 29, 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 hearing a 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:

Pursuant to 21 U.S.C. 958(i), prior to issuing a regulation under 21 U.S.C. 952(a)(2)(B) authorizing the importation of a controlled substance in schedule I or II, DEA is required to provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing. Additionally, pursuant to 21 CFR 1301.34(a), DEA shall, upon the filing of an application for registration to import a controlled substance in schedule I or II under 21 U.S.C. 952(a)(2)(B), provide notice and the opportunity to request a hearing to manufacturers holding registrations for the bulk manufacture of the substance and to applicants for such registrations.