

Controlled substance	Drug code	Schedule
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Fentanyl .....	9801	II

The company plans to manufacture the above-listed controlled substances in bulk for distribution to its customers. In reference to drug codes 7360 marihuana and 7370 tetrahydrocannabinols the company plans to bulk manufacture both as synthetic substances. No other activity for these drug codes is authorized for this registration.

Dated: December 19, 2016.

**Louis J. Milione,**  
Assistant Administrator.

[FR Doc. 2016-31279 Filed 12-27-16; 8:45 am]

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**DEPARTMENT OF JUSTICE**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Application: Cambridge Isotope Laboratories**

**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 in accordance with 21 CFR 1301.33(a) on or before February 27, 2017.

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

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated her 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

September 1, 2016, Cambridge Isotope Laboratories, Inc., 50 Frontage Road, Andover, Massachusetts 01810 applied to be registered as a bulk manufacturer of morphine (9300), a basic class of controlled substance listed in schedule II:

The company plans to utilize small quantities of the listed controlled substance for use in product development of analytical reference standards, for distribution to its customers.

Dated: December 19, 2016.

**Louis J. Milione,**  
Assistant Administrator.

[FR Doc. 2016-31271 Filed 12-27-16; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Application: Wildlife Laboratories, 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 in accordance with 21 CFR 1301.34(a) on or before January 27, 2017. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before January 27, 2017.

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

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated her

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 September 20, 2016, Wildlife Laboratories, Inc., 1230 W. Ash Street, Suite D, Windsor, Colorado 80550-8055 applied to be registered as an importer of the following basic class of controlled substances.

Controlled substance	Drug code	Schedule
Thiafentanil .....	9729	II

The company plans to import the listed controlled substance for sale to its customers.

Dated: December 20, 2016.

**Louis J. Milione,**  
Assistant Administrator.

[FR Doc. 2016-31272 Filed 12-27-16; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Application: Cayman Chemical Company**

**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 in accordance with 21 CFR 1301.33(a) on or before February 27, 2017.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement