Billing Code 4410-09-P

DEPARTMENT OF JUSTICE

Bulk Manufacturer of Controlled Substances Application: Cambridge Isotope Laboratories

[Docket No. DEA-392]

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 [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette 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.

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