Dated: August 20, 2013.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[Docket No. DEA-2013-0067; FR Doc. 2013–20724 Filed 8–23–13; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances; Notice of Application; Cambridge Isotope Lab

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 01, 2013, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, made application by renewal to the Drug Enforcement Administration (DEA) 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 in the preparation of analytical standards.

Any other such applicant, and any person who is presently registered with DEA to manufacture such a substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than October 25, 2013.


Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[Docket No. DEA-2013-0068; FR Doc. 2013–20725 Filed 8–23–13; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances; Notice of Application; Morton Grove Pharmaceuticals

By Notice dated March 12, 2013, and published in the Federal Register on March 20, 2013, 78 FR 17231, Morton Grove Pharmaceuticals, 6451 Main Street, Morton Grove, Illinois 60053–2633, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Gamma Hydroxybutyric Acid (2010), a basic class of controlled substance listed in schedule I.

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

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Morton Grove Pharmaceuticals to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Morton Grove Pharmaceuticals to ensure that the company’s registration is consistent with the public interest.

The investigation has included inspection and testing of the company’s physical security systems; verification of the company’s compliance with state and local laws; and a review of the company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.


Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[Docket No. DEA-2013-0070; FR Doc. 2013–20761 Filed 8–23–13; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Tin T. Win, M.D., Dismissal of Proceeding

On February 27, 2013, I, the Administrator of the Drug Enforcement Administration, issued an Order to Show Cause and Immediate Suspension of Registration to Tin T. Win, M.D. (hereinafter, Registrant), of Lake Havasu, Arizona. GX 10, at 1. Among various charges, the Order alleged that Registrant issued numerous controlled substance prescriptions after the Arizona Medical Board had prohibited her “from prescribing controlled substances” and thus violated both the Board’s order and federal law. Id. at 1–3 (citing Ariz. Rev. Stat. § 32–1401(27)(r); 21 U.S.C. 841). The Order also notified Registrant of her right to either request a hearing on the allegations or submit a written statement of position in lieu of a hearing within thirty (30) days of her receipt of the Order, the procedure for electing either option, and the consequence of failing to elect either option.

On March 6, 2013, the Order was personally served on Registrant by a DEA Special Agent and a Diversion Investigator. See GX 11. On May 20, 2013, the Government filed a Motion for Final Agency Action, which sought the revocation of Registrant’s...