Billing Code 4410-09-M

DEPARTMENT OF JUSTICE

DRUG ENFORCEMENT ADMINISTRATION

Manufacturer of Controlled Substances

Notice of Application

Austin Pharma, Llc.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 28, 2013, Austin Pharma, LLC., 811 Paloma Drive, Suite C, Round Rock, Texas 78665-2402, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Marihuana (7360), a basic class of controlled substance listed in schedule I.

The company plans to manufacture bulk active pharmaceutical ingredients (APIs) for distribution to its customers.

In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol as a synthetic intermediate. No other activity for this drug code is authorized for this registration.

Any other such applicant, and any person who is presently registered with DEA to manufacture such 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 [INSERT DATE 60 DAYS FROM DATE OF PUBLICATION].

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

DATED: May 14, 2013

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