



This document is scheduled to be published in the Federal Register on 01/26/2015 and available online at <http://federalregister.gov/a/2015-01312>, and on [FDsys.gov](http://FDsys.gov)

## **DEPARTMENT OF JUSTICE**

### **Drug Enforcement Administration**

**[Docket No. DEA-392]**

#### **Bulk Manufacturer of Controlled Substances Application: Euticals, 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 1301.33(a) on or before [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152.

#### **SUPPLEMENTARY INFORMATION:**

The Attorney General has delegated his 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, and dispensers of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (“Deputy Assistant Administrator”) pursuant to section 7 of 28 CFR pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.33(a), this is notice that on May 29, 2014, Euticals, Inc., 2460 W. Bennett Street, Springfield, Missouri 65807-1229, applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

<b><u>Controlled Substance</u></b>	<b><u>Schedule</u></b>
Gamma Hydroxybutyric Acid (2010)	I
Amphetamine (1100)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Phenylacetone (8501)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Oripavine (9330)	II
Tapentadol (9780)	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers.

In reference to Amphetamine (1100), the company plans to acquire the listed controlled substance in bulk from a domestic source in order to manufacture other controlled substances in bulk for distribution to its customers.

Dated: January 9, 2015.

Joseph T. Rannazzisi,  
*Deputy Assistant Administrator.*