

Dated: July 20, 2017.  
**Demetra Ashley,**  
*Acting Assistant Administrator.*  
 [FR Doc. 2017-15693 Filed 7-25-17; 8:45 am]  
**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**  
**Drug Enforcement Administration**  
**[Docket No. DEA-392]**

**Bulk Manufacturer of Controlled Substances Application: AMRI Rensselaer, 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 September 25, 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 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, 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 1301.33(a), this is notice that on March 30, 2017, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I
Amphetamine .....	1100	II
Lisdexamfetamine .....	1205	II
Methylphenidate .....	1724	II
Pentobarbital .....	2270	II
4-Anilino-N-phenethyl-4-piperidine (ANPP) .....	8333	II
Codeine .....	9050	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Hydrocodone .....	9193	II
Meperidine .....	9230	II
Morphine .....	9300	II
Fentanyl .....	9801	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers.

In reference to drug codes 7360 (marihuana) and 7370 (THC), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: July 19, 2017.  
**Demetra Ashley,**  
*Acting Assistant Administrator.*  
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**DEPARTMENT OF JUSTICE**  
**Drug Enforcement Administration**  
**[Docket No. DEA-392]**

**Bulk Manufacturer of Controlled Substances Registration**

**ACTION:** Notice of registration.

**SUMMARY:** Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as bulk manufacturers of various classes of controlled substances.

**SUPPLEMENTARY INFORMATION:** The companies listed below applied to be registered as manufacturers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted for these notices.

Company	FR docket	Published
Cedarburg Pharmaceuticals .....	82 FR 19083 .....	April 25, 2017.
Siegfried USA, LLC .....	82 FR 19084 .....	April 25, 2017.
Sigma Aldrich Research Biochemicals, Inc .....	82 FR 19085 .....	April 25, 2017.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international

treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company’s maintenance of effective controls against diversion by inspecting and testing each company’s physical security systems, verifying each

company’s compliance with state and local laws, and reviewing each company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a