

**SUMMARY:** Siegfried USA, LLC applied to be registered as an importer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Siegfried USA, LLC registration as an importer of those controlled substances.

**SUPPLEMENTARY INFORMATION:** By notice dated January 11, 2016, and published in the **Federal Register** on January 19, 2016, 81 FR 2910, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070 applied to be registered as an importer of certain basic classes of controlled substances. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417, (January 25, 2007).

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Siegfried USA, LLC to import the 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 the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above-named company is granted registration as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Opium, raw (9600) .....	II
Poppy Straw Concentrate (9670)	II

The company plans to import the listed controlled substances to manufacture bulk active pharmaceutical ingredients (API) for distribution to its customer.

Dated: April 28, 2016.

**Louis J. Milione,**

*Deputy Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Manufacturer of Controlled Substances Registration: AMRI Rensselaer, Inc.**

**ACTION:** Notice of registration.

**SUMMARY:** AMRI Rensselaer, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants AMRI Rensselaer, Inc. registration as a manufacturer of those controlled substances.

**SUPPLEMENTARY INFORMATION:** By notice dated November 30, 2015, and published in the **Federal Register** on December 8, 2015, 80 FR 76312, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of AMRI Rensselaer, Inc. to manufacture the 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 the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing 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 following basic classes of controlled substances:

Controlled substance	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
Meperidine (9230) .....	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 activity for these drug codes are authorized for this registration.

Dated: April 28, 2016.

**Louis J. Milione,**

*Deputy Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Manufacturer of Controlled Substances Registration: Johnson Matthey, Inc.**

**ACTION:** Notice of registration.

**SUMMARY:** Johnson Matthey, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Johnson Matthey, Inc. registration as a manufacturer of those controlled substances.

**SUPPLEMENTARY INFORMATION:** By notice dated December 9, 2015, and published in the **Federal Register** on December 17, 2015, 80 FR 78765, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066-1742 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Johnson Matthey, Inc. to manufacture the 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 the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing 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 following basic classes of controlled substances: