

Dated: January 11, 2016.  
**Louis J. Milione,**  
*Deputy Assistant Administrator.*  
 [FR Doc. 2016-00789 Filed 1-15-16; 8:45 am]  
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**DEPARTMENT OF JUSTICE**

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

**Manufacturer of Controlled Substances Registration: American Radiolabeled Chemicals, Inc.**

**ACTION:** Notice of registration.

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

**SUPPLEMENTARY INFORMATION:** By notice dated October 2, 2015, and published in the **Federal Register** on October 13, 2015, 80 FR 61469, American Radiolabeled Chemicals, Inc., 101 Arc Drive, Saint Louis, Missouri 63146 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 American Radiolabeled Chemicals, 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
Gamma Hydroxybutyric Acid (2010).	I
ibogaine (7260) .....	I
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370) .....	I

Controlled substance	Schedule
Dimethyltryptamine (7435) .....	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470).	I
Dihydromorphine (9145) .....	I
Heroin (9200) .....	I
Normorphine (9313) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Amobarbital (2125) .....	II
Phencyclidine (7471) .....	II
Phenylacetone (8501) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Ecgonine (9180) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Metazocine (9240) .....	II
Methadone (9250) .....	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300) .....	II
Oripavine (9330) .....	II
Thebaine (9333) .....	II
Oxymorphone (9652) .....	II
Phenazocine (9715) .....	II
Carfentanil (9743) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture small quantities of the listed controlled substances as radiolabeled compounds for biochemical research.

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**DEPARTMENT OF JUSTICE**

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

**Importer of Controlled Substances Application: Siegfried USA, LLC**

**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 1301.34(a) on or before February 18, 2016. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before February 18, 2016.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrisette Drive, Springfield, Virginia

22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417, (January 25, 2007).

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated her 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 Deputy Assistant Administrator of the DEA Office of Diversion Control ("Deputy Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on November 12, 2015, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070 applied to be registered 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.

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**DEPARTMENT OF JUSTICE**

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

**Manufacturer of Controlled Substances Registration: Apertus Pharmaceuticals**

**ACTION:** Notice of registration.