

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.

SUMMARY: Apertus Pharmaceuticals applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Apertus Pharmaceuticals 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 61470, Apertus Pharmaceuticals, 331 Consort Drive, Ballwin, Missouri 63011 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 Apertus Pharmaceuticals 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
Remifentanyl (9739)	II

The company plans to manufacture the above-listed controlled substances in bulk for distribution to its customers. In reference to drug codes 7360 marihuana and 7370 tetrahydrocannabinols the company plans to bulk manufacture both as synthetic substances. No other activity for these drug codes is authorized for this registration.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Manufacturer of Controlled Substances Registration: Rhodes Technologies

ACTION: Notice of registration.

SUMMARY: Rhodes Technologies applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Rhodes Technologies registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated August 21, 2015, and published in the **Federal Register** on August 31, 2015, 80 FR 52511, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816 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 Rhodes Technologies 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
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Methylphenidate (1724)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II

Controlled substance	Schedule
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for conversion and sale to dosage form manufacturers.

In reference to drug code 7370 the company plans to bulk manufacture synthetic tetrahydrocannabinols. No other activity for this drug code is authorized for this registration.

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Deputy Assistant Administrator.

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DEPARTMENT OF JUSTICE

[OMB Number 1121-0321]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision of a Currently Approved Collection: National Institute of Justice Compliance Testing Program

AGENCY: Office of Justice Programs, Department of Justice.

ACTION: 60-day notice.

SUMMARY: The Department of Justice (DOJ), Office of Justice Programs, National Institute of Justice (NIJ), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995.

DATES: Comments are encouraged and will be accepted for 60 days until *March 21, 2016*.

FOR FURTHER INFORMATION CONTACT: If you have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact: Michael O'Shea (202) 305-7954, National Institute of Justice (NIJ), Office of Justice Programs, U.S. Department of Justice, 810 Seventh Street NW., Washington, DC 20531 or Jamie.phillips@usdoj.gov.

SUPPLEMENTARY INFORMATION: Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should