

Controlled substance	Schedule
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II
Ecgonine (9180) .....	II
Hydrocodone (9193) .....	II
Levorphanol (9220) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Methadone intermediate (9254) ...	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Opium tincture (9630) .....	II
Opium, powdered (9639) .....	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Remifentanil (9739) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture the listed controlled substances for internal use and for distribution to other companies.

Dated: January 9, 2015.

**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Manufacturer of Controlled Substances Registration: Chattem Chemicals, Inc.**

**ACTION:** Notice of registration.

**SUMMARY:** Chattem Chemicals, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The DEA grants Chattem Chemicals, Inc. registration as a manufacturer of those controlled substances.

**SUPPLEMENTARY INFORMATION:** By notice dated August 27, 2014, and published in the **Federal Register** on September 4, 2014, 79 FR 52764, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409, applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted to this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Chattem 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 basic classes of controlled substances listed:

Controlled substance	Schedule
Gamma Hydroxybutyric Acid (2010).	I
4-Methoxyamphetamine (7411) ...	I
Dihydromorphone (9145) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Lisdexamfetamine (1205) .....	II
Methylphenidate (1724) .....	II
Pentobarbital (2270) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Meperidine intermediate-A (9232)	II
Meperidine intermediate-B (9233)	II
Meperidine intermediate-C (9234)	II
Methadone (9250) .....	II
Methadone intermediate (9254) ...	II
Morphine (9300) .....	II
Oripavine (9330) .....	II
Thebaine (9333) .....	II
Opium tincture (9630) .....	II
Opium, powdered (9639) .....	II
Opium, granulated (9640) .....	II
Oxymorphone (9652) .....	II
Noroxymorphone (9668) .....	II
Alfentanil (9737) .....	II
Remifentanil (9739) .....	II
Sufentanil (9740) .....	II
Tapentadol (9780) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers. Regarding (9640) the company plans to manufacture another controlled substance for sale to its customers.

Dated: January 9, 2015.

**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Manufacturer of Controlled Substances Registration: Wildlife Laboratories, Inc.**

**ACTION:** Notice of registration.

**SUMMARY:** Wildlife Laboratories, Inc. applied to be registered as a manufacturer of a basic class of controlled substance. The DEA grants Wildlife Laboratories, Inc. registration as a manufacturer of this controlled substance.

**SUPPLEMENTARY INFORMATION:** By notice dated May 28, 2014, and published in the **Federal Register** on June 3, 2014, 79 FR 31985, Wildlife Laboratories, Inc., 1230 W. Ash Street, Suite D, Windsor, Colorado 80550, applied to be registered as a manufacturer of a certain basic class of controlled substance. No comments or objections were submitted to this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Wildlife Laboratories, Inc. to manufacture the basic class of controlled substance 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 Carfentanil (9743), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the above listed controlled substance for sale to veterinary pharmacies, zoos, and other animal and wildlife applications.

Dated: January 9, 2015.

**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator.*

[FR Doc. 2015-01294 Filed 1-23-15; 8:45 am]

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