

Procedures and Penalties Act, 15 U.S.C. § 16, including making copies available to the public of this Final Judgment, the Competitive Impact Statement, and any comments thereon and the United States's responses to comments. Based upon the record before the Court, which includes the Competitive Impact Statement and any comments and response to comments filed with the Court, entry of this Final Judgment is in the public interest.

Date:

Court approval subject to procedures of Antitrust Procedures and Penalties Act, 15 U.S.C. § 16.

United States District Judge
[FR Doc. 2014-21102 Filed 9-3-14; 8:45 am]

BILLING CODE 4410-11-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Application: Fisher Clinical Services,
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 1301.34(a) on or before October 6, 2014. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before October 6, 2014.

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.

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 importers, 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 pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on December 13, 2013, Fisher Clinical Services, Inc., 700A-C Nestle Way, Breinigsville, Pennsylvania 18031-1522 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Methylphenidate (1724)	II
Levorphanol (9220)	II
Noroxymorphone (9668)	II
Tapentadol (9780)	II

The company plans to import the listed substances for analytical research and testing and clinical trials. This authorization does not extend to the import of a finished FDA approved or non-approved dosage form for commercial distribution in the United States.

The company plans to import an intermediate form of Tapentadol (9780) to bulk manufacture Tapentadol for distribution to its customers.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.
[FR Doc. 2014-21056 Filed 9-3-14; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Application: Cody Laboratories, Inc.**

ACTION: Notice of application.

SUMMARY: On July 3, 2014, Cody Laboratories, Inc., Cody, Wyoming, applied to be registered as an importer of basic classes of controlled substances.

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 October 6, 2014. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before October 6, 2014.

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 request 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 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 importers 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 pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on July 3, 2014, Cody Laboratories, Inc., 601 Yellowstone Avenue, Cody, Wyoming 82414-9321, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Phenylacetone (8501)	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780)	II

The company plans to import narcotic raw materials for manufacturing and further distribution to its customers. The company is registered with the DEA as a manufacturer of several controlled substances that are manufactured from poppy straw concentrate.

The company plans to import an intermediate form of tapentadol (9780), to bulk manufacture tapentadol for distribution to its customers.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.
[FR Doc. 2014-21058 Filed 9-3-14; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Registration: Mylan Technologies, Inc.**

ACTION: Notice of registration.

SUMMARY: Mylan Technologies, Inc., applied to be registered as an importer of certain basic classes of controlled substances. The DEA grants Mylan Technologies, Inc., registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated May 28, 2014, and published in the **Federal Register** on June 4, 2014, 79 FR 32316, Mylan Technologies, Inc., 110 Lake Street, Saint Albans, Vermont 05478, applied to be registered as an importer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Mylan Technologies, Inc., 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 basic classes of controlled substances listed:

Controlled substance	Schedule
Methylphenidate (1724)	II
Fentanyl (9801)	II

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to allow the company to export domestically-manufactured FDF to foreign markets.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

[FR Doc. 2014-21052 Filed 9-3-14; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Registration: Arizona Department of
Corrections, Aspc-Florence**

ACTION: Notice of registration.

SUMMARY: Arizona Department of Corrections, ASPC-Florence applied to be registered as an importer of a certain basic class of controlled substance. The DEA grants Arizona Department of Corrections, ASPC-Florence registration as an importer of this controlled substance.

SUPPLEMENTARY INFORMATION: By notice dated May 28, 2014, and published in the **Federal Register** on June 4, 2014, 79 FR 32317, Arizona Department of Corrections, ASPC-Florence, 1305 E. Butte Avenue, Florence, Arizona 85132, applied to be registered as an importer of a certain basic class of controlled substance. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Arizona Department of Corrections, ASPC-Florence to import 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above-named company is granted registration as an importer of Pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The facility intends to import the above listed controlled substance for legitimate use. Supplies of this particular controlled substance are inadequate and are not available in the form needed within the current domestic supply of the United States.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

[FR Doc. 2014-21083 Filed 9-3-14; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

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

ACTION: Notice of registration.

SUMMARY: Wildlife Laboratories, Inc., applied to be registered as an importer of certain basic classes of controlled substances. The DEA grants Wildlife Laboratories, Inc., registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated May 28, 2014, and published in the **Federal Register** on June 4, 2014, FR 79 32318, Wildlife Laboratories, Inc., 1230 West Ash Street, Suite D, Windsor, Colorado 80550, applied to be registered as an importer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Wildlife Laboratories, Inc., 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 basic classes of controlled substances:

Controlled substance	Schedule
Etorphine (except HCl) (9056)	I
Etorphine HCl (9059)	II

The company plans to import the listed controlled substances for sale to its customers.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

[FR Doc. 2014-21087 Filed 9-3-14; 8:45 am]

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