

may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 7, 2014.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 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 manufacturers, distributors, and dispensers 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 sec. 7(g) of 28 CFR part 0, subpart R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on March 20, 2014, Actavis Pharma, Inc., 2455 Wardlow Road, Corona, California 92880–2882, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II

The company plans to import the listed controlled substances for analytical testing and clinical trials.

The import of the above listed basic classes of controlled substances will be granted only for analytical testing and clinical trials. This authorization does not extend to the import of a finished Food and Drug Administration approved or non-approved dosage form for commercial distribution in the United States.

Dated: May 28, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Penick Corporation

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 July 7, 2014. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 7, 2014.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 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 manufacturers, distributors, and dispensers 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 sec. 7(g) of 28 CFR part 0, subpart R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on February 26, 2014, Penick Corporation, 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 controlled substance intermediates for sale to its customers.

Comments and requests for any hearings on applications to import narcotic raw material are not

appropriate. 72 FR 3417 (January 25, 2007).

Dated: May 28, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Mylan Technologies, 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 July 7, 2014. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 7, 2014.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 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 manufacturers, distributors, and dispensers 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 sec. 7(g) of 28 CFR pt. 0, subpt. R, App.

In accordance with 21 CFR 1301.34(a), this is notice that on December 12, 2013, Mylan Technologies, Inc., 110 Lake Street, Saint Albans, Vermont 05478, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled Substance	Schedule
Methylphenidate (1724)	II