

Drug	Schedule
Meperidine (9230) .....	II
Meperidine intermediate-A (9232) .....	II
Meperidine intermediate-B (9233) .....	II
Meperidine intermediate-C (9234) .....	II
Metazocine (9240) .....	II
Methadone (9250) .....	II
Methadone intermediate (9254) .....	II
Methamphetamine (1105) .....	II
Methylphenidate (1724) .....	II
Metopon (9260) .....	II
Moramide intermediate (9802) .....	II
Morphine (9300) .....	II
Nabilone (7379) .....	II
Opium, raw (9600) .....	II
Opium extracts (9610) .....	II
Opium fluid extract (9620) .....	II
Opium tincture (9630) .....	II
Opium poppy/Poppy Straw (9650) .....	II
Oripavine (9330) .....	II
Poppy Straw Concentrate (9670) .....	II
Opium, granulated (9640) .....	II
Oxycodone (9143) .....	II
Oxymorphone (9652) .....	II
Pentobarbital (2270) .....	II
Phenazocine (9715) .....	II
Phencyclidine (7471) .....	II
Phenmetrazine (1631) .....	II
Phenylacetone (8501) .....	II
Piminodine (9730) .....	II
Powdered opium (9639) .....	II
Racemethorphan (9732) .....	II
Racemorphan (9733) .....	II
Remifentanil (9739) .....	II
Secobarbital (2315) .....	II
Sufentanil (9740) .....	II
Tapentadol (9780) .....	II
Thebaine (9333) .....	II

The company plans to import small quantities of the listed controlled substances for the National Institute on Drug Abuse (NIDA) for research activities.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (2007).

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule I or II, which fall under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may, in the circumstances set forth in 21 U.S.C. 958(i), file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR § 1301.43 and in such form as prescribed by 21 CFR § 1316.47.

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, **Federal Register** Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than August 30, 2013.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR § 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, 40 FR 43745–46, all applicants for registration to import a basic class of any controlled substances in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: July 23, 2013.  
**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration; Meda Pharmaceuticals, Inc.**

By Notice dated February 8, 2013, and published in the **Federal Register** on February 21, 2013, 78 FR 12101, Meda Pharmaceuticals, Inc., 705 Eldorado Street, Decatur, Illinois 62523, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Nabilone (7379), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance as a finished drug product in dosage form for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Meda Pharmaceuticals Inc., 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. DEA has investigated Meda Pharmaceuticals Inc., to ensure that the company’s registration is consistent with the public interest. The investigation has included inspection and testing of the company’s physical security systems, verification of the company’s compliance with state and local laws, and a review of 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 class of controlled substance listed.

Dated: July 23, 2013.  
**Joseph T. Rannazzisi,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. 2013–18332 Filed 7–30–13; 8:45 am]  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration; Almac Clinical Services, Inc.**

By Notice dated April 10, 2013, and published in the **Federal Register** on April 19, 2013, 78 FR 23594, Almac Clinical Services, Inc., (ACSI), 25 Fretz Road, Souderton, Pennsylvania 18964, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Tapentadol (9780) .....	II
Fentanyl (9801) .....	II

The company plans to import small quantities of the listed controlled substances in dosage form to conduct 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 FDA approved or non-approved dosage forms for commercial distribution in the United States.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Almac Clinical Services, Inc., (ACSI) 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. DEA has investigated Almac Clinical Services, Inc., (ACSI) to ensure that the company’s registration is consistent with the public interest. The investigation has included inspection and testing of the company’s physical security systems, verification of the company’s compliance with state and