

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
Mephedrone (4-Methyl-N-methylcathinone) (1248).	I
1-Pentyl-3-(1-naphthoyl)indole (7118).	I
5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl-phenol] (7297).	I
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
4-Bromo-2,5-dimethoxyamphetamine (7391).	I
3,4-Methylenedioxymethamphetamine (7405).	I
Dimethyltryptamine (7435)	I
Psilocyn (7438)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Oxycodone (9143)	II
Thebaine (9333)	II
Fentanyl (9801)	II

The company plans to import the listed controlled substances in dosage form to distribute to researchers.

In reference to drug codes 7360 and 7370, the company plans to import a synthetic cannabidiol and a synthetic tetrahydrocannabinol. No other activity for these drug codes is authorized for this registration.

The import of the above listed basic classes of controlled substances would 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: September 26, 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
Registration: Meda Pharmaceuticals,
Inc.**

ACTION: Notice of registration.

SUMMARY: Meda Pharmaceuticals, Inc., applied to be registered as an importer of a certain basic class of controlled

substance. The DEA grants Meda Pharmaceuticals, Inc., registration as an importer of this controlled substance.

SUPPLEMENTARY INFORMATION: By notice dated June 10, 2014, and published in the **Federal Register** on June 17, 2014, 79 FR 34552, Meda Pharmaceuticals, Inc., 705 Eldorado Street, Decatur, Illinois 62523, 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 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. 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 nabilone (7379), a basic class of controlled substance listed in schedule II.

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

Dated: September 26, 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: Chattem Chemicals, 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 November 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 November 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 June 23, 2014, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409, applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Methamphetamine (1105)	II
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Phenylacetone (8501)	II
Thebaine (9333)	II
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II
Tapentadol (9780)	II

The company plans to import the listed controlled substances to manufacture bulk controlled substances for sale to its customers. The company plans to import an intermediate form of Tapentadol (9780), and Thebaine (9333), for the manufacture of other bulk controlled substances and distribution to its customers.

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

Dated: September 26, 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
Registration: Alltech Associates, Inc.

ACTION: Notice of registration.

SUMMARY: Alltech Associates, Inc., applied to be registered as an importer of certain basic classes of controlled substances. The DEA grants Alltech Associates, 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 3, 2014, 79 FR 31986, Alltech Associates, Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, 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 Alltech Associates, 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
Gamma Hydroxybutyric Acid (2010).	I
Lysergic acid diethylamide (7315)	I
Heroin (9200)	I
Cocaine (9041)	II
Codeine (9050)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Morphine (9300)	II

The company plans to import these controlled substances for the manufacture of reference standards.

Dated: September 25, 2014.
Joseph T. Rannazzisi,
Deputy Assistant Administrator.
[FR Doc. 2014–23830 Filed 10–6–14; 8:45 am]
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DEPARTMENT OF LABOR

Occupational Safety and Health
Administration

[Docket No. OSHA–2011–0189]

Servicing Multi-Piece and Single Piece
Rim Wheels; Extension of the Office of
Management and Budget's (OMB)
Approval of Information Collection
(Paperwork) Requirements

AGENCY: Occupational Safety and Health Administration, Labor.
ACTION: Request for public comments.

SUMMARY: OSHA solicits public comments concerning its proposal to extend the Office of Management and Budget's (OMB) approval of the information collection requirements specified in the Standard on Servicing Multi-Piece and Single Piece Rim Wheels (29 CFR 1910.177). The paperwork provisions of the Standard includes a requirement that the manufacturer or a Registered Professional Engineer certify that repaired restraining devices and barriers meet the strength requirements specified in the Standard and a requirement that defective wheels and wheel components be marked or tagged.

DATES: Comments must be submitted (postmarked, sent, or received) by December 8, 2014.

ADDRESSES: *Electronically:* You may submit comments and attachments electronically at <http://www.regulations.gov>, which is the Federal eRulemaking Portal. Follow the instructions online for submitting comments.

Facsimile: If your comments, including attachments, are not longer than 10 pages you may fax them to the OSHA Docket Office at (202) 693–1648.

Mail, hand delivery, express mail, messenger, or courier service: When using this method, you must submit a copy of your comments and attachments to the OSHA Docket Office, Docket No. OSHA–2011–0189, Occupational Safety and Health Administration, U.S. Department of Labor, Room N–2625, 200 Constitution Avenue NW., Washington, DC 20210. Deliveries (hand, express mail, messenger, and

courier services) are accepted during the Department of Labor's and Docket Office's normal business hours, 8:15 a.m. to 4:45 p.m., e.t.

Instructions: All submissions must include the Agency name and OSHA docket number (OSHA–2011–0189) for the Information Collection Request (ICR). All comments, including any personal information you provide, are placed in the docket without change and may be made available online at <http://www.regulations.gov>. For further information on submitting comments see the "Public Participation" heading in the section of this notice titled "SUPPLEMENTARY INFORMATION."

Docket: To read or download comments or other materials in the docket, go to <http://www.regulations.gov> or the OSHA Docket Office at the address above. All documents in the docket (including this **Federal Register** notice) are listed in the <http://www.regulations.gov> index; however, some information (e.g., copyrighted material) is not publicly available to read or download from the Web site. All submissions, including copyrighted material, are available for inspection and copying at the OSHA Docket Office. You may also contact Theda Kenney at the address below to obtain a copy of the ICR.

FOR FURTHER INFORMATION CONTACT: Theda Kenney or Todd Owen, Directorate of Standards and Guidance, OSHA, U.S. Department of Labor, Room N–3609, 200 Constitution Avenue NW., Washington, DC 20210; telephone (202) 693–2222.

SUPPLEMENTARY INFORMATION:

I. Background

The Department of Labor, as part of its continuing effort to reduce paperwork and respondent (i.e., employer) burden, conducts a preclearance consultation program to provide the public with an opportunity to comment on proposed and continuing information collection requirements in accord with the Paperwork Reduction Act of 1995 (PRA–95) (44 U.S.C. 3506(c)(2)(A)). This program ensures that information is in the desired format, reporting burden (time and costs) is minimal, collection instruments are clearly understood, and OSHA's estimate of the information collection burden is accurate. The Occupational Safety and Health Act of 1970 (the OSH Act) (29 U.S.C. 651 et seq.) authorizes information collection by employers as necessary or appropriate for enforcement of the OSH Act or for developing information regarding the causes and prevention of occupational injuries, illnesses, and