

Administration (DEA) to be registered as an importer of Phenylacetone (8501), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance to bulk manufacture amphetamine.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class of controlled substance may 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 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 July 23, 2009.

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 the basic class of any controlled substance 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: June 15, 2009.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E9–14729 Filed 6–22–09; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II and prior to issuing a regulation under 21 U.S.C. 952(a)(2) authorizing the importation of such substances, provide manufacturers holding registrations for the bulk

manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on May 28, 2009, Aptuit (Allendale) Inc., 75 Commerce Drive, Allendale, New Jersey 07401, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the basic classes of controlled substances listed in schedule II.

Drug	Schedule
Lisdexamfetamine (1205)	II
Noroxymorphone (9668)	II

The company plans to import the basic classes of controlled substances for clinical trials and research.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may 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 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 July 23, 2009.

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 the basic classes 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: June 15, 2009.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E9–14724 Filed 6–22–09; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 27, 2009, Boehringer Ingelheim Chemicals, Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, made application by renewal to the Drug Enforcement Administration (DEA) as a bulk manufacturer of the basic classes of controlled substances listed in schedule II:

Drug	Schedule
Amphetamine (1100)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers for formulation into finished pharmaceuticals.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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 24, 2009.

Dated: June 15, 2009.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E9–14707 Filed 6–22–09; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 22, 2009, Noramco Inc., Division of Ortho-McNeil, Inc., 1440 Olympic Drive, Athens, GA 30601, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Tapentadol

(9780), a basic class of controlled substance listed in schedule II.

The company plans to bulk manufacture the above listed controlled substance for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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, VA 22152; and must be filed no later than August 24, 2009.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E9-14705 Filed 6-22-09; 8:45 am]

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DEPARTMENT OF JUSTICE

DRUG ENFORCEMENT ADMINISTRATION

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 1, 2009, Noramco Inc., Division of Ortho-McNeil, Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Noroxymorphone (9668), a basic class of controlled substance listed in schedule II.

The company plans to bulk manufacture the above listed controlled substance for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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, VA 22152; and must be filed no later than August 24, 2009.

Dated: June 15, 2009.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E9-14704 Filed 6-22-09; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 22, 2009, Noramco Inc., Division of Ortho-McNeil, Inc., 500 Swedes Landing Road, Wilmington, Delaware 19801, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Tapentadol (9780), a basic class of controlled substance listed in schedule II.

The company plans to bulk manufacture the above listed controlled substance for distribution to its customers.

Any other such applicant, and any person who is presently registered with DEA to manufacture such a substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

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, VA 22152; and must be filed no later than August 24, 2009.

Dated: June 15, 2009.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E9-14695 Filed 6-22-09; 8:45 am]

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DEPARTMENT OF LABOR

Office of the Secretary; "Promotion of Labor Rights and Employment in Pakistan"

June 17, 2009.

AGENCY: Bureau of International Labor Affairs, Department of Labor.

Announcement Type: New. Notice of Availability of Funds and Solicitation for Cooperative Agreement Applications. The full announcement is posted on <http://www.Grants.gov>.

Funding Opportunity Number: SGA-09-07.

Key Dates: The closing date for receipt of applications is July 31, 2009 via Grants.gov. Applications may also be submitted by mail.

Funding Opportunity Description

The U.S. Department of Labor (USDOL), Bureau of International Labor Affairs (ILAB), announces the availability of funds in the amount of U.S. \$1,600,000 to be awarded by Cooperative Agreement to a qualifying organization for the purpose of implementing a worker rights and employment demonstration project that assists unemployed Pakistanis with vocational and entrepreneurial training to obtain employment or pursue professions and acquire an understanding of their fundamental rights as workers. The project will take place in the Federally Administered Tribal Areas, the Northwest Frontier Province and Baluchistan. Participants provided with vocational training will also receive integrated worker rights training in order to ensure that they are able to fully benefit from their new skills and professions. Similarly, new entrepreneurs may become employers for the first time and will also receive worker rights training to understand their responsibilities as employers. ILAB is authorized to award and administer this program by the Department of Labor Appropriations Act, 2009, Public Law 111-08, 123 Stat. 524 (2009). The Cooperative Agreement awarded under this initiative will be managed by ILAB's Office of Trade and Labor Affairs. The duration of the project funded by this solicitation is three years. The start date of program activities will be negotiated upon award of the Cooperative Agreement, but will be no later than September 30, 2009.

The full Solicitation for Grant Applications is posted on <http://www.Grants.gov> under U.S. Department of Labor/ILAB. If you need to speak to a person concerning these grants, or if you have issues regarding access to the Grants.gov Web site, you may telephone Georgette Nkpa at 202-693-4584 (not a toll-free number).

Signed at Washington, DC this 17th day of June 2009.

Cassandra Mitchell,

Grant Officer.

[FR Doc. E9-14648 Filed 6-22-09; 8:45 am]

BILLING CODE 4510-28-P