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 10, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–11689 Filed 7–21–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated March 6, 2006 and published in the **Federal Register** on March 13, 2006, (71 FR 12714), ISP Freetown Fine Chemicals, 238 South Main Street, Assonet, Massachusetts 02702, made application to the Drug Enforcement 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 Phenylacetone to manufacture Amphetamine.

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 ISP Freetown Fine Chemicals 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, at this time. DEA has investigated ISP Freetown Fine Chemicals 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 10, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–11694 Filed 7–21–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 18, 2006 and published in the **Federal Register** on April 25, 2006, (71 FR 23949), Johnson Matthey Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey 08066, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in Schedule II:

Drug	Schedule
Phenylacetone (8501)	II
Raw Opium (9600)	II
Concentrate of Poppy Straw (9670).	II

The company plans to import the controlled substances as raw materials for use in the manufacture of bulk controlled substances 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 Johnson Matthey Inc. to import the basic class 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, at this time. DEA has investigated Johnson Matthey 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 substances listed.

Dated: July 10, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–11692 Filed 7–21–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 18, 2006 and published in the **Federal Register** on April 25, 2006, (71 FR 23949–23950), Noramco Inc., 500 Old Swedes Landing Road, Wilmington, Delaware 19801, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in Schedule II:

Drug	Schedule
Raw Opium (9600)	II
Concentrate of Poppy Straw (9670).	II

The company plans to import the listed controlled substances to manufacture other controlled substances.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. Sections 823(a) and 952(a) and determined that the registration of Noramco Inc. to import the basic class 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, at this time. DEA has investigated Noramco 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 substances listed.

Dated: July 10, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–11693 Filed 7–21–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 20, 2006, and published in the **Federal Register** on March 27, 2006, (71 FR 15219), Organichem Corporation, 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedules I and II:

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Dextropropoxyphene (9273)	II
Fentanyl (9801)	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers. In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabindiol as a synthetic intermediate. This controlled substance will be further synthesized to bulk manufacture a synthetic THC (7370). No other activity for this drug code is authorized for this registration.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Organichem Corporation to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Organichem Corporation 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: July 10, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–11690 Filed 7–21–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

National Institute of Corrections

Advisory Board Meeting

Time and Date: 8 a.m. to 4:30 p.m. on Monday, September 25, 2006. 8 a.m. to 4:30 p.m. on Tuesday, September 26, 2006.

Place: Courtyard by Marriott Detroit, 333 E. Jefferson Avenue, Detroit, Michigan 48226, Phone: 313–222–7700.

Status: Open.

Matters to be Considered: Site Visit to Michigan Department of Corrections; Observation of Michigan Prisoner ReEntry Initiative; Faith Based; Evidence-based practices, Institutional culture work; and pubic/private funding partnerships; PREA Update; Agency Reports.

For Further Information Contact: Larry Solomon, Deputy Director, 202–307–3106, ext. 44254.

Morris L. Thigpen,

Director.

[FR Doc. 06–6427 Filed 7–21–06; 8:45 am]

BILLING CODE 4410–36–M

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

[Docket No. NRTL3–92]

TUV Rheinland of North America, Inc., Application for Expansion of Recognition

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Notice.

SUMMARY: This notice announces the application of TUV Rheinland of North America, Inc., (TUV) for expansion of its recognition to use additional test standards, and presents the Agency's preliminary finding to grant this request for expansion. This preliminary finding does not constitute an interim or temporary approval of this application.

DATES: You must submit information or comments, or any request for extension of the time to comment, by the following dates:

• *Hard copy:* postmarked or sent by August 8, 2006.

• *Electronic transmission or facsimile:* sent by August 8, 2006.

ADDRESSES: You may submit information or comments to this notice—identified by docket number NRTL3–92—by any of the following methods:

• *Federal eRulemaking Portal: http://www.regulations.gov*. Follow the instructions for submitting comments.

• OSHA Web site: *http://ecomments.osha.gov*. Follow the instructions for submitting comments on OSHA's Web page.

• *Fax:* If your written comments are 10 pages or fewer, you may fax them to the OSHA Docket Office at (202) 693–1648.

• *Regular mail, express delivery, hand delivery and courier service:* Submit three copies to the OSHA Docket Office, Docket No. NRTL3–92, U.S. Department of Labor, 200 Constitution Avenue, NW., Room N–2625, Washington, DC 20210; telephone (202) 693–2350. (OSHA's TTY number is (877) 889–5627). OSHA Docket Office hours of operation are 8:15 a.m. to 4:45 p.m., EST.

Instructions: All comments received will be posted without change to *http://dockets.osha.gov*, including any personal information provided. OSHA cautions you about submitting personal information such as social security numbers and birth dates.

Docket: For access to the docket to read background documents or comments received, go to *http://dockets.osha.gov*. Contact the OSHA Docket Office for information about materials not available through the OSHA Web page and for assistance in using the Web page to locate docket submissions.

Extension of Comment Period: Submit requests for extensions concerning this notice to the Office of Technical Programs and Coordination Activities, NRTL Program, Occupational Safety and Health Administration, U.S. Department of Labor, 200 Constitution Avenue, NW., Room N–3655, Washington, DC 20210. Or, fax to (202) 693–1644.

FOR FURTHER INFORMATION CONTACT: Director, Office of Technical Programs and Coordination Activities, NRTL Program, Occupational Safety and Health Administration, U.S. Department of Labor, 200 Constitution Avenue, NW., Room N–3655, Washington, DC 20210, or phone (202) 693–2110.

SUPPLEMENTARY INFORMATION:

Notice of Application

The Occupational Safety and Health Administration (OSHA) hereby gives notice that TUV Rheinland of North