

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and OMA intends to file additional written notifications disclosing all changes in membership.

On March 18, 1998, OMA filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on December 31, 1998 (63 FR 72333).

The last notification was filed with the Department on January 18, 2007. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on February 26, 2007 (72 FR 8401).

**J. Robert Kramer, II,**

*Director of Operations, Antitrust Division.*

[FR Doc. E8-14596 Filed 6-26-08; 8:45 am]

**BILLING CODE 4410-11-M**

## DEPARTMENT OF JUSTICE

### Antitrust Division

#### **Notice Pursuant to the National Cooperative Research and Production Act of 1993—International SAE Consortium Ltd. (Formerly Known as SAE Consortium Ltd.)**

Notice is hereby given that, on May 21, 2008, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* ("the Act"), International SAE Consortium Ltd. ("ISAEC") has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Daiichi Sankyo, Inc., Edison, NJ; Takeda Global Research and Development Center, Inc., Deerfield, IL; and The Wellcome Trust, London, UNITED KINGDOM have been added as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and ISAEC intends to file additional written notification disclosing all changes in membership.

On September 27, 2007, ISAEC filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to section

6(b) of the Act on November 7, 2007 (72 FR 62867).

The last notification was filed with the Department of Justice on January 25, 2008. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on March 4, 2008 (73 FR 11680).

**J. Robert Kramer, II,**

*Director of Operations, Antitrust Division.*

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## DEPARTMENT OF JUSTICE

### Antitrust Division

#### **Notice Pursuant to the National Cooperative Research and Production Act of 1993—Testing of Methods for Measuring Hydrocarbon Dew Points in Natural Gas Streams**

Notice is hereby given that, on May 13, 2008, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* ("the Act"), SwRI: Testing of Methods for Measuring Hydrocarbon Dew Points in Natural Gas Streams has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its nature and objective. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, the period of performance has been extended to July 31, 2008.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and SwRI intends to file additional written notifications disclosing all changes in membership.

On March 20, 2007, SwRI: Testing of Methods for Measuring Hydrocarbon Dew Points in Natural Gas Streams filed its original notification pursuant to section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to section 6(b) of the Act on April 16, 2007 (72 FR 19023).

The last notification was filed with the Department on October 30, 2007. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on December 20, 2007 (72 FR 72389).

**J. Robert Kramer, II,**

*Director of Operations, Antitrust Division.*

[FR Doc. E8-14598 Filed 6-26-08; 8:45 am]

**BILLING CODE 4410-11-M**

## 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 registration 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 13, 2008, 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 Noroxymorphone (9668), a basic class of controlled substance listed in schedule II.

The company plans to import the basic class of controlled substance for clinical trials and research.

Any manufacturer who 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 written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to 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 28, 2008.

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 substance listed 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 19, 2008.

**Joseph T. Rannazzisi,**

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

[FR Doc. E8-14586 Filed 6-26-08; 8:45 am]

**BILLING CODE 4410-09-P**

## 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 registration under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on May 8, 2008, Aptuit, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Marihuana (7360), a basic class of controlled substance listed in schedule I.

The company plans to import a finished pharmaceutical product containing cannabis extracts in dosage form for packaging for a clinical trial study.

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 being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, **Federal Register** Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, **Federal Register** Representative (ODL), 8701 Morrisette

Drive, Springfield, VA 22152; and must be filed no later than July 28, 2008.

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 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 19, 2008.

**Joseph T. Rannazzisi,**

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

[FR Doc. E8-14590 Filed 6-26-08; 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 19, 2008 and published in the **Federal Register** on March 28, 2008, (73 FR 16718), Penick Corporation, 33 Industrial Park Road, Pennsville, New Jersey 08070, 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
Coca Leaves (9040) .....	II
Opium, raw (9600) .....	II
Poppy Straw (9650) .....	II
Concentrate of Poppy Straw (9670).	II

The company plans to import the listed controlled substances to manufacture bulk controlled substance intermediates for sale 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 Penick Corporation 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, at this time. DEA has investigated Penick

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. 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.

Dated: June 19, 2008.

**Joseph T. Rannazzisi,**

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

[FR Doc. E8-14584 Filed 6-26-08; 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 27, 2008 and published in the **Federal Register** on April 2, 2008, (73 FR 18000), Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Coca Leaves (9040), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance for the manufacture of a bulk controlled substance for distribution to its customer.

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 Stepan Company 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 Stepan Company 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