

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.*

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## 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.*

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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 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