

DEPARTMENT OF JUSTICE**Antitrust Division****Notice Pursuant to the National Cooperative Research and Production Act of 1993—Cooperative Research Group on Development and Evaluation of a Gas Chromatograph Testing Protocol**

Notice is hereby given that, on August 26, 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”), Southwest Research Institute—Cooperative Research Group on Development and Evaluation of a Gas Chromatograph Testing Protocol (“GCTP”) 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, EffeTech, Ltd., Staffordshire, UNITED KINGDOM has 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 GCTP intends to file additional written notifications disclosing all changes in membership.

On March 6, 2008, GCTP 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 7, 2008 (73 FR 18813).

The last notification was filed with the Department on June 4, 2008. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on July 16, 2008 (73 FR 40882).

Patricia A. Brink,

Deputy 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—IMS Global Learning Consortium, Inc.**

Notice is hereby given that, on August 28, 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”)/IMS Global Learning Consortium, Inc. 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, CREDU Co., Ltd., Seoul, REPUBLIC OF KOREA; DaulSoft Co., Ltd., Seoul, REPUBLIC OF KOREA; Laureate Online Education, Baltimore, MD; Miami-Dade College—Virtual College, Miami, FL; and Utah Valley University, Orem, UT have been added as parties to this venture. Also, Intrallect, Scotland, UNITED KINGDOM has withdrawn 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 IMS Global Learning Consortium, Inc. intends to file additional written notifications disclosing all changes in membership.

On April 7, 2008, IMS Global Learning Consortium, Inc. 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 September 13, 2008 (65 FR 55283).

The last notification was filed with the Department on June 10, 2008. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on July 21, 2008 (73 FR 42367).

Patricia A. Brink,

Deputy 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—Digital Entertainment Content Ecosystem (DECE) LLC**

Notice is hereby given that, on August 25, 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”), Digital Entertainment Content Ecosystem (DECE) LLC (“DECE”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the name and principal place of business of the

standards development organization and (2) the nature and scope of its standards development activities. The notifications were filed for the purpose of invoking the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the name and principal place of business of the standards development organization is: Digital Entertainment Content Ecosystem (DECE) LLC, Culver City, CA. The nature and scope of DECE’s standards development activities are: (1) To enable the delivery of digital entertainment content in a manner that allows for interoperability among digital formats and digital rights management systems; and (2) to develop specifications accordingly.

Patricia A. Brink,

Deputy 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 II, and prior to issuing a regulation 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 August 18, 2008, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004–1412, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to import small quantities of ioflupane, in the form of three separate analogues of Cocaine, to validate production and QC systems; for a reference standard; and for producing material for future investigational new drug (IND) submission.

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