under specified circumstances. Specifically, Sound Technology (S.Z.) Co. Ltd., Shenzhen, PEOPLE'S REPUBLIC OF CHINA; Vulcan Inc., Seattle, WA; Yusan Industries, Ltd., Hong Kong, HONG KONG-CHINA; and Zentek Technology Japan, Inc., Tokyo, JAPAN have been added as parties to this venture.

Also, AWIND, Inc., Taipei, TAIWAN; Bestguide Group Limited, Kowloon, HONG KONG-CHINA; Clevo Co., Taipei, TAIWAN; Coretronic Corporation, Miao-Li, TAIWAN; Cosmic Digital Technology, Ltd., Hong Kong, HONG KONG-CHINA; Daewoo Electronics Corporation, Seoul, REPUBLIC OF KOREA; Dahaam E-Tec Co., Seoul, REPUBLIC OF KOREA; Disctronics Texas, Inc. dba DiscUSA, Plano, TX; Ever Best Industrial (H.K.) Limited, Kowloon, HONG KONG-CHINA; Giant Video Electronics Co., Ltd., Yueh Long, HONG KONG-CHINA; Hansong (Nanjing) Electronic Ltd., Nanjing, PEOPLE'S REPUBLIC OF CHINA; Hing Lung Technology (HK) Company Limited, Hong Kong, HONG KONG-CHINA; KRCD India PVT Ltd., Mumbai, INDIA; Leadtek Research, Inc., Taipei, TAIWAN; Link Concept Technology Ltd., Kowloon, HONG KONG-CHINA; Linpus Technologies, Inc., Taipei, TAIWAN; Major Digital Technology Co., Ltd., Jiangxi, PEOPLE'S REPUBLIC OF CHINA; ODS Optical Disc Service GmbH, Dassow, GERMANY; Premium Disc Corp., Mississauga, Ontario, CANADA; Princeton Technology Corp., Taipei, TAIWAN; Prof ilo Telra Elektronic San. Ve Tic. A.S., Istanbul, TURKEY; SKC Co. Ltd., Seoul, REPUBLIC OF KOREA; Zhongshan Dingcai AV Technology Ltd., Zhongshan, PEOPLE'S REPUBLIC OF CHINA; and Ziova Corporation Pty Ltd., Lonsdale, South Australia, AUSTRALIA have withdrawn as parties to this venture.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 DVD CCA intends to file additional written notifications disclosing all changes in membership.

On April 11, 2001, DVD CCA 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 August 3, 2001 (66 FR 40727).

The last notification was filed with the Department on September 10, 2008. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on October 21, 2008 (73 FR 62541)

Patricia A. Brink,

Deputy Director of Operations, Antitrust Division.

[FR Doc. E9–757 Filed 1–16–09; 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—Wireless Industrial Technology Konsortium Inc.

Notice is hereby given that, on December 2, 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'') Wireless Industrial Technology Konsortium Inc. (''WITK'') 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, Softing AG, Haar, GERMANY; and Cooper Industries, Houston, TX have been added as parties to this venture. Also, Airsprite Technologies, Inc., Marlborough, MA 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 WITK intends to file additional written notifications disclosing all changes in membership.

On August 8, 2008, WITK 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 18, 2008 (73 FR 54170).

Patricia A. Brink,

Deputy Director of Operations, Antitrust Division.

[FR Doc. E9–758 Filed 1–16–09; 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 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 November 26, 2008, Kenco VPI, Division of Kenco Group, Inc., 350 Corporate Place, Chattanooga, Tennessee 37419, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Nabilone (7379), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance for distribution to its customers.

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 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than February 20, 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), 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: January 9, 2009.

Joseph T. Rannazzisi,

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

[FR Doc. E9–1051 Filed 1–16–09; 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 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 § 1301.34(a), this is notice that on October 23, 2008, Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, has made letter to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic class Thebaine (9333), a controlled substance listed in schedule II.

The company plans to import analytical reference standards for distribution to its customers for research purposes.

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), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than February 20, 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 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: January 9, 2009.

Joseph T. Rannazzisi,

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

[FR Doc. E9–1052 Filed 1–16–09; 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 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 September 4, 2008, Medical Isotopes Inc., 100 Bridge Street, Pelham, New Hampshire 03076, made application 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
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Diprenorphine (9058)	II
Etorphine HCL (9059)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Methadone (9250)	II
Morphine (9300)	II
Thebaine (9333)	II
Opium, powdered (9639)	II
Levo-alphacetylmethadol (9648)	II
Oxymorphone (9652)	II
Fentanyl (9801)	II

The company plans to import small quantities of the listed controlled substances as reference standards for distribution for research and analytical purposes only.

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 being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than February 20, 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 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: January 9, 2009.

Joseph T. Rannazzisi,

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

[FR Doc. E9–1055 Filed 1–16–09; 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 November 5, 2008, Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003