

Information Systems, Massy CEDEX, FRANCE; and International Computers Limited, Bracknell, Berks, ENGLAND.

New Allied Sponsors of OSF, i.e., a group of affiliated members which share a single vote, are NEC Corporation, Tokyo, JAPAN; Siemens Nixdorf Informationssysteme AG, Munchen, GERMANY; Silicon Graphics Computer Systems, Mountain View, CA; Sony Corporation, Tokyo, JAPAN; and Transarc Corporation, Pittsburgh, PA.

The previous notification filed on May 11, 1994 is hereby corrected to show a change in address as follows: Persetel (Pty) Ltd., Sandton, Johannesburg, SOUTH AFRICA; USAF ESD, Hanscom AFB, MA; and University of Bilkent, Ankara, TURKEY; and to add Unilever PLC/NV, London, ENGLAND as a member of OSF.

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

On May 11, 1994, OSF 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 31, 1994 (59 Fed. Reg. 45009).

**Constance K. Robinson,**

*Director of Operations, Antitrust Division.*  
[FR Doc. 95-12873 Filed 5-24-95; 8:45 am]  
BILLING CODE 4410-01-M

**Notice Pursuant to the National Cooperative Research and Production Act of 1993; SI Diamond Technology, Inc.**

Notice is hereby given, that on March 21, 1995, pursuant to the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* ("the Act"), SI Diamond Technology, Inc., for itself and on behalf of its members, has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the identities of the parties and (2) the nature and objectives of the venture. The notifications were filed for the purpose 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 identities of the parties are: SI Diamond Technology, Inc., Houston, TX; and Supertex, Inc., Sunnyvale, CA. The purpose of this joint venture is to develop and demonstrate diamond diode field

emission display process technology needed for production of a 10", full color, VGA, flat panel display. The activities of this Joint Venture project will be partially funded by an award from the Advanced Technology Program, National Institute of Standards and Technology, Department of Commerce.

**Constance K. Robinson,**

*Director of Operations, Antitrust Division.*  
[FR Doc. 95-12874 Filed 5-24-95; 8:45 am]  
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**Drug Enforcement Administration**

**Importation of Controlled Substances; Notice of Application**

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (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 Section 1002(a) 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 Section 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on February 17, 1994, Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121-4340, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370) .....	I
Mescaline (7381) .....	I
Amphetamine (1100) .....	II
Phencyclidine (7471) .....	II
Phenylacetone (8501) .....	II
Cocaine (9041) .....	II

The firm plans to import small quantities of the listed controlled substances to make reagents for distribution to the biomedical research commodity.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in

such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 26, 1995.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42 (b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745-43746 (September 23, 1975), all applicants for registration to import basic classes 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 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated May 18, 1995.

**Gene R. Haislip,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
[FR Doc. 95-12782 Filed 5-24-95; 8:45 am]  
BILLING CODE 4410-09-M

**Importer of Controlled Substances; Notice of Registration**

By Notice dated April 25, 1994, and published in the **Federal Register** on May 4, 1994, (59 FR 23081), Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Dimethyltryptamine (7435) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Benzoylcegonine (9180) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Fentanyl (9801) .....	II

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of

Federal Regulations, Section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: May 18, 1995.

**Gene R. Haislip,**

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

[FR Doc. 95-12779 Filed 5-24-95; 8:45 am]

BILLING CODE 4410-09-M

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated October 20, 1994, and published in the **Federal Register** on October 28, 1994, (59 FR 54219), Hoffman-LaRoche, Inc., 340 Kingsland Street, Nutley, New Jersey 07110, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Levorphanol (9220), a basic class of controlled substance listed in Schedule II.

No comments or objections have been received. Therefore, pursuant to Section 303 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 and Title 21, Code of Federal Regulations, Section 1301.54(e), the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: May 17, 1995.

**Gene R. Haislip,**

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

[FR Doc. 95-12780 Filed 5-24-95; 8:45 am]

BILLING CODE 4410-09-M

**Manufacturer of Controlled Substances; Notice of Correction**

In the **Federal Register** (FR Doc. 95-8920) Vol. 60, No. 70 at page 18618, dated April 12, 1995, the listing of controlled substances should have included Oxycodone (9143), Hydromorphone (9150), Diphenoxylate (9170) and Noroxymorphone (9668) for Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147.

Dated: May 17, 1995.

**Gene R. Haislip,**

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

[FR Doc. 95-12778 Filed 5-24-95; 8:45 am]

BILLING CODE 4410-09-M

**Importation of Controlled Substances; Notice of Application**

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (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 Section 1002(a) 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 Section 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on January 18, 1995, North Pacific Trading Company, 1505 SE Gideon Street, Portland, Oregon 97202, made application to the Drug Enforcement Administration to be registered as an importer of Marihuana (7360) a basic class of controlled substance in Schedule I.

This application is exclusively for the importation of marithuana seed which will be rendered non-viable and used as bird seed.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in such form as prescribed by 21 CFR 1305.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justices, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 26, 1995.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43747-46 (September 23, 1975), all applicants for registration to import basic classes 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 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: May 18, 1995.

**Gene R. Haislip,**

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

[FR Doc. 95-12783 Filed 5-24-95; 8:45 am]

BILLING CODE 4410-09-M

**Importer of Controlled Substances; Application Withdrawal for Nycomed Incorporated**

By letter dated April 17, 1995, Nycomed Inc., 33 Riverside Avenue, Rensselaer, New York 12144, withdrew their request to be registered as an importer of Meperidine (9230).

Therefore, the Notice dated February 14, 1995, in **Federal Register** (FR Doc. 95-3627), Vol. 60, No. 30 at page 8414 is hereby withdrawn.

Dated: May 17, 1995.

**Gene R. Haislip,**

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

[FR Doc. 95-12781 Filed 5-24-95; 8:45 am]

BILLING CODE 4410-09-M

**Importation of Controlled Substances; Notice of Application**

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (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 Section 1002(a) 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 Section 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 22, 1995, Research Biochemicals, Limited Partnership, One Strathmore Road, Natick, Massachusetts 01760, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Methaqualone (2565) .....	I
Ibogaine (7260) .....	I
Tetrahydrocannabinol (7370) .....	I
Bufotenine (7433) .....	I
Dimethyltryptamine (7435) .....	I
Etorphine (except HC1) (9056) ....	I
Methylphenidate (1724) .....	II
Etorphine Hydrochloride (9059) ...	II
Diphenoxylate (9170) .....	II