have been made in either the membership, corporate name, or planned activities of this group research project. Membership in the project remains open, and the Semiconductor Research Corporation intends to file additional written notifications disclosing all changes in membership.

On January 7, 1985, the Semiconductor Research Corporation 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 January 30, 1985 (50 FR 4281). The last notification was filed with the Department on September 16, 1997. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on October 31, 1997 (62 FR 58983).

Constance K. Robinson,

Director of Operations, Antitrust Division.

[FR Doc. 98–3594 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 8, 1997, Johnson & Johnson Pharmaceutical Partners, HC–02 State Road 933, KMO.1 Mamey Ward HC–02 Box 19250, Gurabo, Puerto Rico 00778–9629, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of sufentanil (9740), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture sufentanil for bulk distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than April 13, 1998.

Dated: January 21, 1998.

John H. King,

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

[FR Doc. 98–3611 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21, of the Code of Federal Regulations (CFR), this is notice that on November 4, 1997, Knoll Pharmaceutical Company, 30 North Jefferson Road, Whippany, New Jersey 07981, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Dihydromorphine (9145)	I
Hydromorphone (9150)	II

The firm plans to produce bulk product and finished dosage units for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than April 13, 1998.

Dated: January 21, 1998.

John H. King,

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

[FR Doc. 98–3612 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Registration

By Notice dated October 3, 1997, and published in the **Federal Register** on October 22, 1997, (62 FR 54856), Norac Company, Inc., 405 S. Motor Avenue, Azusa, California 91702, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of tetrahydrocannabinols (7370), a basic class of controlled substance listed in Schedule I.

The firm plans to manufacture medication for the treatment of AIDS wasting syndrome and as an antiemetic.

DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Norac Company, Inc. to manufacture tetrahydrocannabinols is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, 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: January 21, 1998.

John H. King,

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

[FR Doc. 98–3608 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Registration

By Notice dated October 6, 1997, and published in the **Federal Register** on October 22, 1997, (62 FR 54857), Novartis Pharmaceuticals Corp., Attn: Compliance, 59 Route 10, East Hanover, 556 Morris Avenue, Summit, New Jersey 07901, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of methylphenidate (1724), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture the finished product for distribution to its customers.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Novartis Pharmaceuticals Corp. to manufacture methylphenidate is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, 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: January 21, 1998.

John H. King,

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

[FR Doc. 98–3607 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Registration

By Notice dated October 3, 1997, and published in the **Federal Register** on October 22, 1997, (62 FR 54857), Nycomed, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of meperidine (9230), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture the bulk product for distribution to its customers.

DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Nycomed, Inc. to manufacture meperidine is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, 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: January 21, 1998.

John H. King,

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

[FR Doc. 98–3606 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that by letter dated October 2, 1997, which was received for processing on October 26, 1997, Nycomed, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of methylphenidate (1724) a basic class of controlled substance listed in Schedule II.

This bulk manufacture of methylphenidate is being conducted in conjunction and coordination with another bulk manufacturer.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than April 13, 1998.

Dated: January 8, 1998.

John H. King,

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

[FR Doc. 98–3613 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 9, 1997, Orpharm, Inc., 728 West 19th Street, Houston, Texas 77008, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methadone (9250)	II
Methadone-intermediate (9254) ..	II
Levo-alphacetylmethadol (9648)	II

The firm plans manufacture methadone and methadone-intermediate for production of LAAM.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than April 13, 1998.

Dated: January 21, 1998.

John H. King,

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

[FR Doc. 98–3609 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that Roche Diagnostic Systems, Inc., 1080 U.S. Highway 202, Somerville, New Jersey 08876–3771, made application to the Drug Enforcement Administration (DEA) by letter dated December 17, 1997, for registration as a bulk manufacturer of ecgonine (9180), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture small quantities of ecgonine which will be further converted into derivatives for incorporation in drug of abuse detection kits.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than April 13, 1998.

Dated: January 21, 1998.

John H. King,

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

[FR Doc. 98–3610 Filed 2–11–98; 8:45 am]

BILLING CODE 4410–09–M