proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g. permitting electronic submission of responses.

Overview of this information collection:

(1) Type of Information Collection: Existing Collection in Use Without an OMB Control Number.

(2) Title of the Form/Collection: Notice of Entry of Appearance as Attorney or representative Before the Immigration Court, Executive Office for Immigration Review.

(3) Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection: Form EOIR–28, Executive Office for immigration Review, U.S. Department of Justice.

(4) Affected public who will be asked to respond, as well as a brief abstract: The information collected on EOIR–28 will be used: (i) To determine whether or not a responding attorney or representative is duly authorized to represent aliens before the Immigration Court, (ii) to provide the responding represented party an opportunity to expressly consent to such representation and to release of Executive Office for Immigration Review records to the representative as required by law, and (iii) to notify the Immigration and Naturalization Service and the Executive Office for immigration Review of such representation.

(5) An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: 77,000 responses per year at 6 minutes per response.

(6) An estimate of the total public burden (in hours) associated with the collection: 7,700 annual burden hours.

If additional information is required contact: Mr. Robert B. Briggs, Clearance Officer, United States Department of Justice, Information Management and Security Staff, Justice Management Division, Suite 850, Washington Center, 1001 G Street, NW, Washington, DC 20530.

Dated: July 1, 1999.

Robert B. Briggs,

Clearance Officer, Department of Justice.

[FR Doc. 99–17220 Filed 7–6–99; 8:45 am]

BILLING CODE 4410–30–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 April 21, 1999, American Radiolabeled Chemicals, Inc., 11624 Bowling Green Drive, St. Louis, Missouri 63146, 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
Dimethyltryptamine (7435)	I
Dihydromorphine (9145)	I
Cocaine (9041)	II
Codeine (9050)	II
Benzoylecgonine (9180)	II
Meperidine (9230)	II
Metazocine (9240)	II
Morphine (9300)	II
Oxymorphone (9652)	II

The firm plans to bulk manufacture small quantities of the listed controlled substances as radiolabeled compounds.

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 September 7, 1999.

Dated: June 23, 1999.

John H. King,

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

[FR Doc. 99–17097 Filed 7–6–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; 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 § 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on April 1, 1999, Applied Science Labs, Inc., A Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Heroin (9200)	I
Cocaine (9041)	II
Codeine (9050)	II
Meperidine (9230)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to import these controlled substances for the manufacture of reference standards.

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 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, 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 then August 6, 1999.

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 at 40 FR 43745-46 (September 23, 1975), all applicants for registration to import the 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 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: June 23, 1999.

John H. King,

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

[FR Doc. 99–17095 Filed 7–6–99; 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 April 27, 1999, Damocles10, 3529 Lincoln Highway, Thorndale, Pennsylvania 19372, 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
Codeine-N-oxide (9053)	I
Heroin (9200)	I
Morphine-N-oxide (9307)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Phencyclidine (7471)	II
Codeine (9050)	II
Morphine (9300)	II

The firm plans to manufacture the listed controlled substances for the purpose of deuterium labeled internal standards for distribution to analytical laboratories.

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 September 7, 1999.

Dated: June 23, 1999.

John H. King,

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

[FR Doc. 99–17096 Filed 7–6–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Registration

By Notice dated February 23, 1999, and published in the **Federal Register** on March 5, 1999, (64 FR 10724), Ganes Chemicals, Inc., Industrial Park Road, Pennsville, New Jersey 08070, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methylphenide (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Methadone (9250)	II
Methadone-intermediate (9254)	II
Dextropropoxphene, bulk (non-dosage forms) (9273).	II

The firm plans to manufacture the listed controlled substances for distribution as bulk products to its customers.

DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Ganes Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Ganes Chemicals, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. 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 classes of controlled substances listed above is granted.

Dated: June 23, 1999.

John H. King,

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

[FR Doc. 99–17094 Filed 7–6–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Registration

By notice dated March 19, 1999, and published in the **Federal Register** on April 9, 1999, (64 FR 17417), Roberts Laboratories, Inc., 4 Industrial Way East, Eatontown, New Jersey 07724, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of propiram (9649), a basic class of controlled substance listed in Schedule I.

The firm plans to import the propiram for product development.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Roberts Laboratories, Inc. to import propiram is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Roberts Laboratories, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: June 23, 1999.

John H. King,

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

[FR Doc 99–17093 Filed 7–6–99; 8:45 am]

BILLING CODE 4410–09–M