

## DEPARTMENT OF THE INTERIOR

## National Park Service

**Availability of Plan of Operations and Environmental Assessment; Continuing Operation of Four Gas Wells, Saltwater Disposal Well and Production Facility; Vector Energy Corporation Padre Island National Seashore, Kleberg County, TX**

Notice is hereby given in accordance with Section 9.52(b) of Title 36 of the Code of Federal Regulations that the National Park Service has received from Vector Energy Corporation a Plan of Operations for the continuing operation of four gas wells, a saltwater disposal well, and production facility located within Padre Island National Seashore, in Kleberg County, Texas.

The Plan of Operations and Environmental Assessment are available for public review and comment for a period of 30 days from the publication date of this notice in the Office of the Superintendent, Padre Island National Seashore, 20301 Park Road 22, Corpus Christi, Texas. Copies are available from the Superintendent, Padre Island National Seashore, Post Office Box 181300, Corpus Christi, Texas 78480-1300, and will be sent upon request.

If you wish to comment, you may submit your comments by mailing them to the post office address provided above, or, you may hand-deliver comments to the park at the street address provided above. Our practice is to make comments, including names and home addresses of respondents, available for public review during regular business hours. Individual respondents may request that we withhold their home address from the decisionmaking record, which we will honor to the extent allowable by law. There also may be circumstances in which we would withhold from the decisionmaking record a respondent's identity, as allowable by law. If you wish us to withhold your name and/or address, you must state this prominently at the beginning of your comment. However, we will not consider anonymous comments. We will make all submissions from organizations or businesses, and from individuals identifying themselves as representatives or officials of organizations or businesses, available for public inspection in their entirety.

Dated: October 8, 1999.

**Jock F. Whitworth,**  
Superintendent, Padre Island National Seashore.  
[FR Doc. 99-27120 Filed 10-15-99; 8:45 am]  
BILLING CODE 4310-70-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 September 1, 1999, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810, 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
Methaqualone (2565) .....	I
Dimethyltryptamine (7435) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Pentobarbital (2270) .....	II
Secobarbital (2315) .....	II
Phencyclidine (7471) .....	II
Phenylacetone (8501) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Benzoylcegonine (9180) .....	II
Methadone (9250) .....	II
Dextropropoxyphene, bulk (non-dosage forms) (9273) .....	II
Morphine (9300) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture small quantities of the listed controlled substances to produce isotope labeled standards for drug analysis.

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 December 17, 1999.

Dated: October 8, 1999.

**John H. King,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.  
[FR Doc. 99-27098 Filed 10-15-99; 8:45 am]  
BILLING CODE 4410-09-M

## 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 August 12, 1999, Cedarburg Laboratories, Inc., 870 Bandger Circle, Grafton, Wisconsin 53024, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of propiram (9649), a basic class of controlled substance listed in Schedule I.

The firm plans will manufacture propiram in the process of manufacturing other targeted test compounds for another firm.

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 December 17, 1999.

Dated: October 8, 1999.

**John H. King,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.  
[FR Doc. 99-27099 Filed 10-15-99; 8:45 am]  
BILLING CODE 4410-09-M

## DEPARTMENT OF JUSTICE

## Drug Enforcement Administration

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

By Notice dated July 6, 1999, and published in the **Federal Register** on August 2, 1999, (64 FR 41969), Chiragene, Inc., 7 Powder Horn Drive, Warren, New Jersey 07059, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The firm plans to import the phenylacetone to manufacture amphetamine.

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 Chiragene, Inc. to import phenylacetone 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 Chiragene, 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, Section 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: October 8, 1999.

**John H. King,**

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

[FR Doc. 99-27097 Filed 10-15-99; 8:45 am]

BILLING CODE 4410-09-M

## DEPARTMENT OF JUSTICE

### 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 § 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on June 21, 1999, Chirex Technology Center, Inc., DBA Chirex Cauldron, 383 Phoenixville Pike, Malvern, Pennsylvania 19355, made application to the Drug Enforcement Administration to be registered as an importer of phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The firm plans to import the phenylacetone for the manufacture of amphetamine.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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 than November 17, 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 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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: October 8, 1999.

**John H. King,**

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

[FR Doc. 99-27100 Filed 10-15-99; 8:45 am]

BILLING CODE 4410-09-M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is, notice that on June 21, 1999, Chirex Technology Center, Inc., DBA Chirex Cauldron, 383 Phoenixville Pike, Malvern, Pennsylvania 19355, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Amphetamine (1100), a basic class of controlled substance listed Schedule II.

The firm plans to bulk manufacture amphetamine salts for product development.

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 December 17, 1999.

Dated: October 8, 1999.

**John H. King,**

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

[FR Doc. 99-27101 Filed 10-15-99; 8:45 am]

BILLING CODE 4410-09-M

## DEPARTMENT OF JUSTICE

### 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 is 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 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on August 25, 1999, Glaxo Wellcome Inc., Attn: Jeffrey A. Weiss, 1011 North Arendell Avenue, P.O. Box 1271, Zebulon, North Carolina 27597-2309, made application by renewal to the Drug Enforcement Administration to be registered as an importer of remifentanyl (9739), a basic class of controlled substance listed in Schedule II.

The remifentanyl is being imported for the production of Ultiva dosage forms and for research and new product development.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the