

Drug	Schedule
Methamphetamine (1105) .....	II
Amobarbital (2125) .....	II
Secobarbital (2315) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Benzoylcegonine (9180) .....	II
Hydrocodone (9193) .....	II
Levorphanol (9220) .....	II
Methadone (9250) .....	II
Dextropropoxyphene, bulk (non-dosage forms) (9273) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Oxymorphone (9652) .....	II
Alfentanil (9737) .....	II
Fentanyl (9801) .....	II

The firm plans to import small reference standard quantities of finished commercial product from its sister company in Switzerland for sale to its customers for drug testing and pharmaceutical research and development.

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, DC 20537, Attention DEA Federal Register Representative (CCR), and must be filed no later than March 10, 2003.

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: January 27, 2003.  
**Laura M. Nagel,**  
*deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. 03-2913 Filed 2-5-03; 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 July 24, 2002, National Center for Natural Products Research-NIDA MProject University of Mississippi, 135 Coy Waller Complex, University, Mississippi 38677, 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
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	I

The firm will cultivate marijuana for the National Institute of Drug Abuse for research approved by the Department of Health and Human Services.

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 7, 2003.

Dated: January 27, 2003.  
**Laura M. Nagel,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. 03-2918 Filed 2-5-03; 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 July 10, 2002, Norac Company, Inc., 405 S. Motor Avenue, Azusa, California 91702, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of tetrahydrocannabinols (7370), a basic class of controlled substance listed in Schedule I.

The firm plans to manufacture bulk tetrahydrocannabinols for formulation into pharmaceutical products.

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 comment 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 7, 2003.

Dated: January 27, 2003.  
**Laura M. Nagel,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. 03-2915 Filed 2-5-03; 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 October 1, 2002, Noramco, Inc., 1440 Olympic Drive, Athens, Georgia 30601, 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
Amphetamine (1100) .....	II
Oxycodone (9143) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The firm plans to support its other manufacturing facility with manufacturing and analytical testing.

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 7, 2003.

Dated: January 27, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-2920 Filed 2-5-03; 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 September 24, 2002, OraSure Technologies, Inc., 1745 Eaton Avenue, Bethlehem, Pennsylvania 18018, 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
Alphamethadol (9605) .....	I
Benzoylecgonine (9180) .....	II
Morphine (9300) .....	II

The firm plans to bulk manufacture the listed controlled substances to be used in-house to manufacture other controlled substances.

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 7, 2003.

Dated: January 27, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-2917 Filed 2-5-03; 8:45 am]

**BILLING CODE 4410-09-M**

**DEPARTMENT OF JUSTICE**

**Office of Justice Programs**

**[OJP(OJP)-1372]**

**Meeting of the Public Safety Officer Medal of Valor Review Board**

**AGENCY:** Office of Justice Programs (OJP), Justice.

**ACTION:** Notice of meeting.

**SUMMARY:** This is an announcement of a meeting of the Public Safety Officer Medal of Valor Review Board to review and discuss the timetable for carrying out the Board's responsibilities for 2003.

**DATES:** The meeting will take place on Thursday, February 13, 2003, from 2 p.m. to 5 p.m. E.S.T.

**ADDRESSES:** The meeting will take place at the Office of Justice Programs, 810 7th St., NW., Washington, DC 20531; Phone: 202-307-5933.

**FOR FURTHER INFORMATION CONTACT:**

Tracy A. Henke, Principal Deputy Assistant Attorney General, Office of Justice Programs, 810 7th Street NW., Sixth Floor, Washington, DC 20531; Phone: (202) 307-5933 (note: this not a toll free number).

**SUPPLEMENTARY INFORMATION:** This meeting will be open to the public and registrations will be accepted on a space available basis. Members of the public who wish to attend the meeting must register at least seven (7) days in advance of the meeting by contacting Ms. Henke at the above address. Access to the meeting will not be allowed without registration. All attendees will be required to sign in at the meeting registration desk. Please bring photo identification and allow extra time prior to the meeting.

Anyone requiring special accommodations should contact Ms. Henke at least seven (7) days in advance of the meeting.

**Authority**

The Public Safety Officer Medal of Valor Review Board is authorized to carry out its advisory function under 42 U.S.C. 15202. 42 U.S.C. 15201 authorizes the President to award the Public Safety Officer Medal of Valor, the

highest national award for valor by a public safety officer.

**Tracy A. Henke,**

*Principal Deputy Assistant Attorney General, Office of Justice Programs.*

[FR Doc. 03-2814 Filed 2-5-03; 8:45 am]

**BILLING CODE 4410-18-P**

**DEPARTMENT OF LABOR**

**Office of the Secretary**

**Privacy Act of 1974; Publication of A New System of Records**

**AGENCY:** Office of the Secretary, Labor.

**ACTION:** Notice of a new system of records.

**SUMMARY:** The Privacy Act of 1974 requires that each agency publish notice of all of the systems of records that it maintains. This document adds a new system of records to the Department's current systems of records. With the addition of the new system of records, the Department will be maintaining 148 systems of records.

**DATES:** Persons wishing to comment on this new system of records may do so by March 18, 2003.

**EFFECTIVE DATE:** Unless there is a further notice in the **Federal Register**, this new system of records will become effective on April 2, 2003.

**ADDRESSES:** Written comments may be mailed or delivered to Robert A. Shapiro, Associate Solicitor, Division of Legislation and Legal Counsel, 200 Constitution Avenue, NW., Room N-2428, Washington, DC 20210 or by e-mail to *miller-miriam@dol.gov*.

**FOR FURTHER INFORMATION CONTACT:** Miriam McD. Miller, Co-Counsel for Administrative Law, Office of the Solicitor, Department of Labor, 200 Constitution Avenue, NW., Room N-2428, Washington, DC 20210, telephone (202) 693-5500.

**SUPPLEMENTARY INFORMATION:** Pursuant to section three of the Privacy Act of 1974 (5 U.S.C. 552a(e)(4)), hereinafter referred to as the Act, the Department hereby publishes notice of a new system of records currently maintained pursuant to the Act. On April 8, 2002, in volume 67 at page 16816 of the **Federal Register**, the Department published a notice of 147 systems of records which are maintained under the Act. The new system of records presented herein, established by the Office of the 21st Century Workforce, is entitled DOL/21st CENTURY-1, *Correspondents With the Office of the 21st Century Workforce*. This system contains information necessary to