such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on August 16, 2007, Clinical Supplies Management, Inc., 4733 Amber Valley Parkway, Fargo, North Dakota 58104, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule I and II:

| Drug | Schedule |
|---|---|
| Tetrahydrocannabinols (7370) | I |
| Sufentanil (9740) | II |

The company plans to import the listed controlled substances for clinical trials, research, analytical purposes, and distribution to its customers.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class of controlled substance may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, VA 22152; and must be filed no later than December 31, 2007.

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 published in the **Federal Register** on September 23, 1975, (40 FR 43745–46), all applicants for registration to import a basic class 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(b), (c), (d), (e), and (f) are satisfied.

Dated: November 19, 2007.

**Joseph T. Rannazzisi,**

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

[FR Doc. E7–23188 Filed 11–29–07; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Importer of Controlled Substances; Notice of Application

Pursuant to 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 registration under 21 U.S.C. 952(a)(2) 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 Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on October 12, 2007, Lipomed Inc., One Broadway, Cambridge, Massachusetts 02142, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule I:

| Drug | Schedule |
|---|---|
| Methcathinone (1237) | I |
| N-ethylamphetamine (1475) | I |
| Gamma Hydroxybutyric Acid (2010). | I |
| 2,5-Dimethoxy-4-[n]-propylthiophenethylamine (7348). | I |

The company plans to import the listed controlled substances for clinical trials, research, analytical purposes, and distribution to its customers.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, VA. 22152; and must be filed no later than December 31, 2007.

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 published in the **Federal Register** on September 23, 1975, (40 FR 43745–46), all applicants for registration to import a basic class 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(b), (c), (d), (e), and (f) are satisfied.

Dated: November 20, 2007.

**Joseph T. Rannazzisi,**

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

[FR Doc. E7–23184 Filed 11–29–07; 8:45 am]

**BILLING CODE 4410–09–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 November 1, 2007, Norac Inc., 405 S. Motor Avenue, P.O. Box 577, Azusa, California 91702–3232, 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 company plans to manufacture the listed controlled substance in bulk for formulation into the pharmaceutical controlled substance Marinol®.

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 pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administrator, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to

Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than January 29, 2008.

Dated: November 20, 2007.

**Joseph T. Rannazzisi,**

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

[FR Doc. E7–23185 Filed 11–29–07; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Office of Justice Programs

**[OJP (OJP) Docket No. 1475]**

### 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 applications for the 2006–2007 Medal of Valor Awards and to discuss upcoming activities. Due to the late scheduling of this meeting, publication 15 days prior to the meeting was not possible. This notice will be published less than 15 days prior to the meeting pursuant to 41 CFR 102–3.105(b). The meeting time and location are located below.

**DATES:** December 13, 2007, 10 a.m. to 3 p.m.

**ADDRESSES:** The meeting will take place at the Office of Justice Programs, 810 7th Street, NW., Washington, DC 20531.

**FOR FURTHER INFORMATION CONTACT:** Greg Joy, Policy Advisor, Bureau of Justice Assistance, Office of Justice Programs, 810 7th Street, NW., Washington, DC 20531, by telephone at (202) 514–1369, toll free (866) 859–2687, or by e-mail at *gregorgy.joy@usdoj.gov.*

**SUPPLEMENTARY INFORMATION:** The Public Safety Officer Medal of Valor Review Board carries out those advisory functions specified in 42 U.S.C. 15202. Pursuant to 42 U.S.C. 15201, the President of the United States is authorized to award the Public Safety Officer Medal of Valor, the highest national award for valor by a public safety officer. The purpose of this meeting is to review applications for the 2006–2007 Public Safety Officer Medal of Valor Awards and to discuss upcoming activities related thereto.

This meeting will be open to the public. For security purposes, members of the public who wish to attend must register at least five (5) days in advance of the meeting by contacting Mr. Joy. All attendees will be required to sign in at the front desk. **Note:** Photo identification will be required for admission. Additional identification documents may be required.

Access to the meeting will not be allowed without prior registration. Anyone requiring special accommodations should contact Mr. Joy at least five (5) days in advance of the meeting.

Dated: November 26, 2007.

**Cybele K. Daley,**

*Acting Assistant Attorney General, Office of Justice Programs.*

[FR Doc. E7–23240 Filed 11–29–07; 8:45 am]

**BILLING CODE 4410–18–P**

## DEPARTMENT OF LABOR

### Office of the Secretary

### Proposed Collection; Comment Request

**ACTION:** Notice.

**SUMMARY:** The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden conducts a pre-clearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995 [44 U.S.C. 3506(c)(2)(A)]. This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. Currently, the Civil Rights Center within the Office of the Assistant Secretary for Administration and Management is soliciting comments concerning the proposed extension of the collection of the Compliance Information Report—29 CFR part 31 (Title VI of the Civil Rights Act), Nondiscrimination—Disability—29 CFR part 32 (section 504 of the Rehabilitation Act), and Nondiscrimination—Workforce Investment Act—29 CFR part 37 (section 188 of the Workforce Investment Act). A copy of the proposed information collection request (ICR) can be obtained by contacting the office listed below in the addresses section of this notice. In addition, a copy of the ICR in alternate formats of large print and electronic file on computer disk are available upon request.

**DATES:** Written comments must be submitted to the office listed in the addresses section below on or before January 30, 2008.

**ADDRESSES:** Comments should be sent to Annabelle T. Lockhart, Director of the Civil Rights Center. Electronic mail is the preferred method of submittal of comments. Comments by electronic mail must be clearly identified as pertaining to the ICR and sent to *civilrightscenter@dol.gov.* Brief comments (maximum of five pages), clearly identified as pertaining to the ICR, may be submitted by facsimile machine (Fax) to (202) 693–6505. Where necessary, hard copies of comments, clearly identified as pertaining to the ICR, may also be delivered to the Civil Rights Center Director at the U.S. Department of Labor, 200 Constitution Ave., NW., Room N–4123, Washington, DC 20210. Because of problems with U.S. Postal Service mail delivery, the Civil Rights Center suggests that those submitting comments by means of the U.S. Postal Service should place those comments in the mail well before the deadline by which comments must be received.

Receipt of submissions, whether by U.S. Postal Service, e-mail, fax transmittal, or other means will not be acknowledged; however, the sender may request confirmation that a submission has been received, by telephoning the Civil Rights Center at the telephone numbers listed below.

Comments received will be available for public inspection during normal business hours at the above address. Persons who need assistance to review the comments will be provided with appropriate aids such as readers or print magnifiers. Copies of the ICR will be made available, upon request, in large print or electronic file on computer disk. Provision of the rule in other formats will be considered upon request. To schedule an appointment to review the comments and/or obtain the ICR in an alternate format contact the Civil Rights Center at (202) 693–6500 (Voice) or (202) 693–6515/16 (TTY). Please note that these are not toll free telephone numbers.

**FOR FURTHER INFORMATION CONTACT:** Julia Tamakloe-Mankata, Civil Rights Center, (202) 693–6519 (Voice) or (202) 693–6515/16 (TTY). Please note that these are not toll free telephone numbers.

**SUPPLEMENTARY INFORMATION:**

#### I. Background

The Compliance Information Report and its information collection is designed to ensure that programs or activities funded in whole or in part by