

California, a federally recognized Indian tribe, which Grant Deed was approved by the United States of America on October 9, 2003, and recorded on October 9, 2003, in the official records of Contra Costa County Recorder Office, Contra Costa, California, as Document No. 2003-0506433-00 (Trust Lands).

The Trust Lands are proclaimed to be an addition to and part of the reservation of the Lytton Rancheria of California under sections 5 and 7 of the Act of June 18, 1934 (48 Stat. 985; 25 U.S.C. § 467). The Trust Lands are further proclaimed to be held in trust and part of the reservation of the Lytton Rancheria of California before October 17, 1988.

This proclamation does not affect title to the land described above, nor does it affect any valid existing easements for public roads and highways, public utilities and for railroads and pipelines and any other rights-of-way or reservations of record.

Dated: June 29, 2004.

**Aurene M. Martin,**

*Principal Deputy Assistant Secretary—Indian Affairs.*

[FR Doc. 04-15820 Filed 7-12-04; 8:45 am]

**BILLING CODE 4310-W7-P**

## INTERNATIONAL TRADE COMMISSION

[USITC SE-04-005]

### Sunshine Act Meeting

**AGENCY HOLDING THE MEETING:** United States International Trade Commission.

**ORIGINAL DATE:** July 13, 2004.

**ORIGINAL TIME:** 11 a.m.

**NEW DATE:** July 15, 2004.

**NEW TIME:** 11 a.m.

**PLACE:** Room 101, 500 E Street SW., Washington, DC 20436, Telephone: (202) 205-2000.

**STATUS:** Open to the public.

In accordance with 19 CFR 201.35(d)(1), the Commission has determined to change the day and time for the meeting of 11 a.m., July 13, 2004 to 11 a.m., July 15, 2004.

Issued: July 9, 2004.

By order of the Commission.

**Marilyn R. Abbott,**

*Secretary to the Commission.*

[FR Doc. 04-15976 Filed 7-9-04; 2:36 pm]

**BILLING CODE 7020-02-P**

## DEPARTMENT OF JUSTICE

### Bureau of Alcohol, Tobacco, Firearms and Explosives

#### Agency Information Collection Activities: Proposed Collection; Comments Requested

**ACTION:** 30-day notice of information collection under review: application to register as an importer of U.S. Munitions Import List articles.

The Department of Justice (DOJ), Bureau of Alcohol, Tobacco, Firearms, and Explosives (ATF) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register**, volume 69, number 24, on page 5578 on February 5, 2004, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until August 12, 2004. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395-5806.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and
- 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:* Reinstatement, without change, of a previously approved collection.

(2) *Title of the Form/Collection:* Application to Register as an Importer of U.S. Munitions Import List Articles.

(3) *Agency Form Number, if Any, and the Applicable Component of the Department of Justice Sponsoring the Collection: Form Number:* ATF F 4587 (5330.4). Bureau of Alcohol, Tobacco, Firearms and Explosives.

(4) *Affected Public Who Will be Asked or Required to Respond, as Well as a Brief Abstract: Primary:* Business or other for-profit. *Other:* None. The purpose of this information collection is to allow ATF to determine if the registrant qualifies to engage in the business of importing a firearm or firearms, ammunition, and the implements of war, and to facilitate the collection of registration fees.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* There will be an estimated 300 respondents, who will complete the form within approximately 30 minutes.

(6) *An estimate of the total burden (in hours) associated with the collection:* There are an estimated 150 total burden hours associated with this collection.

**FOR FURTHER INFORMATION CONTACT:** Brenda E. Dyer, Clearance Officer, United States Department of Justice, Policy and Planning Staff, Justice Management Division, Suite 1600, Patrick Henry Building, 601 D Street, NW., Washington, DC 20530.

Dated: July 8, 2004.

**Brenda E. Dyer,**

*Clearance Officer, United States Department of Justice.*

[FR Doc. 04-15834 Filed 7-12-04; 8:45 am]

**BILLING CODE 4410-FY-P**

## 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 April 29, 2004, Cedarburg Pharmaceuticals, Inc.,

870 Badger Circle, Grafton, Wisconsin 53024, 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
Tetrahydrocannabinols (7370) .....	I
Dihydromorphine (9145) .....	I
Hydromorphone (9150) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture the listed controlled substances 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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than September 13, 2004.

Dated: June 28, 2004.

**William J. Walker,**

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

[FR Doc. 04-15771 Filed 7-12-04; 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 April 29, 2004, Eli-Elsohly Laboratories, Inc., Mahmoud A. Elsohly Ph.D., 5 Industrial Park Drive, Oxford, Mississippi 38655, 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
Tetrahydrocannabinols (7370) .....	I
Cocaine (9041) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II

The firm plans to manufacture the controlled substances for use in analysis and drug test standards.

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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than September 13, 2004.

Dated: June 28, 2004.

**William J. Walker,**

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

[FR Doc. 04-15772 Filed 7-12-04; 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 April 22 and 28, 2004, Noramco, Inc., 1440 Olympic Drive, Athens, Georgia 30601-1645, made application by letter 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
Morphine-N-Oxide (9307) .....	I
Hydromorphone-N-Oxide (9150) ..	II

The firm plans to manufacture small quantities of the Schedule I products for internal testing; the Schedule II product will be manufactured for distribution to a customer.

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: Federal Register Representative, Office

of Chief Counsel (CCD) and must be filed no later than September 13, 2004.

Dated: June 28, 2004.

**William J. Walker,**

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

[FR Doc. 04-15773 Filed 7-12-04; 8:45 am]

**BILLING CODE 4410-09-M**

**DEPARTMENT OF LABOR**

**Mine Safety and Health Administration**

**Proposed Information Collection Request Submitted for Public Comment and Recommendations; Records of Results of Examinations of Self-Rescuers**

**ACTION:** Notice.

**SUMMARY:** The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden conducts a preclearance 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 (PRA95) (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.

**DATES:** Submit comments on or before September 13, 2004.

**ADDRESSES:** Send comments to Melissa Stoehr, Acting Chief, Records Management Branch, 1100 Wilson Boulevard, Room 2134, Arlington, VA 22209-3939. Commenters are encouraged to send their comments on computer disk, or via e-mail to [stoehr.melissa@dol.gov](mailto:stoehr.melissa@dol.gov). Ms. Stoehr can be reached at (202) 693-9827 (voice), or (202) 693-9801 (facsimile).

**FOR FURTHER INFORMATION CONTACT:** Contact the employee listed in the **ADDRESSES** section of this notice.

**SUPPLEMENTARY INFORMATION:**

**I. Background**

The Self-Rescue devices are subjected to harsh in-mine conditions that may result in damage to the device which could cause the device to malfunction or provide less than adequate protection. The 90-day examination of the device is necessary in order to provide for early detection of potential