

requested, the Commission will transmit its report to the Committee by May 14, 2004. The Committee indicated that it intends to make the report public.

**FOR FURTHER INFORMATION CONTACT:** Industry-specific information may be obtained from Jonathan Coleman, Project Leader (202-205-3465 or [jcoleman@usitc.gov](mailto:jcoleman@usitc.gov)) or Warren Payne, Deputy Project Leader (202-205-3317 or [wpayne@usitc.gov](mailto:wpayne@usitc.gov)), Office of Industries, U.S. International Trade Commission, Washington, DC 20436. For information on legal aspects of this investigation, contact William Gearhart of the Office of General Counsel (202-205-3091 or [wgearhart@usitc.gov](mailto:wgearhart@usitc.gov)). Hearing impaired individuals are advised that information on this matter can be obtained by contacting the TDD terminal on (202-205-1810).

### Public Hearing

A public hearing in connection with the investigation will be held at the U.S. International Trade Commission Building, 500 E Street SW, Washington, DC beginning at 9:30 a.m. on December 4, 2003. All persons shall have the right to appear, by counsel or in person, to present information and to be heard. Requests to appear at the public hearing should be filed with the Secretary, United States International Trade Commission, 500 E Street SW, Washington, DC 20436, no later than 5:15 p.m., November 20, 2003. Any prehearing briefs (original and 14 copies) should be filed not later than 5:15 p.m., November 24, 2003, the deadline for filing post-hearing briefs or statements is 5:15 p.m., December 18, 2003. In the event that, as of the close of business on November 20, 2003, no witnesses are scheduled to appear at the hearing, the hearing will be canceled. Any person interested in attending the hearing as an observer or non-participant may call the Secretary (202-205-1806) after November 20, 2003, to determine whether the hearing will be held.

### Written Submissions

In lieu of or in addition to participating in the hearing, interested parties are invited to submit written statements concerning the investigation. Commercial or financial information that a submitter desires the Commission to treat as confidential must be submitted on separate sheets of paper, each clearly marked "Confidential Business Information" at the top. All submissions requesting confidential treatment must conform with the requirements of section 201.6 of the *Commission's Rules of Practice and Procedure* (19 CFR 201.6). All written

submissions, except for confidential business information, will be made available for inspection by interested parties. The Senate Committee on Finance has requested that the Commission prepare a public report (containing no confidential business information). Accordingly, any confidential business information received by the Commission in this investigation and used in preparing the report will not be published in a manner that would reveal the operations of the firm supplying the information. To be assured of consideration by the Commission, written statements relating to the Commission's report should be submitted to the Commission at the earliest practical date and should be received no later than the close of business on December 18, 2003. All submissions should be addressed to the Secretary, United States International Trade Commission, 500 E Street SW, Washington, DC 20436. The Commission's rules do not authorize filing submissions with the Secretary by facsimile or electronic means, except to the extent permitted by section 201.8 of the Commission's Rules, as amended, 67 FR 8036 (Nov. 8, 2002). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at <http://edis.usitc.gov>. Hearing-impaired individuals are advised that information on this matter can be obtained by contacting our TDD terminal on (202) 205-1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202-205-2000.

### List of Subjects

Milk proteins, government intervention, tariffs, and imports.

Issued: June 5, 2003.

By order of the Commission.

**Marilyn R. Abbott,**  
Secretary.

[FR Doc. 03-14792 Filed 6-10-03; 8:45 am]

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## 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: extension of a currently approved collection; Application and Permit for Temporary

### Importation of Firearms and Ammunition by Nonimmigrant Aliens.

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 68, Number 51, and page 12716 on March 17, 2002, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until July 11, 2003. 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) 395-7285.

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:

(1) 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;

(2) 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;

(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.

**Overviews of This Information Collection**

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Application and Permit for Temporary Importation of Firearms and Ammunition by Nonimmigrant Aliens.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: ATF F 6NIA (5330.3D). 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: Individuals or households. Other: none. Abstract: This information collection is needed to determine if the firearms or ammunition listed on the application qualify for importation and to certify that a nonimmigrant alien is in compliance with 18 U.S.C. 922(g)(5)(B). This application will also serve as the authorization for importation.

(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 15,000 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 7,500 estimated total burden hours associated with this collection.

If additional information is required contact: Ms. Brenda E. Dyer, Deputy Clearance Officer, United States Department of Justice, Information Management and Security Staff, Justice Management Division, Suit 1600, Patrick Henry Building, 601 D Street, NW., Washington, DC 20530.

Dated: June 5, 2003.

**Brenda E. Dyer,**

*Deputy Clearance Officer, United States Department of Justice.*

[FR Doc. 03-14620 Filed 6-10-03; 8:45 am]

**BILLING CODE 4410-FB-M**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Pending Registration**

By Notice dated January 27, 2003, and published in the **Federal Register** on February 6, 2003, (68 FR 6183), Houba, Inc., 16235 State Road 17, Culver, Indiana 46511, made application by renewal to the Drug Enforcement Administration (DEA) for registration as

a bulk manufacturer of the basic classes of Schedule II controlled substances Oxycodone (9143) and Hydrocodone (9193). The firm plans to bulk manufacture the controlled substances for sale to its customers for the production of finished dosage form products.

Comments and Objections have been filed with the Drug Enforcement Administration and are currently under review. A final decision regarding the firm's renewal application as a bulk manufacturer is hereby being held pending investigation and resolution of issues raised.

Houba, Inc. is authorized to continue operating under the manufacturer registration issued pursuant to the Notice of Registration published June 13, 2002 (67 FR 40752) pending final approval or denial of the renewal application.

Dated: May 23, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-14737 Filed 6-10-03; 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 October 24, 2002, Noramco, Inc. (formerly Noramco of Delaware, Inc.), 500 Swedes Landing Road, Wilmington, Delaware 19801, made application by renewal and on December 4 and 26, 2002, by letters to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Morphine-N-Oxide .....	I
Codeine-N-Oxide (9053) .....	I
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Oxymorphone (9652) .....	II

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

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 August 11, 2003.

Dated: May 16, 2003.

**Laura M. Nagel,**

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

[FR Doc. 03-14755 Filed 6-10-03; 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 October 3, 2002, Varian, Inc., Lake Forest, 25200 Commercentre Drive, Lake Forest, California 92630-8810, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Phencyclidine (7470) .....	II
1-Piperidincyclohexane-carbonitrile (8603).	II
Benzoylcegonine (9180) .....	II

The firm plans to manufacture small quantities of controlled substances for use in diagnostic products.

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 (CCD) and must be filed no later than August 11, 2003.