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 mintues.

(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-carbonitrille (8603).	II
Benzoylecgoonine (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.