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 permanent exportation of firearms.

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 74, page 19009 on April 17, 2003, 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 7, 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, or facsimile (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:

(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 agency's 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.

Overview 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 Permanent Exportation of Firearms.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection: Form Number:* ATF F 9 (5320.9). Bureau of Alcohol, Tobacco, Firearms and Explosives, United States Department of Justice.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Business or other for-profit. Other: Individual or households. The form is used to obtain permission to export firearms and serves as a vehicle to allow either the removal of the firearm from registration in the National Firearms Registration and Transfer Record or collection of an excise tax. It is used by Federal firearms licensees and others to obtain a benefit.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that 70 respondents will complete a 18 minute form.

(6) *An estimate of the total public burden (in hours) associated with the collection:* There are an estimated 1,050.

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

Dated: July 1, 2003.

Brenda E. Dyer,

Deputy Clearance Officer, Department of Justice.

[FR Doc. 03–17120 filed 7–7–03; 8:45 am]

BILLING CODE 4410–FB–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 March 18, 2003, Applied Science Labs, Division of Alltech Associates Inc., 2701 Carolean Industrial Drive, PO Box 440, State College, Pennsylvania 16801, made renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic class of controlled substances listed below:

Drug	Schedule
Methcathinone (1237)	I
N-Ethylamphetamine (1475)	I
N,N-Dimethylamphetamine (1480)	I
4-Methylaminorex (cis isomer) (1590).	I
Lysergic acid diethylamide (7315)	I
Mescaline (7381)	I
3,4-Methylenedioxyamphetamine (7400).	I
N-Hydroxy-3, 4-methylenedioxyamphetamine (7402).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3/4-Methylenedioxymethamphetamine (7405).	I
N-Ethyl-1- phenylcyclohexylamine (7455).	I
1-(1-Phenylcyclohexyl) pyrrolidine (7458).	I
1-[1- (2-Thienyl) cyclohexyl] piperidine (7470).	I
Dihydromorphine (9145)	I
Normorphine (9313)	I
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexanecarbonitrile (8603).	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Benzoylecgonine (9180)	II
Morphine (9300)	II
Noroxymorphone (9668)	II

The firm plans to manufacture small quantities of the listed controlled substances for reference 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 Assistance 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 8, 2003.

Dated: June 20, 2003

Laura M. Nagel,

Deputy Assistance Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 03–17124 Filed 7–7–03; 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 May 8, 2003, Cody Laboratories, Inc., 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:

Diphenoxylate (9170)—Schedule II
Meperidine (9230)—Schedule II
Oxymorphone (9652)—Schedule II
Sufentanil (9740)—Schedule II

The firm plans to manufacture bulk material 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:* DEA Federal Register Representative (CCD) and must be filed no later than September 8, 2003.

Laura M. Nagel,

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

[FR Doc. 03–17123 Filed 7–7–03; 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 April 21, 2003, Lilly Del Caribe, Inc., Chemical Plant, Kilometer 146.7, State Road 2, Mayaguez, Puerto Rico 00680, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Dextropropoxyphene (9273), a basic class of controlled substance listed in Schedule II.

The firm plans to bulk manufacture product 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 8, 2003.

Dated: June 20, 2003.

Laura M. Nagel,

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

[FR Doc. 03–17125 Filed 7–7–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 27, 2003, and published in the **Federal Register** on February 6, 2003, (68 FR 6184), Noramco, Inc., 1440 Olympic Drive, Athens, GA 30601, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Amphetamine	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.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Noramco, Inc., to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Noramco, Inc., to ensure that the company's registration is consistent with the public interest. This investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: June 20, 2003.

Laura M. Nagel,

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

[FR Doc. 03–17126 Filed 7–7–03; 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 March 14, 2003, Pressure Chemical Company, 3419 Smallman Street, Pittsburgh, Pennsylvania 15201, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of 2, 5-Dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The firm plans to manufacturer the substance for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacturer 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 8, 2003.

Dated: June 20, 2003.

Laura M. Nagel,

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

[FR Doc. 03–17122 Filed 7–7–03; 8:45 am]

BILLING CODE 4410–09–M