previously published in **Federal Register** on December 6, 2002, Volume 67, Number 235, Pages 72701–72702, allowing for a 60 day public comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until March 10, 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.

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.

Overview of This Information Collection

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

(2) *Title of the Form/Collection:* Application for Individual Manufacturing Quota for a Basic Class of Controlled Substances.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: DEA Form 189. Office of Diversion Control, Drug Enforcement Administration, U.S. 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: None.

Abstract: Title 21, United States Code, section 826, and title 21, Code of Federal Regulations (CFR) 1303.22 require that any person who is registered to manufacture any basic class of controlled substances listed in Schedule I or II and who desires to manufacture a quantity of such class must apply on DEA Form 189 for a manufacturing quota for such quantity of such class.

(5) An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: There are an estimated 264 responses, provided by 44 respondents. The estimated time required for the average respondent to respond is 30 minutes.

(6) An estimate of the total public burden (in hours) associated with the collection: There are 132 annual burden hours associated with this collection.

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

Dated: February 3, 2003.

Robert B. Briggs,

Department Clearance Officer, Department of Justice.

[FR Doc. 03–3078 Filed 2–6–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Registration

By Notice dated June 28, 2002, and published in the **Federal Register** on August 7, 2002, (67 FR 51294), Applied Science Labs, Division of Alltech Associates, Inc., 2701 Carolean Industrial Drive, State College, Pennsylvania 16801, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes 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-methylenedioxy-amphetamine (7402).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymeth-amphetamine (7405).	I
N-Ethyl-1-phenylcyclohexylamine (7455).	I
1-(1-Phenylcyclohexyl)pyrrolidine (7458).	I
1-[-(2-Thienyl)cyclohexyl]piperidine (7470).	I
Dihydromorphine (9145)	I
Normorphine (9313)	I
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexane-carbonitrile (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.

No comments or objections were received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Applied Science Labs to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Applied Science Labs on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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 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: January 27, 2003.

Laura M. Nagel,

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

[FR Doc. 03–3049 Filed 2–6–03; 8:45 am]

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