manufacturers and sellers of Swiss watches and watch parts engaged in a conspiracy ''to restrict, eliminate and discourage the manufacture of watches and watch parts in the United States, and to restrain United States imports and exports of watches and watch parts for manufacturing and repair purposes.'' *Id.* The United States also charged that these companies agreed to fix minimum prices for watches and maximum prices for repair parts, regulate the use and distribution of watches and repair parts, boycott those who violated these restrictions. *Id.* The conspiracy came about through the adoption and enforcement of an agreement known as the Collective Convention of the Swiss Watch Industry. ''The purpose of the Collective Convention was to protect, develop and stablize the Swiss watch industry and to impede the growth and competitive watch industries outside of Switzerland.'' *United States* v. *The Watchmakers of Switzerland Information Center, Inc.,* 1963–1 Trade Cas. (CCH) ¶70,600, at 77,426 (S.D.N.Y. Dec. 20, 1962).

The AWA was named as a defendant because, as a trade association whose members included most of the defendant manufacturers and importers, there was concern that the AWA could aid the alleged conspiracy by policing members' conduct and influencing members to participate in the cartel.

Foote was named as a defendant in the Complaint, becuase as an advertising agency and an agent for some of the defendants, there was concern that Foote, similar to the AWA, was policing the alleged conspiracy and thus aiding the defendants in the enforcement of the cartel.

On March 9, 1960, prior to trial, the United States and the defendant importers (not the AWA since it is a trade association, nor Foote since it is an advertising agency) named in the complaint agreed to enter into the Watchmakers Final Judgment in lieu of going to trial. *United States* v. *The Watchmakers of Switzerland Information Center, Inc.,* Trade Reg. Rep. (CCH) ¶69,655 (S.D.N.Y. Mar. 9, 1960). Also on March 9, 1960, the United States and Defendants AWA and Foote agreed to enter into the AWA Final Judgment and the Foote Final Judgment, respectively, in lieu of going to trial. *Id.* Most of the restrictions in the AWA and Foote Final Judgments prohibit conduct that each company, respectively, could have taken to facilitate the conspiracy.

The Department has filed with the Court a memorandum setting forth the reasons why the United States believes that termination of the AWA and Foote Final Judgments would serve the public interest. Copies of the AWA's and Foote's joint motion to terminate, the stipulation containing the United States' tentative consent, the United States' memorandum, and all further papers filed with the Court in connection with the AWA's and Foote's joint motion will be available for inspection at the Antitrust Documents Group, Antitrust Division, Room 215, 325 7th Street, NW., Washington, DC 20004, and at the Office of the Clerk of the United States District Court for the Southern District of New York. Copies of these materials may be obtained from the Antitrust Division upon request and payment of the copying fee set by Department regulations.

Interested persons may submit comments regarding the proposed termination of the AWA and Foote Final Judgments to the United States. Such comments must be received by the Antitrust Division within sixty (60) days and will be filed with the Court by the United States. Comments should be addressed to John R. Read, Chief, Litigation III Section, Antitrust Division, U.S. Department of Justice, 325 7th Street, NW., Suite 300, Washington, DC 20530.

Dorothy B. Fountain,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 06–6625 Filed 8–1–06; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 20, 2006 and published in the **Federal Register** on March 24, 2006, (71 FR 14948), Cerilliant API Services LLC, 811 Paloma Drive, Suite A, Round Rock, Texas 78664, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II; and by letter to modify its name to Austin Pharma LLC. Subsequent to the publication of the Notice of Application, by letter, the company has also requested to withdraw thirty-five drug codes from their initial application request.

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Alphamethadol (9605)	I
Methadone (9250)	II
Methadone intermediate (9254)	II
Levo-alphacetylmethadol (9648)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol as a synthetic intermediate. This controlled substance will be further synthesized to bulk manufacture a synthetic THC (7370). No other activity for this drug code is authorized for this registration.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cerilliant API Services LLC to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cerilliant API Services LLC to ensure that the company's registration is consistent with the public interest. The 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: July 26, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12478 Filed 8–1–06; 8:45 am]

BILLING CODE 4410–09–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 June 22, 2006, Clariant LSM (Missouri) Inc., 2460 W. Bennett Street, or (P.O. Box 1246, zip 65801), Springfield, Missouri 65807–1229, 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 in Schedules II:

Drug	Schedule
Methylphenidate (1724)	II
Phenylacetone (8501)	II
Methadone intermediate (9254)	II

The company plans to manufacture the listed controlled substance in bulk for sale to its customer.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC. 20537, Attention: DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA **Federal Register** Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

Dated: July 26, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12455 Filed 8–1–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) Title 21 of the Code of Federal Regulations (CFR), this is notice that on June 27, 2006, Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of basic drug code (1724) methylphenidate.

The company plans to bulk manufacture methylphenidate for a customer to use in the production of a controlled substance product.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC. 20537, Attention: DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

Dated: July 26, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12457 Filed 8–1–06; 8:45 am]

BILLING CODE 4410–09–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 February 19, 2006, Orasure Technologies, Inc., Lehigh University, Seeley G. Mudd-Building 6, Bethlehem, Pennsylvania 18015, made application by renewal, and by letter, to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II:

Drug	Schedule
Lysergic acid diethylamide (LSD) (7315).	I
4–Methoxyamphetamine (7411)	I
Normorphine (9313)	I
Tetrahydrocannabinols (THC) (7370).	I
Alphamethadol (9605)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Cocaine (9041)	II
Hydromorphone (9150)	II
Benzoylecgonine (9180)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Oxycodone (9143)	II
Meperidine (9230)	II
Methadone (9250)	II
Oxymorphone (9652)	II

The company plans to manufacture the listed controlled substances in bulk to manufacture controlled substance derivatives. These derivatives will be used in diagnostic products created specifically for internal use only.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC. 20537, Attention: DEA **Federal Register** Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA **Federal Register** Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

Dated: July 26, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12459 Filed 8–1–06; 8:45 am]

BILLING CODE 4410–09–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 January 10, 2006, Siegfried (USA), Inc., Industrial Park Road, Pennsville, New Jersey 08070, 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 in Schedule II:

Drug	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the