factors in Title 21, United States Code, Section 823(a) and determined that the registration of Pharmacia & Upjohn Company to manufacture 2,5-Dimethoxyamphetamine is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 CFR 0.100 and 0.104, the Acting 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 class of controlled substance listed above is granted.

Dated: March 31, 1997.

Terrance W. Woodworth,

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

[FR Doc. 97–11925 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 19, 1996, and published in the **Federal Register** on December 24, 1996, (61 FR 67853), Radian International LLC, 8501 North Mopac Blvd., P.O. Box 201088, Austin, Texas 78720, made application by letter to the Drug Enforcement Administration (DEA) to be registered to manufacture small quantities of controlled substances for drug reference standards as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Alpha-Ethyltryptamine (7249)	I
3,4,5-Trimethoxyamphetamine (7390).	I
4-Bromo-2,5-dimethoxyamphetamine (7391).	I
4-Bromo-2,5-dimethoxyphenethylamine (7392).	I
4-Methyl-2,5-dimethoxyamphetamine (7395).	I
2,5-Dimethoxyamphetamine (7396)	I
2,5-Dimethoxy-4-ethylamphetamine (7399).	I
5-Methoxy-3,4-methylenedioxyamphetamine (7401).	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402).	I
Bufotenine (7433)	I
Codeine-N-oxide (9053)	I
Heroin (9200)	I
Morphine-N-oxide (9307)	I
Pholcodine (9314)	I
Alphamethadol (9605)	I
Betcetylmethadol (9607)	I
Betamethadol (9609)	I
Norlevorphanol (9634)	I
Para-Fluorofentanyl (9812)	I
Alpha-methylfentanyl (9814)	I
Acetyl-alpha-methylfentanyl (9815)	I
Beta-hydroxyfentanyl (9830)	I
Beta-hydroxy-3-methylfentanyl (9831).	I
Alpha-Methylthiofentanyl (9832)	I
3-Methylthiofentanyl (9833)	I
Thiofentanyl (9835)	I
Phenmetrazine (1631)	II
Glutethimide (2550)	II
Cocaine (9041)	II
Codeine (9050)	II
Levomethorphan (9210)	II
Levorphanol (9220)	II

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 Radian International LLC to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 CFR 0.100 and 0.104, the Acting 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: March 31, 1997.

Terrance W. Woodworth,

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

[FR Doc. 97–11926 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 17, 1997, Roche Diagnostic Systems, Inc., 1080 U.S. Highway 202, Somerville, New Jersey 08876–3771, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Phencyclidine (7471)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to manufacture small quantities of the listed controlled substances for incorporation in drug of abuse detection kits.

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 above application.

Any such comments or objections may be addressed, in quintuplicate, to the Acting Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than July 7, 1997.

Dated: March 31, 1997.

Terrance W. Woodworth,

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

[FR Doc. 97–11927 Filed 5–7–97; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Immigration and Naturalization Service

Agency Information Collection Activities: Proposed Collection; Comment Request

ACTION: Extenstion of existing collection; Affidavit of support.

The Office of Management and Budget (OMB) approval is being sought for the information collection listed below. This proposed information collection was previously published in the **Federal Register** on March 3, 1997, at 62 FR 9452, allowing for a 60-day public comment period. No comments were received by the Immigration and Naturalization Service. The purpose of this notice is to allow an additional 30 days for public comments. Comments are encouraged and will be accepted until June 9, 1997. This process is conducted in accordance with 5 CFR Part 1320.10.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Regulatory Affairs, Attention: Department of Justice Desk Officer, Washington, DC 20530. Additionally,