

Drug	Schedule
4-Methyl-2,5-dimethoxyamphetamine (7395).	I
2-5-Dimethoxy-4-ethylamphetamine (7399).	I
3,4-Methylenedioxyamphetamine (7400).	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxy-methamphetamine (7405).	I
Alpha-methyltryptamine (AMT) (7432).	I
Bufotenine (7433)	I
Diethyltryptamine (7434)	I
Dimethyltryptamine (7435)	I
Psilocybin (7437)	I
Psilocyn (7438)	I
5-methoxy-N,N-diisopropyltryptamine (5-MeO-DIPT) (7439).	I
N-Ethyl-1-phenylcyclohexylamine (7455).	I
1-(1-Phenylcyclohexyl)pyrrolidine (PCPy) (7458).	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470).	I
Dihydromorphine (9145)	I
Normorphine (9313)	I
Methamphetamine (1105)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
1-Piperidinocyclohexanecarbonitrile (8603).	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Benzoylcegonine (9180)	II
Ethylmorphine (9190)	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 substance 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 2, 2004.

Dated: May 18, 2004.  
**William J. Walker,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. 04-12460 Filed 6-2-04; 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 1, 2004, American Radiolabeled Chemicals, Inc., 101 Arc Drive, St. Louis, Missouri 63146, 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
Gamma hydroxybutyric acid (2010).	I
Dimethyltryptamine (7435)	I
Dihydromorphine (9145)	I
Cocaine (9041)	II
Codeine (9050)	II
Dihydromorphine (9150)	II
Benzoylcegonine (9180)	II
Ecgonine (9180)	II
Meperidine (9230)	II
Metazocine (9240)	II
Morphine (9300)	II
Oxymorphone (9652)	II

The firm plans to bulk manufacture small quantities of the listed controlled substances as radiolabeled compounds.

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 2, 2004.

Dated: May 21, 2004.  
**William J. Walker,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. 04-12467 Filed 6-2-04; 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 March 17, 2004, Boehringer Ingelheim Chemicals Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, 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
Amphetamine (1100)	II
Methylphenidate (1724)	II
Methadone (9250)	II
Methadone Intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Levo-alphaacetylmethadol (9648)	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances for formulation into finished pharmaceuticals.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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 2, 2004.

Dated: May 5, 2004.  
**William J. Walker,**  
*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*  
 [FR Doc. 04-12461 Filed 6-2-04; 8:45 am]

**BILLING CODE 4410-09-M**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated February 4, 2004, and published in the **Federal Register** on February 18, 2004, (69 FR 7655), Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50619, made application by renewal to the Drug