New Mexico MPS), NM 4 over Rio Grande, San Ildefonso vicinity, 97000730
Sierra County, Percha Creek Bridge (Historic Highway Bridges of New Mexico MPS), NM 90 over Percha Cr., Hillsboro, 97000731
Taos County, Rio Grande Gorge Bridge (Historic Highway Bridges of New Mexico MPS), NM 111 over Rio Grande Gorge, Taos vicinity, 97000733

North Carolina

Pitt County, Greenville Tobacco Warehouse Historic District, Roughly bounded by Twelfth, Clark, Ficklen, and Washington Sts., Greenville, 97000726

South Carolina

Anderson County, Boone—Douthit House, 1000 Milwee Creek Rd., Pendleton vicinity, 97000742
Greenville County, Carolina Supply Company, 35 W Court St., Greenville, 97000743
Sumter County, Lenoir Store, 3240 Horatio Rd., Horatio, 97000744
Mason, Charles T., House, 111 Mason Croft, Sumter, 97000745

Tennessee

Giles County, Pulaski Courthouse Square Historic District (Boundary Increase), 114 E. Jefferson St., Pulaski, 97000746

Vermont

Windsor County, Quechee Historic Mill District, Roughly along High, Quechee Main, River, and School Sts., and River, Waterman Hill, Deweys Mill, and Cemetery Rds., Hartford, 97000747

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33 of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 13, 1997, Damocles10, 3529 Lincoln Highway, Thorndale, Pennsylvania 19372, 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
Heroin (9200)	I
Codeine (9050)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Morphine (9300)	II

The firm plans to manufacture the listed controlled substances for the purpose of deuterium labeled internal standards for distribution to analytical laboratories.

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, D.C. 20537, Attention: DEA Federal Register Representative, (CCR), and must be filed no later than August 18, 1997.

Dated: June 9, 1997.

John H. King,

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

[FR Doc. 97–15843 Filed 6–16–97; 8:45 am]
BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33 of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 16, 1997, Dupont Pharmaceuticals, The Dupont Merck Pharmaceutical Co., 1000 Steward Avenue, Garden City, New York 11530, 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
Oxycodone (9143)	II
Hydrocodone (9193)	II
Oxymorphone (9652)	II

The firm plans to manufacture the listed controlled substances to make finished products.

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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than August 18, 1997.

Dated: June 9, 1997.

John H. King,

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

[FR Doc. 97–15844 Filed 6–16–97; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for hearing.

Therefore, in accordance with Section 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on May 19, 1997, Wildlife Laboratories, Inc., 1401 Duff Drive, Suite 600, Ft. Collins, Colorado 80524, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Etorphien Hydrochloride (9059)	II
Carfentanil (9743)	II

The firm plans to import the listed controlled substances to produce finished products for distribution to its customers. There is no domestic source of etorphein hydrochloride or carfentanil.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in