Janesville Sts., Village of Oregon, 00000699
Waukesha County
Needham, Enoch Gardner and Mary Caroline Koch, House, 12713 W. Greenfield Ave., New Berlin, 00000700

A request for *Removal* has been made for the following resources:

Kansas

Kiowa County
Belvidere Medicine River Bridge, (Masonary Arch Bridges of Kansas TR) 0.25 mi. N of Belvidere, Belvidere vicinity, 85001418
Wyandotte County
Huron Building, 905 N. 7th St., Kansas City, 84001243

[FR Doc. 00–13376 Filed 5–26–00; 8:45 am]

BILLING CODE 4310–70–P

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 21, 2000, American Radiolabeled Chemical, Inc., 11624 Bowling Green Drive, St. Louis, Missouri 63146, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of gamma hydroxybutyric acid (2010), a basic class of controlled substance listed in Schedule I.

The firm plans to bulk manufacture small quantities of the listed controlled substance as radiolabeled compound.

Any other such applicant and any person who is prsently 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than July 31, 2000.

Dated: May 19, 2000.

John H. King,

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

[FR Doc. 00–13439 Filed 5–26–00; 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 10, 2000, and published in the **Federal Register** on February 17, 2000, (65 FR 8206), Ansys Diagnostics, Inc., 25200 Commercentre Drive, Lake Forest, California 92630, 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
Phencyclidine (7471)	II
1–Piperid-ino-cyclo-hex-ane-carbo-nitrile (PCC) (8603).	II
Benzoylecgonine (9180)	II

The firm plans to manufacture the listed controlled substances to produce standards and controls for in-vitro diagnostic drug testing systems.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Ansys Diagnostics, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Ansys Diagnostics, Inc. 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby order 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: May 19, 2000.

John H. King,

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

[FR Doc. 00–13437 Filed 5–26–00; 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 February 29, 2000, Chemic Laboratories, Inc., 480 Neponset Street, Building 7C, Canton, Massachusetts 02021, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of cocaine (9041), a basic class of controlled substance listed in Schedule II.

The firm plans to bulk manufacture small quantities of cocaine derivative for a customer.

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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than July 31, 2000.

Dated: May 19, 2000.

John H. King,

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

[FR Doc. 00–13438 Filed 5–26–00; 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 10, 2000, and published in the **Federal Register** on February 17, 2000, (65 FR 8207), Noramco of Delaware, Inc., Division of McNeilab, Inc., which has changed its name to Noramco of Delaware, Inc., Division of Ortho-McNeil, Inc., 500 Old Swedes Landing Road, Wilmington, Delaware 19801, 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
Codeine (9050)	II
Oxycodone (9143)	II