FOR FURTHER INFORMATION CONTACT: The Offices of the Mojave National Preserve, Barstow, California 92311. Tel. (760) 255–8801.

SUPPLEMENTARY INFORMATION: In order to resolve the encroachment of a private residence and ranch headquarters on federal land, it is necessary for the National Park Service to effect a land exchange at Mojave National Preserve, San Bernardino California. Comments received in response to the Federal Register Notice of Realty Action, included requests for a public hearing on the exchange. National Park Service staff will brief those in attendance at the hearing on the history, purpose, and procedures involved in the exchange. For more detailed information on the proposed exchange, see **Federal Register** Notice of Realty Action published April 18, 2000 (Volume 65, Number 75, pages 20831–20832).

Dated: September 12, 2000.

Cynthia L. Ip,

Acting Regional Director, Pacific West Region.

[FR Doc. 00–24503 Filed 9–22–00; 8:45 am]

BILLING CODE 4310–70–P

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 a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on June 19, 2000, Abbott Laboratories, 1776 North Centennial Drive, McPherson, Kansas 67460–1247, made application to the Drug Enforcement Administration to be registered as an importer of remifentanil (9739), a basic class of controlled substance listed in Schedule II.

The firm plans to import the remifentanil to manufacture Ultiva for the U.S. market.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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 accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing 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 October 25, 2000.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import a basic class of any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(a), (b), (c), (d), (e), and (f) are satisifed.

Dated: September 8, 2000.

John H. King,

Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 00–24556 Filed 9–22–00; 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 July 24, 2000, American Radiolabeled Chemical, Inc., 11624 Bowling Green Drive, St. Louis, Missouri 63146, made application by renewal to the Drug Enforcement Administration (DEA) for registratin as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Gamma hydroxybutyric acid (2010).	I
Lysergic acid diethylamide (7315)	I
Dimethyltryptamine (7435)	I
Dihydromorphine (9145)	I
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Hydromorphone (9150)	II
Benzoylecgonine (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 compound.

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: DEA Federal Register Representative (CCR), and must be filed no later than November 24, 2000.

Dated: September 6, 2000.

John H. King,

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

[FR Doc. 00–24560 Filed 9–22–00; 8:45 am]

BILLING CODE 4410–09–M

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 a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on July 5, 2000, Glaxo Wellcome Inc., Attn: Jeffrey A. Weiss, 1011 North Arendell Avenue, P.O. Box 1217, Zebulon, North Carolina 27597–2309, made application by renewal to the Drug Enforcement Administration to be registered as an importer of remifentanil (9739), a basic class of controlled substance listed in Schedule II.

The remifentanil is being imported for the production of Ultiva dosage forms and for research and new product development.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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 accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing 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: DEA Federal Register Representative (CCR), and must be filed no later than October 25, 2000.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic class of any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: September 1, 2000.

John H. King,

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

[FR Doc. 00–24557 Filed 9–22–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated April 21, 2000, and published in the **Federal Register**on April 28, 2000, (65 FR 24986), Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application by letter to the Drug Enforcement Administration (DEA) to be registered ax a bulk manufacturer of gamma hydroxybutyric acid (2010), a basic class of controlled substance listed in Schedule I.

The firm plans to bulk manufacture gamma hydroxybutyric acid for distribution to its customers.

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 Lonza Riverside to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Lonza Riverside 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 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: September 1, 2000.

John H. King,

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

[FR Doc. 00–24561 Filed 9–22–00; 8:45 am]

BILLING CODE 4410–09–M

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 a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 17, 2000, Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Phenylacetone (8501)	II
Fentanyl (9801)	II

The firm plans to import phenylacetone for the production of amphetamine and fentanyl for seed material for the manufacture of fentanyl base.

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 accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing 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: DEA Federal Register Representative (CCR), and must be filed no later than October 25, 2000.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34 (b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic class of any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: September 5, 2000.

John H. King,

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

[FR Doc. 00–24555 Filed 9–22–00; 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 July 28,