States International Trade Commission (Commission) determines, pursuant to section 751(b) of the Tariff Act of 1930 (19 U.S.C. 1675d(b)) (the Act), that revocation of the antidumping duty orders covering certain frozen warmwater shrimp and prawns from India and Thailand would be likely to lead to continuation or recurrence of material injury to an industry in the United States. Certain frozen warmwater shrimp and prawns from India and Thailand are provided for in subheadings 0306.13.00 and 1605.20.10 of the Harmonized Tariff Schedule of the United States.

Background

On December 17, 2004, the Department of Commerce determined that imports of certain frozen and canned warmwater shrimp and prawns from India and Thailand are being sold in the United States at less than fair value (LTFV) within the meaning of section 731 of the Act (19 U.S.C. 1673) (69 FR 76916, 76918, December 23, 2004); and on January 6, 2005 the Commission determined, pursuant to section 735(b)(1) of the Act (19 U.S.C. 1673d(b)(1)), that an industry in the United States was materially injured by reason of imports of such LTFV merchandise. Accordingly, Commerce ordered that antidumping duties be imposed on such imports (70 FR 5143, February 1, 2005).

On January 6, 2005, when the Commission conducted its vote in the original investigations, it stated that it was concerned about the possible impact of the December 26, 2004, tsunami on the shrimp industries of India and Thailand. The tsunami occurred prior to the closing of the record in the original investigations on December 27, 2004. At the time the record closed, however, factual information as to any impact of the tsunami on the ability of producers in India or Thailand to produce and export shrimp was not available. On February 8, 2005, the Commission published a **Federal Register** notice (70 FR 6728) inviting comments from the public on whether changed circumstances exist sufficient to warrant the institution of changed circumstances reviews of the Commission's affirmative determinations concerning certain frozen warmwater shrimp and prawns from India and Thailand.

The Commission instituted the subject investigations (investigation Nos. 751–TA–28–29), effective May 5, 2005, after having reviewed the comments it received in response to that request, and having determined that it had received information which showed changed circumstances sufficient to warrant instituting review investigations and that there was good cause for instituting such review investigations within two years after publication of the orders. Notice of the scheduling of the Commission's investigations and of a public hearing to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** of May 5, 2005 (70 FR 23884). The hearing was held in Washington, DC, on September 14, 2005, and all persons who requested the opportunity were permitted to appear in person or by counsel.

The Commission transmitted its determinations in these investigations to the Secretary of Commerce on November 21, 2005. The views of the Commission are contained in USITC Publication 3813 (November 2005), entitled *Certain Frozen Warmwater Shrimp and Prawns from India and Thailand: Investigation Nos. 751–TA–28–29.*

Issued: November 21, 2005.

By order of the Commission.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E5–6593 Filed 11–28–05; 8:45 am]

BILLING CODE 7020–02–P

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 3, 2005, 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 in Schedules I and II:

Drug	Schedule
Gamma hydroxybutyric acid (2010).	I
Dimethyltryptamine (7435)	I
Dihydromorphine (9145)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Lysergic acid diethylamide (7315).	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylecgonine (9180)	II
Ecgonine (9180)	II
Meperidine (9230)	II
Metazocine (9240)	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II

The company plans to manufacture in bulk, 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 a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Acting Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than January 30, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E5–6609 Filed 11–28–05; 8:45 am]

BILLING CODE 4410–09–P

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 May 6, 2005, Chemic Laboratories, Inc., 480 Neponset Street, Building 7C, Canton, Massachusetts 02021, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in Schedule II.

The company plans to manufacture small quantities of a cocaine derivative for distribution to its customers for the purpose of research.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance

may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than January 30, 2006.

Dated: November 18, 2005.

Joseph T. Rannazzisi,

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

[FR Doc. E5–6602 Filed 11–28–05; 8:45 am]

BILLING CODE 4410–09–P

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 April 18, 2005, Dade Behring Inc., 100 GBE Drive, MS514, Post Office Box 6101, Attention: RA/QS, Newark, Delaware 19714–6101, 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 in Schedules I and II:

Drug	Schedule
Tetrahydrocannabionols (7370)	I
Benzoylecgonine (9180)	II
Morphine (9300)	II

The company plans to produce the listed controlled substances in bulk to be used in the manufacture of reagents and drug calibrator/controls for DEA exempt products.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters,

Attention: DEA Federal Register Representative/ODL, 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than January 30, 2006.

Dated: November 18, 2005.

Joseph T. Rannazzisi,

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

[FR Doc. E5–6603 Filed 11–28–05; 8:45 am]

BILLING CODE 4410–09–P

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 May 13, 2005, Dade Behring, Inc., Regulatory Affairs, Quality Systems, 20400 Mariani Avenue, Cupertino, California 95014, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II:

Drug	Schedule
Tetrahydrocannabionols (7370)	I
Benzoylecgonine (9180)	II
Morphine (9300)	II

The company plans to produce the listed controlled substances in bulk to be used in the manufacture of reagents and drug calibrator/controls for DEA exempt products.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than January 30, 2006.

Dated: November 18, 2005.

Joseph T. Rannazzisi,

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

[FR Doc. E5–6605 Filed 11–28–05; 8:45 am]

BILLING CODE 4410–09–P

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 August 10, 2005, ISP, Freetown Fine Chemicals, Inc., 238 South Main Street, Assonet, Massachusetts 02702, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed in Schedules I and II:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396).	I
Amphetamine (1100)	II
Phenylacetone (8501)	II

The company plans to manufacture phenylacetone to be used in the manufacture of amphetamine for distribution to its customers. The bulk 2,5-dimethoxyamphetamine will be used for conversion into non-controlled substances.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Acting Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than January 30, 2006.