record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov.* Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810.

SUPPLEMENTARY INFORMATION: On May 31, 2006, the Commission instituted an investigation based on a complaint filed by Ajinomoto Heartland LLC (''Heartland'') of Chicago, Illinois under section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337 (71 FR 30958, May 31, 2006). The complaint, as amended and supplemented, alleges violations of section 337 in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain L-lysine feed products and genetic constructs for production thereof by reason of infringement of claims 13, 15–19, and 21–22 of U.S. Patent No. 5,827,698 and claims 1, 2, 15, and 22 of U.S. Patent No. 6,040,160. The complaint further alleges that an industry in the United States exists as required by subsection (a)(2) of section 337. Global Bio-Chem Technology Group Company Ltd.; Changchun Dacheng Bio-Chem Engineering Development Co., Limited; Changchun Baochen Bio-Chem Development Co., Ltd; Changchun Dahe Bio Technology Development Co. Ltd., all of China, and Bio-Chem Technology (HK) Limited of Hong Kong (collectively ''Bio-Chem'') were named respondents in the investigation. Id.

On June 29, 2006, complainant Heartland filed a motion to amend the complaint to add its parent company, Ajinomoto, Inc. (''Ajinomoto'') as a complainant. The motion was supported by the Commission investigative attorney and Bio-chem. On July 11, 2006, the ALJ granted complainant's motion, finding that complainant had demonstrated good cause for adding Ajinomoto as a complainant at this time. No petitions for review of the ID were filed.

This action is taken under the authority of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) and section 210.42(h) of the Commission's Rules of Practice and Procedure (19 CFR 210.42(h)).

Issued: July 25, 2006.

By order of the Commission.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E6–12144 Filed 7–28–06; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Inv. No. 337–TA–560]

In the Matter of Certain Nor and Nand Flash Memory Devices and Products Containing Same; Notice of Correction

AGENCY: U.S. International Trade Commission.

ACTION: Correction of the notice of investigation for the above-captioned investigation.

SUMMARY: On February 13, 2006, the Commission published the notice of investigation for the above-captioned investigation under Section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337). 71 FR 77576. The Commission hereby gives notice of the following corrections to that notice: (1) In the section labeled **SUMMARY,** ''flash memory devices'' should read ''flash memory devices and products containing same,'' and (2) in the section labeled Scope of Investigation, ''flash memory devices'' should read ''flash memory devices or products containing same.'' The Commission expects that the administrative law judge will extend the target date for completion of the investigation to the extent necessary to avoid any prejudice to any of the parties.

FOR FURTHER INFORMATION CONTACT: Clint Gerdine, Esq., telephone 202–708–2310, Office of the General Counsel, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436. Copies of all nonconfidential documents filed in connection with this investigation are or will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (*http://www.usitc.gov*). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS–ON–LINE) at *http://edis.usitc.gov.* Hearing-impaired persons are advised that information on the matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810.

Issued: July 24, 2006.

By order of the Commission.

Marilyn R. Abbott.

Secretary to the Commission.

[FR Doc. E6–12143 Filed 7–28–06; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Project Management Institute

Notice is hereby given that, on June 14, 2006, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), Project Management Institute (''PMI'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the name and principal place of business of the standards development organization, and (2) the nature and scope of its standards development activities. The notifications were filed for the purpose of invoking the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the name and principal place of business of the standards development organization is: Project Management Institute, Newtown Square, PA. The nature and scope of PMI's standards development activities are to develop standards for the project management profession that are valued by PMI members, the marketplace and other stakeholders. More details regarding PMI's standards development activities can be found at *http://www.pmi.org.*

Dorothy B. Fountain,

Deputy Director of Operations Antitrust Division.

[FR Doc. 06–6569 Filed 7–28–06; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 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 21 U.S.C. 952(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 21 CFR 1301.34(a), this is notice that on May 25, 2006, Aptuit, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, made application by letter to the Drug

Enforcement Administration (DEA) for registration as an importer of Marihuana (7360), a basic class of controlled substance in Schedule I.

The company plans to import a finished pharmaceutical product containing cannabis extracts in dosage form for packaging for a clinical trial study.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/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 August 30, 2006.

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 published in the **Federal Register** on September 23, 1975, (40 FR 43745–46), all applicants for registration to import a basic class of any controlled substance listed 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 301.34(b), (c), (d), (e) and (f) are satisfied.

Dated: July 25, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12171 Filed 7–28–06; 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 October 28, 2005, MGI Pharma, 6611 Tributary Street, Baltimore, Maryland 21224, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in Schedules II.

The company plans to manufacture a cocaine derivative to be used in domestic and foreign clinical research studies.

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 should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/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 September 29, 2006.

Dated: July 25, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12172 Filed 7–28–06; 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 March 22, 2006, Penick Corporation, 33 Industrial Park Road, Pennsville, New Jersey 08070, 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 Schedule II:

Drug	Schedule
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II

The company plans to manufacture the listed controlled substances as bulk controlled substance intermediates for distribution to its customers for further manufacture or to manufacture pharmaceutical dosage forms.

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 should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/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 September 29, 2006.

Dated: July 25, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12173 Filed 7–28–06; 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 January 26, 2006, Roche Diagnostics Operations Inc., Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, 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 Schedule I and II:

Drug	Schedule
Lysergic Acid Diethylamide (7315)	I
Tetrahydrocannabinol (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Ecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II