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