filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Keithley Instruments, Inc., Solon, OH; and PLX Technology, Sunnyvale, CA have been added as parties to this venture. Also, Mapsuka Industries Co., Ltd., Taipei, TAIWAN has withdrawn as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and PXI Systems Alliance, Inc. intends to file additional written notifications disclosing all changes in membership.

On November 22, 2000, PXI Systems Alliance, Inc. filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on march 8, 2001 (66 FR 13971).

The last notification was filed with the Department on October 5, 2006. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on November 22, 2006 (71 FR 67642).

Patricia A. Brink,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 07–319 Filed 1–24–07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Correction to Notice of Application

The Drug Enforcement Administration (DEA) is hereby correcting a notice of application that appeared in the **Federal Register** on January 23, 2006 (71 FR 3545). That document announced the application of Cody Laboratories, Inc., to be registered as an importer of raw opium, poppy straw, and concentrate of poppy straw.

The January 23, 2006, notice of application incorrectly stated that ''[a]ny 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.'' Correctly stated, under the Controlled Substances Act (CSA) and DEA regulations, applications to import narcotic raw materials, including raw opium, poppy straw, and concentrate of poppy straw, are not required to be published in the **Federal Register**. Further, the notice of application, although not required to be published at all, should have stated that ''bulk manufacturers'' of raw opium, poppy straw, or concentrate of poppy straw may file a written request for a hearing. As explained in the Correction to Notice of Application pertaining to Rhodes Technologies published today, since there are no domestic bulk manufacturers of narcotic raw materials registered with DEA, no registrant has a statutory or regulatory right to a hearing on the application. For the reasons set forth therein, I correct the Notice of Application dated January 23, 2006. I direct the Administrative Law Judge to remove from the agency's administrative docket the hearing on the application of Cody Laboratories, Inc. to be registered as an importer of narcotic raw materials.

Dated: January 18, 2007.

Michele M. Leonhart,

Deputy Administrator.

[FR Doc. E7–1052 Filed 1–24–07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Correction to Notice of Application

The Drug Enforcement Administration (DEA) is hereby correcting a notice of application that appeared in the **Federal Register** on April 17, 2006 (71 FR 20729). That document announced the application of Rhodes Technologies to be registered as an importer of raw opium and concentrate of poppy straw. This is the second correction to the original notice of application. This document augments the correction which was published in the **Federal Register** on May 22, 2006 (71 FR 29354).

The April 17, 2006, notice of application incorrectly stated that ''[a]ny 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.'' Correctly stated, under the Controlled Substances Act (CSA) and DEA regulations, applications to import narcotic raw materials, including raw opium and concentrate of poppy straw, are not required to be published in the **Federal Register**. Further, the notice of application, although not required to be published at all, should have stated that ''bulk manufacturers'' of raw opium or concentrate of poppy straw may file a written request for a hearing. As explained below, since there are no domestic bulk manufacturers of narcotic raw materials registered with DEA, no registrant has a statutory or regulatory right to a hearing on the application.

In response to the notice, several importers of narcotic raw materials who also hold manufacturing registrations (but not as ''bulk manufacturers'' of narcotic raw materials) requested a hearing on the application. DEA's Administrative Law Judge (ALJ) accepted the requests for hearings and placed the case on DEA's administrative hearing docket. This correction notifies the applicant, the public, and those importers/manufacturers that requested a hearing that DEA is denying the requests for hearing and dismissing the case on the agency's administrative docket.

Statutory and Regulatory Provisions

As set forth in 21 U.S.C. 958(i), the Attorney General (by delegation, the Administrator and Deputy Administrator of DEA) [1] shall, prior to issuing an importer registration 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.'' (Emphasis added.) Thus, the CSA contemplates that only ''bulk manufacturers'' shall be entitled to hearing on an application to import a schedule I or II controlled substance and, further, that only those who are registered to bulk manufacture the particular substance that the applicant seeks to import. Accordingly, if no one is registered to bulk manufacture the substance that the applicant seeks to import, no one is entitled to a hearing on that application.

DEA's registration database confirms that no person holds a registration as a bulk manufacturer of raw opium, concentrate of poppy straw, or any of the other narcotic raw materials listed in 21 U.S.C. 952(a)(1).[2] Accordingly, the

[1] 21 U.S.C. 871(a); 28 CFR 0.100(b) and 0.104, appendix to subpart R, sec. 12.

[2] When applying for registration, manufacturers are required to complete DEA Form-225, which

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