

Register pursuant to Section 6(b) of the Act on June 28, 2017 (82 FR 29328).

**Patricia A. Brink,**

*Director of Civil Enforcement, Antitrust Division.*

[FR Doc. 2017-19791 Filed 9-15-17; 8:45 am]

BILLING CODE P

## DEPARTMENT OF JUSTICE

### Antitrust Division

#### Notice Pursuant to the National Cooperative Research and Production Act of 1993—*fd.io* Project, Inc.

Notice is hereby given that, on August 24, 2017, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), *fd.io* Project, Inc. (“*fd.io*”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, ARM Ltd., Cambridge, UNITED KINGDOM; Linaro Limited, Cambridge, UNITED KINGDOM; Rubicon Communications LLC dba Netgate, Austin, TX; and CENGN (Centre of Excellence in Next Generation Networks), Ottawa, CANADA, have been added as parties to this venture.

Also, Brocade Communications Systems, Inc., San Jose, CA, has withdrawn as a party to this venture.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 *fd.io* intends to file additional written notifications disclosing all changes in membership.

On May 4, 2016, *fd.io* 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 June 9, 2016 (81 FR 37211).

The last notification was filed with the Department on May 30, 2017. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on June 28, 2017 (82 FR 29329).

**Patricia A. Brink,**

*Director of Civil Enforcement, Antitrust Division.*

[FR Doc. 2017-19789 Filed 9-15-17; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

#### Bulk Manufacturer of Controlled Substances Application: AMPAC Fine Chemicals LLC

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before November 17, 2017.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on August 16, 2017, AMPAC Fine Chemicals LLC, Highway 50 and Hazel Avenue, Building 05001, Rancho Cordova, California 95670 applied to be registered as a bulk manufacturer of levomethorphan (9210), a basic class of controlled substance in schedule II.

The company plans to manufacture the listed controlled substance as an intermediate in the bulk manufacture of other controlled substances for distribution to its customers.

Dated: September 11, 2017.

**Demetra Ashley,**

*Acting Assistant Administrator.*

[FR Doc. 2017-19831 Filed 9-15-17; 8:45 am]

BILLING CODE 4410-09-P

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

#### Importer of Controlled Substances Application: Catalent Centers, LLC

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before October 18, 2017. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before October 18, 2017.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on July 25, 2017, Catalent Centers, LLC, 10245 Hickman Mills Drive, Kansas City, Missouri 64137 applied to be registered as an importer of the following basic classes of controlled substances: