Dated: April 3, 2018.

**Elizabeth K. Appel,**

*Director, Office of Regulatory Affairs and Collaborative Action—Indian Affairs.*

[FR Doc. 2018–07122 Filed 4–6–18; 8:45 am]

**BILLING CODE 4337–15–P**

## DEPARTMENT OF JUSTICE

### Antitrust Division

### Notice Pursuant to the National Cooperative Research and Production Act of 1993—UHD Alliance, Inc.

Notice is hereby given that, on March 8, 2018, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), UHD Alliance, Inc. (''UHD Alliance'') 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, Google, Inc., Mountain View, CA; Teledyne LeCroy, Elgin, IL; and Synaptics, San Jose, CA, have been added as parties to this venture.

Also, HDAnywhere Ltd., Malvern, UNITED KINGDOM; Quantum Data, Inc., Elgin, IL; and Sky UK Ltd., Isleworth, UNITED KINGDOM, have withdrawn as parties to this venture.

In addition, the following members have changed their names: Koninklijke Philips N.V. to Philips International B.V.–IP&S, Eindhoven, NETHERLANDS; and DTS, Inc., to Xperi Corporation, Calabasas, CA.

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 UHD Alliance intends to file additional written notifications disclosing all changes in membership.

On June 17, 2015, UHD Alliance 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 July 17, 2015 (80 FR 42537).

The last notification was filed with the Department on December 15, 2017. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on February 12, 2018 (83 FR 6051).

**Patricia A. Brink,**

*Director of Civil Enforcement, Antitrust Division.*

[FR Doc. 2018–07129 Filed 4–6–18; 8:45 am]

**BILLING CODE 4410–11–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–392]**

### Importer of Controlled Substances Application: United States

**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 on or before May 9, 2018. Such persons may also file a written request for a hearing on the application on or before May 9, 2018.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All request for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrissette 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 delegated 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 March 1, 2018, United States Pharmacopeial Convention, 12601 Twinbrook Parkway, Rockville, MD, 20852 applied to be registered as an importer of the following basic classes of controlled substances:

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Cathinone | 1235 | I |
| Methaqualone | 2565 | I |
| Lysergic acid diethylamide | 7315 | I |
| 4-Methyl-2,5-dimethoxyamphetamine | 7395 | I |
| 4-Methoxyamphetamine | 7411 | I |
| Codeine-N-oxide | 9053 | I |
| Difenoxin | 9168 | I |
| Heroin | 9200 | I |
| Morphine-N-oxide | 9307 | I |
| Norlevorphanol | 9634 | I |
| Methamphetamine | 1105 | II |
| Phenmetrazine | 1631 | II |
| Methylphenidate | 1724 | II |
| Amobarbital | 2125 | II |
| Pentobarbital | 2270 | II |
| Secobarbital | 2315 | II |
| Glutethimide | 2550 | II |
| Phencyclidine | 7471 | II |
| Phenylacetone | 8501 | II |
| Alphaprodine | 9010 | II |
| Anileridine | 9020 | II |
| Cocaine | 9041 | II |