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Ltd., Malvern, UNITED KINGDOM; Quantum Data, Inc., Elgin, IL; and Sky UK Ltd., Isleworth, UNITED KINGDOM, have withdrawn as parties to this venture.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

Controlled substance	Drug code	Schedule
Dihydrocodeine	9120	II
Diphenoxylate	9170	II
Levomethorphan	9210	II
Levorphanol	9220	II
Meperidine	9230	II
Dextropropoxyphene, bulk (non-dosage forms)	9273	II
Thebaine	9333	II
Noroxymorphone	9668	II
Alfentanil	9737	II
Sufentanil	9740	II

The company plans to import the bulk controlled substances for distribution of analytical reference standards to its customers for research and analytical purposes.

Dated: April 2, 2018.

Susan A. Gibson,

Deputy Assistant Administrator.

[FR Doc. 2018–07167 Filed 4–6–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: AMRI Rensselaer, Inc.

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. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

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 21, 2018, AMRI Rensselaer, 33 Riverside Ave., Rensselaer, NY 12144, applied to be registered as an importer of the following basic class of controlled substance:

Controlled substance	Drug code	Schedule
Poppy Straw Concentrate	9670	II

The company plans to import the listed controlled substance to manufacture bulk controlled substance for distribution to its customers.

Dated: April 2, 2018.

Susan A. Gibson,

Deputy Assistant Administrator.

[FR Doc. 2018–07165 Filed 4–6–18; 8:45 am]

BILLING CODE 4410–09–P

NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

National Endowment for the Arts

Arts Advisory Panel Meetings

AGENCY: National Endowment for the Arts, National Foundation on the Arts and Humanities.

ACTION: Notice of meetings.

SUMMARY: Pursuant to the Federal Advisory Committee Act, as amended, notice is hereby given that 2 meetings of the Arts Advisory Panel to the National Council on the Arts will be held by teleconference unless otherwise noted.

DATES: See the **SUPPLEMENTARY INFORMATION** section for individual meeting times and dates. All meetings are Eastern time and ending times are approximate:

ADDRESSES: National Endowment for the Arts, Constitution Center, 400 7th St. SW, Washington, DC 20506.

FOR FURTHER INFORMATION CONTACT: Further information with reference to these meetings can be obtained from Ms. Sherry P. Hale, Office of Guidelines & Panel Operations, National Endowment for the Arts, Washington, DC 20506; *hales@arts.gov,* or call 202/682–5696.

SUPPLEMENTARY INFORMATION: The closed portions of meetings are for the purpose of Panel review, discussion, evaluation, and recommendations on financial assistance under the National Foundation on the Arts and the Humanities Act of 1965, as amended, including information given in confidence to the agency. In accordance with the determination of the Chairman of July 5, 2016, these sessions will be closed to the public pursuant to subsection (c)(6) of section 552b of title 5, United States Code.

The upcoming meetings are:

Literature Fellowships: Translation Projects (review of applications): This meeting will be closed. *Date and time:* May 16, 2018; 3:00 p.m. to 5:00 p.m.