(individual member), Sydney, AUSTRALIA; Douglas McGee (individual member), Columbus, OH; and Nick Ryan (individual member), London, UNITED KINGDOM, have been added as parties to this venture.

Also, Ad-ID, New York, NY; CNN/Turner Broadcasting System, Atlanta, GA; Masstech Group, Inc., Markham, Ontario, CANADA; Video Stream Networks (VSN), Barcelona, SPAIN; and SDVI Corporation, Menlo Park, CA, have withdrawn as parties to this venture.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 Advanced Media Workflow Association, Inc. intends to file additional written notifications disclosing all changes in membership.

On March 28, 2000, Advanced Media Workflow Association, 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 June 29, 2000 (65 FR 40127).

The last notification was filed with the Department on December 23, 2015. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on January 22, 2016 (81 FR 3823).

Patricia A. Brink,

Director of Civil Enforcement, Antitrust Division.

[FR Doc. 2016–08804 Filed 4–15–16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Cody Laboratories, Inc.

ACTION: Notice of registration.

SUMMARY: Cody Laboratories, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Cody Laboratories, Inc. registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated December 4, 2015, and published in the **Federal Register** on December 10, 2015, 80 FR 76709, Cody Laboratories, Inc., 601 Yellowstone Avenue, Cody, Wyoming 82414 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cody Laboratories, Inc. to manufacture the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above-named company is granted registration as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Dihydromorphine (9145)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
4-Anilino-N-phenethyl-4-piperidine (ANPP) (8333).	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers.

Dated: April 11, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–08843 Filed 4–15–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration: Research Triangle Institute

ACTION: Notice of registration.

SUMMARY: Research Triangle Institute applied to be registered as an importer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Research Triangle Institute registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated July 29, 2015, and published in the **Federal Register** on August 4, 2015, 80 FR 46330, Research Triangle Institute, Kenneth S. Rehder, Hermann Building East Institute Drive, Room 106, Research Triangle Park, North Carolina 27709–2194 applied to be registered as an importer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Research Triangle Institute to import the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above-named company is granted registration as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
AB–PINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)1-pentyl-1H-indazole-3-carboxamide) (7023)	I
AB–CHMINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-(cyclohexylmethyl)-1H-indazole-3-carboxamide (7031)	I
AM–2201 (1-(5-Fluoropentyl)-3-(1-naphthoyl) indole) (7201)	I