National Fire Protection Association ("NFPA") has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing additions or changes to its standards development activities. 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, NFPA has provided an updated and current list of its standards development activities, related technical committee and conformity assessment activities. Information concerning NFPA regulations, technical committees, current standards, standards development and conformity assessment activities are publicly available at nfpa.org.

On September 20, 2004, NFPA 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 October 21, 2004 (69 FR 61869).

The last notification was filed with the Department on October 10, 2018. A notice was published in the **Federal** **Register** pursuant to Section 6(b) of the Act on October 22, 2018 (83 FR 53297).

Suzanne Morris,

Chief, Premerger and Division Statistics Unit, Antitrust Division.

[FR Doc. 2019–01489 Filed 2–6–19; 8:45 am] BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: IsoSciences, 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 on or before April 8, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 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 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 November 1, 2017, IsoSciences, LLC, 340 Mathers Road, Ambler, Pennsylvania 19002, applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Cathinone 1235 Methcathinone 1237 Lysergic acid diethylamide 7315 Marihuana 7360 Tetrahydrocannabinols 7370 3.4-Methylenedioxyamphetamine 7400 3.4-Methylenedioxy-N-ethylamphetamine 7404 5-Methoxy-N-N-drimethyltryptamine 7431 Alpha-methyltryptamine 7432 Bufotenine 7433 Diethyltryptamine 7434 Psilocybin 7435 Psilocybin 7438 P-Methoxy-N,N-diisopropyltryptamine 7437 Psilocybin 7438 P-Methoxy-N,N-diisopropyltryptamine 7438 Inhydromorphine 9145 Heroin 9200 Nicocodeine 9309 Nicomorphine 9312 Normorphine 9315	Controlled substance	Drug code	Schedule
Lysergic acid diethylamide	Cathinone	1235	1
Lysergic acid diethylamide	Methcathinone	1237	1
Tetrahydrocannabinols 7370 1 3,4-Methylenedioxyamphetamine 7400 1 3,4-Methylenedioxy-N-ethylamphetamine 7404 1 5-Methoxy-N-U-dimethyltryptamine 7431 1 5-Methoxy-N-N-U-dimethyltryptamine 7432 1 1 1 1 1 1 1 1 1		7315	1
Tetrahydrocannabinols 7370 1 3,4-Methylenedioxyamphetamine 7400 1 3,4-Methylenedioxy-N-ethylamphetamine 7404 1 5-Methoxy-N-U-dimethyltryptamine 7431 1 5-Methoxy-N-N-U-dimethyltryptamine 7432 1 1 1 1 1 1 1 1 1		7360	1
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5-Methoxy-N-N-Idmethyltryptamine 7431 Alpha-methyltryptamine 7432 Bufotenine Diethyltryptamine 7433 I Dimethyltryptamine 7434 I Dimethyltryptamine 7435 I Psilocybin 7437 I Psilocyn 7438 I 5-Methoxy-N,N-diisopropyltryptamine 7439 I Dihydromorphine 9145 I Heroin 9200 I Nicocodeine 9309 I Nicomorphine 9312 I Normorphine 9313 I Thebacon 9315 I Normethadone 9335 I Acryl fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylacrylamide) 9811 I Para-Fluorofentanyl 9813 I Alpha-methylfentanyl 9813 I Alpha-methylfentanyl 9814 I Acetyl-alpha-methylfentanyl 9815 I N-(2-fluorophenyl)-N-(1-phenethylpiperidin-4-yl)-n-phenylacetamide) 9815 I Butryl Fentanyl 9821 I Butryl Fentanyl 9822 I 4-Fluoroisobutryl fentanyl (N-(4-fluorophenyl)-N-(1-phenethylpiperidin-4-yl)isobutyramide) 9824 I 2-methoxy-N-(1-phenethylpiperidin-4-yl)-N-phenylaceta	3,4-Methylenedioxy-N-ethylamphetamine	7404	1
Alpha-methyltryptamine 7432 1	5-Methoxy-N-N-dimethyltryptamine	7431	1
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Furanyl fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylfuran-2-carboxamide)			

Controlled substance	Drug code	Schedule
Beta-hydroxythiofentanyl	9836	I
N-(1-phenethylpiperidin-4-yl)-N-phenyltetrahydrofuran-2-carboxamide	9843	1
Amphetamine	1100	П
Methamphetamine	1105	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	П
Hydromorphone	9150	II
Hydrocodone	9193	II
Isomethadone	9226	II
Methadone	9250	II
Methadone intermediate	9254	II
Morphine	9300	II
Thebaine	9333	II
Levo-alphacetylmethadol	9648	II
Oxymorphone	9652	II
Thiafentanil	9729	II
Alfentanil	9737	П
Sufentanil	9740	II
Carfentanil	9743	П
Fentanyl	9801	II

The company plans to manufacture the listed controlled substances to make reference standards which will be distributed to their customers.

Dated: December 21, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019-01510 Filed 2-6-19; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: The registrant listed below has applied for and been granted a

registration by the Drug Enforcement Administration (DEA) as an importer of schedule I and II controlled substances.

SUPPLEMENTARY INFORMATION: The company listed below applied to be registered as an importer of various basic classes of controlled substances. Information on the previously published notice is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for this notice.

Company	FR Docket	Published
Sigma Aldrich Co., LLC	83 FR 60903	November 27, 2018.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrant to import the applicable basic classes of schedule I and II 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 system, 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 DEA has granted a registration as an importer for schedule I and II controlled substances to the above listed company.

Dated: January 3, 2019.

John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01521 Filed 2–6–19; $8{:}45~\mathrm{am}]$

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Research Triangle Institute

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 March 11, 2019. Such persons may also file a written request for a

hearing on the application on or before March 11, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 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 requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 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