of the listed controlled substances in bulk and to conduct analytical testing in support of the company's primary manufacturing facility in West Deptford, New Jersey. The controlled substances manufactured in bulk at this facility will be distributed to the company's customers.

Dated: November 25, 2014.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2014–29117 Filed 12–10–14; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Sigma Aldrich Research Biochemicals, Inc.

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: Notice of registration.

SUMMARY: Sigma Aldrich Research Biochemicals, Inc., applied to be registered as a manufacturer of certain basic classes of controlled substances. The DEA grants Sigma Aldrich Research Biochemicals, Inc. registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated January 15, 2014, and published in the **Federal Register** on February 4, 2014, 79 FR 6633, Sigma Aldrich Research Biochemicals, Inc. 1–3 Strathmore Road, Natick, Massachusetts 01760–2447, applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted to this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Sigma Aldrich Research Biochemicals, 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 basic classes of controlled substances listed:

Controlled substance	Schedule
Cathinone (1235)	I
Methcathinone (1237)	I
Mephedrone (4-Methyl-N-methylcathinone) (1248).	I
Aminorex (1585)	I
Alpha-ethyltryptamine (7249)	I
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
4-Bromo-2,5-dimethoxyamphetamine (7391).	I
4-Bromo-2,5-dimethoxyphenethylamine (7392).	I
4-Methyl-2,5-dimethoxyamphetamine (7395).	I
2,5-Dimethoxyamphetamine (7396).	I
3,4-Methylenedioxyamphetamine (7400).	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymethamphetamine (MDMA) (7405).	I
Dimethyltryptamine (7435)	I
Psilocybin (7437)	I
5-Methoxy-N, N-diisopropyltryptamine (7439).	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (TCP) (7470).	I
N-Benzylpiperazine (BZP) (7493)	I
MDPV (3,4-Methylenedioxypyrovalerone) (7535).	I
Methylone (3,4-Methylenedioxy-N-methylcathinone) (7540).	I
Heroin (9200)	I
Normorphine (9313)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Nabilone (7379)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Ecgonine (9180)	II
Levomethorphan (9210)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Metazocine (9240)	II
Methadone (9250)	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphacetylmethadol (9648)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Carfentanil (9743)	II
Fentanyl (9801)	II

The company plans to manufacture reference standards.

Dated: November 25, 2014.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2014–29119 Filed 12–10–14; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Employee Benefits Security Administration

Proposed Extension of Information Collection Request Submitted for Public Comment; Revisions to Coverage of Certain Preventive Services Under the Affordable Care Act

AGENCY: Employee Benefits Security Administration, Department of Labor.

ACTION: Notice.

SUMMARY: The Department of Labor (the Department), in accordance with the Paperwork Reduction Act of 1995 (PRA 95) (44 U.S.C. 3506(c)(2)(A)), provides the general public and Federal agencies with an opportunity to comment on proposed and continuing collections of information. This helps the Department assess the impact of its information collection requirements and minimize the reporting burden on the public and helps the public understand the Department's information collection requirements and provide the requested data in the desired format. Currently, the Employee Benefits Security Administration is soliciting comments on the revision of the Coverage of Certain Preventive Services Under the Affordable Care Act information collection to reflect the new option of notifying the Department of Health and Human Services of the respondents' objections to providing coverage in response to the Supreme Court of the United States' interim order in connection with an application for an injunction in the pending case of *Wheaton College* v. *Burwell.* A copy of the information collection request (ICR) may be obtained by contacting the office listed in the **ADDRESSES** section of this notice.

DATES: Written comments must be submitted to the office shown in the Addresses section on or before February 9, 2015.

ADDRESSES: Direct all written comments regarding the information collection request and burden estimates to G. Christopher Cosby, Office of Policy and Research, Employee Benefits Security Administration, U.S. Department of Labor, 200 Constitution Avenue NW., Room N–5718, Washington, DC 20210. Telephone: (202) 693–8410; Fax: (202)