SUPPLEMENTARY INFORMATION: In light of the ongoing Coronavirus (COVID–19) outbreak, the Commission has determined to extend postponement of all section 337 in-person hearings until June 10, 2020. Commission Administrative Law Judges (''ALJ'') are directed to notify all affected parties and to schedule new dates for the hearings as appropriate. ALJs may otherwise conduct their investigations in accordance with their established procedures.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in Part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: April 15, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020–08388 Filed 4–20–20; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–596]

Bulk Manufacturer of Controlled Substances Application: Cedarburg Pharmaceuticals

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before June 22, 2020.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 3, 2018, Cedarburg Pharmaceuticals, 870 Badger Circle, Grafton, Wisconsin 53024–0000 applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols ..	7370	I
Methylphenidate	1724	II
Nabilone	7379	II
4-Anilino-N-phenethyl-4-piperidine (ANPP).	8333	II
Fentanyl	9801	II

The company plans to manufacture bulk active pharmaceutical ingredients (API) for distribution to its customers. In reference to drug code 7370 (Tetrahydrocannabinols) the company plans to bulk manufacture as synthetic. No other activity for this drug code is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–08352 Filed 4–20–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–606]

Importer of Controlled Substances Application: Sigma Aldrich Co., LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturer of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before May 21, 2020. Such persons may also file a written request for a hearing on the application on or before May 21, 2020.

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 a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a 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: In accordance with 21 CFR 1301.34(a), this is notice that on January 27, 2020, Sigma Aldrich Co., LLC, 3500 DeKalb Street, Saint Louis, Missouri 63118 applied to be registered as an importer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Cathinone	1235	I
Methcathinone	1237	I
Mephedrone (4-Methyl-N-methylcathinone)	1248	I
Gamma Hydroxybutyric Acid	2010	I
Tetrahydrocannabinols	7370	I
4-Bromo-2,5-dimethoxyamphetamine	7391	I
4-Bromo-2,5-dimethoxyphenethylamine	7392	I
2,5-Dimethoxyamphetamine	7396	I
3,4-Methylenedioxyamphetamine	7400	I
3,4-Methylenedioxy-N-ethylamphetamine	7404	I
3,4-Methylenedioxymethamphetamine	7405	I
4-Methoxyamphetamine	7411	I
Dimethyltryptamine	7435	I
N-Benzylpiperazine	7493	I
Heroin	9200	I
Normorphine	9313	I
Amobarbital	2125	II
Secobarbital	2315	II
Nabilone	7379	II
Phencyclidine	7471	II
Ecgonine	9180	II
Ethylmorphine	9190	II

Controlled substance	Drug code	Schedule
Levorphanol	9220	II
Meperidine	9230	II
Thebaine	9333	II
Opium, powdered	9639	II
Levo-alphacetylmethadol	9648	II

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis. In reference to drug code 7370 (Tetrahydrocannabinols) the company plans to import synthetic tetrahydrocannabinols. No other activities for these drug codes are authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–08354 Filed 4–20–20; 8:45 am]

BILLING CODE 4410–09–P

NATIONAL ARCHIVES AND RECORDS ADMINISTRATION

[NARA–2020–036]

Freedom of Information Act (FOIA) Advisory Committee; Meeting

AGENCY: Office of Government Information Services (OGIS), National Archives and Records Administration (NARA).

ACTION: Notice of Federal Advisory Committee meeting.

SUMMARY: We are announcing an upcoming Freedom of Information Act (FOIA) Advisory Committee meeting in accordance with the Federal Advisory Committee Act and the second United States Open Government National Action Plan.

DATES: The meeting will be on May 1, 2020, from 10:00 a.m. to 1:30 p.m. EST. You must register by midnight EST April 28, 2020, to attend the meeting.

Location: This meeting will be held virtually. Instructions on how to access will be sent to those who register according to instructions below.

FOR FURTHER INFORMATION CONTACT: Kirsten Mitchell, Designated Federal Officer for this committee, by email at *foia-advisory-committee@nara.gov* or by telephone at 202.741.5770.

SUPPLEMENTARY INFORMATION:

Agenda and meeting materials: This is the eighth meeting of the third committee term. The Committee will discuss the report of the committee and consider additional recommendations from the Committee's three subcommittees, focusing on FOIA vision, time/volume, and records management. We will post meeting materials online at *https://www.archives.gov/ogis/foia-advisory-committee/2018-2020-term/meetings.*

Procedures: This virtual meeting is open to the public. You must register through Eventbrite in advance if you wish to attend, and you must provide an email address so that we can provide information to you on accessing the meeting online. Because of scheduling complications related to the COVID–19 situation and notice processing time, we have not been able to provide the full 15-day notice before this advisory committee meeting. We apologize for the inconvenience. To request additional accommodations (*e.g.,* a transcript), email *foia-advisory-committee@nara.gov* or call 202.741.5775. Members of the media who wish to register, those who are unable to register online, and those who require special accommodations, should contact Kirsten Mitchell (contact information listed above).

Maureen MacDonald,

Designated Committee Management Officer.

[FR Doc. 2020–08357 Filed 4–20–20; 8:45 am]

BILLING CODE 7515–01–P

NATIONAL ARCHIVES AND RECORDS ADMINISTRATION

[NARA–20–0010; NARA–2020–035]

Records Schedules; Availability and Request for Comments

AGENCY: National Archives and Records Administration (NARA).

ACTION: Notice of availability of proposed records schedules; request for comments

SUMMARY: The National Archives and Records Administration (NARA) publishes notice of certain Federal agency requests for records disposition authority (records schedules). We publish notice in the **Federal Register** and on regulations.gov for records schedules in which agencies propose to dispose of records they no longer need to conduct agency business. We invite public comments on such records schedules.

DATES: We must receive comments by June 5, 2020.

ADDRESSES: You may submit comments by either of the following methods. You must cite the control number, which appears on the records schedule in parentheses after the name of the agency that submitted the schedule.

- *Federal eRulemaking Portal: http://www.regulations.gov.*
- *Mail:* Records Appraisal and Agency Assistance (ACR); National Archives and Records Administration; 8601 Adelphi Road; College Park, MD 20740–6001.

FOR FURTHER INFORMATION CONTACT: Kimberly Keravuori, Regulatory and External Policy Program Manager, by email at *regulation_comments@nara.gov.* For information about records schedules, contact Records Management Operations by email at *request.schedule@nara.gov,* by mail at the address above, or by phone at 301–837–1799.

SUPPLEMENTARY INFORMATION:

Public Comment Procedures

We are publishing notice of records schedules in which agencies propose to dispose of records they no longer need to conduct agency business. We invite public comments on these records schedules, as required by 44 U.S.C. 3303a(a), and list the schedules at the end of this notice by agency and subdivision requesting disposition authority.

In addition, this notice lists the organizational unit(s) accumulating the records or states that the schedule has agency-wide applicability. It also provides the control number assigned to each schedule, which you will need if you submit comments on that schedule. We have uploaded the records schedules and accompanying appraisal memoranda to the regulations.gov docket for this notice as ''other'' documents. Each records schedule contains a full description of the records at the file unit level as well as their proposed disposition. The appraisal memorandum for the schedule includes information about the records.

We will post comments, including any personal information and attachments, to the public docket unchanged. Because comments are public, you are responsible for ensuring that you do not include any confidential or other information that you or a third