ACTION: Notice of Federal Advisory Committee meeting; amended.

SUMMARY: This notice amends the location of the partially closed meeting of the Advisory Committee on Actuarial Examinations previously announced in the **Federal Register** of June 14, 2024.

DATES: July 11, 2024, from 9 a.m. to 5 p.m., and July 12, 2024, from 9:30 a.m. to 3 p.m.

FOR FURTHER INFORMATION CONTACT: Elizabeth Van Osten, Designated Federal Officer, Advisory Committee on Actuarial Examinations, at (202) 312–3648 or *Elizabeth.jvanosten@irs.gov.*

SUPPLEMENTARY INFORMATION: As published in the **Federal Register** of June 14, 2024 (89 FR 50634), the meeting was to be held at the Internal Revenue Service, 1111 Constitution Avenue NW, Washington, DC 20224. However, due to an unexpected building closure precluding an in-person meeting, the meeting will be held by teleconference instead. There are no other changes to the meeting. Because the circumstances necessitating the change to the venue of the meeting are beyond the control of the Joint Board or the Enrollment of Actuaries, it is unable to provide public notification about the changes, as required by 41 CFR 102–3.150(a).

Dated: July 3, 2024.

Thomas V. Curtin, Jr.,

Executive Director, Joint Board for the Enrollment of Actuaries.

[FR Doc. 2024–14991 Filed 7–5–24; 8:45 am]

BILLING CODE 4830–01–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1389]

Bulk Manufacturer of Controlled Substances Application: Curia Missouri Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Curia Missouri Inc. has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before September 6, 2024. Such persons may also file a written request for a hearing on the application on or before September 6, 2024.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on May 29, 2024, Curia Missouri Inc., 2460 West Bennett Street, Springfield, Missouri 65807–1229, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Phenylacetone	8501	II
Tapentadol	9780	II

The company plans to bulk manufacture the listed controlled substances for internal use intermediates or for sale to its customers. No other activities for these drug codes are authorized for this registration.

Marsha L. Ikner,

Acting Deputy Assistant Administrator.

[FR Doc. 2024–14926 Filed 7–5–24; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1385]

Bulk Manufacturer of Controlled Substances Application: S&B Pharma LLC DBA Norac Pharma

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: S&B Pharma LLC DBA Norac Pharma has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before September 6, 2024. Such persons may also file a written request for a hearing on the application on or before September 6, 2024.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on April 30, 2024, S&B Pharma LLC DBA Norac Pharma, 405 South Motor Avenue, Azusa, California 91702, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I

The company plans to manufacture the above listed controlled substance for internal research and for development purposes as part of the process in seeking Food and Drug Administration approval prior to distribution to customers. No other activity for this drug code is authorized for this registration.

Marsha L. Ikner,

Acting Deputy Assistant Administrator.

[FR Doc. 2024–14910 Filed 7–5–24; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1388]

Importer of Controlled Substances Application: Arizona Department of Corrections

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Arizona Department of Corrections has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before August 7, 2024. Such persons may also file a written request for a hearing on the application on or before August 7, 2024.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must 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. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on June 3, 2024, Arizona Department of Corrections, 1305 East Butte Avenue, ASPC-Florence, Florence, Arizona 85132–9221, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Pentobarbital	2270	II

The facility intends to import the above-listed controlled substance for legitimate needs. This particular controlled substance is not available for the intended legitimate need within the current domestic supply of the United States. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Marsha L. Ikner,

Acting Deputy Assistant Administrator.

[FR Doc. 2024–14918 Filed 7–5–24; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Consent Decree Under the Clean Air Act

On July 1, 2024, the Department of Justice lodged a proposed consent decree with the United States District Court for the Southern District of New York in the lawsuit entitled *United States* v. *Gristede's Foods NY, Inc.,* Civil Action No. 24 Civ. 4981.

The United States filed this lawsuit seeking injunctive relief and civil penalties for violations of the Clean Air Act against defendant Gristede's Foods NY, Inc. (''Gristedes'') for violations of the United States Environmental Protection Agency's (''EPA'') Recycling and Emissions Reduction Rule, 40 CFR part 82, subpart F, for failing to take actions necessary to monitor, prevent, leak, and record refrigerant emissions.

The consent decree requires Gristedes to implement a new Refrigerant Compliance Management Plan; to reduce its company-wide refrigerant leak rates; to repair or replace specified appliances; to convert certain stores to using more advanced refrigerants; and to pay a $400,000 civil penalty to the United States.

The publication of this notice opens a period for public comment on the consent decree. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States* v. *Gristede's Foods NY, Inc.,* D.J. Ref. No. 90–5–2–1–12759. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

To submit comments:	*Send them to:*
By email	*pubcomment-ees.enrd@usdoj.gov.*