Sterling Wisconsin LLC., W130N10497 Washington Drive, Germantown, Wisconsin 53022–4448, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Lysergic Acid Diethylamide.	7315	I
Marihuana Extract	7350	1
Marihuana	7360	1
Tetrahydrocannabinols	7370	1
Mescaline	7381	1
5-Methoxy-N-N-	7431	1
Dimethyltryptamine.		
Psilocybin	7437	1
Oliceridine	9245	П
Thebaine	9333	П
Alfentanil	9737	II

The company plans to bulk manufacture the listed controlled substances to be commercially sold to registered manufacturers/suppliers. In reference to drug codes 7350 (Marihuana Extract), 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

#### Matthew Strait,

Deputy Assistant Administrator. [FR Doc. 2023–03839 Filed 2–23–23; 8:45 am] BILLING CODE P

#### **DEPARTMENT OF JUSTICE**

#### **Drug Enforcement Administration**

[Docket No. DEA-1153]

# Importer of Controlled Substances Application: S&B Pharma LLC

**AGENCY:** Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

**SUMMARY:** S&B Pharma LLC 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 March 27, 2023. Such persons may also file a written request for a hearing on the application on or before March 27, 2023.

**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/OALI, 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 January 10, 2023, S&B Pharma LLC, 405 South Motor Avenue, Azusa, California 91702, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
4-Anilino-N-phenethyl-4-piperidine (ANPP).	8333	II
Tapentadol	9780	II

The company plans to import intermediate forms of Tapentadol (9780) for further manufacturing prior to distribution to its customers. The company plans to import ANPP (8333) to bulk manufacture other controlled substances for distribution to its customers. 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.

#### Matthew Strait,

Deputy Assistant Administrator. [FR Doc. 2023–03829 Filed 2–23–23; 8:45 am] BILLING CODE P

#### **DEPARTMENT OF JUSTICE**

#### **Drug Enforcement Administration**

[Docket No. DEA-1150]

# Bulk Manufacturer of Controlled Substances Application: Scottsdale Research Institute

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

SUMMARY: Scottsdale Research Institute, 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 April 25, 2023. Such persons may also file a written request for a hearing on the application on or before April 25, 2023.

**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 January 10, 2023, Scottsdale Research Institute, 12815 North Cave Creek Road, Phoenix, Arizona 85022, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

The company plans to bulk manufacture the listed controlled substances to support clinical trials and distribution to their customers for research purposes. No other activities for these drug codes are authorized for this registration.

#### Matthew Strait,

Deputy Assistant Administrator. [FR Doc. 2023–03826 Filed 2–23–23; 8:45 am] BILLING CODE P

#### **LEGAL SERVICES CORPORATION**

#### **Sunshine Act Meetings**

TIME AND DATE: The Search Committee for LSC Inspector General (Search Committee) of the Legal Services Corporation Board of Directors will meet virtually on Tuesday, February 28, 2023. The meeting will commence at 11:00 a.m. EST and will continue until the conclusion of the Committee's agenda.

**PLACE:** Public notice of virtual meetings. LSC will conduct the February 28, 2023 meeting via Zoom.

STATUS: Closed.

## MATTERS TO BE CONSIDERED:

#### **Closed Session**

- 1. Approval of Agenda.
- 2. Discuss the interviews of candidates for the position of Legal Services Corporation Inspector General.
- 3. Decide which candidates, if any, to consider further.
- 4. Determine whether to conduct further interviews of selected candidates, discuss the questions the Search Committee would like to ask, and determine the dates of such further interviews.
  - 5. Adjourn.

#### **CONTACT PERSON FOR MORE INFORMATION:**

Cheryl DuHart, Administrative Coordinator, Office of Legal Affairs, at (202) 295–1621. Questions may also be sent by electronic mail to *duhartc@lsc.gov*.

Dated: February 21, 2023.

#### Stefanie Davis,

Senior Associate General Counsel for Regulations.

[FR Doc. 2023–03923 Filed 2–22–23; 8:45 am]

BILLING CODE 7050-01-P

#### POSTAL REGULATORY COMMISSION

# [Docket Nos. MC2023-112 and CP2023-115]

#### **New Postal Products**

**AGENCY:** Postal Regulatory Commission. **ACTION:** Notice.

**SUMMARY:** The Commission is noticing a recent Postal Service filing for the Commission's consideration concerning a negotiated service agreement. This notice informs the public of the filing, invites public comment, and takes other administrative steps.

**DATES:** Comments are due: February 28, 2023.

ADDRESSES: Submit comments electronically via the Commission's Filing Online system at http://www.prc.gov. Those who cannot submit comments electronically should contact the person identified in the FOR FURTHER INFORMATION CONTACT section by telephone for advice on filing alternatives.

**FOR FURTHER INFORMATION CONTACT:** David A. Trissell, General Counsel, at 202–789–6820.

## SUPPLEMENTARY INFORMATION:

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I. Introduction
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# I. Introduction

The Commission gives notice that the Postal Service filed request(s) for the Commission to consider matters related to negotiated service agreement(s). The request(s) may propose the addition or removal of a negotiated service agreement from the Market Dominant or the Competitive product list, or the modification of an existing product currently appearing on the Market Dominant or the Competitive product list.

Section II identifies the docket number(s) associated with each Postal Service request, the title of each Postal Service request, the request's acceptance date, and the authority cited by the Postal Service for each request. For each request, the Commission appoints an officer of the Commission to represent the interests of the general public in the proceeding, pursuant to 39 U.S.C. 505 (Public Representative). Section II also establishes comment deadline(s) pertaining to each request.

The public portions of the Postal Service's request(s) can be accessed via the Commission's website (http://www.prc.gov). Non-public portions of the Postal Service's request(s), if any, can be accessed through compliance

with the requirements of 39 CFR 3011.301.1

The Commission invites comments on whether the Postal Service's request(s) in the captioned docket(s) are consistent with the policies of title 39. For request(s) that the Postal Service states concern Market Dominant product(s), applicable statutory and regulatory requirements include 39 U.S.C. 3622, 39 U.S.C. 3642, 39 CFR part 3030, and 39 CFR part 3040, subpart B. For request(s) that the Postal Service states concern Competitive product(s), applicable statutory and regulatory requirements include 39 U.S.C. 3632, 39 U.S.C. 3633, 39 U.S.C. 3642, 39 CFR part 3035, and 39 CFR part 3040, subpart B. Comment deadline(s) for each request appear in section II.

#### II. Docketed Proceeding(s)

1. Docket No(s).: MC2023–112 and CP2023–115; Filing Title: USPS Request to Add Priority Mail Express International, Priority Mail International, First-Class Package International Service & Commercial ePacket Contract 13 to Competitive Product List and Notice of Filing Materials Under Seal; Filing Acceptance Date: February 17, 2023; Filing Authority: 39 U.S.C. 3642, 39 CFR 3040.130 through 3040.135, and 39 CFR 3035.105; Public Representative: Jennaca D. Upperman; Comments Due: February 28, 2023.

This Notice will be published in the **Federal Register**.

#### Erica A. Barker,

Secretary.

# **SMALL BUSINESS ADMINISTRATION**

[Disaster Declaration #17767 and #17768; California Disaster Number CA-00368]

# Presidential Declaration Amendment of a Major Disaster for Public Assistance Only for the State of California

**AGENCY:** U.S. Small Business Administration. **ACTION:** Amendment 3.

**SUMMARY:** This is an amendment of the Presidential declaration of a major disaster for Public Assistance Only for the State of California (FEMA–4683–DR), dated 01/26/2023.

*Incident:* Severe Winter Storms, Flooding, Landslides, and Mudslides.

<sup>&</sup>lt;sup>1</sup> See Docket No. RM2018–3, Order Adopting Final Rules Relating to Non-Public Information, June 27, 2018, Attachment A at 19–22 (Order No. 4679).