

for a hearing on the application on or before August 8, 2022.

**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 February 2, 2022, Stepan Company, 100 West Hunter Avenue, Maywood, New Jersey 07607–1021, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Cocaine .....	9041	II
Ecgonine .....	9180	II

The company plans to bulk manufacture the listed controlled substances for the internal use intermediates or for sale to its customers.

**Kristi O'Malley,**

*Assistant Administrator.*

[FR Doc. 2022–12198 Filed 6–6–22; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA–1021]

#### Bulk Manufacturer of Controlled Substances Application: Pisgah Laboratories Inc.

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

**ACTION:** Notice of application.

**SUMMARY:** Pisgah Laboratories 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 August 8, 2022. Such persons may also file a written request for a hearing on the application on or before August 8, 2022.

**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 March 9, 2022, Pisgah Laboratories Inc., 3222 Old Hendersonville Highway, Pisgah Forest, North Carolina 28768, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
4-Bromo-2,5-dimethoxyphenethylamine .....	7392	I
Methylone (3,4-Methylenedioxy-N-methylcathinone) .....	7540	I
Difenoxin .....	9168	I
Diphenoxylate .....	9170	II
Levorphanol .....	9220	II
Meperidine .....	9230	II
Methadone .....	9250	II
Remifentanyl .....	9739	II
Tapentadol .....	9780	II

The company plans to bulk manufacture the above-listed controlled substances in bulk for distribution to its customers. No other activities for these drug codes are authorized for this registration.

**Kristi O'Malley,**

*Assistant Administrator.*

[FR Doc. 2022–12197 Filed 6–6–22; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Notice of Proposed Interim Settlement and Crediting Agreement Under Comprehensive Environmental Response, Compensation and Liability Act

As of May 25, 2022, the United States Fish and Wildlife Service (“USFWS”), on behalf of the Department of the Interior, the National Oceanic and Atmospheric Administration (“NOAA”), on behalf of the Department of Commerce, (collectively, the “Trustees”), the Department of Justice, and potentially responsible party (“PRP”) BASF Corporation (“BASF”) signed a proposed non-judicial Interim Settlement and Crediting Agreement

concerning early natural resource restoration work under the Comprehensive Environmental Response, Compensation, and Liability Act (“CERCLA”), 42 U.S.C. 9601 *et seq.*, for a five-acre property in East Newark, New Jersey. The United States contends BASF and other PRPs are liable for natural resource damages under Section 107(a) of CERCLA, 42 U.S.C. 9607(a), concerning the Diamond Alkali Superfund Site, including Newark Bay and the upstream 17 mile stretch of the Passaic River, and the Berry's Creek Study Area, Bergen County, New Jersey (collectively “the Sites”). The proposed agreement facilitates early natural resource restoration work, and provides for credit for accomplished early restoration work, in advance of the