DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–635]

Bulk Manufacturer of Controlled Substances Application: Research Triangle Institute

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 July 6, 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 March 4, 2020, Research Triangle Institute, 3040 East Cornwallis Road, Hermann Building, Room 106, Research Triangle Park, North Carolina 27709, 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

The purpose for the bulk manufacturing of the controlled substance is for the preparation and the sale of small quantities of Tetrahydrocannabinols (7370), which will be manufactured by synthesis for use by customers as analytical reference standards.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–09555 Filed 5–4–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–636]

Bulk Manufacturer of Controlled Substances Application: Patheon API Manufacturing, Inc.

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 July 6, 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 February 21, 2020, Patheon A PI Manufacturing, Inc, 309 Delaware Street, Greenville, South Carolina 29605, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
Alpha-methyltryptamine	7432	I
Thebaine	9333	II
Noroxymorphone	9668	II

The company plans to bulk manufacture the above-listed controlled substances as an Active Pharmaceutical Ingredient (API) for distribution to its customers. No other activities for these drug codes are authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–09556 Filed 5–4–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–626]

Importer of Controlled Substances Application: Alcami Carolinas Corporation

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 4, 2020. Such persons may also file a written request for a hearing on the application on or before June 4, 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 request 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 March 13, 2020, Alcami Carolinas Corporation, 1726 North 23rd Street, Wilmington, North Carolina 28405–1822, applied to be registered as an importer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Psilocybin	7437	I
Psilocyn	7438	I
Thebaine	9333	II
Pentobarbital	2270	II

The company plans to import the listed controlled substances in bulk for the manufacturing of capsules/tablets for Phase II clinical trials. Approval of permit applications will occur only when the registrant's activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA-approved or non-approved finished dosage forms for commercial sale.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–09552 Filed 5–4–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–611]

Importer of Controlled Substances Application: Unither Manufacturing LLC

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 4, 2020. Such persons may also file a written request for a hearing on the application on or before June 4, 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 February 27, 2020, Unither Manufacturing LLC, 331 Clay Road, Rochester, New York 14623, applied to be registered as an importer of the following basic class(es) of a controlled substance:

Controlled substance	Drug code	Schedule
Methylphenidate	1724	II

The company plans to import the listed controlled substance solely for updated analytical testing purposes for European customer requirements. This analysis is required to allow the company to export domestically-manufactured finished dosage forms to foreign markets. Approval of permit applications will occur only when the registrant's activity is consistent with what is authorized under to 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA-approved or non-approved finished dosage forms for commercial sale.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–09514 Filed 5–4–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–634]

Bulk Manufacturer of Controlled Substances Application: Absolute Standards, Inc.

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 July 6, 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 March 12, 2020, Absolute Standards, Inc., 44 Rossotto Drive, Hamden, Connecticut 06514–1335, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled Substance	Drug Code	Schedule
Pentobarbital	2270	II

The company plans to bulk manufacture the above-listed controlled substance for distribution to customers.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–09553 Filed 5–4–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

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

On April 29, 2020, the Department of Justice lodged a proposed Consent Decree with the United States District Court for the Northern District of Illinois in the lawsuit entitled *United States* v. *American Zinc Recycling Corp.,* Case No. 1:20–cv–02582.

The United States filed a Complaint seeking civil penalties and injunctive relief from Defendant American Zinc Recycling Corp. (''AZR'') for alleged violations of the Clean Air Act, 42 U.S.C. 7401–7671q, at its electric arc furnace flue dust recycling facility in Chicago (the ''Facility''). Among other things, the United States alleges that AZR has violated statutory and regulatory requirements limiting particulate matter emissions from the Facility, as well as corresponding requirements in AZR's Clean Air Act permits for the Facility.

When the Complaint was filed, the United States also lodged a proposed Consent Decree that would settle the claims asserted in the Complaint. The proposed Consent Decree would require that AZR implement appropriate injunctive relief to control air pollutant emissions from the Facility, including upgrading multiple bag collectors that filter and remove particulate matter from air exhausted from the Facility. The Consent Decree also assess a $1,054,000 civil penalty. $654,000 of the penalty assessment would be payable on discounted basis under AZR's 2016 Chapter 11 bankruptcy reorganization plan. The remaining $400,000 would be paid in full.

The publication of this notice opens a period for public comment on the proposed Consent Decree. Comments should be addressed to the Principal Deputy Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States* v. *American Zinc Recycling Corp.,* D.J. Ref. No. 90–5–2–1–11205. 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.*
By mail	Principal Deputy Assistant Attorney General U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044–7611.

During the public comment period, the proposed Consent Decree may be examined and downloaded at this Justice Department website: *https://www.justice.gov/enrd/consent-decrees.*

We will provide a paper copy of the proposed Consent Decree upon written request and payment of reproduction costs. Please mail your request and payment to: Consent Decree Library U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044–7611.

Please enclose a check or money order for $18.00 (25 cents per page reproduction cost) payable to the United States Treasury.

Patricia A. McKenna,

Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.

[FR Doc. 2020–09595 Filed 5–4–20; 8:45 am]

BILLING CODE 4410–15–P

DEPARTMENT OF LABOR

Bureau of Labor Statistics

Information Collection Activities; Comment Request

AGENCY: Bureau of Labor Statistics, Department of Labor.

ACTION: Notice of information collection; request for comment.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a pre-clearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995. This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and