

Public Availability of Comments

Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time. While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

Authority

The authority for this action is section 4(f) of the Endangered Species Act of 1973, as amended (16 U.S.C. 1531 et seq.).

Robyn Thorson,

Regional Director, U.S. Fish and Wildlife Service.

[FR Doc. 2021-16521 Filed 8-2-21; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-874]

Bulk Manufacturer of Controlled Substances Application: Purisys, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Purisys, LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

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 October 4, 2021. Such persons may also file a written request for a hearing on the application on or before October 4, 2021.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on June 30, 2021, 1550 Olympic Drive, Athens, Georgia 30601-1602, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Table with 3 columns: Controlled substance, Drug code, Schedule. Rows include Opium, powdered (9639, II) and Opium, granulated (9640, II).

The company plans to bulk manufacture the listed controlled substances as 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. 2021-16500 Filed 8-2-21; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-873]

Bulk Manufacturer of Controlled Substances Application: Cerilliant Corporation

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cerilliant Corporation has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to SUPPLEMENTAL INFORMATION listed below for further drug information.

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 October 4, 2021. Such persons may also file a written request for a hearing on the application on or before October 4, 2021.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on June, 24, 2021, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665-2402, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Large table with 3 columns: Controlled substance, Drug code, Schedule. Lists various controlled substances like 3-Fluoro-N-methylcathinone, Cathinone, Methcathinone, etc., with their respective drug codes and schedules.