membership, which is bound by Indian Preference Act of 1990 (25 U.S.C. 472).

5. Basis for Nominations

If you wish to nominate someone for appointment to the Advisory Board, please do not make the nomination until the person has agreed to have his or her name submitted to the BIE for this purpose. A person can also self-nominate.

6. Nomination Application

Please submit a complete application form and a copy of the nominee's resume or curriculum vitae to the DFO by Wednesday, January 31, 2024. The nomination application form can be found on the BIE website at *https://www.bie.edu/sites/default/files/inline-files/Advisory-Board-Membership-Nomination-Form%20%28Expires%206-30-24%29.pdf.*

7. Information Collection

This collection of information is authorized by OMB Control Number 1076–0179, ''Solicitation of Nominations for the Advisory Board for Exceptional Children,'' with a June 30, 2024, expiration date.

(Authority: 5 U.S.C. ch. 10; 20 U.S.C. 1400 *et seq.*)

Bryan Newland,

Assistant Secretary—Indian Affairs.

[FR Doc. 2023–26518 Filed 12–1–23; 8:45 am]

BILLING CODE 4337–15–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1304]

Bulk Manufacturer of Controlled Substances Application: Element Materials Technology Santa Rosa

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Element Materials Technology Santa Rosa 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 February 2, 2024. Such persons may also file a written request for a hearing on the application on or before February 2, 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 November 1, 2023, Element Materials Technology Santa Rosa, 3331 Industrial Drive, Suite B, Santa Rosa, California 95403–2062, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Oxymorphone	9652	II
Fentanyl	9801	II

The company is a contract manufacturer. At the request of the company's customers, it manufactures derivatives of the listed controlled substances in bulk form. No other activities for these drug codes are authorized for this registration.

Claude Redd,

Acting Deputy Assistant Administrator.

[FR Doc. 2023–26553 Filed 12–1–23; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1300]

Importer of Controlled Substances Application: Caligor Coghlan Pharma Services

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Caligor Coghlan Pharma Services, 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 January 3, 2024. Such persons may also file a written request for a hearing on the application on or before January 3, 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 October 13, 2023, Caligor Coghlan Pharma Services, 1500 Business Park Drive, Unit B, Bastrop, Texas 78602, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Dimethyltryptamine	7435	I

The company plans to import the listed controlled substance as finished dosage units for use in clinical trials. 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-