health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

(i) explain how the articles potentially subject to the requested remedial orders are used in the United States;

(ii) identify any public health, safety, or welfare concerns in the United States relating to the requested remedial orders;

(iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;

(iv) indicate whether complainant, complainant’s licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to the requested exclusion order and/or a cease and desist order within a commercially reasonable time; and

(v) explain how the requested remedial orders would impact United States consumers.

Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the Federal Register. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business, eight calendar days after publication of this notice in the Federal Register. Complainant may file replies to any written submissions no later than by close of business, eight calendar days after the date of publication of any reply that was filed in opposition to the initial determination. Any responses to any replies must be filed by no later than the close of business, eight calendar days after the date of publication of any response filed in opposition to the initial determination in the Federal Register. No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this Investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel, solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary on EDIS. Any person desiring to submit a confidential submission or a document containing confidential information and documents for which confidential treatment is properly sought will be directed to the Secretary to the Commission in confidence must request confidential treatment.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission’s Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission.


Lisa Barton,
Secretary to the Commission.

[FR Doc. 2023–11774 Filed 6–1–23; 8:45 am]

BILLING CODE 7020–02–P


2 All contract personnel will sign appropriate nondisclosure agreements.


DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1208]

Importer of Controlled Substances Application: Veranova, L.P.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Veranova, L.P. 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 July 3, 2023. Such persons may also file a written request for a hearing on the application on or before July 3, 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/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 April 19, 2023, Veranova, L.P., 2003 Nolte Drive, West Deptford, New Jersey 08066–1727, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Veranova, L.P. 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.
### Controlled substance

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coca Leaves ..........</td>
<td>9040</td>
<td>II</td>
</tr>
<tr>
<td>Thebaine .............</td>
<td>9333</td>
<td>II</td>
</tr>
<tr>
<td>Opium, raw ..........</td>
<td>9600</td>
<td>II</td>
</tr>
<tr>
<td>Noroxymorphone ......</td>
<td>9668</td>
<td>II</td>
</tr>
<tr>
<td>Poppy Straw Concentrate.</td>
<td>9670</td>
<td>II</td>
</tr>
<tr>
<td>Fentanyl ............</td>
<td>9801</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import Coca Leaves (9040), Opium, raw (9600), and Poppy Straw Concentrate (9670) in order to bulk manufacture Active Pharmaceutical Ingredients (API) for distribution to its customers. The company plans to also import Thebaine (9333), Noroxymorphone (9668), and Fentanyl (9801) to use as analytical reference standards, both internally and to be sold to their customers to support testing of Veranova, L.P. APIs only. 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. 2023–11740 Filed 6–1–23; 8:45 am]

### BILLING CODE P

## DEPARTMENT OF JUSTICE

[OMB Number 1110–0046]

| Agency Information Collection Activities: Proposed eCollection eComments Requested: Revision of a Currently Approved Collection; Friction Ridge Cards: Arrest and Institution FD–249; Applicant FD–258; Identity History Summary Request FD–1164; FBI Standard Palm Print FD–884; Supplemental Finger and Palm Print FD–884a; Voluntary Appeal File Fingerprint FD–1212; Firearm-Related Challenge Fingerprint FD–1211
|---|

### AGENCY:

Federal Bureau of Investigation, Criminal Justice Information Services Division, Department of Justice.

### ACTION:

30-Day notice.

### SUMMARY:

Department of Justice (DOJ), Federal Bureau of Investigation (FBI), Criminal Justice Information Services (CJIS) Division will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection was previously published in the Federal Register, Volume 88 pages 16664–16665, on March 20, 2023, allowing a 60-day comment period.

### DATES:

The DOJ encourages public comment and will accept input until July 3, 2023.

## FOR FURTHER INFORMATION CONTACT:

If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact: Larry E. Cotton-Zinn, Management and Program Analyst, FBI, CJIS, Criminal History Information and Policy Unit, BTC–3, 1000 Caster Hollow Road, Clarksburg, WV 26306; phone: 304–425–5300 or email fbi.iii@fbi.gov.

### SUPPLEMENTARY INFORMATION:

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected; and/or
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology. e.g., permitting electronic submission of responses.

Written comments and recommendations for this information collection should be submitted within 30 days of the publication of this notice on the following website www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function and entering either the title of the information collection or the OMB Control Number 1110–0046. This information collection request may be viewed at www.reginfo.gov. Follow the instructions to view Department of Justice, information collections currently under review by OMB.

DOJ seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOJ notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

## Overview of This Information Collection

1. **Type of Information Collection:** Revision of a currently approved collection.
2. **Title of the Form/Collection:** Friction Ridge Cards: Arrest and Institution; Applicant; Identity History Summary Request; FBI Standard Palm Print; Supplemental Finger and Palm Print; Voluntary Appeal File Fingerprint; Firearm-Related Challenge Fingerprint.
3. **Agency form number, if any, and the applicable component of the Department sponsoring the collection:** Agency form number: Forms FD–249 (Arrest and Institution), FD–258 (Applicant), and FD–1164 (Identity History Summary Request); FD–884 (FBI Standard Palm Print); FD–884a (Supplemental Finger and Palm Print); FD–1212 (Voluntary Appeal File Fingerprint); FD–1211 (Firearm-Related Challenge Fingerprint) encompassed under OMB 1110–0046; CJIS Division, FBI, DOJ.

4. **Affected public who will be asked or required to respond, as well as a brief abstract:** Primary: City, county, state, federal and tribal law enforcement agencies; civil entities requesting security clearance and background checks. This collection is needed to collect information on individuals requesting background checks, security clearance, or those individuals who have been arrested for or accused of criminal activities. Acceptable data is stored as part of the Next Generation Identification System (NGI) of the FBI.

5. **Obligation to Respond:** Required to obtain or retain a benefit.
6. **Total Estimated Number of Respondents:** 460,762.
7. **Total Estimated Number of Responses:** 69,200,000.
8. **Estimated Time per Respondent:** 10 minutes.
9. **Frequency:** On occasion.
10. **Total Estimated Annual Time Burden:** 11,500,000 hours.
11. **Total Estimated Annual Other Costs Burden:** $0.