6, 10, 12, and 13 of the '007 patent; and whether an industry in the United States exists as required by subsection (a)(2) of section 337;

(2) Pursuant to section 210.10(b)(1) of the Commission's Rules of Practice and Procedure, 19 CFR 210.10(b)(1), the plain language description of the accused products or category of accused products, which defines the scope of the investigation, is ''robotic vacuums and wet/dry mops, their docking stations, and associated parts and components (including software)'';

(3) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

(a) The complainant is:
iRobot Corporation
8 Crosby Drive
Bedford, MA 01730

(b) The respondents are the following entities alleged to be in violation of section 337, and are the parties upon which the complaint is to be served:
SharkNinja Operating LLC, 89 A St. #100, Needham, MA 02494
SharkNinja Management LLC, 89 A St. #100, Needham, MA 02494
SharkNinja Management Co., 89 A St. #100, Needham, MA 02494
SharkNinja Sales Co., 89 A St. #100, Needham, MA 02494
EP Midco LLC, 89 A St. #100, Needham, MA 02494
SharkNinja Hong Kong Co. Ltd., 238 Des Voeux Road Central, Sheung Wan, Central & Western District—Hong Kong Island, Hong Kong

(4) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

The Office of Unfair Import Investigations will not participate as a party in this investigation.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with section 210.13 of the Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), as amended in 85 FR 15798 (March 19, 2020), such responses will be considered by the Commission if received not later than 20 days after the date of service by the complainant of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: February 25, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021–04294 Filed 3–1–21; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–801]

Importer of Controlled Substances Application: Stepan Company

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Stepan Company has applied to be registered as an importer 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 April 1, 2021. Such persons may also file a written request for a hearing on the application on or before April 1, 2021.

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 08, 2021, Stepan Company, 100 West Hunter Avenue, Maywood, New Jersey 07607–1021, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug Code	Schedule
Coca Leaves	9040	II

The company plans to import the listed controlled substance in bulk for the manufacture of controlled substances for distribution to its customers. No other activity for this drug code is 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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–04245 Filed 3–1–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Federal Bureau of Investigation

[OMB Number 1110–0039]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision of a Currently Approved Collection Bioterrorism Preparedness Act: Entity/Individual Information

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

ACTION: 60-Day notice.

SUMMARY: The 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 is published to obtain comments from the public and affected agencies.

DATES: Comments are encouraged and will be accepted for 60 days until May 3, 2021.

FOR FURTHER INFORMATION CONTACT: If you have additional 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 James J. Sheets, Biometric Services Section, Identity Management Unit, Federal Bureau of Investigation, CJIS Division, Biometric Technology Center, 1000 Custer Hollow Road, Clarksburg, West Virginia 26306.

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:

—Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Bureau of Justice Statistics, including whether the information will have practical utility;
—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;
—Evaluate whether and if so how the quality, utility, and clarity of the information to be collected can be enhanced; and
—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.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a currently approved collection.

2. *The Title of the Form/Collection:* Federal Bureau of Investigation Bioterrorism Preparedness Act: Entity/Individual Information.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* The form is FD–961. The applicable component within the Criminal Justice Information Services Division, Department of Justice (DOJ), Federal Bureau of Investigation (FBI).

4. *Affected public who will be asked or required to respond, as well as a brief abstract:.*

Primary: City, country, state, federal, individuals, business or other for profit, and not-for-profit institute. This collection is needed to receive names and other identifying information submitted by individuals requesting access to specific agents or toxins, and consult with appropriate officials of the Department of Health and Human Services and the Department of Agriculture as to whether certain individuals specified in the provisions should be denied access to or granted limited access to specific agents.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* There are approximately 3,655 (FY 2020) and respondents at 1 hour 30 minutes for the FD–961 form.

6. *An estimate of the total public burden (in hours) associated with the collection:* The estimated public burden associated with this collection is approximately 5,483 hours annual burden, associated with this information collection.

If additional information is required contact: Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, 3E.405A, Washington, DC 20530.

Dated: February 24, 2021.

Melody Braswell,

Department Clearance Officer for PRA, U.S. Department of Justice.

[FR Doc. 2021–04208 Filed 3–1–21; 8:45 am]

BILLING CODE 4410–02–P

DEPARTMENT OF LABOR

Employee Benefits Security Administration

State All Payer Claims Databases Advisory Committee

AGENCY: Employee Benefits Security Administration, Department of Labor.

ACTION: Notice.

SUMMARY: This notice announces the establishment of the State All Payer Claims Databases Advisory Committee (Committee) and also solicits nominations for members to be appointed by the Secretary of Labor (Secretary). The No Surprises Act, enacted as part of the Consolidated Appropriations Act, 2021, requires the Department of Labor (DOL) to establish (and periodically update) a standardized reporting format for voluntary reporting by group health plans to State All Payer Claims Databases and provide guidance to States on collecting data. The Act also directs the Secretary to convene an advisory committee.

DATES: Nominations for membership will be considered if they are received by March 16, 2021.

ADDRESSES: Send nominations to *SAPCDACnominations@dol.gov* to the attention of Elizabeth Schumacher, Office of Health Plan Standards and Compliance Assistance, Employee Benefits Security Administration, DOL.

FOR FURTHER INFORMATION CONTACT: Elizabeth Schumacher, Employee Benefits Security Administration, DOL at 202–693–8335. For press inquiries contact Grant Vaught, Office of Public Affairs, DOL at 202–693–4672.

SUPPLEMENTARY INFORMATION:

I. Background

Section 735 of the Employee Retirement Income Security Act of 1974, as added by section 115(b) of the No Surprises Act, enacted as part of the Consolidated Appropriations Act, 2021, div. BB, tit. I, Public Law 116–260 (Dec. 27, 2020), requires the DOL to establish (and periodically update) a standardized reporting format for voluntary reporting by group health plans to State All Payer Claims Databases and provide guidance to States on collecting data. The Act also directs the Secretary to convene an advisory committee consisting of no more than 15 members to advise the Secretary regarding the format and guidance for voluntary reporting by group health plans to State All Payer Claims Databases. The Committee will remain in existence from the time of its convention until it submits its report with its recommendations to the Secretary, the Committee on Health, Education, Labor, and Pensions of the Senate, and the Committee on Energy and Commerce and the Committee on Education and Labor of the House of Representatives. The Committee is governed by the provisions of the Federal Advisory Committee Act, 5 U.S.C. App. 2.

II. Charter, General Responsibilities, and Composition of the State All Payer Claims Databases Advisory Committee

A. Charter Information and General Responsibilities

On February 17, 2021, the Acting Secretary approved the charter establishing the Committee. The Committee will advise the Secretary regarding the standardized reporting format for the voluntary reporting by group health plans to State All Payer Claims Databases. Reporting will include medical claims, pharmacy claims, dental claims, and eligibility and provider files collected from private