respect to the asserted '026 mark. No party petitioned for review of the ID.

On September 19, 2022, the Commission determined not to review the RID. 87 FR 58130–1 (Sept. 23, 2022). On November 15, 2022, the Commission issued a final determination finding a violation, issuing a GEO prohibiting the unlicensed importation of chocolate milk powder and packaging thereof that infringe the '026 mark, and terminating the investigation. 87 FR 70864–65. On the same day, the Commission issued an opinion explaining the basis for its final determination.

On October 9, 2023, Meenaxi filed a complaint requesting that the Commission institute an enforcement proceeding under Commission Rule 210.75 to investigate alleged violations of the General Exclusion Order by four proposed enforcement respondents: (1) Organic Ingredients; (2) New India; (3) Bharat Bazar; and (4) Coconut Hill Inc. of Sunnyvale, California. Meenaxi asserts that the four proposed enforcement respondents continue to import, sell for importation, advertise, market, distribute, offer to sell ''Bournvita'' products that infringe the '026 mark. Meenaxi also alleges that the four proposed enforcement respondents are in continuing violation of the GEO and as a result, it is sustaining ''immediate and irreparable harm.'' None of the respondents answered Meenaxi's enforcement complaint.

Having examined the enforcement complaint and the supporting documents, the Commission has determined to institute a formal enforcement proceeding, pursuant to Commission Rule 210.75(a) (19 CFR 210.75(a)), to determine whether violations of the GEO, issued on November 15, 2022 in the above-referenced investigation, have occurred and to determine what, if any, enforcement measures are appropriate. The named respondents are: (1) Organic Ingredients Inc. d/b/a Namaste Plaza Indian Super Market; (2) New India Bazar Inc.; (3) Bharat Bazar Inc.; and (4) Coconut Hill Inc. d/b/a Coconut Hill. OUII is also named as a party.

In the Order issued concurrently herewith, the Commission has delegated this enforcement proceeding to the CALJ for designation of a presiding Administrative Law Judge to conduct any necessary proceedings, issue an Enforcement Initial Determination, and make a recommendation on appropriate enforcement measures, if any.

The Commission's vote on this determination took place on November 9, 2023.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: November 9, 2023.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2023–25279 Filed 11–15–23; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–683 and 731–TA–1594–1596 (Final)]

Paper File Folders From China, India, and Vietnam

Determinations

On the basis of the record [1] developed in the subject investigations, the United States International Trade Commission (''Commission'') determines, pursuant to the Tariff Act of 1930 (''the Act''), that an industry in the United States is materially injured by reason of imports of paper file folders from China, India, and Vietnam, provided for in subheading 4820.30.00 of the Harmonized Tariff Schedule of the United States, that have been found by the U.S. Department of Commerce (''Commerce'') to be sold in the United States at less than fair value (''LTFV''), and to be subsidized by the government of India.[2]

Background

The Commission instituted these investigations, effective October 12, 2022, following receipt of petitions filed with the Commission and Commerce by the Coalition of Domestic Folder Manufacturers, Hastings, Minnesota and Naperville, Illinois. The final phase of the investigations was scheduled by the Commission following notification of preliminary determinations by Commerce that imports of paper file folders from India were subsidized within the meaning of section 703(b) of the Act (19 U.S.C. 1671b(b)) and imports of paper file folders from China, India, and Vietnam were sold at LTFV within the meaning of 733(b) of the Act (19 U.S.C. 1673b(b)). Notice of the scheduling of the final phase of the Commission's investigations and of a public hearing to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the **Federal Register** on June 8, 2023 (88 FR 37579). The public hearing in connection with the investigations, originally scheduled for October 3, 2023, was cancelled.[3]

The Commission made these determinations pursuant to §§ 705(b) and 735(b) of the Act (19 U.S.C. 1671d(b) and 19 U.S.C. 1673d(b)). It completed and filed its determinations in these investigations on November 13, 2023. The views of the Commission are contained in USITC Publication 5472 (November 2023), entitled *Paper File Folders from China, India, and Vietnam: Investigation Nos. 701–TA–683 and 731–TA–1594–1596 (Final).*

By order of the Commission.

Issued: November 13, 2023.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2023–25331 Filed 11–15–23; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0023]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Import/Export Declaration for List I and List II Chemical

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 30-Day notice.

SUMMARY: The Drug Enforcement Administration (DEA), Department of Justice (DOJ), 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. This proposed information collection was previously published in the **Federal Register** on September 11, 2023, allowing for a 60-day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until December 18, 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 Scott A. Brinks, Regulatory Drafting and Policy Support Section,

[1] The record is defined in § 207.2(f) of the Commission's Rules of Practice and Procedure (19 CFR 207.2(f)).

[2] 88 FR 69130, 88 FR 69134, 88 FR 69138, and 88 FR 69141, October 5, 2023.

[3] 88 FR 68670, October 4, 2023.

Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (571) 362–3261, email: *scott.a.brinks@dea.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:

—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;
—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;
—Enhance the quality, utility, and clarity of the information to be collected; and/or
—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 1117–0023. 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:* Extension of a currently approved collection.

2. *Title of the Form/Collection:* Import/Export Declaration for List I and List II Chemicals.

3. *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* DEA Forms: 486, 486A. The applicable component within the Department of Justice is the Drug Enforcement Administration, Diversion Control Division.

4. *Affected public who will be asked or required to respond, as well as a brief abstract: Affected public (Primary):* Business or other for-profit. Affected public (Other): Not-for-profit institutions; Federal, State, local, and tribal governments.

Abstract: Section 1018 of the Controlled Substances Import and Export Act (CSIEA) (21 U.S.C. 971) and Title 21 Code of Federal Regulations (21 CFR) Part 1313 require any persons who import, export, or conduct international transactions involving list I and list II chemicals are required to establish a system of recordkeeping and report certain information regarding those transactions to DEA. The chemicals subject to control are used in the clandestine manufacture of controlled substances. The reports of domestic, import, and export regulated transactions in listed chemicals are submitted electronically through the Diversion Control Division secure network application. Any person who desires to import non-narcotic substances in schedules III, IV, and V must electronically file their return information. Any person who desires to export non-narcotic substances in schedules III and IV and any other substance in schedule V is also required to electronically file a controlled substances import declaration/ controlled substance export invoice.

5. *Obligation to Respond:* Mandatory per 21 CFR 1313.

6. *Total Estimated Number of Respondents:* 631.

7. *Estimated Time per Respondent:* 11 minutes for DEA–486 Import, DEA–486 International, and DEA–486A Import, and 12 minutes for DEA–486 Export.

8. *Frequency:* 1 for DEA–486 Import and DEA–486A Import, 4 for DEA–486 International, and 77 for DEA–486 Export.

9. *Total Estimated Annual Time Burden:* Ex: 4,134 hours.

10. *Total Estimated Annual Other Costs Burden:* $0.

If additional information is required, contact: Darwin Arceo, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, United States Department of Justice, Two Constitution Square, 145 N Street NE, 4W–218 Washington, DC 20530.

Dated: November 7, 2023.

Darwin Arceo,

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

[FR Doc. 2023–25326 Filed 11–15–23; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0013]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Application for Permit To Import Controlled Substances for Domestic and/or Scientific Purposes

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 30-Day notice.

SUMMARY: The Drug Enforcement Administration (DEA), Department of Justice (DOJ), 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. This proposed information collection was previously published in the **Federal Register** on September 11, 2023, allowing for a 60-day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until December 18, 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 Scott A. Brinks, Regulatory Drafting and Policy Support Section, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152; Telephone: (571) 362–3261, email: *scott.a.brinks@dea.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:

—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;
—Evaluate the accuracy of the agency's estimate of the burden of the