

Commission's TDD terminal on (202) 205-1810.

SUPPLEMENTARY INFORMATION: The Commission has received a complaint and a submission pursuant to § 210.8(b) of the Commission's Rules of Practice and Procedure filed on behalf of Crocs, Inc. on June 8, 2021. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain casual footwear and packaging thereof. The complainant names as respondents: Cape Robbin Inc. of Pomona, CA; Bijora, Inc., d/b/a Akira of Chicago, IL; Carol Wright Enterprise LLC of Bloomfield, NJ; Dr. Leonard's Healthcare Corp. of Edison, NJ; Crocsky of Austin, TX; Fullbeauty Brands Inc. d/b/a Kingsize of New York, NY; Hawkins Footwear, Sports, Military & Dixie Store of Brunswick, GA; Hobibear Shoes and Clothing Ltd. of Brighton, CO; Hobby Lobby Stores, Inc. of Oklahoma City, OK; Ink Tee of Los Angeles, CA; La Modish Boutique of West Covina, CA; Legend Footwear, Inc., d/b/a Wild Diva of City of Industry, CA; Loeffler Randall Inc. of New York, NY; Maxhouse Rise Ltd. of Hong Kong; New Genesis Online LLC of Newcastle, WA; PW Shoes, Inc. a/k/a P&W of Maspeth, NY; SG Footwear Meser Grp. Inc. a/k/a S. Goldberg & Co. of Hackensack, NJ; Shoe-Nami, Inc. of Gretna, LA; Sketchers USA, Inc. of Manhattan Beach, CA; Star Bay Group Inc. of Hackensack, NJ; Yoki Fashion International LLC of New York, NY; Quanzhou ZhengDe Network Corp., d/b/a Amoji of China; 718Closeouts of Brooklyn, NY; Royal Deluxe Accessories, LLC of New Providence, NJ; and Fujian Huayuan Well Import and Export Trade Co., Ltd. of China. The complainant requests that the Commission issue a general exclusion order, or in the alternative a limited exclusion order, and cease and desist orders.

Proposed respondents, other interested parties, and members of the public are invited to file comments on any public interest issues raised by the complaint or § 210.8(b) filing. Comments should address whether issuance of the relief specifically requested by the complainant in this investigation would affect the public 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 three calendar days after the date on which any initial submissions were due. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number ("Docket No. 3551") in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, Electronic Filing Procedures¹). Please note the Secretary's Office will accept only electronic filings during this time. Filings must be made through the Commission's Electronic Document Information System (EDIS, <https://edis.usitc.gov>.) No in-person paper-based filings or paper copies of any

electronic filings will be accepted until further notice. Persons with questions regarding filing should contact the Secretary at EDIS3Help@usitc.gov.

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,² solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.³

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.

Issued: June 8, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021-12310 Filed 6-10-21; 8:45 am]

BILLING CODE 7020-02-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-848]

Importer of Controlled Substances Application: Adiramedica, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

² All contract personnel will sign appropriate nondisclosure agreements.

³ Electronic Document Information System (EDIS): <https://edis.usitc.gov>.

¹ Handbook for Electronic Filing Procedures: https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf.

SUMMARY: Adiramedica, LLC. 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 file written comments on or objections to the issuance of the proposed registration on or before July 12, 2021. Such persons may also file a written request for a hearing on the application on or before July 12, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on May 17, 2021, Adiramedica, LLC., 585 Turner Industrial Way, Aston, Pennsylvania 19014, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Tapentadol	9780	II

The company plans to import Tapentadol (9780) in dosage form for clinical trials. 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. 2021-12290 Filed 6-10-21; 8:45 am]

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NATIONAL ARCHIVES AND RECORDS ADMINISTRATION

[NARA-2021-026]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: National Archives and Records Administration (NARA).

ACTION: Notice of proposed extension request.

SUMMARY: We are proposing to request an extension from the Office of Management and Budget (OMB) of a currently approved information collection, "Use of NARA Official Seals and Logos." Members of the public and other Federal agencies provide information under this collection as part of their requests to use our official seal(s) and logo(s). We invite you to comment on this proposed information collection pursuant to the Paperwork Reduction Act of 1995.

DATES: We must receive written comments on or before August 10, 2021.

ADDRESSES: Send comments by email to tamee.fechhelm@nara.gov. Because our buildings are temporarily closed during the COVID-19 restrictions, we are not able to receive comments by mail during this time.

FOR FURTHER INFORMATION CONTACT:

Tamee Fechhelm, Paperwork Reduction Act Officer, by email at tamee.fechhelm@nara.gov or by telephone at 301.837.1694 with requests for additional information or copies of the proposed information collection and supporting statement.

SUPPLEMENTARY INFORMATION: Pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104-13), we invite the public and other Federal agencies to comment on proposed information collections. If you have comments or suggestions, they should address one or more of the following points: (a) Whether the proposed information collection is necessary for NARA to properly perform its functions; (b) our estimate of the burden of the proposed information collection and its accuracy; (c) ways we could enhance the quality, utility, and clarity of the information we collect; (d) ways we could minimize the burden on respondents of collecting the information, including through information technology; and (e) whether this collection affects small businesses.

We will summarize any comments you submit and include the summary in our request for OMB approval. All comments will become a matter of public record.

In this notice, we solicit comments concerning the following information collection:

Title: Use of NARA Official Seals and Logos.

OMB number: 3095-0052.

Agency form number: N/A.

Type of review: Regular.

Affected public: Business or other for-profit, not-for-profit institutions, Federal Government.

Estimated number of respondents: 37.

Estimated time per response: 15 minutes.

Frequency of response: On occasion.

Estimated total annual burden hours: 9 hours.

Abstract: The authority for this information collection is contained in 36 CFR 1200.8. NARA's three official seals are the National Archives and Records Administration seal; the National Archives seal; and the National Archives Trust Fund Board seal. The official seals are used to authenticate various copies of official records in our custody and for other official NARA business. We also have an official NARA logo, and other official program and office logos (such as the Federal Register logo, Presidential library logos, Controlled Unclassified Information logo, National Historical Publications and Records Center logo, and more). Occasionally, when criteria are met, we will permit the public or other Federal agencies to use our official seals and logos. The requestor must submit a written request, that includes certain information outlined in 36 CFR 1200, to use the official seals and logos. We approve or deny the request using specific criteria, also outlined in the regulation.

Swarnali Haldar,

Executive for Information Services/CIO.

[FR Doc. 2021-12268 Filed 6-10-21; 8:45 am]

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NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

National Endowment for the Arts

National Council on the Arts 203rd Meeting

AGENCY: National Endowment for the Arts, National Foundation on the Arts and the Humanities.

ACTION: Notice of meeting.

SUMMARY: Pursuant to section 10(a)(2) of the Federal Advisory Committee Act, as amended, notice is hereby given that a meeting of the National Council on the Arts will be held open to the public by videoconference or teleconference.