

number of persons wishing to speak, and the time available, the time for individual comments may be limited. Any written comments received will be provided to the AMWG members.

Before including your address, phone number, email address, or other personal identifying information in your comment, you should be aware that your entire comment—including your personal identifying information—may be made publicly available at any time.

While you can ask us in your comment to withhold your personal identifying information from public review, we cannot guarantee that we will be able to do so.

Authority: 5 U.S.C. appendix 2.

Lee Traynham,

*Chief, Adaptive Management Group,
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BILLING CODE 4332-90-P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731-TA-1557 (Final)]

Certain Mobile Access Equipment and Subassemblies Thereof From China

Determination

On the basis of the record¹ developed in the subject investigation, 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 threatened with material injury by reason of imports of certain mobile access equipment and subassemblies thereof (“mobile access equipment”) from China, provided for in subheadings 8427.10.80, 8427.20.80, 8427.90.00, and 8431.20.00 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”).²

Background

The Commission instituted this investigation effective February 26, 2021, following receipt of antidumping and countervailing duty petitions filed with the Commission and Commerce by the Coalition of American Manufacturers of Mobile Access Equipment (“CAMMAE” or “the

Coalition”).³ The Commission scheduled the final phase of these investigations following notification of a preliminary determination by Commerce that imports of mobile access equipment from China were being subsidized within the meaning of section 703(b) of the Act (19 U.S.C. 1671b(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** of August 12, 2021 (86 FR 44402). In light of the restrictions on access to the Commission building due to the COVID-19 pandemic, the Commission conducted its hearing through written testimony and video conference on October 12, 2021. All persons who requested the opportunity were permitted to participate.

The investigation schedules became staggered when Commerce did not align its countervailing duty investigation with its antidumping duty investigation. Following notification of a final determination by Commerce that imports of mobile access equipment from China were being subsidized within the meaning of section 705(a) of the Act (19 U.S.C. 1671d(a)),⁴ on December 3, 2021, the Commission issued a final affirmative determination in its countervailing duty investigation of mobile access equipment from China.⁵ Following notification of a final determination by Commerce that imports of mobile access equipment from China were being sold at LTFV within the meaning of section 735(a) of the Act (19 U.S.C. 1673d(a)),⁶ notice of the supplemental scheduling of the final phase of the Commission’s antidumping duty investigation 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** of March 2, 2022 (87 FR 11730).

The Commission made this determination pursuant to § 735(b) of the Act (19 U.S.C. 1673d(b)). It completed and filed its determination in this investigation on April 8, 2022. The views of the Commission are contained in USITC Publication 5317 (April 2022),

entitled *Certain Mobile Access Equipment and Subassemblies Thereof from China: Investigation No. 731-TA-1557 (Final)*.

By order of the Commission.
Issued: April 8, 2022.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2022-07912 Filed 4-12-22; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation No. 337-TA-1238]

Certain Plant-Derived Recombinant Human Serum Albumins (“rHSA”) and Products Containing Same; Notice of Request for Submissions on the Public Interest

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that on April 7, 2022, the presiding administrative law judge (“ALJ”) issued an Initial Determination on Violation of Section 337. The ALJ also issued a Recommended Determination on remedy and bonding should a violation be found in the above-captioned investigation. The Commission is soliciting submissions on public interest issues raised by the recommended relief should the Commission find a violation. This notice is soliciting comments from the public only.

FOR FURTHER INFORMATION CONTACT: Ronald A. Traud, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone (202) 205-3427. Copies of non-confidential documents filed in connection with this investigation may be viewed on the Commission’s electronic docket (EDIS) at <https://edis.usitc.gov>. For help accessing EDIS, please email EDIS3Help@usitc.gov. General information concerning the Commission may also be obtained by accessing its internet server at <https://www.usitc.gov>. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission’s TDD terminal on (202) 205-1810.

SUPPLEMENTARY INFORMATION: Section 337 of the Tariff Act of 1930 provides that, if the Commission finds a violation, it shall exclude the articles concerned from the United States: Unless, after considering the effect of such exclusion upon the public health and welfare, competitive conditions in

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

² 87 FR 9576 (February 22, 2022).

³ The Coalition is composed of JLG Industries, Inc. (“JLG”), Hagerstown, Maryland and Terex Corp. (“Terex”), Redmond, Washington.

⁴ 86 FR 57809 (October 19, 2021).

⁵ 86 FR 70147 (December 9, 2021).

⁶ 87 FR 9576 (February 22, 2022).

the United States economy, the production of like or directly competitive articles in the United States, and United States consumers, it finds that such articles should not be excluded from entry. 19 U.S.C. 1337(d)(1). A similar provision applies to cease and desist orders. 19 U.S.C. 1337(f)(1).

The Commission is soliciting submissions on public interest issues raised by the recommended relief should the Commission find a violation, specifically: A limited exclusion order directed to certain plant-derived recombinant human serum albumins ("rHSA") and products containing imported, sold for importation, and/or sold after importation by respondents Wuhan Healthgen Biotechnology Corp. of Wuhan, China; Aspira Scientific, Inc. of Milpitas, California ("Aspira"); eEnzyme LLC of Gaithersburg, Maryland ("eEnzyme"); and ScienCell Research Laboratories, Inc., of Carlsbad, California ("ScienCell"); and cease and desist orders directed to Aspira, eEnzyme, and ScienCell. Parties are to file public interest submissions pursuant to 19 CFR 210.50(a)(4).

The Commission is interested in further development of the record on the public interest in this investigation. Accordingly, members of the public are invited to file submissions of no more than five (5) pages, inclusive of attachments, concerning the public interest in light of the ALJ's Recommended Determination on Remedy and Bonding issued in this investigation on April 7, 2022. Comments should address whether issuance of the recommended remedial orders in this investigation, should the Commission find a violation, 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 recommended remedial orders are used in the United States;
- (ii) identify any public health, safety, or welfare concerns in the United States relating to the recommended 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 recommended orders within a commercially reasonable time; and

(v) explain how the recommended orders would impact consumers in the United States.

Written submissions must be filed no later than by close of business on May 9, 2022.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. The Commission's paper filing requirements in 19 CFR 210.4(f) are currently waived. 85 FR 15798 (Mar. 19, 2020). Submissions should refer to the investigation number ("Inv. No. 337-TA-1238") in a prominent place on the cover page and/or the first page. (See *Handbook for Electronic Filing Procedures*, https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf). Persons with questions regarding filing should contact the Secretary (202-205-2000).

Any person desiring to submit a document to the Commission in confidence must request confidential treatment by marking each document with a header indicating that the document contains confidential information. This marking will be deemed to satisfy the request procedure set forth in Rules 201.6(b) and 210.5(e)(2) (19 CFR 201.6(b) & 210.5(e)(2)). Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. Any non-party wishing to submit comments containing confidential information must serve those comments on the parties to the investigation pursuant to the applicable Administrative Protective Order. A redacted non-confidential version of the document must also be filed simultaneously with any confidential filing and must be served in accordance with Commission Rule 210.4(f)(7)(ii)(A) (19 CFR 210.4(f)(7)(ii)(A)). 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 contract personnel will sign appropriate nondisclosure agreements. All nonconfidential written submissions will be available for public inspection on EDIS.

This action is taken under the authority of 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: April 8, 2022.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2022-07904 Filed 4-12-22; 8:45 am]

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DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request, the Substance Use-Disorder Prevention That Promotes Opioid Recovery and Treatment for Patients and Communities (SUPPORT) Act Grants Evaluation, New Collection

AGENCY: Office of the Assistant Secretary for Policy, Chief Evaluation Office, Department of Labor.

ACTION: Notice of information collection; request for comment.

SUMMARY: The Department of Labor (DOL), as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and federal agencies with an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA95). This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents is properly assessed. Currently, the Department of Labor is soliciting comments concerning the collection of data about the SUPPORT Act Grant Program Evaluation. A copy of the proposed Information Collection Request (ICR) can be obtained by contacting the office listed below in the addressee section of this notice.

DATES: Written comments must be submitted to the office listed in the **ADDRESSES** section below on or before June 13, 2022.

ADDRESSES: You may submit comments by either one of the following methods: