

UNITED STATES INTERNATIONAL TRADE COMMISSION
Washington, D.C.

In the Matter of

**CERTAIN PNEUMATIC COMPRESSION
DEVICES AND COMPONENTS THEREOF**

Investigation No. 337-TA-1316

**NOTICE OF A COMMISSION DETERMINATION NOT TO REVIEW
AN INITIAL DETERMINATION TERMINATING THE INVESTIGATION DUE TO A
SETTLEMENT AGREEMENT AND CONSENT ORDER; TERMINATION OF THE
INVESTIGATION**

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission (“Commission”) has determined not to review an initial determination (“ID”) (Order No. 16) issued by the presiding administrative law judge (“ALJ”) terminating the above-captioned investigation due to a settlement agreement, consent order, and consent order stipulation. The investigation is hereby terminated.

FOR FURTHER INFORMATION CONTACT: Carl P. Bretscher, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street S.W., Washington, D.C. 20436, telephone (202) 205-2382. 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: The Commission instituted this investigation on June 6, 2022, based on a complaint, as supplemented, filed by Precision Holdings USA Inc. of Rocklin, California, and Innovamed Health LLC of San Antonio, Texas (collectively, “Complainants”). 87 FR 34299-300 (June 6, 2022). The complaint, as supplemented, alleges violations of section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, in the importation into the United States, sale for importation, or sale in the United States after importation of certain pneumatic compression devices and components thereof by reason of infringement of certain claims of U.S. Patent Nos. 10,058,475 and 10,912,704. *Id.* The complaint further alleges that a domestic industry exists. *Id.*

The Commission’s notice of investigation names the following respondents: ManaMed, Inc. of Las Vegas, Nevada (“ManaMed”); Grandway Healthcare Ltd. of Hong Kong S.A.R., China (“Grandway”); Medline Industries, Inc. of Northfield, Illinois (“Medline”); and Vive

Health LLC (d/b/a Coretech) of Naples, Florida (“Vive Health”). *Id.* at 34300. The Office of Unfair Import Investigations (“OUII”) is also a party to this investigation. *Id.*

On November 1, 2022, the Commission terminated the investigation with respect to ManaMed, Medline, and Grandway due to a settlement agreement. Order No. 14 (Oct. 7, 2022), *unreviewed by Comm’n Notice* (Nov. 1, 2022).

On November 14, 2022, Complainants and the remaining respondent, Vive Health, jointly moved to terminate the investigation due to a settlement agreement, consent order, and consent order stipulation. On November 17, 2022, OUII filed a response in support of the joint motion to terminate.

On November 18, 2022, the presiding ALJ issued the subject ID granting the joint motion to terminate the investigation. The ID finds that the consent order and consent order stipulation satisfy the requirements of Commission Rule 210.21(c)(3), (4) (19 CFR 210.21(c)(3), (4)). The presiding ALJ also found that the settling parties satisfy the requirements of Commission Rule 210.21(a), (b) (19 CFR 210.21(a), (b)) by providing a copy of the settlement agreement and stipulating that there are no other agreements, oral or written, express or implied, between Complainants and Vive Health regarding the subject matter of the investigation. The ID further finds that public policy generally favors settlement to preserve public and private resources, and that settlement would not harm the public health and welfare, competitive conditions in the U.S. economy, U.S. consumers, or the production of like or directly competitive articles in the United States.

No party filed a petition for review of the subject ID.

The Commission has determined not to review the subject ID. The above-captioned investigation is hereby terminated.

The Commission vote for this determination took place on December 8, 2022.

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.



Katherine M. Hiner
Acting Secretary to the Commission

Issued: December 8, 2022