

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337-TA-1022]

Certain Sleep-Disordered Breathing Treatment Mask Systems and Components Thereof; Institution of Investigation**AGENCY:** U.S. International Trade Commission.**ACTION:** Notice.

SUMMARY: Notice is hereby given that a complaint was filed with the U.S. International Trade Commission on August 17, 2016, under section 337 of the Tariff Act of 1930, as amended, on behalf of ResMed Corp. of San Diego, California; ResMed Inc. of San Diego, California; and ResMed Ltd. of Australia. The complaint alleges violations of section 337 based upon the importation into the United States, the sale for importation, and the sale within the United States after importation of certain sleep-disordered breathing treatment mask systems and components thereof by reason of infringement of certain claims of U.S. Patent No. 8,960,196 (“the ‘196 patent”) and U.S. Patent No. 9,119,931 (“the ‘931 patent”). The complaint further alleges that an industry in the United States exists as required by subsection (a)(2) of section 337.

The complainants request that the Commission institute an investigation and, after the investigation, issue a limited exclusion order and cease and desist orders.

ADDRESSES: The complaint, except for any confidential information contained therein, is available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW., Room 112, Washington, DC 20436, telephone (202) 205-2000. Hearing impaired individuals are advised that information on this matter can be obtained by contacting the Commission’s TDD terminal on (202) 205-1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at (202) 205-2000. General information concerning the Commission may also be obtained by accessing its internet server at <https://www.usitc.gov>. The public record for this investigation may be viewed on the Commission’s electronic docket (EDIS) at <https://edis.usitc.gov>.

FOR FURTHER INFORMATION CONTACT: The Office of Docket Services, U.S. International Trade Commission, telephone (202) 205-1802.

Authority: The authority for institution of this investigation is contained in section 337 of the Tariff Act of 1930, as amended, and in section 210.10 of the Commission’s Rules of Practice and Procedure, 19 CFR 210.10 (2016).

Scope of Investigation: Having considered the complaint, the U.S. International Trade Commission, on September 16, 2016, *ordered that—*

(1) Pursuant to subsection (b) of section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine whether there is a violation of subsection (a)(1)(B) of section 337 in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain sleep-disordered breathing treatment mask systems and components thereof by reason of infringement of one or more of claims 23–86 of the ‘196 patent and claims 1, 5–8, 11–14, 18–22, 25, 26, 28–31, 33–37, 40, 41, 43, 46, 48, 49, 51, 53–55, 57, 58, 60–65, 69–71, 77, and 78 of the ‘931 patent, and whether an industry in the United States exists as required by subsection (a)(2) of section 337;

(2) 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 complainants are:
ResMed Corp., 9001 Spectrum Center Drive, San Diego, CA 92123.
ResMed Inc., 9001 Spectrum Center Drive, San Diego, CA 92123.
ResMed Ltd., 1 Elizabeth Macarthur Drive, Bella Vista NSW 2153, Australia.

(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:

Fisher & Paykel Healthcare Limited, 15 Maurice Paykel Place, East Tamaki, Auckland 2013, P.O. Box 14 348, Panmure, Auckland 1741, New Zealand.
Fisher & Paykel Healthcare, Inc., 173 Technology Drive, Suite 100, Irvine, CA 92618.
Fisher & Paykel Healthcare Distribution Inc., 173 Technology Drive, Suite 100, Irvine, CA 92618.

(3) 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 be a party to 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), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission 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: September 19, 2016.

Lisa R. Barton,*Secretary to the Commission.*

[FR Doc. 2016-22865 Filed 9-21-16; 8:45 am]

BILLING CODE 7020-02-P**NATIONAL AERONAUTICS AND SPACE ADMINISTRATION****Notice of Information Collection**

AGENCY: National Aeronautics and Space Administration (NASA); *Notice:* (16-067).

ACTION: Notice of information collection.

SUMMARY: The National Aeronautics and Space Administration, as part of its continuing effort to reduce paperwork and respondent burden, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. 3506(c)(2)(A)).

DATES: All comments should be submitted within 30 calendar days from the date of this publication.

ADDRESSES: Interested persons are invited to submit written comments regarding the proposed information collection to the Office of Information and Regulatory Affairs, Office of