June 22, 2021

Dear Sir/Madam,

We are contacting you to provide important information relating to ResMed and our devices in light of the Philips voluntary recall and field safety notice announced on June 14, 2021, relating to sound abatement foam used in Philips' CPAP, Bi-Level and Ventilator devices. ResMed's first priority is patient safety and the assurance that patients have access to safe and effective devices for treatment. As such, ResMed would like to provide the following information to you regarding our devices.

ResMed devices are not subject to Philips' June 14, 2021 Field Safety Notice. ResMed devices have a different design and use a different type of foam material for sound abatement than that reported by Philips.

ResMed's devices are extensively tested as part of rigorous verification and validation activities during product development, including tests for particulates and volatile organic compounds (VOC). This testing has not detected any unsafe levels of particulates or chemicals, including those mentioned by Philips in its recall and field safety notice. Furthermore, ResMed continues to evaluate the safety of our products as we learn new information, consistent with our global regulatory obligations, including analysis of complaints. ResMed concludes that all of our devices continue to be safe for patients to use, when following the directions for use.

ResMed is committed to doing everything we can to support the global patient population and maximize availability of our devices to our customers. We will continue to communicate with our customers as the situation develops. For more information, you can visit our FAQ page here, or contact your local ResMed representative.

Regards,
Daniel Judson

VP, Global Product Quality
Quality Assurance and Regulatory Affairs

ResMed Pty Limited 1 Elizabeth Macarthur Drive Bella Vista NSW 2153 Australia
T +61 (2) 8884 1000 **F** +61 (2) 8884 2000 **E** contact@resmed.com.au **ABN** 30 003 765 142 **ResMed.com**