

PHILIPS RESPIRONICS **RECALL INFORMATION**

We want to make you aware of an important development related to certain CPAPs, BiLevel PAPs, and ventilators (including non-invasive and invasive ventilation) manufactured by Philips Respironics. Please be sure to read this information in detail.

On June 14, 2021, Philips Respironics announced a voluntary recall for Continuous and Non-Continuous Ventilators (certain CPAP, BiLevel PAP and Ventilator Devices) due to two issues related to the polyester-based polyurethane (PE-PUR) sound abatement foam used in these devices.

For information on the Recall Notice, a complete list of impacted products, and potential health risks, visit:

<https://www.usa.philips.com/healthcare/e/sleep/communications/src-update>

Our teams are in communication with Philips about this recall and we will do our best to communicate information to you as it becomes available. We encourage you to contact your physician with any additional questions you may have regarding your therapy.

You can access the Philips Q&A at:

[https://www.usa.philips.com/healthcare/e/sleep/communications/src-update#questions and answers](https://www.usa.philips.com/healthcare/e/sleep/communications/src-update#questions_and_answers)

We realize this is unwelcome news, but please be assured that the Advanced Care Network's primary focus is on your safety and wellbeing.