PHILIPS RESPIRONICS RECALL INFORMATION

Philips wants to make you aware of the RECALL referenced above regarding certain CPAPs, Bilevel PAPs, and Ventilators (including non-invasive and invasive ventilation) manufactured by Philips Respironics. PLEASE 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 polyurethan (PE-PUR) sound abatement foam used in these devices. Attached is a copy of the notice provided by Philips.

For additional information on the Recall Notice, a complete list of impacted products, and potential health risks, please visit **www.philips.com/src-update.**

Philips has also provided a Question & Answer resource located at: **https://www.usa.philips.com/healthcare/e/sleep/communications/src-update#questions_and_answers.**

Please visit **www.philips.com/src-update** for the most up-to-date information or call the Philips' toll-free recall support hotline at 1-877-907-7508.