INTEGRATED
HOME CARE SERVICES

Philips Respironics has announced a voluntary Urgent Medical Device 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 a complete list of impacted products, please see the two Recall Notices issued by Philips Respironics.

For additional information regarding the recall, impacted products, and potential health risks, visit Philips.com/src-update.

Please contact our office at 954-381-7957 with any questions.