Update: Philips Respironics Recall

One of the vendors ThedaCare At Home has used to support the sleep and ventilation needs of our customers has issued the below recall notification. Before proceeding with any action, please verify the device you are using is one manufactured by **Philips Respironics**. This can be found on the front of your device. Those devices by other Sleep and Ventilator vendors are not impacted by this recall notification and no further action is necessary.

> Philips Respironics announced a voluntary recall for Continuous and Non-Continuous Ventilators (certain CPAP, Bi-level 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, potential health risks, and the vendor guidance on continuing use of your affected device, visit philips.com/src-update.

We are taking steps with Philips Respironics to resolve this issue as quickly as possible. Further information regarding actions will be updated on this site as soon as available. Should you have any immediate questions, you may contact our office directly with any questions.