

Medical Device Recall Alert

06/14/2021

Please visit our website for medical recalls and alerts that may affect your PAP therapy at www.redriversleep.com. Click on Forms & Policies > Look under “Recalls & Alerts”.

As of 06/14/2021, 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 philips.com/src-update.

The recall notice states that there may be added health risk with ozone-related cleaning devices breaking down components of their PAP devices over time. Philips is recommending halt of use of ozone-related cleaning devices with their PAP devices.

Red River Sleep Center primarily dispenses ResMed PAP devices, however, as a safety precaution; Red River Sleep Center is recommending the following:

1. Review the recall notice posted by Philips and the information posted by ResMed.
2. Temporarily discontinue using ozone-related cleaning products with PAP devices until safety guidance is released by the involved manufacturers.
3. If you have a device subject to the Philips recall, notify your DME supplier to begin the process of replacing your PAP therapy device.
4. If you are not using an ozone cleaning device, but are using a Philips PAP device, contact your DME provider for guidance on replacing your PAP device.
5. If you have a device that is recalled and need assistance to determine if you should continue to use your device, please contact Red River Sleep Center at (318) 443-1684.
6. Clean your PAP device and equipment according to manufacture recommendations.
7. Monitor our Facebook page at <https://www.facebook.com/redriversleep> for updates.

Please contact our office with any additional questions.

Thank you,

Red River Sleep Center, Inc.