AmeriHealth Caritas
District of Columbia

**To:** AmeriHealth Caritas DC Providers

**Date:** February 3, 2022

**Subject:** **Philips Respironics Medical Device Recall**

**Summary**: Certain Philips Respironics devices manufactured before April 26, 2021, have been recalled. Please contact your Provider Account Executive to notify them of enrollees who have been prescribed one of the devices listed below, and contact your patients who may be affected by the recall.

We are writing to notify you that on June 14, 2021, Philips Respironics recalled the CPAP, Bi-Level PAP, and ventilator devices listed below. In these devices manufactured before April 26, 2021, the polyester-based polyurethane (PE-PUR) foam used to reduce sound and vibration in the device may break down and potentially enter the device's air pathway. If this occurs, black debris from the foam or certain chemicals released into the device's air pathway may be inhaled or swallowed by the person using the device.

**We request your assistance in identifying enrollees who were prescribed a recalled device. Please contact your Provider Account Executive to let them know of any enrollees who were prescribed one of the devices below.** If you do not know who your Account Executive is, call Provider Services at 202-408-2237 or visit https://www.amerihealthcaritasdc.com/pdf/provider/contact-provider-account-executive.pdf.

Please visit the U.S. Food and Drug Administration (FDA) website for more information on the recall: https://www.fda.gov/medical-devices/safety-communications/certain-philips-respironics-ventilators-bipap-and-cpap-machines-recalled-due-potential-health-risks.

## Recalled Philips Respironics CPAP and BiPAP devices

| Device type | Model name and number (all serial numbers) |
|---|---|
| Continuous ventilator, minimum ventilatory support, facility use | E30 (emergency use authorization) |
| Continuous ventilator, non-life supporting | DreamStation ASV |
| | DreamStation ST, AVAPS |
| | SystemOne ASV4 |
| | C-Series ASV |
| | C-Series S/T and AVAPS |
| | OmniLab Advanced+ |
| Noncontinuous ventilator | SystemOne (Q-Series) |
| | DreamStation |
| | DreamStation Go |
| | Dorma 400 |
| | Dorma 500 |
| | REMstar SE Auto |

## Recalled Philips Respironics ventilators

| Device type | Model name and number (all series numbers) |
|---|---|
| Continuous ventilator | Trilogy 100 |
| | Trilogy 200 |
| | Garbin Plus, Aeris, LifeVent |
| Continuous ventilator, minimum ventilatory support, facility use | A-Series BiPAP Hybrid A30 (not marketed in U.S.) |
| | A-Series BiPAP V30 Auto |
| Continuous ventilator, non-life supporting | A-Series BiPAP A40 |
| | A-Series BiPAP A30 |

If you have any questions about this communication, please contact your Provider Account Executive or call Provider Services at 202-408-2237.