



HEALTH ALERT

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Certain Philips Respironics Ventilators, BiPAP, CPAP Machines Recalled Due to Potential Health Risks

The Office of Developmental Programs is issuing this Health Alert to bring awareness of a recall affecting certain Philips Respironics Ventilators, BiPAP, and CPAP machines due to potential health risks. The Food and Drug Administration recently issued a Safety Communication regarding this: [Certain Philips Respironics Ventilators, BiPAP, and CPAP Machines Recalled Due to Potential Health Risks: FDA Safety Communication | FDA](#). In their communication, the FDA provides guidance for people who use the affected devices and their caregivers. The devices involved are listed below.

If you use one of these affected devices (see table below), in conjunction with the recommendations provided by the FDA, talk to your health care provider to decide on a suitable treatment for your condition and along with the recommendations provided by the FDA.

CPAP and BiPAP Devices

<u>Device Type</u>	<u>Model Name and Number (All Serial Numbers)</u>
Continuous Ventilator, Minimum Ventilatory Support, Facility Use	<ul style="list-style-type: none"> E30 (Emergency Use Authorization)
Continuous Ventilator, Non-life Supporting	<ul style="list-style-type: none"> 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**

Ventilators

<u>Device Type</u>	<u>Model Name and Number (All Serial Numbers)</u>
Continuous Ventilator	<ul style="list-style-type: none">• Trilogy 100• Trilogy 200• Garbin Plus, Aeris, LifeVent
Continuous Ventilator, Minimum Ventilatory Support, Facility Use	<ul style="list-style-type: none">• A-Series BiPAP Hybrid A30 (not marketed in US)• A-Series BiPAP V30 Auto
Continuous Ventilator, Non-life Supporting	<ul style="list-style-type: none">• A-Series BiPAP A40• A-Series BiPAP A30

Where to Learn More:

- FDA: <https://www.fda.gov/medical-devices/safety-communications/certain-philips-respironics-ventilators-bipap-and-cpap-machines-recalled-due-potential-health-risks>
- Philips Recall Letter: <https://www.philips.com/c-dam/b2bhc/master/landing-pages/src/update/documents/philips-recall-letter-2021-05-a-2021-06-a.pdf>