State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

California Device Recall Information Sheet

Philips Issues Correction For Trilogy Evo Platform Ventilators

Recall Date	Product Description	Recalling Firm	Recall Reason
4/15/2026	**Trilogy Evo** Software Version 1.05.15.00. Continuous home-use ventilator device	**PHILIPS RESPIRONICS, INC.** Murrysville, Pennsylvania	Using non-pneumatic nebulizers with Trilogy Evo Platform ventilators may result in a discrepancy between the set tidal volume and the tidal volume received by the patient.
4/15/2026	**Trilogy Evo O2** Software Version 1.05.15.00. Continuous home-use ventilator device	**PHILIPS RESPIRONICS, INC.** Murrysville, Pennsylvania	Tidal Volume Discrepancy
4/15/2026	**Trilogy Ev300** Software Version 1.05.15.00. Continuous home-use ventilator device	**PHILIPS RESPIRONICS, INC.** Murrysville, Pennsylvania	Tidal Volume Discrepancy
4/15/2026	**Trilogy Evo Universal** Software Version 1.05.15.00. Continuous home-use ventilator device	**PHILIPS RESPIRONICS, INC.** Murrysville, Pennsylvania	Tidal Volume Discrepancy

Recall Class	Product Identification	Distribution	Affected Dates
I	**Trilogy Evo** DS2110X11B UDI: 00606959051942	113,717 Units Nationwide	April 2026 and prior

I	**Trilogy Evo O2** DS2100X11B UDI: 00606959051997	113717 Units Nationwide	April 2026 and prior
I	**Trilogy EV300** DS2200X11B UDI: 00606959052017	113717 Units Nationwide	April 2026 and prior
I	**Trilogy Evo Universal** DS2000X11B UDI: 00606959052000	113,717 Units Nationwide	April 2026 and prior

For additional information, please visit the FDA website