California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Philips Respironics, Inc. Recalls Philips Respironics Trilogy

Recall Date	Product Description	Recalling Firm	Recall Reason
3/26/2026	**Product 1** Philips Respironics Trilogy Evo Universal, Software Version 1.05.15.00. Continuous home-use ventilator device	Philips Respironics, Inc. Murrysville, PA	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.
3/26/2026	**Product 2** Philips Respironics Trilogy EV300, Software Version 1.05.15.00. Continuous home-use ventilator device	Philips Respironics, Inc. Murrysville, PA	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.

Recall Class	Product Identification	Distribution	Affected Dates
I	**Product 1** All Model No.; All UDI; All Serial No.	113,717 Units Worldwide	March 2026 and prior
I	**Product 2** All Model No.; All UDI; All Serial No.	113,717 Units Worldwide California	March 2026 and prior

For additional information, please visit Product 1 (Evo Universal) , Product 2 (EV300)