California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Ventec Life Systems, Inc. VOCSN Multi-Function Ventilators VOCSN+Pro and V+Pro

Recall Date	Product Description	Recalling Firm	Recall Reason
02/26/2025	VOCSN Multi-Function Ventilators: VOCSN+Pro (V+O+C+S+N+Pro, English), REF: PRT-00490-001 V+Pro (V+Pro, English), REF: PRT-01185-000, PRT-01185-002	Ventec Life Systems, Inc.	Multi-Function Ventilators were serviced using incorrect parts which have the potential to cause unexpected shutdown, or when using an active circuit could result in inaccurate tidal volume monitor, not triggering on patient efforts, less inspiratory volume, less inspiratory pressure, and less PEEP delivered.

Recall Class	Product Identification	Distribution	Affected Dates
I	REF: PRT-00490-001 UDI-DI: 00855573007792 Serial Numbers: 123627, 121892, 119639 REF: PRT-01185-000 UDI-DI: 00855573007877 Serial Numbers: 5038421, 116700, 117933, 5038112,	30 units in California, Missouri, and New York	February and prior

	5038619, 5038717, 5038953, 5039086, 5039091, 5039361, 5039499, 5039503, 5039541, 5039636, 5039938, 5040019, 5040640, 5040762, 5041060, 5041078, 5041534, 5042558, 117344, 118464, 118509, 5039387 REF: PRT-01185-002 UDI-DI: 00850018761154 Serial Numbers: 5037655		

For additional information, please visit FDA Website.