



California Medical Device Recall Information



Recall Name

Covidien Recalls Puritan Bennett 980 Ventilator System Due to a Software Error

Recall Date	Product Description	Recalling Firm	Recall Reason
7/16/15	Puritan Bennett 980 Ventilator System	Covidien Boulder, CO	<i>When the ventilator is in neonatal Volume Control Plus (VC+) mode, a software error may cause the amount of air being delivered to the patient to be lower than the amount programmed by the clinician.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<p>All Lots with the following Product Codes are affected:</p> <ul style="list-style-type: none"> • 980U1ENDIUU • 980U1ENDIUUS • 980U3ENDIUU • 980U3ENDIUUS • 980N1ENDIUU • 980N1ENDIUUS • 980N3ENDIUU • 980N3ENDIUUS 	CA , nationwide	<p>Manufactured and distributed between:</p> <p>March 1, 2014 to June 17, 2015</p>

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm460955.htm>