California Medical Device Recall Information

Recall Name
CareFusion Recalls AVEA® Ventilators Due to Incorrect Tidal Volume

Recall Date	Product Description	Recalling Firm	Recall Reason
9/03/13	**AVEA® Ventilators**	**CareFusion** San Diego, CA	*AVEA ventilators may experience an underreporting of tidal volume if used in conjunction with the **neonatal** hotwire flow sensor.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	**All** Models	**CA**, nationwide	N/A

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm370509.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery