California Medical Device Recall Information

Recall Name
Medtronic Recalls the Battery Pack in Covidien Oridion Labeled Capnostream™ 20 and Capnostream™ 20p Patient Monitors Due to a Potential for Thermal Damage

Recall Date	Product Description	Recalling Firm	Recall Reason
4/20/16	**Battery Pack** in • Capnostream 20 • Capnostream 20p	**Medtronic, Inc.** Dublin, Ireland	*A battery manufacturing defect that may increase the risk of thermal damage in the battery pack.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Capnostream: Battery Pack Suspect Model Numbers: • **010520** • **016400**	**CA**, nationwide	Manufactured between: **April 2014** and **February 2016**

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm497049.htm