California Medical Device Recall Information

Recall Name
Medtronic Recalls SynchroMed II/EL Implantable Drug Infusion Pumps Due to Possible Motor Stall

Recall Date	Product Description	Recalling Firm	Recall Reason
11/09/12	SynchroMed II / EL Implantable Drug Infusion Pumps	**Medtronic Neuromodulation, Inc.** Minneapolis, MN	*Possible motor stall when used with unapproved drugs*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	SynchroMed II Implantable Drug Infusion Pump • Models: 8637-40, 8637-20 SynchroMed EL Implantable Drug Infusion Pump • Models: 8626-10, 8626L-10, 8626-18, 8626L-18, 8627-10, 8627L-10, 8627-18, 8627L-18 All Lots Recalled	**CA**, nationwide	Manufactured May 1998 through November 2012

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm333231.htm