California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Medtronic Neuromodulation myPTM Software Application

Recall Date	Product Description	Recalling Firm	Recall Reason
04/01/2025	A820 myPTM Software Application associated with Medtronic SynchroMed Pump and Infusion System	Medtronic Neuromodulation	Product complaints were received describing the A820 myPTM app taking longer than expected for patients to interact with their implantable pump.

Recall Class	Product Identification	Distribution	Affected Dates
II	version v.2x UDI/DI: 00763000632793	729 units in California	January and prior

For additional information, please visit the FDA Website.