MEDICAL DEVICE RECALL LISTING MARCH 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
16/03/2024	MDA/Recall/P0248-76933609-2024	Voluntary Recall	CLEARVIEW® INTRACORONARY SHUNT	GB2174023-125091	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
16/03/2024	MDA/Recall/P0249-39416580-2024	Voluntary Recall	MC2® TWO STAGE VENOUS CANNULA	GB2190423-128552	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
16/03/2024	MDA/Recall/P0250-42161663-2024	Voluntary Recall	DLP® SINGLE STAGE VENOUS CANNULA	GB2321923-125105	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
16/03/2024	MDA/Recall/P0251-10714053-2024	Voluntary Recall	DLP® LEFT HEART VENT CATHETER	GB2414723-125057	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
16/03/2024	MDA/Recall/P0252-38526905-2024	Voluntary Recall	DLP® ONE-PIECE PEDIATRIC ARTERIAL CANNULA	GB5559423-128558	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
16/03/2024	MDA/Recall/P0253-53959033-2024	Voluntary Recall	DLP® CARDIAC SUCTION TUBE	GB6549023-127517	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
16/03/2024	MDA/Recall/P0254-30232634-2024	Voluntary Recall	DLP® SINGLE STAGE VENOUS CANNULA	GB6879723-129857	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123

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16/03/2024	MDA/Recall/P0255-29355891-2024	Voluntary Recall	VC2™ ATRIAL CAVAL VENOUS CANNULA	GB8384823-129855	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
20/03/2024	MDA/Recall/P0256-16071109-2024	Voluntary Recall	KLARO™ IN VIVO SURGICAL LIGHTING	GB6647721-65910	Class II	A02: Manufacturing, Packaging or Shipping Problem	GOODMAN PLT.	MDA-3604-WDP122
25/03/2024	MDA/Recall/P0258-42286569-2024	Voluntary Recall	GRAFTON™ DBM	GD9615023-127511	Class III	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123

* The information contained in the Medical Device Authority Recall database is released under Regulation 7(8) and Regulation 8 (5) of Malaysia's Medical Device Regulations 2019.