MEDICAL DEVICE RECALL LISTING JUNE 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
01/06/2023	MDA/Recall/P0171-13423663-2023	Voluntary Recall	VERITAS VISION SYSTEM	GC2538221-71446	Class III	A26: Insufficient Information	JOHNSON & JOHNSON SDN BHD	MDA-4880-WDP123
02/06/2023	MDA/Recall/P0170-25517312-2023	Voluntary Recall	PATIENT CONNECTIONS	GB491791268519	Class II	A02: Manufacturing, Packaging or Shipping Problem	KL MED SUPPLIES (M) SDN BHD	MDA-2434-WDP121
14/06/2023	MDA/Recall/P0176-61388126-2023	Voluntary Recall	ROTATING SPINE POSITIONERS	GA9655122-88079	Class II	A23: Use of Device Problem	WELCH ALLYN MALAYSIA SDN BHD	MDA-310-W21615
29/06/2023	MDA/Recall/P0182-83668044-2023	Voluntary Recall	MAHURKAR ACUTE DUAL AND TRIPLE LUMEN CATHETER KIT	GD10956523-131786	Class I	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.