MEDICAL DEVICE RECALL LISTING MARCH 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
15/03/2023	MDA/Recall/P0128-60818328-2023	Voluntary Recall	ZENITH BRANCH ENDOVASCULAR GRAFT - ILIAC BIFURCATION	GD6890667816	Class II	A05: Mechanical Problem	COOK ASIA (MALAYSIA) SDN BHD	MDA-1987-WDP121
24/03/2023	MDA/Recall/P0130-62798032-2023	Voluntary Recall	InnoQ Ultrasound Gel	GA10900121-67932	Class III	A04: Material Integrity Problem	IDS MEDICAL SYSTEMS (M) SDN BHD	MDA-2377-WDP121
24/03/2023	MDA/Recall/P0133-57936436-2023	Voluntary Recall	PERIPHERAL VASCULAR CATHETHER	GB8366674516	Class III	A22: Human-Device Interface Problem	B. BRAUN MEDICAL INDUSTRIES SDN BHD	MDA-4298-K123
23/03/2023	MDA/Recall/P0134-82763169-2023	Voluntary Recall	SHILEY ADULT FLEXIBLE TRACHEOSTOMY TUB	GC94666774518	Class II	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-0074-WDP7414
30/03/2023	MDA/Recall/P0135-75622318-2023	Voluntary Recall	TOSOH AUTOMATED GLYCOHEMOGLOBIN ANALYZER HLC-723GX	IVDB2953711516	Class III	A02: Manufacturing, Packaging or Shipping Problem	ALL EIGHTS (M) SDN BHD	MDA-4259-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.

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