MEDICAL DEVICE RECALL LISTING APRIL 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
04/04/2024	MDA/Recall/P0260-60513733-2024	Voluntary Recall	F&P AIRVO 2 HUMIDIFIER	GB13073385917	Class II	A07: Electrical /Electronic Property Problem	EMERGO MALAYSIA SDN. BHD.	MDA-5078-W123
09/04/2024	MDA/Recall/P0261-18637955-2024	Voluntary Recall	COBAS® HCV GT	IVDC84546154618	Class III	A09: Output Problem	ROCHE DIAGNOSTICS (M) SDN. BHD.	MDA-5585-WDP124
26/04/2024	MDA/Recall/P0263-33396706-2024	Voluntary Recall	ORTHOPAEDIC IMPLANTS	GC86706621718	Class II	A02: Manufacturing, Packaging or Shipping Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4250-W123
30/04/2024	MDA/Recall/P0270-78379448-2024	Voluntary Recall	ARTIS SYMBIOSE	GC5714520-45735	Class II	A26: Insufficient Information (require revision)	TRANSMEDIC HEALTHCARE SDN. BHD.	MDA-4762-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.