



MEDICAL DEVICE RECALL LISTING MAY 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
06/05/2024	MDA/Recall/P0271-76917725-2024	Voluntary Recall	HUMIDIFIER MYAIRVO2	GB197511053618	Class II	A07: Electrical /Electronic Property Problem	EMERGO MALAYSIA SDN. BHD.	MDA-5078-W123
08/05/2024	MDA/Recall/P0274-20416296-2024	Voluntary Recall	LAPAROSCOPY SYSTEM	GB26809989718	Class III	A27: Appropriate Term/Code Not Available (require revision_	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
14/05/2024	MDA/Recall/P0277-92317170-2024	Voluntary Recall	TACTOSET INJECTABLE BONE SUBSTITUTE SYSTEM	GD5361723-124925	Class II	A25: No Apparent Adverse Event	CIMED HEALTHCARE SDN. BHD.	MDA-3498-W122
15/05/2024	MDA/Recall/P0278-91305682-2024	Voluntary Recall	PDS™ II (POLYDIOXANONE) STERILE SYNTHETIC, ABSORBABLE SUTURE	GD4349722-103519	Class III	A02: Manufacturing, Packaging or Shipping Problem	JOHNSON & JOHNSON SDN BHD.	MDA-4880-WDP123
15/05/2024	MDA/Recall/P0279-39693928-2024	Voluntary Recall	CASCADE IOMAX	GC5241323-144670	Class III	A02: Manufacturing, Packaging or Shipping Problem	T T MEDICAL MANAGEMENT SDN. BHD.	MDA-2373-W121

* 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.