

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	02/03/2026	MDA/Recall/P0514-86310994-2026	Establishment (Voluntary Recall)	SUPERPULSE SYSTEM	GC81659199517	Class II: Moderate Risk	A05: Mechanical Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
2.	13/03/2026	MDA/Recall/P0515-34376841-2026	Establishment (Voluntary Recall)	KIWI VACUUM DELIVERY SYSTEM	GB71670789518	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	TRANSMEDIC HEALTHCARE SDN BHD	MDA-4762-WDP123
3.	19/03/2026	MDA/Recall/P0520-26538397-2026	Establishment (Voluntary Recall)	CENTROFLO LONG TERM HEMODIALYSIS CATHETER	GD62284755018	Class II: Moderate Risk	A23: Use of Device Problem	MERIT MEDICAL MALAYSIA SDN. BHD.	MDA-5946-W124

* 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.