MEDICAL DEVICE RECALL LISTING JANUARY 2026

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	13/01/2026	MDA/Recall/P0476-79011959-2026	Establishment (Voluntary Recall)	VIZISHOT 2 FLEX SINGLE USE ASPIRATION NEEDLE	GB78398709818	Class II :Moderate Risk	A05: Mechanical Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
2.	13/01/2026	MDA/Recall/P0478-41002997-2026	Establishment (Voluntary Recall)	HIGH FREQUENCY SPHINCTEROTOME	GC726531024218	Class II :Moderate Risk	A05: Mechanical Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
3.	19/01/2026	MDA/Recall/P0479-76694841-2026	Establishment (Voluntary Recall)	DA VINCI SI SURGICAL SYSTEM, ENDOSCOPIC INSTRUMENT CONTROL SYSTEM, MODEL IS3000	GC74543334017	Class II :Moderate Risk	A27: Appropriate Term/Code Not Available	DTG MEDICAL SDN BHD	MDA-4091-WDP122
4.	19/01/2026	MDA/Recall/P0480-11682079-2026	Establishment (Voluntary Recall)	DA VINCI SI SURGICAL SYSTEM, ENDOSCOPIC INSTRUMENT CONTROL SYSTEM, MODEL IS3000	GC74543334017	Class II :Moderate Risk	A27: Appropriate Term/Code Not Available	DTG MEDICAL SDN BHD	MDA-4091-WDP122
5.	15/01/2026	MDA/Recall/P0481-25326490-2026	Establishment (Voluntary Recall)	HICKMAN CENTRAL VENOUS CATHETERS	GD78617561018	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123

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6.	15/01/2026	MDA/Recall/P0482-88125165-2026	Establishment (Voluntary Recall)	BROVIAC CENTRAL VENOUS CATHETER	GD6710123-133101	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123
7.	19/01/2026	MDA/Recall/P0483-15942341-2026	Establishment (Voluntary Recall)	DA VINCI XI SURGICAL SYSTEM, ENDOSCOPIC INSTRUMENT CONTROL SYSTEM, MODEL IS4000	GC3810021-64842	Class II :Moderate Risk	A27: Appropriate Term/Code Not Available	DTG MEDICAL SDN BHD	MDA-4091-WDP122
8.	19/01/2026	MDA/Recall/P0484-95973794-2026	Establishment (Voluntary Recall)	DA VINCI XI SURGICAL SYSTEM, ENDOSCOPIC INSTRUMENT CONTROL SYSTEM, MODEL IS4000	GC3810021-64842	Class II :Moderate Risk	A27: Appropriate Term/Code Not Available	DTG MEDICAL SDN BHD	MDA-4091-WDP122
9.	27/01/2026	MDA/Recall/P0488-52812230-2026	Establishment (Voluntary Recall)	AQUASIL PUTTY IMPRESSION MATERIAL	GMD61708268217A	Class III :Low Risk	A25: No Apparent Adverse Event	DENTSPLY SIRONA MALAYSIA SDN. BHD.	MDA-4163-WDP122

* 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.