MEDICAL DEVICE RECALL LISTING FEBRUARY 2025

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason for Recall	Recalling Establishment	Establishment License
1.	04/02/2025	MDA/Recall/P0361-82066130-2025	Voluntary Recall	UNIFLOW COAXIAL BREATHING SYSTEMS	GB447531163118	Class II: Moderate Risk	A12: Connection Problem	INTERSURGICAL SDN BHD	MDA-5088-WDP123
2.	04/02/2025	MDA/Recall/P0363-37699536-2025	Voluntary Recall	PIPELINE VANTAGE EMBOLIZATION DEVICE WITH SHIELD TECHNOLOGY	GD8603721-68288	Class I: High Risk	A23: Use of Device Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
3.	06/02/2025	MDA/Recall/P0364-37080000-2025	Voluntary Recall	MĪAR™ AORTIC ROOT CANNULA	GB8326523-125097	Class III :Low Risk	A02: Manufacturing, Packaging or Shipping Problem		
4.	06/02/2025	MDA/Recall/P0365-80908849-2025	Voluntary Recall	DLP® AORTIC ROOT CANNULA	GB5806223-125793	Class III :Low Risk	A02: Manufacturing, Packaging or Shipping Problem		

5.	17/02/2025	MDA/Recall/P0366-97884014-2025	Voluntary Recall	AESCULAP ORTHOPILOT® SYSTEM	GB91243575018	Class III :Low Risk	A04: Material Integrity Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4250-W123
6.	20/02/2025	MDA/Recall/P0368-61499460-2025	Voluntary Recall	TB STAINS AND REAGENTS	IVDB19407246518	Class III :Low Risk	A02: Manufacturing, Packaging or Shipping Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123
7.	24/02/2025	MDA/Recall/P0369-68831920-2025	Voluntary Recall	PERICARDIAL SUCKER	GB630391339119	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	NUFA AWANA SDN BHD	MDA-3974-W122

* 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.