MEDICAL DEVICE RECALL LISTING JULY 2025

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	24/01/2025	MDA/Recall/P0415-39645057-2025	Voluntary Recall	NASOGEL TUBE	GB5801523-154871	Class III	A02: Manufacturing, Packaging or Shipping Problem	JETPHARMA SDN BHD	MDA-5334-WDP123
2.	08/07/2025	MDA/Recall/P0419-30602785-2025	Voluntary Recall	CAROTID WALLSTENT MONORAIL	GD94941309717	Class III	A23: Use of Device Problem	BOSTON SCIENTIFIC (MALAYSIA) SDN BHD	MDA-5810-WD124
3.	16/07/2025	MDA/Recall/P0422-69098658-2025	Voluntary Recall	VENCLOSE RADIOFREQUENCY SYSTEM	GC5750923-148658	Class II	A07: Electrical /Electronic Property Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123
4.	24/07/2025	MDA/Recall/P0425-41063047-2025	Voluntary Recall	CELSITE® IMPLANTABLE VASCULAR ACCESS SYSTEMS AND ACCESSORIES	GD59669269417	Class III	A15: Activation, Positioning or Separation Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4298-K123

* 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.