MEDICAL DEVICE RECALL LISTING AUGUST 2025

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	15/08/2025	MDA/Recall/P0429-13072711-2025	Voluntary Recall	ENDOTRACHEAL TUBES SYSTEM	GB6823123-143999	Class I :High Risk	A01: Patient Device Interaction Problem	VENTICARE MEDICAL (M) SDN BHD	MDA-5498-K124
2.	01/08/2025	MDA/Recall/P0430-62799712-2025	Voluntary Recall	NEEDLE HOLDER	GMD56025205417A	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123
3.	05/08/2025	MDA/Recall/P0431-17249514-2025	Voluntary Recall	ENCORE 26 INFLATION DEVICE	GA3949581516	Class III :Low Risk	A04: Material Integrity Problem	BOSTON SCIENTIFIC (MALAYSIA) SDN BHD	MDA-5810-WD124
4.	08/08/2025	MDA/Recall/P0433-31831757-2025	Voluntary Recall	CONSTELLATION VISION SYSTEM	GC96073327017	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	ALCON LABORATORIES (M) SDN. BHD.	MDA-5021-W123
5.	12/08/2025	MDA/Recall/P0436-55913975-2025	Voluntary Recall	Sterile Syringe	GA10626625-193082	Class II :Moderate Risk	A09: Output Problem	ADVENTA HEALTHCARE SDN. BHD.	MDA-5667-WDP124
6.	13/08/2025	MDA/Recall/P0437-16677350-2025	Voluntary Recall	HINOTORI SURGICAL ROBOT SYSTEM	GC8683624-179851	Class II :Moderate Risk	A02: Manufacturing, Packaging or	SYSMEX (MALAYSIA) SDN. BHD.	MDA-4997-WDP123

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							Shipping Problem		
7.	14/08/2025	MDA/Recall/P0438-38655840-2025	Voluntary Recall	VIZISHOT 2 FLEX SINGLE USE ASPIRATION NEEDLE	GB78398709818	Class II :Moderate Risk	A05: Mechanical Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
8.	18/08/2025	MDA/Recall/P0439-21973484-2025	Voluntary Recall	STRATAFIX SPIRAL PDS PLUS KNOTLESS TISSUE CONTROL DEVICE	GD2913522-110161	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	JOHNSON & JOHNSON SDN BHD	MDA-4880-WDP123
9.	20/08/2025	MDA/Recall/P0440-55937739-2025	Voluntary Recall	DLP® LEFT HEART VENT CATHETER	GB2414723-125057	Class II :Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
10.	20/08/2025	MDA/Recall/P0441-27160344-2025	Voluntary Recall	AESCULAP POWER SYSTEM	GB68825829018	Class I :High Risk	A21: Labelling, Instructions for Use or Training Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4298-K123
11.	29/08/2025	MDA/Recall/P0443-57749742-2025	Voluntary Recall	INSTRUMENTS	GC48560748218	Class I :High Risk	A04: Material Integrity Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4298-K123