MEDICAL DEVICE RECALL LISTING NOVEMBER 2025

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	07/11/2025	MDA/Recall/P0453-43804394-2025	Establishment (Voluntary Recall)	GM85 (C-TYPE, F-TYPE, FIT-TYPE)	GC5394423-134640	Class II: Moderate Risk	A04: Material Integrity Problem	LAC MEDICAL SUPPLIES SDN BHD	MDA-5984-WDP124
2.	05/11/2025	MDA/Recall/P0456-47674455-2025	Establishment (Voluntary Recall)	LEGION KNEE SYSTEM	GC19512623018	Class III: Low Risk	A21: Labelling, Instructions for Use or Training Problem	SMITH & NEPHEW HEALTHCARE SDN BERHAD	MDA-4767-WP123
3.	6/11/2025	MDA/Recall/P0460-20270414-2025	Establishment (Voluntary Recall)	INFUSION SETS & ACCESSORIES	GB411691095818	Class II: Moderate Risk	A05: Mechanical Problem	COMCORDE MEDICAL (M) SDN BHD	MDA-4949-K123
4.	24/11/2025	MDA/Recall/P0463-23956658-2025	Establishment (Voluntary Recall)	FLEXOR CHECK-FLO INTRODUCER	GB5300419-34174	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	COOK ASIA (MALAYSIA) SDN BHD	MDA-5123-WDP123
5.	26/11/2025	MDA/Recall/P0464-18511740-2025	Establishment (Voluntary Recall)	BIOSURE REGENESORB INTERFERENCE SCREW	GD74672799818	Class III: Low Risk	A21: Labelling, Instructions for Use or Training Problem	SMITH & NEPHEW HEALTHCARE SDN BERHAD	MDA-4767-WP123

* 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.