MEDICAL DEVICE RECALL LISTING SEPTEMBER 2025

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	30/09/2025	MDA/Recall/P0257-57406180-2024	Establishment (Voluntary Recall)	QUO-LAB A1C SYSTEM	IVDB8128921-67256	A02: Manufacturing, Packaging or Shipping Problem	Class II: Moderate Risk	UBISSON SDN BHD	MDA-5314-W123
2.	11/09/2025	MDA/Recall/P0442-55829451-2025	Establishment (Voluntary Recall)	SAMARITAN PAD 350P/360P/500P	GC1448998616	A02: Manufacturing, Packaging or Shipping Problem	Class III: Low Risk	MURSMEDIC MALAYSIA SDN. BHD.	MDA-5600-W124
3.	05/09/2025	MDA/Recall/P0444-23689701-2025	Establishment (Voluntary Recall)	EURODEFIPADS	GC10719224-185696	A02: Manufacturing, Packaging or Shipping Problem	Class II: Moderate Risk	JUST NU MEDICARE SDN BHD	MDA-5938-WDP124
4.	10/09/2025	MDA/Recall/P0445-37674373-2025	Establishment (Voluntary Recall)	COBAS® HCV	IVDD3403524-176110	A02: Manufacturing, Packaging or Shipping Problem	Class III: Low Risk	ROCHE DIAGNOSTICS (M) SDN. BHD.	MDA-5585-WDP124
5.	11/09/2025	MDA/Recall/P0446-96203019-2025	Establishment (Voluntary Recall)	CONFIRM ANTI-ESTROGEN RECEPTOR (ER) (SP1) RABBIT MONOCLONAL	IVDC19194190418	A04: Material Integrity Problem	Class III: Low Risk	ROCHE DIAGNOSTICS (M) SDN. BHD.	MDA-5585-WDP124

				PRIMARY ANTIBODY					
6.	14/09/2025	MDA/Recall/P0447-84556323-2025	Establishment (Voluntary Recall)	STA® - LIATEST® D-DI PLUS	IVDC5347522-90795	A04: Material Integrity Problem	Class III: Low Risk	ALL EIGHTS (M) SDN BHD	MDA-4259-WDP123
7.	25/09/2025	MDA/Recall/P0448-71268790-2025	Establishment (Voluntary Recall)	SIGNIA SMALL DIAMETER RELOADS	GD9181722-85855	A27: Appropriate Term/Code Not Available	Class III: Low Risk	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123

* 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.