



MEDICAL DEVICE RECALL LISTING AUGUST 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
07/08/2024	MDA/Recall/P0318-83004726-2024	Voluntary Recall	SUTURES	GD62950129617	Class III	A02: Manufacturing, Packaging or Shipping Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4298-K123
09/08/2024	MDA/Recall/P0321-53506257-2024	Voluntary Recall	DAILIES TOTAL 1 MULTIFOCAL SOFT CONTACT LENSES	GB31347275217	Class II	A02: Manufacturing, Packaging or Shipping Problem	ALCON LABORATORIES (M) SDN. BHD.	MDA-5021-W123
15/08/2024	MDA/Recall/PX0320-45427297-2024	Voluntary Recall	DISPOSABLE BLOOD GIVING SET	GB24805760818	Class III	A23: Use of Device Problem	MYMEDIC INNOVATION SDN BHD	MDA-5449-K124
16/08/2024	MDA/Recall/P0329-80390932-2024	Voluntary Recall	REGENETEN BIOINDUCTIVE IMPLANT	GD8137521-61832	Class II	A18: Contamination / decontamination Problem	SMITH & NEPHEW HEALTHCARE SDN BERHAD	MDA-4767-WP123
22/08/2024	MDA/Recall/P0332-95573555-2024	Voluntary Recall	INTERVENTIONAL PAIN MANAGEMENT	GC19187572718	Class II	A18: Contamination / decontamination Problem	GLOBAL HEALTHCARE LOGISTICS (M) SDN BHD	MDA-5941-WDP124
27/08/2024	MDA/Recall/P0333-54768955-2024	Voluntary Recall	ARCHITECT HBSAG	IVDD89016121618	Class III	A18: Contamination / decontamination Problem	ABBOTT LABORATORIES (MALAYSIA) SDN. BHD.	MDA-5104-W123

* 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.



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