MEDICAL DEVICE RECALL LISTING JUNE 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
07/06/2024	MDA/Recall/P0285-10572756-2024	Voluntary Recall	DR WOUND NANO SILVER SPRAY	GC3497719-26935	Class II	A18: Contamination / decontamination Problem	FARMASIA SDN BHD	MDA-5666-K124
21/06/2024	MDA/Recall/P0298-31421182-2024	Voluntary Recall	LINQ II™ INSERTABLE CARDIAC MONITOR	GD1544422-94694	Class I	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
18/06/2024	MDA/Recall/P0300-44004172-2024	Voluntary Recall	Cobas HBV	IVDD5502023-131998	Class III	A02: Manufacturing, Packaging or Shipping Problem	ROCHE DIAGNOSTICS (M) SDN. BHD.	MDA-5585-WDP124
24/06/2024	MDA/Recall/P0280-79157745-2024	Voluntary Recall	AUTOSTAINER LINK 48	IVDA321495216	Class III	A07: Electrical /Electronic Property Problem	BITA LIFESCIENCE SDN. BHD.	MDA-4664-WDP123
27/06/2024	MDA/Recall/P0301-93746175-2024	Mandatory Recall	TRANSMISSION GEL	GA13780404317	Class II	A18: Contamination / decontamination Problem	MYMEDIC INNOVATION SDN BHD	MDA-5449-K124

* 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.