MEDICAL DEVICE RECALL LISTING MAY 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
03/05/2023	MDA/Recall/P0152-94493240-2023	Voluntary Recall	TELEFLEX WECK METAL LIGATING CLIPS (HORIZON)	GD84037633018	Class II	A02: Manufacturing, Packaging or Shipping Problem	TELEFLEX MEDICAL SDN.BHD.	MDA-3059-K121
05/05/2023	MDA/Recall/P0154-79475888-2023	Voluntary Recall	QUICKCLIP-SINGLE USE ROTATABLE CLIP FIXING	GB58901283217	Class II	A02: Manufacturing, Packaging or Shipping Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
17/05/2023	MDA/Recall/P0158-74970035-2023	Voluntary Recall	CELL-DYN EMERALD 22	IVDB5774422-110093	Class II	A10: Temperature Problem	ABBOTT LABORATORIES (MALAYSIA) SDN. BHD.	MDA-1685-W121
19/05/2023	MDA/Recall/P0160-78943884-2023	Voluntary Recall	UROPASS URETERAL ACCESS SHEATH SET	GB32940617118	Class II	A04: Material Integrity Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
24/05/2023	MDA/Recall/P0159-92691485-2023	Voluntary Recall	GELITA-SPON	GD97157451717	Class III	A25: No Apparent Adverse Event	IDS MEDICAL SYSTEMS (M) SDN. BHD.	MDA-2377-WDP121
24/05/2023	MDA/Recall/P0161-74043748-2023	Voluntary Recall	TELEFLEX MEDICAL RUSCH BRONCHOPART DOUBLE LUMEN BRONCHIAL TUBE	GB16263304517	Class II	A18: Contamination / decontamination Problem	TELEFLEX MEDICAL SDN.BHD	MDA-3059-K121

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24/05/2023	MDA/Recall/P0164-62777011-2023	Voluntary Recall	TELEFLEX RUSCH TRACHEAL TUBES (UNCUFFED)	GB449871169418	Class II	A18: Contamination / decontamination Problem	TELEFLEX MEDICAL SDN.BHD	MDA-3059-K121
24/05/2023	MDA/Recall/P0166-67878571-2023	Voluntary Recall	ENDOTRACHEAL TUBE	GB87665877718	Class II	A18: Contamination / decontamination Problem	TELEFLEX MEDICAL SDN.BHD	MDA-3059-K121

* 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.