MEDICAL DEVICE RECALL LISTING NOVEMBER 2022

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
11/11/2022	MDA/Recall/P0080-34550611-2022	Voluntary recall	TELEFLEX RUSCH URETERAL CATHETERS	GB46067877818	Class II	A01: Patient Device Interaction Problem	TELEFLEX MEDICAL SDN.BHD.	MDA-3058-W121
16/11/2022 (Late reporting)	MDA/Recall/P0085-15057098-2022	Voluntary recall	ALTERNA® BASEPLATE	GMD343681007019A	Class I	A21: Labelling, Instructions for Use or Training Problem	DKSH MALAYSIA SDN. BHD.	MDA-0023-WDP2314
16/11/2022 (Late reporting)	MDA/Recall/P0089-77559257-2022	Voluntary recall	ALTERNA® BASEPLATE	GMD343681007019A	Class I	A21: Labelling, Instructions for Use or Training Problem	DKSH MALAYSIA SDN. BHD.	MDA-0023-WDP2314
21/11/2022	MDA/Recall/P0090-20968288-2022	Voluntary recall	AQUACEL AG ADHESIVE/NON-ADHESIVE FOAM HYDROFIBER DRESSING	GD12979451117	Class III	A18: Contamination / decontamination Problem	CONVATEC MALAYSIA SDN. BHD.	MDA-1913-W121
22/11/2022	MDA/Recall/P0092-28460710-2022	Voluntary recall	VENTANA PD-L1 (SP142) ASSAY	IVDC77381365919	Class II	A02: Manufacturing, Packaging or Shipping Problem	ROCHE DIAGNOSTICS (M) SDN. BHD.	MDA-1674-WDP121

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24/11/2022	MDA/Recall/P0093-55791877-2022	Voluntary recall	VLIWAKTIV ABSORBENT ACTIVATED CHARCOAL DRESSING	GB2547620-42323	Class III	A02: Manufacturing, Packaging or Shipping Problem	NYPRAX BUSINESS SOLUTIONS	MDA-400-WDP30415

* 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.