MEDICAL DEVICE RECALL LISTING DECEMBER 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
03/12/2024	MDA/Recall/P0348-41448449-2024	Voluntary Recall	CENTRIFUGE	GA84458804218	Class III	A05: Mechanical Problem	INTERSCIENCE SDN BHD	MDA-5690-W124
05/12/2024	MDA/Recall/P0350-82624946-2024	Voluntary Recall	STEALTHSTATION S7 TREATMENT GUIDANCE SYSTEM (SPINAL)	GD2205223-131802	Class II	A23: Use of Device Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
10/12/2024	MDA/Recall/P0353-70976527-2024	Voluntary Recall	SELDINGER CHEST DRAINAGE KITS	GB2435122-96118	Class III	A02: Manufacturing, Packaging or Shipping Problem	SOMNOTEC (M) SDN BHD	MDA-4866-WDP123
18/12/2024	MDA/Recall/P0354-13401169-2024	Voluntary Recall	MECHANICAL LITHOTRIPTOR	GA65102107116	Class II	A21: Labelling, Instructions for Use or Training Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-6344-WDP124

* 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.