MEDICAL DEVICE RECALL LISTING JANUARY 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
09/01/2024	MDA/Recall/P0234-84890215-2024	Voluntary Recall	PK INSTRUMENTS	GC34084617018	Class II	A05: Mechanical Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121
16/01/2024	MDA/Recall/P0235-17535027-2024	Voluntary Recall	SUPRASORB® A + AG	GD6355223-124001	Class III	A02: Manufacturing, Packaging or Shipping Problem	NYPRAX BUSINESS SOLUTIONS	MDA-5323-WDP123
26/01/2024	MDA/Recall/P0236-57177993-2024	Voluntary Recall	DUET EXTERNAL DRAINAGE AND MONITORING SYSTEM	GD29959666618	Class I	A23: Use of Device Problem	MEDTRONIC MALAYSIA SDN. BHD.	MDA-4793-WDP123

* 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.