MEDICAL DEVICE RECALL LISTING FEBRUARY 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
2 February 2022	MDA/Recall/P0115-69489709-2023	Voluntary recall	STERILIZATION CONTAINER SYSTEM	GMD48453111917A	Class I	A17: Compatibility Problem	BECTON DICKINSON SDN BHD	MDA-0033-W3314
7 February 2022	MDA/Recall/P0117-22967415-2023	Voluntary recall	INSULIN SYRINGE	GB56396803618	Class II	A18: Contamination / decontamination Problem	BECTON DICKINSON SDN BHD	MDA-0033-W3314
10 February 2022	MDA/Recall/P0119-94336789-2023	Voluntary recall	GRAFTON DBM	GD67453892418	Class III	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-0074-WDP7414

* 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.