MEDICAL DEVICE RECALL LISTING SEPTEMBER 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
03/09/2024	MDA/Recall/P0336-70507259-2024	Voluntary Recall	BARD MAGNUM DISPOSABLE CORE TISSUE BIOPSY NEEDLE TO BE USED WITH BARD MAGNUM INSTRUMENT	GB62221655118	Class II	A18: Contamination / decontamination Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123
10/09/2024	MDA/Recall/PX0330-18226750-2024	Voluntary Recall	SURGICAL CONNECTING TUBE	GA94267171917	Class II	A02: Manufacturing, Packaging or Shipping Problem	HOSPITECH MANUFACTURING SERVICES SDN. BHD.	MDA-4689-K123

* 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.