MEDICAL DEVICE RECALL LISTING NOVEMBER 2024

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
11/11/2024	MDA/Recall/P0346-47411217-2024	Voluntary Recall	HYDROPHOBIC ACRYLIC INTRAOCULAR LENS	GC7143222-113629	Class III	A02: Manufacturing, Packaging or Shipping Problem	MY IOL SDN BHD	MDA-6395-K124
11/11/2024	MDA/Recall/P0347-99297066-2024	Voluntary Recall	BD BBL™ SENSI-DISC™ SUSCEPTIBILITY TEST DISCS	IVDB6578123-148932	Class II	A23: Use of Device Problem	BECTON DICKINSON SDN BHD	MDA-5083-W123
26/11/2024	MDA/Recall/P0349-25322329-2024	Voluntary Recall	3M UNITEK TRANSBOND PLUS SELF ETCHING PRIMER	GB71798433017	Class III	A02: Manufacturing, Packaging or Shipping Problem	3M MALAYSIA SDN BHD	MDA-5752-WP124

* 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.