



MEDICAL DEVICE RECALL LISTING APRIL 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
06/04/2023	MDA/Recall/P0142-26605188-2023	Voluntary Recall	HOYA VIVINEX TORIC PRELOADED INTRAOCULAR LENS	GC372051170718	Class II	A26: Insufficient Information	MEDFRAY MEDICAL SDN. BHD.	MDA-3605-WDP122
11/04/2023	MDA/Recall/P0146-18665945-2023	Voluntary Recall	HYDRASYS 2	IVDB53611300018	Class I	A02: Manufacturing, Packaging or Shipping Problem	UTAS MAJU SDN. BHD.	MDA-0974-WDP120
18/04/2023	MDA/Recall/P0150-50911713-2023	Voluntary Recall	EVIS EXERA III DUODENOVideoscope TJF-Q190V	GB4632321-70352	Class II	A22: Human-Device Interface Problem	OLYMPUS (MALAYSIA) SDN. BHD.	MDA-2218-WDP121

* 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.