



MEDICAL DEVICE RECALL LISTING JANUARY 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
13 January 2023	MDA/Recall/P0106-84504416-2023	Voluntary recall	COVERA PLUS COVERED VASCULAR STENT	GD8549319-32164	Class III	A05: Mechanical Problem	BECTON DICKINSON SDN. BHD.	MDA-0033-W3314
25 January 2023	MDA/Recall/P0111-68100620-2023	Voluntary recall	IRVINE SCIENTIFIC MEDIA FOR ASSISTED REPRODUCTIVE	GD56583937918	Class III	A04: Material Integrity Problem	LABIVF (M) SDN. BHD.	MDA-3218-WDP122
30 January 2023	MDA/Recall/P0113-95103752-2023	Voluntary recall	EXTENSION FLUID LINES FOR PRESSURE AND GRAVITY INFUSIONS	GB92776178817	Class II	A22: Human-Device Interface Problem	B. BRAUN MEDICAL INDUSTRIES SDN. BHD.	MDA-4250-W123

* 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.