MEDICAL DEVICE RECALL LISTING OCTOBER 2023

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
05/10/2023	MDA/Recall/P0197-79659247-2023	Voluntary Recall	NEX-LOAD SYSTEM SP	GC3790323-117204	Class III	A05: Mechanical Problem	EYE NATION MEDICAL SDN BHD	MDA-2018-WDP121
31/10/2023	MDA/Recall/P0215-43251362-2023	Voluntary Recall	ALINITY M HCV	IVDD3475320-42033	Class III	A02: Manufacturing, Packaging or Shipping Problem	ABBOTT LABORATORIES (MALAYSIA) SDN. BHD.	MDA-5104-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.