



## MEDICAL DEVICE RECALL LISTING JANUARY 2025

No	Date Received	Reference Number	Recall Type	Product Name	Product Registration	Recall Class	Reason for Recall	Recalling Establishment	Establishment License
1	03/01/2025	MDA/Recall /P0299-16637967-2024	Establishment (Voluntary Recall)	ACROBAT-I VACUUM STABILIZER SYSTEM	GB3805023-133245	Class II: Moderate Risk	A17: Compatibility Problem	RBD HEALTHCARE SDN BHD	MDA-4488-WDP123
2	09/01/2025	MDA/Recall /P0357-66712436-2025	Establishment (Voluntary Recall)	ZIMMER NEXGEN COMPLETE KNEE SOLUTION LEGACY CONSTRAINED CONDYLAR KNEE (LCCK) SYSTEM	GC85953379417	Class III: Low Risk	A05: Mechanical Problem	ZIMMER MALAYSIA (MALAYSIA) SDN BHD	MDA-6538-WDP124
3	13/01/2025	MDA/Recall /P0359-67093794-2025	Establishment (Voluntary Recall)	PRELOADED INTRAOCULAR LENS	GC2656025716	Class III: Low Risk	A02: Manufacturing, Packaging or Shipping Problem	CARL ZEISS SDN BHD	MDA-4561-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.