MEDICAL DEVICE RECALL LISTING APRIL 2026

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	03/04/2026	MDA/Recall/P0521-59732809-2026	Establishment (Voluntary Recall)	TENOR	GA6312419-29083	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	BEST CONTACT (M) SDN BHD	MDA-5525-WDP124
2.	02/04/2026	MDA/Recall/P0524-40215251-2026	Establishment (Voluntary Recall)	ZIMMER BONE CEMENT ACCESSORIES	GB619831022718	Class III: Low Risk	A02: Manufacturing, Packaging or Shipping Problem	ZIMMER MEDICAL (MALAYSIA) SDN BHD	MDA-6538-WDP124
3.	06/04/2026	MDA/Recall/P0525-72702643-2026	Establishment (Voluntary Recall)	SYRINGE	GB6979824-157861	Class III: Low Risk	A23: Use of Device Problem	H&A MEDICAL SUPPLY SDN BHD	MDA-6010-WDP124
4.	13/04/2026	MDA/Recall/P0527-66202214-2026	Establishment (Voluntary Recall)	INZONE DETACHMENT SYSTEM	GC54327161017	Class II: Moderate Risk	A07: Electrical /Electronic Property Problem	STRYKER CORPORATION (MALAYSIA) SDN. BHD.	MDA-5657-WDP124

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5.	22/04/2026	MDA/Recall/P0530-34168771-2026	Establishment (Voluntary Recall)	DLP® RETROGRADE CORONARY SINUS PERFUSION CANNULA	GB2266423-128554	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	MEDTRONIC MALAYSIA SDN BHD	MDA-4793-WDP123
6.	28/04/2026	MDA/Recall/P0533-16589695-2026	Establishment (Voluntary Recall)	GENTEAL LUBRICANT EYE DROPS	GB35237660118	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	ALCON LABORATORIES (M) SDN. BHD.	MDA-5021-W123
7.	28/04/2026	MDA/Recall/P0534-75693836-2026	Establishment (Voluntary Recall)	GENTEAL LUBRICANT EYE GEL	GB67890515817	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	ALCON LABORATORIES (M) SDN. BHD.	MDA-5021-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.