MEDICAL DEVICE RECALL LISTING FEBRUARY 2026

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	06/02/2026	MDA/Recall/P0495-17131337-2026	Establishment (Voluntary Recall)	SUPRASORB® A + AG	GD6355223-124001	Class III: Low Risk	A02: Manufacturing, Packaging or Shipping Problem	NYPRAX BUSINESS SOLUTIONS	MDA-5323-WDP123
2.	26/02/2026	MDA/Recall/P0513-38427720-2026	Establishment (Voluntary Recall)	INSTINCT PLUS™ ENDOSCOPIC CLIPPING DEVICE	GC7168824-176936	Class II: Moderate Risk	A05: Mechanical Problem	COOK ASIA (MALAYSIA) SDN BHD	MDA-5123-WDP123

* 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.