MEDICAL DEVICE RECALL LISTING JUNE 2026

No.	Date Received	Reference Number	Recall Type	Product Name	Product Registration Number	Recall Class	Reason of Recall	Recalling Establishment	Establishment License
1.	12/06/2026	MDA/Recall/P0538-80158382-2026	Establishment (Voluntary Recall)	GUEDEL AIRWAY	GA529191078418	Class II: Moderate Risk	A02: Manufacturing, Packaging or Shipping Problem	INTERSURGICAL SDN BHD	MDA-5088-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.