**MEDICAL DEVICE RECALL LISTING JULY 2024**

| Date Received | Reference No. | Recall Type | Product Name | Product Registration | Recall Class | Reason of Recall | Recalling Establishment | Establishment License |
|---|---|---|---|---|---|---|---|---|
| **01/07/2024** | MDA/Recall/P0302-75727245-2024 | Voluntary Recall | CAPIO SLIM SUTURE CAPTURING DEVICE | GB71551564718 | Class II | A24: Adverse Event Without Identified Device or Use Problem | BOSTON SCIENTIFIC (MALAYSIA) SDN BHD | MDA-5810-WD124 |
| **03/07/2024** | MDA/Recall/P0303-51703664-2024 | Voluntary Recall | ALINITY C PROTEINS | IVDB5170819-34873 | Class II | A08: Calibration Problem | ABBOTT LABORATORIES (MALAYSIA) SDN. BHD. | MDA-5104-W123 |
| **05/07/2024** | MDA/Recall/P0304-42114480-2024 | Voluntary Recall | VASCUGRAFT® NEO | GC37117108716 | Class III | A21: Labelling, Instructions for Use or Training Problem | B. BRAUN MEDICAL INDUSTRIES SDN. BHD. | MDA-4250-W123 |
| **09/07/2024** | MDA/Recall/P0305-15896132-2024 | Voluntary Recall | DSD EDGE ENDOSCOPE REPROCESSOR | GC16706518418 | Class II | A21: Labelling, Instructions for Use or Training Problem | STERIS MALAYSIA SDN. BHD. | MDA-5302-WDP123 |
| **09/07/2024** | MDA/Recall/P0309-17658684-2024 | Voluntary Recall | NIM-NEURO 3.0 SYSTEM AND ACCESSORIES | GB45802845818 | Class II | A23: Use of Device Problem | MEDTRONIC MALAYSIA SDN BHD | MDA-4793-WDP123 |
| **18/07/2024** | MDA/Recall/P0311-30582816-2024 | Voluntary Recall | MCGRATH MAC LARYNGOSCOPE | GB10147922-107333 | Class III | A27: Appropriate Term/Code Not Available | MEDTRONIC MALAYSIA SDN BHD | MDA-4793-WDP123 |
| **23/07/2024** | MDA/Recall/P0312-10509171-2024 | Voluntary Recall | SUSCEPTIBILITY AND IDENTIFICATION TEST FOR MYCOBACTERIA | IVDC23724200218 | Class II | A23: Use of Device Problem | BECTON DICKINSON SDN BHD | MDA-5083-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