**MEDICAL DEVICE RECALL LISTING FEBRUARY 2024**

| Date Received | Reference No. | Recall Type | Product Name | Product Registration | Recall Class | Reason of Recall | Recalling Establishment | Establishment License |
|---|---|---|---|---|---|---|---|---|
| **13/02/2024** | MDA/Recall/P0242-56360924-2024 | Voluntary Recall | DLP® VESSEL CANNULA | GB2870623-133586 | Class II | A02: Manufacturing, Packaging or Shipping Problem | MEDTRONIC MALAYSIA SDN. BHD. | MDA-4793-WDP123 |
| **19/02/2024** | MDA/Recall/P0246-39637683-2024 | Voluntary Recall | ARTERIAL CATHETERIZATION SET | GB35257208217 | Class II | A01: Patient Device Interaction Problem | TELEFLEX MEDICAL SDN.BHD. | MDA-3058-W121 |

* 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.