**MEDICAL DEVICE RECALL LISTING JULY 2023**

| Date Received | Reference No. | Recall Type | Product Name | Product Registration | Recall Class | Reason of Recall | Recalling Establishment | Establishment License |
|---|---|---|---|---|---|---|---|---|
| **04/07/2023** | MDA/Recall/P0183-98610616-2023 | Voluntary Recall | POWERSPIRAL | GB5498322-96108 | Class II | A27: Appropriate Term/Code Not Available | OLYMPUS (MALAYSIA) SDN. BHD. | MDA-2218-WDP121 |
| **10/07/2023** | MDA/Recall/P0184-81141062-2023 | Voluntary Recall | ACROBAT SUV VACUUM OFF-PUMP SYSTEM | GB3806023-134469 | Class II | A17: Compatibility Problem | RBD HEALTHCARE SDN BHD | MDA-4488-WDP123 |
| **13/07/2023** | MDA/Recall/P0185-35263762-2023 | Voluntary Recall | COBAS X 480 INSTRUMENT | IVDA87215200918 | Class III | A23: Use of Device Problem | ROCHE DIAGNOSTICS (M) SDN. BHD. | MDA-1674-WDP121 |
| **14/07/2023** | MDA/Recall/P0187-62671494-2023 | Voluntary Recall | BenchMark | IVDA7917823-115778 | Class III | A23: Use of Device Problem | ROCHE DIAGNOSTICS (M) SDN. BHD. | MDA-1674-WDP121 |
| **14/07/2023** | MDA/Recall/P0188-89730954-2023 | Voluntary Recall | KARL STORZ SINGLE-USE FLEXIBLE INTUBATION VIDEO ENDOSCOPE | GB5838321-52999 | Class I | A02: Manufacturing, Packaging or Shipping Problem | UMMI SURGICAL SDN BHD | MDA-4833-WDP123 |
| **24/07/2023** | MDA/Recall/P0190-39405912-2023 | Voluntary Recall | SINGLE USE SYRINGES, 2-PIECE | GA1919740916 | Class III | A02: Manufacturing, Packaging or Shipping Problem | B. BRAUN MEDICAL INDUSTRIES SDN. BHD. | MDA-4250-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.