



MEDICAL DEVICE RECALL LISTING APRIL 2022

Date Received	Reference No.	Recall Type	Product Name	Product Registration	Recall Class	Reason Of Recall	Recalling Establishment	Establishment License
1 April 2022	MDA/PMSV/R2022-019	Voluntary Recall	Exeter V40 Cemented Hip Stem	GC23592916318 GC8157019-32104	Class III	A21: Labelling, Instructions for Use or Training Problem	Stryker Corporation (Malaysia) Sdn Bhd	MDA-542-WDP44515
5 April 2022	MDA/PMSV/R2022-020	Voluntary Recall	Disposable Grasping Forceps [Model No: FG-51D]	GB267901024018	Class II	A05: Mechanical Problem	Olympus (Malaysia) Sdn Bhd	MDA-2218-WDP121
1 April 2022	MDA/PMSV/R2022-021	Voluntary Recall	Triathlon® Tritanium® Tibial Component (Size 6) – 5536-B-600 Triathlon® Tritanium® Tibial Component (Size 5) – 5536-B-500	GD42222611618	Class II	A21: Labelling, Instructions for Use or Training Problem	Stryker Corporation (Malaysia) Sdn Bhd	MDA-2123-WDP121
7 April 2022	MDA/PMSV/R2022-022	Voluntary Recall	Cobas cuvette	IVDA9186284817	Class III	A02: Manufacturing, Packaging or Shipping Problem	Roche Diagnostics (M) Sdn Bhd	MDA-1674-WDP121
1 April 2022	MDA/PMSV/R2022-023	Voluntary Recall	Reagent: Albumin in Urine/CSF (Microalbumin)	IVDB8526619-35899	Class II	A04: Material Integrity Problem	Symex Malaysia Sdn Bhd	MDA-515-WDP41815
13 April 2022	MDA/PMSV/R2022-024	Voluntary Recall	Arm and Tray Cover Sterile	GC82278603718	Class IIII	A02: Manufacturing, Packaging or Shipping Problem	Johnson & Johnson Sdn Bhd	MDA-0081-WDP415
14 April 2022	MDA/PMSV/R2022-025	Voluntary Recall	17731; 17732 - Gibeck® Humid-Vent® 2S Flex DC	GB439621028018 GB57567332017 GB628421055618	Class II	A12: Connection Problem A14: Infusion or flow problem	Teleflex Malaysia Sdn Bhd	MDA-3059-K121

* 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.



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			19061; 19062 - Gibeck® Humid-Vent® Filter Compact S with Smooth-Flo® / Swivel 19812 - Gibeck® Humid-Vent® Filter Compact S with Flex Tube 20011 - Flex Tube 22511; 22512 - Smooth-Flo® Flex with Double Swivel 22552 - Smooth-Flo® Flex					
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