



DQS CERTIFICATION (M) SDN. BHD.

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CAB REGISTRATION NUMBER: **MDA/CAB-006**
VALIDITY: **12/09/2020 - 11/09/2023**

SCOPE OF REGISTRATION

Conformity Assessment of Quality Management System (QMS) and Post-Market Surveillance System (PMSS)		
1	ISO 13485	Quality Management System for Medical Devices – Requirements for Regulatory Purpose
2	GDPMO	Good Distribution Practice for Medical Devices

Conformity Assessment of Technical Documentation		
3	MD 0101	Non-active devices for anesthesia, emergency and intensive care
4	MD 0102	Non-active devices for injection, infusion, transfusion and dialysis
5	MD 0103	Non-active orthopaedic and rehabilitation devices
6	MD 0106	Non-active instruments
7	MD 0107	Contraceptive medical devices
8	MD 0202	Non-active orthopaedic implants
9	MD 0301	Bandages and wound dressings
10	MD 0403	Dental implants
11	MD 1101	Devices for extra-corporal circulation, infusion and haemopheresis

Conformity Assessment by Way of Verification		
12	VERIFICATION	Conformity Assessment by Way of Verification

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Section 10(1), Medical Device Act 2012 (Act 737) and Regulation 8, Medical Device Regulations 2012