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Validity **MDA/CAB-001**: **21/11/2019 - 20/11/2022**

SCOPE OF REGISTRATION

Conformity Assessment of Quality Management System (QMS)		
(1)	GDPMD	Good Distribution Practice for Medical Devices
(2)	ISO 13485	Quality Management Systems for Medical Devices – Requirements for Regulatory Purpose
Conformity Assessment of Medical Device Technical Areas		
(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 0104	Non-active medical devices with measuring function
(7)	MD 0106	Non-active instruments
(8)	MD 0107	Contraceptive medical devices
(9)	MD 0109	Non-active devices for in vitro fertilization (IVF) and assisted reproductive technologies (ART)
(10)	MD 0301	Bandages and wound dressings
(11)	MD 0303	Other medical devices for wound care
(12)	MDS 7005	Medical devices referencing the Directive 89/686/EEC on personal protective equipment (PPE)
Conformity Assessment of Medical Device Technical Areas		
(13)	VERIFICATION	Conformity Assessment by Way of Verification

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