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CAB REGISTRATION NUMBER: **MDA/CAB-008**
VALIDITY: **12/09/2020 – 11/09/2023**

PERSON RESPONSIBLE:
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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	GDPMD	Good Distribution Practice for Medical Devices

Conformity Assessment of Technical Documentation		
3	MD 0106	Non-active instruments
4	MD 0107	Contraceptive medical devices
5	MD 0108	Non-active medical devices for disinfecting, cleaning, rinsing
6	MD 1111	Software
7	MD 1301	Monitoring devices of non-vital physiological parameters

Conformity Assessment by Way of Verification		
8	VERIFICATION	Conformity Assessment by Way of Verification

< End of List >

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