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CAB REGISTRATION NUMBER: **MDA/CAB-015**
VALIDITY: 27/02/2019 – 26/02/2022

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 0301 | Bandages and wound dressings |

| Conformity Assessment by Way of Verification | | |
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| 5 | 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