CERTIFICATE

This is to certify that the company

pro med instruments GmbH

Bötzinger Straße 86
79111 Freiburg
Germany

with the organizational units/sites as listed in the annex

has implemented and maintains a **Quality Management System**.

Scope of certification:
Design, development, manufacturing, service and distribution of surgical instruments and retractor systems, headrest systems, skull pins, bipolar electrosurgical forceps and cables.
-AUS (a), BRA, CND, JPN, USA (a,b,c,d)

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of the following standard:

ISO 13485 : 2016

including applicable country-specific regulatory requirements, as indicated below the scope (full references of abbreviations are listed in the annex)

Certificate registration no.	221096 MDSAP16
Certificate unique ID	170779077
Effective date	2022-01-25
Expiry date	2024-06-17
Frankfurt am Main	2022-01-25

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Szymon Kurdyn
Product Manager

Annex to certificate
Certificate registration No.: 221096 MDSAP16
Certificate unique ID: 170779077
Effective date: 2022-01-25

pro med instruments GmbH

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Audited site	REPs FEI No.: site scope and country-specific requirements
pro med instruments GmbH Bötzinger Straße 86 79111 Freiburg Germany	Design, development, manufacturing, service and distribution of surgical instruments and retractor systems, headrest systems, skull pins, bipolar electrosurgical forceps and cables; customer service, incoming inspection, storage and shipping. **- AUS (a), BRA, CND, JPN, USA (a,b,c,d)** **REPs FEI No.: F000625**

Annex to certificate
Certificate registration No.: 221096 MDSAP16
Certificate unique ID: 170779077
Effective date: 2022-01-25

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Full references of country-specific requirements of MDSAP participating Regulatory Authorities

Abbreviation	Jurisdiction	Reference
AUS	Australia	(a) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 1 – Full Quality Assurance Procedure (b) Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 4 – Production Quality Assurance Procedure
BRA	Brazil	RDC ANVISA n. 16/2013 RDC ANVISA n. 23/2012 RDC ANVISA n. 67/2009
CND	Canada	Medical Device Regulations SOR/98-282, Part 1
JPN	Japan	MHLW Ministerial Ordinance No. 169 (2004) as amended by MHLW Ordinance No. 128 (2014), Articles 4 to 68 Japan PMD Act (as applicable)
USA	United States	(a) 21 CFR Part 803 (b) 21 CFR Part 806 (c) 21 CFR Part 807 (d) 21 CFR Part 820 (e) 21 CFR Part 821