CERTIFICATE

This is to certify that the company

RUDOLF

RUDOLF Medical GmbH + Co. KG

Zollerstrasse 1
78567 Fridingen
Germany

with the organizational units/sites as listed in the annex
has implemented and maintains a **Quality Management System**.

Scope of certification:
The development, manufacture, service and sale of surgical instruments, equipment and following medical devices:
Suction systems, irrigation systems, insufflators, light sources, camera systems, image storage/transmission systems, equipment trolleys, self-retaining retractors, HF-surgical equipment, non-active implants, sterilization support systems and endoscopes for: arthroscopy, hysteroscopy, cystoscopy, resectoscopy, nephroscopy, ureterorenoscopy, laparoscopy, thoracoscopy, sinuscopy, otoscopy, neuroendoscopy, ventriculoscopy and microdisectomy.
-AUS (a), 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.	492576 MDSAP16
Certificate unique ID	170774645
Effective date	2021-07-30
Expiry date	2024-07-29
Frankfurt am Main	2021-07-30

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Szymon Kurdyn
Product Manager

Annex to certificate
Certificate registration No.: 492576 MDSAP16
Certificate unique ID: 170774645
Effective date: 2021-07-30

RUDOLF Medical GmbH + Co. KG

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78567 Fridingen
Germany

Audited site	REPs FEI No.: site scope and country-specific requirements
537401 **RUDOLF Medical GmbH + Co. KG** **Manufacture** Tuttlingerstr. 4 78567 Fridingen Germany	The development, manufacture, service and sale of surgical instruments, equipment and following medical devices: Suction systems, irrigation systems, insufflators, light sources, camera systems, image storage/transmission systems, equipment trolleys, self-retaining retractors, HF-surgical equipment, non-active implants, sterilization support systems and endoscopes for: arthroscopy, hysteroscopy, cystoscopy, resectoscopy, nephroscopy, ureterorenoscopy, laparoscopy, thoracoscopy, sinuscopy, otoscopy, neuroendoscopy, ventriculoscopy and microdisectomy. **-AUS (a), CND, JPN, USA (a,b,c,d)** **REPs FEI No.: F001971**
537402 **RUDOLF Medical GmbH + Co. KG** **Head Office** Zollerstrasse 1 78567 Fridingen Germany	The development, manufacture, service and sale of surgical instruments, equipment and following medical devices: Suction systems, irrigation systems, insufflators, light sources, camera systems, image storage/transmission systems, equipment trolleys, self-retaining retractors, HF-surgical equipment, non-active implants, sterilization support systems and endoscopes for: arthroscopy, hysteroscopy, cystoscopy, resectoscopy, nephroscopy, ureterorenoscopy, laparoscopy, thoracoscopy, sinuscopy, otoscopy, neuroendoscopy, ventriculoscopy and microdisectomy. **-AUS (a), CND, JPN, USA (a,b,c,d)** **REPs FEI No.: F000639**

Annex to certificate
Certificate registration No.: 492576 MDSAP16
Certificate unique ID: 170774645
Effective date: 2021-07-30

RUDOLF Medical GmbH + Co. KG

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78567 Fridingen
Germany

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