CERTIFICATE

This is to certify that the company

A.M.I. Agency for Medical Innovations GmbH

Im Letten 1
6800 Feldkirch
Austria

with the organizational units/sites as listed in the annex

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

Scope of certification:
Design and Development, Manufacturing and Distribution of medical devices for Coloproctology, Oncology, Laparoscopy, Morbid Obesity, Gynaecology/ Urogynaecology and Urology
-AUS (a), BRA, CND, 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.	066924 MDSAP16
Certificate unique ID	1000168841
Effective date	2024-06-13
Expiry date	2027-06-12
Frankfurt am Main	2024-04-01

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Marc Goedecke
Product Manager

Annex to certificate
Certificate registration No.: 066924 MDSAP16
Certificate unique ID: 1000168841
Effective date: 2024-06-13

A.M.I. Agency for Medical Innovations GmbH

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Audited site	REPs FEI No.: site scope and country-specific requirements
066924 **A.M.I. Agency for Medical Innovations GmbH** Im Letten 1 6800 Feldkirch Austria	Design and Development, Manufacturing and Distribution of medical devices for Coloproctology, Oncology, Laparoscopy, Morbid Obesity, Gynaecology/ Urogynaecology and Urology **-AUS (a), BRA, CND, USA (a,b,c,d)** **REPs FEI No.: F000632**

Annex to certificate
Certificate registration No.: 066924 MDSAP16
Certificate unique ID: 1000168841
Effective date: 2024-06-13

A.M.I. Agency for Medical Innovations GmbH

Im Letten 1
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Austria

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. 665/2022 RDC ANVISA n. 551/2021 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