



CERTIFICATE



This is to certify that the company

Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH

Brunner Str. 67
1230 Vienna
Austria

with the organizational units/sites as listed in the annex

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

Scope of certification:

Development, production and distribution of in-vitro diagnostics and analysing systems for haemostasis, protein and immune diagnostics.

-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.	543530 MDSAP16
Certificate unique ID	1000265340
Effective date	2026-02-04
Expiry date	2029-02-03
Frankfurt am Main	2025-12-29



DQS Medizinprodukte GmbH

Heinrich von Mettenheim
Managing Director



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DQS Medizinprodukte GmbH is recognized under the Medical Devices Single Audit Program.
The validity of the certification can only be verified by the QR-code.



Annex to certificate
Certificate registration No.: 543530 MDSAP16
Certificate unique ID: 1000265340
Effective date: 2026-02-04

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Brunner Str. 67
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Audited site

543530
Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH
Brunner Str. 67
1230 Vienna
Austria

REPs FEI No.: site scope and country-specific requirements

Development, production and distribution of in-vitro diagnostics and analysing systems for haemostasis, protein and immune diagnostics.
-AUS (a), BRA, CND, JPN, USA (a,b,c,d)
REPs FEI No.: F004534



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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. 665/2022 RDC ANVISA n. 551/2021 RDC ANVISA n. 67/2009
CND	Canada	Medical Devices Regulations – Part 1- SOR 98/282 Medical Devices Regulations – Part 1.1 – SOR 98/282 (as applicable)
JPN	Japan	MHLW Ministerial Ordinance 169, Article 4 to Article 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