



# CERTIFICATE



This is to certify that the company

## Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH

Brunner Str. 67  
1230 Wien  
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	170779833
Effective date	2023-02-04
Expiry date	2026-02-03
Frankfurt am Main	2022-12-11



### DQS Medizinprodukte GmbH

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Product Manager



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DQS Medizinprodukte GmbH is recognized under the Medical Devices Single Audit Program.

Visit <https://www.dqs.de/en/customer-database/> to validate this certificate.

The validity of this certificate can only be verified by the QR-code.



**Annex to certificate**  
**Certificate registration No.: 543530 MDSAP16**  
**Certificate unique ID: 170779833**  
**Effective date: 2023-02-04**

## **Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH**

Brunner Str. 67  
1230 Wien  
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### **Audited site**

**543530**  
**Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH**  
Brunner Str. 67  
1230 Wien  
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**



**Annex to certificate**  
**Certificate registration No.: 543530 MDSAP16**  
**Certificate unique ID: 170779833**  
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### **Full references of country-specific requirements of MDSAP participating Regulatory Authorities**

<b>Abbreviation</b>	<b>Jurisdiction</b>	<b>Reference</b>
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