



# CERTIFICATE



This is to certify that the company

## Ritter Implants GmbH & Co.KG

Freiburger Straße 45  
88400 Biberach  
Germany

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 Dental Implant Systems, Abutments, Surgical Kits, Drills and Instruments.  
-CAN, 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

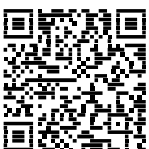
including applicable country-specific regulatory requirements, as indicated below the scope (full references of abbreviations are listed in the annex)

Certificate registration no.	549603 MDSAP16
Certificate unique ID	1000235416
Effective date	2025-08-16
Expiry date	2028-08-15
Frankfurt am Main	2025-08-03



DQS Medizinprodukte GmbH

Heinrich von Mettenheim  
Managing Director



August-Schanz-Straße 21, 60433 Frankfurt am Main,  
Tel. +49 (0) 69 95427-300, [info-med@dqs.de](mailto:info-med@dqs.de)  
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 the certification can only be verified by the QR-code.



Annex to certificate  
Certificate registration No.: 549603 MDSAP16  
Certificate unique ID: 1000235416  
Effective date: 2025-08-16

## Ritter Implants GmbH & Co.KG

Freiburger Straße 45  
88400 Biberach  
Germany

### Audited site

549603  
Ritter Implants GmbH & Co.KG  
Freiburger Str. 45  
88400 Biberach  
Germany

550472  
Ritter Implants GmbH & Co.KG  
Freiburger Str. 32  
88400 Biberach  
Germany

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

Design and development, manufacturing and  
distribution of Dental Implant Systems,  
Abutments, Surgical Kits, Drills and Instruments.  
-CAN, USA (a,b,c,d)  
REPs FEI No.: F006159

Design and development, manufacturing and  
distribution of Dental Implant Systems,  
Abutments, Surgical Kits, Drills and Instruments.  
-CAN, USA (a,b,c,d)  
REPs FEI No.: F006159



Annex to certificate  
Certificate registration No.: 549603 MDSAP16  
Certificate unique ID: 1000235416  
Effective date: 2025-08-16

## Ritter Implants GmbH & Co.KG

Freiburger Straße 45  
88400 Biberach  
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. 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