



CERTIFICATE



This is to certify that the company



Osstell AB

Stampgatan 14
41101 Göteborg
Sweden

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

Scope of certification:

Design, manufacturing and distribution of sterile and non-sterile medical devices for determining stability of human bone-anchored implants.

-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.	543847 MDSAP16
Certificate unique ID	1000213837
Effective date	2025-09-20
Expiry date	2028-09-19
Frankfurt am Main	2025-08-26



DQS Medizinprodukte GmbH

Heinrich von Mettenheim
Managing Director



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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 the certification can only be verified by the QR-code.



Annex to certificate
Certificate registration No.: 543847 MDSAP16
Certificate unique ID: 1000213837
Effective date: 2025-09-20

Osstell AB

Stampgatan 14
41101 Göteborg
Sweden

Audited site

543847
Osstell AB
Stampgatan 14
41101 Göteborg
Sweden

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

Design, manufacturing and distribution of
sterile and non-sterile medical devices for
determining stability of human bone-
anchored implants.

-AUS (a), BRA,CND, JPN, USA (a,b,c,d)

REPs FEI No.: F004960



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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