



CERTIFICATE



This is to certify that the company

Parker Laboratories, Inc.

286 Eldridge Road
Fairfield, NJ 07004
United States of America

with the organizational units/sites as listed in the annex

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

Scope of certification:

Design & Development, Manufacturing and Distribution of Gels, Lotions, Sterile Securement Dressings, and Probe Covers for the Areas of Ultrasound and Electromedical Procedures.
-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

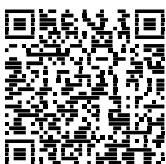
including applicable country-specific regulatory requirements, as indicated below the scope (full references of abbreviations are listed in the annex)

Certificate registration no.	31622677 MDSAP16
Certificate unique ID	1000208822
Effective date	2025-04-14
Expiry date	2028-04-13
Frankfurt am Main	2025-04-14



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.: 31622677 MDSAP16
Certificate unique ID: 1000208822
Effective date: 2025-04-14

Parker Laboratories, Inc.

286 Eldridge Road
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United States of America

Audited site

31622679
Parker Laboratories, Inc.
286 Eldridge Road
Fairfield, NJ 07004
United States of America

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

Design & Development and Manufacturing of Gels, Lotions, Sterile Securement Dressings, and Probe Covers for the Areas of Ultrasound and Electromedical Procedures.
-AUS (a), BRA, CND, JPN, USA (a,b,c,d)
REPs FEI No.: F003631

31622678
Parker Laboratories, Inc.
4 Sperry Road
Fairfield, NJ 07004
United States of America

Design & Development, Manufacturing and Distribution of Gels, Lotions, Sterile Securement Dressings, and Probe Covers for the Areas of Ultrasound and Electromedical Procedures.
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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