



CERTIFICATE



This is to certify that the company

Miele

Miele & Cie. KG

Carl-Miele-Straße 29
33332 Gütersloh
Germany

with the organizational units/sites as listed in the annex

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

Scope of certificate and applicable country-specific requirements:

Design, development, production, distribution and service of appliances for cleaning and disinfection of medical devices, Distribution and service of appliances for sterilization of medical devices.

-AUS (a), 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 (MDSAP Audit Model Edition 2)

including country-specific requirements as shown in the scope
(full references are listed in the annex)

Certificate registration no.	294819 MDSAP16
Certificate unique ID	170714121
Effective date	2018-10-27
Expiry date	2021-10-26
Frankfurt am Main	2018-10-27



DQS Medizinprodukte GmbH

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

Visit <https://www.mydqs.com/en/customers/customer-database.html> to validate this certificate.



Annex to certificate

Certificate registration No.: 294819 MDSAP16

Certificate unique ID: 170714121

Effective date: 2018-10-27

Miele & Cie. KG

Carl-Miele-Straße 29

33332 Gütersloh

Germany

Audited site

DUNS No., site scope and country-specific requirements

Miele & Cie. KG

Carl-Miele-Straße 29

33332 Gütersloh

Germany

Distribution and service of appliances for cleaning and disinfection of medical devices.

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

DUNS No.: 315206441

Miele & Cie. KG

Mielestraße 2

33611 Bielefeld

Germany

Design, development and production of appliances for cleaning and disinfection of medical devices.

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

DUNS No.: 315206441



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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. 16/2013 RDC ANVISA n. 23/2012 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