



CERTIFICATE



This is to certify that the company

Haag-Streit AG

Gartenstadtstrasse 10
3098 Koeniz
Switzerland

with the organizational units/sites as listed in the annex

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

Scope of certification:

Design, development and manufacturing of mechanical, optical and electronic precision instruments including related software for ophthalmologic application.

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

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.	335325 MDSAP16
Certificate unique ID	170776376
Effective date	2022-01-11
Expiry date	2025-01-10
Frankfurt am Main	2021-12-12



DQS Medizinprodukte GmbH

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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.



Annex to certificate

Certificate registration No.: 335325 MDSAP16

Certificate unique ID: 170776376

Effective date: 2022-01-11

Haag-Streit AG

Gartenstadtstrasse 10
3098 Koeniz
Switzerland

Audited site

335325

Haag-Streit AG

Gartenstadtstrasse 10
3098 Koeniz
Switzerland

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

Design, development and manufacturing of
mechanical, optical and electronic precision
instruments including related software for
ophthalmologic application.

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

REPs FEI No.: F000619



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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