



America

CERTIFICATE

No. QS6 084922 0007 Rev. 02

Certificate Holder: Copan Wasp S.r.l.
Via A. Grandi 32
25125 BRESCIA
ITALY

Certification Mark:



Scope of Certificate: Design and Development, Production, Distribution, Installation and Servicing of In-Vitro Diagnostic Instruments and Software for Microbiology;
Design and Development, Production, Distribution of In-Vitro Diagnostic Sterile and Non-Sterile Reagents for Microbiology;
Design and Development, Production, Distribution of In-Vitro Diagnostic Generic Use Consumables for Microbiology Devices

Standard(s): ISO 13485:2016

Regulatory Authority(ies): Australia TGA, Brazil ANVISA, Health Canada, Japan MHLW / PMDA, USA FDA. See attached for listing of specific regulatory requirements.

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. For details and certificate validity see:

www.tuvsud.com/ps-cert?q=cert:QS6_084922_0007_Rev.02

TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

REPs Facility ID: F005494
Report No.: ITA200220001031_CN
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(Renee Walker)
Director, US Certification Body, MHS

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Regulatory Requirements: Audit/Certification Criteria

Australia

Therapeutic Goods (Medical Devices) Regulations 2002
 - Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil

- RDC ANVISA n. 665/2022 - Good Manufacturing Practices
 - RDC ANVISA n. 551/2021
 - RDC ANVISA n. 67/2009 - Vigilance

Canada

- Medical Device Regulations – Part 1- SOR 98/282

Japan

- MHLW Ministerial Ordinance No. 169 (2004), as amended by MHLW Ministerial Ordinance No.60 (2021)
 - Japan PMD Act (as applicable)

United States

- 21 CFR Part 803
 - 21 CFR Part 806
 - 21 CFR Part 807 – Subparts A to D
 - 21 CFR Part 820

Facility(ies):

Copan Wasp S.r.l.

Via A. Grandi 32, 25125 BRESCIA, ITALY

Facility Scopes:

Design and Development, Production, Distribution, Installation and Servicing of In-Vitro Diagnostic Instruments and Software for Microbiology;

Design and Development, Production, Distribution of In-Vitro Diagnostic Sterile and Non-Sterile Reagents for Microbiology;

Design and Development, Production, Distribution of In-Vitro Diagnostic Generic Use Consumables for Microbiology Devices
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