



Certificate

No. Q5 109578 0004 Rev. 00

Holder of Certificate: **Agilent Technologies Singapore
(International) Pte. Ltd.**

No. 1 Yishun Avenue 7
Singapore 768923
SINGAPORE

Certification Mark:



Scope of Certificate: **The Design, Development, Manufacture and Distribution of Antibodies, Reagents and Kits for In-vitro Diagnostics used in Diagnosis and Management of Cancer, Genetic Testing, Immune Status, Disease Status, Autoimmune Status, Protein Metabolism, Blood Analytes, Coagulation, Transmissible Agents, Sexually Transmissible Agents and Immunological Typing The Design, Development, Manufacture, Distribution, Installation and Service of In-vitro Diagnostic Instruments and Software used in the Diagnosis and Management of Cancer, Genetic Testing, Immune Status, Disease Status, Autoimmune Status, Protein Metabolism, Blood Analytes, Coagulation, Transmissible Agents, Sexually Transmissible Agents and for Immunological Typing**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5_109578_0004_Rev._00

Report No.: 72185314

Valid from: 2023-12-15

Valid until: 2026-12-14

Date, 2023-12-15



Christoph Dicks
Head of Certification/Notified Body

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Applied Standard(s): ISO 13485:2016
(EN ISO 13485:2016/AC:2018, EN ISO 13485:2016/A11:2021)
Medical devices - Quality management systems -
Requirements for regulatory purposes

Facility(ies): **Agilent Technologies Singapore (International) Pte.
Ltd.**
No. 1 Yishun Avenue 7, Singapore 768923, SINGAPORE

Management.

Agilent Technologies Denmark ApS
Produktionsvej 42, 2600 Glostrup, DENMARK

See Scope of Certificate.