





Certificate

No. Q5 113968 0003 Rev. 01

Holder of Certificate: Agilent Technologies, Inc.

5301 Stevens Creek Boulevard Santa Clara CA 95051

USA

Certification Mark:



The Design and Development, and Scope of Certificate:

Manufacture of In-vitro Diagnostic Test Kits,

Reagents used in the Diagnosis and/or Management of Cancer, Immune Status, Disease Status, Autoimmune Status, Blood

Analytes, Immunological Typing and

Disease Management

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 113968 0003 Rev. 01

Report No.: 72185439

Valid from: 2023-08-17 Valid until: 2026-08-16

Christoph Dicks 2023-08-18 Date,

Head of Certification/Notified Body





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EN ISO 13485:2016 Applied Standard(s):

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016) DIN EN ISO 13485:2016

Facility(ies): Agilent Technologies, Inc.

5301 Stevens Creek Boulevard, Santa Clara CA 95051, USA

Management and Administration

Agilent Technologies, Inc.

6392 Via Real, Carpinteria CA 93013, USA

Design and Development of In-Vitro Diagnostic Test Kits, Reagents used in the Diagnosis and/or Management of Cancer, Immune Status, Blood Analytes, Immunological Typing and Disease Management

Agilent Technologies, Inc.

1170 Mark Ave., Carpinteria CA 93013, USA

Manufacture of In-Vitro Diagnostic Test Kits, Reagents used in the Diagnosis and/or Management of Cancer, Immune Status, Blood Analytes, Immunological Typing and Disease Management