





**Product Service** 

## **Certificate**

No. Q5 082138 0005 Rev. 05

**Holder of Certificate:** Agilent Biosciences (Hangzhou) Co., Ltd.

Building 4 and Building 3, No. 208, Zhenzhong Road

Xihu District

310030 Hangzhou, Zhejiang PEOPLE'S REPUBLIC OF CHINA

**Certification Mark:** 



Design and Development, Production and Distribution of In Scope of Certificate: Vitro Diagnostic Reagents, Controls, Instruments and

**Consumables for Flow Cytometry and Cytology** 

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, Certification, Validation and Verification Regulations TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5 082138 0005 Rev. 05

Report No.: SH2475201 Valid from: 2024-12-05 Valid until: 2027-12-04

2024-09-05

Christoph Dicks

Head of Certification/Notified Body





Date,







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No. Q5 082138 0005 Rev. 05

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 Biosciences (Hangzhou) Co., Ltd.

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See Scope of Certificate

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