



Certificate

No. Q6 090019 0002 Rev. 01

Holder of Certificate: **MP Biomedicals Germany GmbH**
Thüringer Straße 15
37269 Eschwege
GERMANY

Certification Mark:



Scope of Certificate: **Manufacturing and distribution of in-vitro-diagnostic devices for infectiology, immunology and clinical chemistry**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system (excluding subclause 7.3), 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:Q6 090019 0002 Rev. 01](http://www.tuvsud.com/ps-cert?q=cert:Q6_090019_0002_Rev._01)

Report No.: 713204659

Valid from: 2021-04-13

Valid until: 2024-04-12

Date, 2021-04-08



Christoph Dicks

Head of Certification/Notified Body

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Applied Standard(s):

EN ISO 13485:2016
Medical devices - Quality management systems -
Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

Facility(ies):

MP Biomedicals Germany GmbH
Thüringer Straße 15, 37269 Eschwege, GERMANY

Manufacturing and distribution of in-vitro-diagnostic devices for
infectiology, immunology and clinical chemistry

Parameters: ./.