ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

DAkkS Deutsche Akkreditierungsstelle D-ZM-11321-01-00

TÜV SÜD Product Service

Certificate

No. Q5 068207 0053 Rev. 02

Holder of Certificate: **MP Biomedicals Asia Pacific Pte Ltd**
2 Pioneer Place
Singapore 627885
SINGAPORE

Certification Mark:

Scope of Certificate:

Design & Development, Production and Distribution for In-Vitro Diagnostic Devices for Infectious Diseases, Immunology and Molecular Diagnostics;

Production and Distribution of Instruments and Stand-alone Software Supporting In-Vitro Diagnostics;

Provision of production service of Antigen, Antibodies and Proteins as raw material for In-Vitro Diagnostics Devices;

The provision of Design and Development, Production of In-Vitro Diagnostic Devices

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 068207 0053 Rev. 02

Report No.: SIN_8017068_RA_2023

Valid from: 2023-10-01
Valid until: 2026-09-30

Date, 2023-09-25

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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No. Q5 068207 0053 Rev. 02

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 Asia Pacific Pte Ltd
2 Pioneer Place, Singapore 627885, SINGAPORE

See Scope of Certificate

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