



Malaysia

CERTIFICATE

No. GDPMD 088574 0003 Rev. 02

Holder of Certificate: **QIAGEN Biotechnology Malaysia Sdn Bhd**
Level 9, Tower 8
Avenue 5, Horizon Phase 2
Bangsar South
No.8 Jalan Kerinchi
59200 Kuala Lumpur
MALAYSIA

Facility(ies): QIAGEN Biotechnology Malaysia Sdn Bhd
Level 9, Tower 8, Avenue 5, Horizon Phase 2, Bangsar South,
No.8 Jalan Kerinchi, 59200 Kuala Lumpur, MALAYSIA

Scope of Certificate: **Local Authorized Representative, Import,
Storage and handling, Warehousing,
Distribution (inc. transportation),
Installation, testing & commissioning,
Maintenance and calibration and
Documentation incl. traceability of
In-vitro Diagnostic Devices**

Applied Standard(s): Good Distribution Practice for Medical Device
(GDPMD)
Appendix 4 Schedule 3 Medical Device
Regulation 2012 of Malaysia

The Conformity Assessment Body of TÜV SÜD (Malaysia) Sdn. Bhd. certifies that the company mentioned above has established and is maintaining a quality 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:GDPMD 088574 0003 Rev. 02

Report No.: 5774719-721432054

Valid from: 2023-08-21
Valid until: 2026-08-20

Date: 2023-05-19

Vincent Lam Chee Choong

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TÜV SÜD (Malaysia) Sdn. Bhd. is a Conformity Assessment Body according to Malaysia Medical Device Regulations with CAB Registration Number of MDA/CAB-001.