



Deutsche
Akkreditierungsstelle
D-ZM-11321-01-00



Product Service

Certificate

No. Q5 120404 0002 Rev. 01

Holder of Certificate: **iLife Medical Devices Pvt Ltd**
86, Sector-56, Phase-IV, IE Kundli
Sonapat, Haryana 131028
INDIA

Facility(ies): **iLife Medical Devices Pvt Ltd**
86, Sector-56, Phase-IV, IE Kundli, Sonapat, Haryana 131028,
INDIA

See Scope of Certificate

Certification Mark:



Scope of Certificate: **Design and Development, Manufacture, Sales and Distribution of Sterile Disposable Medical Devices VIZ.I.V. Cannula, Threeway Stop Cock (with and without extension tube), Extension Tubes, I V Set and B T Set.**

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

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 120404 0002 Rev. 01

Report No.: TPS1752_RA
Valid from: 2025-06-03
Valid until: 2028-06-02

Date, 2025-06-02

Christoph Dicks
Head of Certification/Notified Body