





CERTIFICATE

No. QS6 067069 0028 Rev. 05

Certificate Holder: LINET spol. s r.o.

Želevčice 5 274 01 Slaný

CZECH REPUBLIC

Certification Mark:



Scope of Certificate: Design and Development, Manufacture, Distribution,

Servicing and Installation of Hospital Beds, Nursing Beds, Birthing Beds, ICT Solutions, Stretchers, Anti-Decubitus Systems and Mattresses and Furniture for the areas of Hospital Care, Gynecology, Social Care and Home Care

Standard(s): ISO 13485:2016

Regulatory Authority(ies): Australia TGA, Brazil ANVISA, Health Canada, Japan MHLW

/ PMDA, USA FDA. See attached for listing of specific

regulatory requirements.

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:QS6 067069 0028 Rev. 05

TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

REPs Facility ID: F002178
Report No.: 713282314
Effective Date: 2023-06-28
Expiry Date: 2024-09-16

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Date of Issue: 2023-07-25

(Renee Walker)

Director, US Certification Body, MHS





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Regulatory Requirements: Audit/Certification Criteria

Australia

Therapeutic Goods (Medical Devices) Regulations 2002 - Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil

- RDC ANVISA n. 665/2022 - Good Manufacturing Practices

- RDC ANVISA n. 551/2021

- RDC ANVISA n. 67/2009 - Vigilance

Canada

- Medical Device Regulations - Part 1- SOR 98/282

Japan

- MHLW Ministerial Ordinance No. 169 (2004), as amended by MHLW Ministerial Ordinance No.60 (2021)

- Japan PMD Act (as applicable)

United States

- 21 CFR Part 803

- 21 CFR Part 806

- 21 CFR Part 807 – Subparts A to D

- 21 CFR Part 820

Facility(ies): LINET spol. s r.o.

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Politických vězňů 1969, 274 01 Slaný, CZECH REPUBLIC

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