Global Inter Certification

Management System Certification Body No.MSCB-108

CERTIFICATE

No. 22-B-0090 Rev.1/a

This is to certify that the Medical Devices-Quality Management Systems of

Feel Tech Co., Ltd.

4 Floor, Standard Factory 5-dong, 1 Floor, Standard Factory 1-dong, 3, 4 Floor, Standard Factory 2-dong, 15, Jayumuyeok 2-gil, Gunsan-si, Jeollabuk-do, Korea

Company Reg. No.: 401-81-38751

has documented and implemented system in compliance with the requirements of

ISO 13485:2016
Medical Devices-Quality Management Systems

for

Design & Development, Production and Sales of Sterile Single Use Polydioxanone Suture with Needle, Sterile Single Use PLLA (Poly-L-Lactic Acid) Suture with Needle, Sterile Single Use Polycaprolactone Suture with Needle, Disposable Syringe with Needle, Disposable Syringe Needle, Cannula, General-Purpose and Sterile P (LA/CL) (Poly (L-Lactide-co-ε-Caprolactone) suture with needle

Technical Area:
Non-active Medical Devices - General non-active, non-implantable medical devices
Non-active Medical Devices - Devices for wound care

The certificate is issued on the basis of the results mentioned in the pertinent audit report. Validity of the certificate is conditionally limited by positive results of surveillance audits, which the certified company is committed to undergo.

This certificate can be invalid if the certificate holder does not fulfill the conditions set out in the certification agreement.

ISO 13485 CERTIFIED

ACCREDITED Management Systems Certification Body MSCB 108

Initial issue date:	Jul. 09. 2019
Re-issue date:	Nov. 30. 2023
Expire date:	Jul. 08. 2025

Tyrone Dyse
Head of Certification Body

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