



Reg. No. 305/Q-054

CERTIFICATE

This certifies that the Quality management system for medical devices of company

REMEDI Co., Ltd.

Head Office and manufacturing site:

84, Osongsaengmyeong 2-ro, Osong-eup, Heungdeok-gu, Cheongju-si, Chungcheongbuk-do 28161,
Republic of Korea

*has been assessed by 3EC International
and found to be in conformance with the following standard:*

EN ISO 13485:2016

for the following scope:

**DESIGN, DEVELOPMENT, PRODUCTION AND SERVICE OF ACTIVE MEDICAL DEVICES:
ELECTROMAGNETIC STIMULATOR, TRANSCRANIAL MAGNETIC STIMULATOR,
EXTRACORPOREAL SHOCK WAVE THERAPY AND ELECTROSURGICAL UNIT**

Certificate No.: M-0473/25

Date of issuance: August 18th, 2025


Original date of approval: May 25th, 2020

This certificate is valid from August 18th, 2025 to May 24th, 2026 on condition that organization will maintain effective Quality management system for medical devices. To verify the validity of this certificate please contact our office at: +421 (0)2 5831 8343.

This certificate fully supersedes previous certificate No. M-0473/24 issued on June 24th, 2024.

Issuing office: 3EC International a. s., Hraničná 18, 821 05 Bratislava, Slovak Republic




Dr. Katarína Tomin Srdošová
Head of Certification Body 3EC International a. s.

Certification body 3EC International a. s. is accredited by SNAS under registration number 305/Q-054 with accreditation certificate No. Q-054 for certification of Quality management systems for medical devices covered by EA MLA and IAF MLA.