





CERTIFICATE

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

SPL Life Sciences Co., Ltd.

48, Geumgang-ro 2047 beon-gil, Naechon-myeon, Pocheon-city, Gyeonggi-do, 11192, Korea E-beam radiation site: 262, Naejin-ro, Naechon-myeon, Pocheon-si, Gyeonggi-do, 11190, 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, MANUFACTURE AND SALES OF NON ACTIVE MEDICAL DEVICES

AND IN VITRO DIAGNOSTIC MEDICAL DEVICES:

IVF PRODUCTS, CELL CULTURE SLIDE, CELL CULTURE PLATE, CELL CULTURE DISH, CELL CULTURE FLASK, CELL STRAINER, CONICAL TUBE, SNAP TUBE, CRYOVIAL, SEROLOGICAL PIPETTE, CYTO PAP BRUSH, MEDIA BOTTLE, ROLLER BOTTLE

PROVISION E-BEAM IRRADIATION SERVICE OF NON ACTIVE MEDICAL DEVICES AND IN VITRO DIAGNOSTIC MEDICAL DEVICES AS PER ISO11137-1, ISO11137-2, ISO11137-3

Certificate No.: M-0369/24

Date of issuance: September 19th, 2024

Original date of approval: October 7th, 2015

This certificate is valid from October 5th, 2024 to October 4th, 2027 on condition that organization will maintain effective quality management system. To verify the validity of this certificate please contact our office at: +421 (0)2 5831 8343.

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

Dr. Katarina 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.