Certificate of Registration

In accordance with European Communities Council Directive 98/79/EC as amended, concerning In Vitro Medical Devices as transposed into European national law by the member states

Certificate No.
KOR/2015/07/01

Certificate issue date;
1st November 2017

Certificate expiry date;
31st October 2018

We hereby declare that

- **An examination has been made of this organisation's Declaration of Conformity(s) and where appropriate Notified Body certification(s) exist.**
- **The EU Authorised Representative contract has been fulfilled**

And the CE mark may be applied to the products listed below.

Organisation / Client:

SPL Life Sciences Co., Ltd.
26 Geumgang-ro 2047 beon-gil
Naechon-Myeon, Pocheon-si
Gyeonggi-do 487 835
Republic of Korea

Products:

Cell Strainer, Cell Culture Plate, Cell Culture Dish, Cell Culture Slide, Cell Culture Flask & Cryovial
EDMA Code 23051001 Histology/Cytology consumables
See attached EC Declaration of Conformity including product schedule

Competent Authority Information:

IVD Medical Device Directive registration is with the UK Medicines and Healthcare Regulatory Agency (MHRA) and the below registration has been issued.
IVD000839

Authorised Representative Labelling Information:

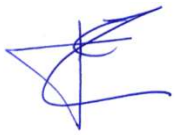
EC REP Advena Ltd. Pure Offices, Plato Close, Warwick CV34 6WE UK.

Advena Limited.

Registered office;
Pure Offices, Plato Close, Tachbrook Park
Warwick CV34 6WE. United Kingdom
Registered in England & Wales No. 3517275

☎ +44 1926 800153
Email; info@advenamedical.com

Authorised Signature:



This certificate is subject to the organisation maintaining their documentation in compliance with the regulations stated in this certificate.