

Certificate of Registration



Client Ref. **GBR/2021/01/34**

Issued To: **Cell Projects Ltd**
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Ashford Road,
Harrietsham, Kent,
ME17 1AB, UK

Legal Manufacturer [SRN: GB-MF-000002266]

Issued By: **Advena Limited**
Tower Business Centre, 2nd Flr, Tower
Street, Swatar, BKR 4013. Malta.

EU-REP [SRN: MT-AR-000000234]

EU Competent Authority: **Malta Medicines Authority (MMA)**
Sir Temi Zammit Buildings, Malta Life
Sciences Park, San Gwann SGN 3000
Malta.
Tel: +356 2343 9000
Email: info.medicinesauthority@gov.mt

Eudamed Actor ID: MT-CA-019

We hereby declare that:

- Device registrations for the medical devices mentioned within this certificate have duly been completed with the Malta Medicines Authority (MMA) the Competent Authority of Malta.
- Due to the 26th May 2021 Date of Application of Regulation (EU) 2017/745 (MDR) the validity of this certificate is subject to the Legal Manufacturer providing satisfactory evidence that any device claiming compliance to Directive 93/42/EEC (MDD) through Article 120 (3) of Regulation (EU) 2017/745 as amended, is legitimately permitted.
- Due to the 26th May 2022 Date of Application of Regulation (EU) 2017/746 (IVDR) the validity of this certificate is subject to the Legal Manufacturer providing satisfactory evidence that any device claiming compliance to Directive 98/79/EC (IVDD) through Article 110(3) of Regulation (EU) 2017/746 as amended, is legitimately permitted.

Anthony Kirby – Managing Director

Date of Issue: 24 March 2026

AR Cover Begins: 01 March 2026

AR Cover Ends: 28 February 2027

Appendix A



Product Details, Names or Trade Names	EU Legislation	Classification	Device Registration Reference(s)
GeneFiX DNA/RNA Collector, Non-Sterile.	IVDR	Class A	DVC-MT-21-06-000377
Isohelix Swab Pack, Non-Sterile	MDR	Class I	MT-MDF03-AA243A
STOOLFIX MICROBIOME DNA COLLECTOR, Non-Sterile	IVDR	Class A	MT-MDF23-AA060



This certificate is subject to the organisation maintaining their documentation in compliance with the EU legislation as indicated in this certificate.

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