

Certificate of Registration



Certificate No. CE/GBR/2020/07/14

Issued To: **Rapid Labs Ltd**
Unit 2 & 2a Hall Farm Business Centre,
Church Road
Little Bentley
Colchester, Essex
CO7 8SD
UK

Legal Manufacturer [SRN: Not yet available]

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

EC-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

We hereby declare that:

- Device registrations for the medical devices mentioned within this certificate have duly been completed with an EU Competent Authority.
- 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 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 is legitimately permitted.

Anthony Kirby – Managing Director

Date of Issue: 1 December 2021

AR Cover Begins: 01 December 2021

AR Cover Ends: 30 November 2022

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

This certificate is for the exclusive use of Advena Ltd's clients and is provided pursuant of the European Authorised Representative agreement (Mandate) between Advena Ltd and the client. Advena's responsibility and liability is limited to the terms and conditions of this agreement. Advena Ltd assumes no liability to any party for any loss, expense or damage occasioned by the use of this certificate and the European Authorised Representative agreement (Mandate). Only the client is authorised to copy or distribute this certificate. Any use of the Advena Ltd name by others who are not covered by the above agreement, or any similar contract, is prohibited. This certificate remains valid until the expiry date has been reached or has been terminated by Advena Limited.

Appendix A



Product Details, Names or Trade Names	EU Legislation	Classification	Device Registration Reference(s)
ELISA kits	IVDD	General IVD	DVC-MT-21-01-000037
Febrile antigens reagents & kits	IVDD	General IVD	DVC-MT-21-01-000038
Latex and slide agglutination kits	IVDD	General IVD	DVC-MT-21-01-000039
Lectins	IVDD	General IVD	DVC-MT-21-01-000040
Rapid Strips & devices	IVDD	General IVD	DVC-MT-21-01-000041
Syphilis reagents & kits	IVDD	General IVD	DVC-MT-21-01-000042
Urinalysis	IVDD	General IVD	DVC-MT-21-01-000043
virusPHIX	IVDD	General IVD	DVC-MT-21-01-000044
Anti-Human Globulin	IVDD	List B	DVC-MT-21-01-000045
Blood grouping reagents	IVDD	List A	DVC-MT-21-01-000046
Culture media	IVDD	General IVD	DVC-MT-21-01-000047
Antibiotic discs & rings	IVDD	General IVD	DVC-MT-21-01-000048

