Certificate of Declaration

CE

We hereby declare that the technical file of product complied with the requirement of directives (98/79/EC) In-Vitro Diagnostic Devices Directive.

Certificate No.: CE-4404

Manufacturer

Name : **KREATIVE TECHNOLABS**

Address : **Ste# 619, 606 S Hill Street, Los Angeles, CA**

Product : **SARS-Cov-2 Real Time PCR Kit Pack Size: 100 Determinations**

The Certification body has performed an audit of the above product quality system covering the design, manufacture and final inspection of the certified product. The quality system has been assessed, approved and is subject to continuous surveillance according to Directive (98/79/EC) In-Vitro Diagnostic Devices Directive.

This certificate is issued under the following conditions:

1. It applies only to the quality system maintained in the manufacture of above referenced models and it does not substitute the design or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.

The CE mark as shown above can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product. It does not imply an assessment of the whole production.

Validity of this certificate can be verified at www.ukcertifications.org.uk/verify

Date of Certification	04th February 2020
1st Surveillance Audit Due	03rd February 2021
2nd Surveillance Audit Due	03rd February 2022
Certificate Expiry (subject to the company maintaining its system to the required standard)	03rd February 2023

Authorised Signatory

UK Certification & Inspection Ltd.