CERTIFICATE OF REGISTRATION



Lorne Laboratories Ltd

Unit 1 Cutbush Park Industrial Estate
Danehill
Lower Earley
Berkshire RG6 4UT UNITED KINGDOM

UL LLC®(UL) issues this certificate to the Firm named above, after assessing the Firm's quality system and finding it in compliance with:

ISO 13485:2016

EN ISO 13485:2016

The manufacture of in vitro diagnostic blood grouping reagents. The purchase for resale of in vitro diagnostic serology test kit.





Authorized by



Michael J. Windler, P.E.

Manager of Global Regulatory Service

Distinguished Member of the Technical Staff

Life and Health Sciences, UL LLC

Check Certificate
Status: here

File Number A12241 Cycle Start May 23, 2020 Certificate Number 1458.200523 Effective Date May 23, 2020 Initial Issue Date June 26, 2018 Expiry Date May 22, 2023

This quality system registration is included in UL's Directory of Registered Firms and applies to the provision of goods and/or services as specified in the scope of registration from the address(es) shown above. By issuance of this certificate the firm represents that it will maintain its registration in accordance with the applicable requirements. This certificate is not transferable and remains the property of UL LLC.



UL LLC 333 Pfingsten Road Northbrook, IL 60062-2096 USA



LORNE LABORATORIES LTD

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Unit 1 Cutbush Park Industrial Estate Danehill, Lower Earley, Berkshire, RG6 4UT United Kingdom

> Phone: +44 (0) 118 921 2264 Fax: +44 (0) 118 986 4518 Email: info@lornelabs.com

CERTIFICATE OF ANALYSIS

DESCRIPTION	LOT NO.	EXPIRY	PRODUCT CODE
RPR Carbon Kit	LO16261	2023-03	044150A

STORAGE: Refrigerated at $2 - 8^{\circ}$ C. Protect from light. Do not freeze.

SHIPPING: This product has data supporting stability tolerance during fluctuations in

ambient shipping temperature.

This product is in compliance with **MEDICAL DEVICES REGULATIONS 2002** for Annex 3 and Self-Certification. This product was manufactured, packaged and tested in accordance with **LORNE QUALITY SYSTEMS** and meets all product specifications.

DEAGENT	SPECIFICATIONS			550111.7
REAGENT	Appearance	Colour	Functionality	RESULT
RPR Carbon Kit	Liquid suspension free of macroscopic particles	Grey	Tested against kit (+) and (-)	PASS
Positive Control Negative Control	Liquid solution	Clear and transparent	controls	PASS

If applicable:

- Components from human origin have been tested and found negative for the presence of antibody to HIV as well as HBsAg and HCV. Handle with caution as potentially infectious.
- This product was tested by methods described in the manufacturers package insert.

• This product is intended for *In Vitro* Diagnostic use only.

We certify that this product has been released as meeting our acceptance criteria

APPROVED BY:

DATE: 14 April 2021

Eddy Velthuis Technical Director

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