

# **CERTIFICATE**OF REGISTRATION

This is to certify that the management system of:

## **Chrono-log Corporation**

(FIN F000776)

Main Site: 2 West Park Road

Havertown, PA 19083 United States

has been registered by Intertek, an MDSAP recognized auditing organization, as conforming to the requirements of:

#### ISO 13485:2016

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6)

Brazil: RDC ANVISA n. 665/2022, RDC ANVISA n. 551/2021, RDC ANVISA n. 67/2009

Canada: Medical Devices Regulations - Part 1- SOR 98/282

United States: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 (Subparts A to D)

#### The management system is applicable to:

Design, manufacture, installation, service and distribution of platelet aggregometers, reagents for platelet aggregation, ristocetin cofactor assay kits and cuvettes.

**Certificate Number:** 

0085301-02

**Initial Certification Date:** 

2018-12-18

**Date of Certification Decision:** 

2024-12-03

**Certification Effective Date:** 

2024-12-17

**Certification Expiry Date:** 

2027-12-17



### intertek

Calin Moldovean

President, Business Assurance

Intertek Testing Services NA, Inc. 4700 Broadmoor SE, Suite 200 Kentwood, MI, USA, 49512



In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate validation@intertek.com or by scanning the code to the right with a smartphone. The certificate remains the property of Intertek, to whom it must be returned upon request. Validity of this certificate may be verified at



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