

CERTIFICATE OF REGISTRATION

This is to certify that the management system of:

Clearside Biomedical, Inc.

(FIN F007259)

Main Site: 900 North Point Parkway, Suite 200

Alpharetta, Georgia, 30005, 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)

Japan: MHLW Ministerial Ordinance 169, Article 4 to Article 68; PMD Act (as applicable)

The management system is applicable to:

The design, development, and manufacture of sterile piston syringes, needles, and associated accessories for the area of ophthalmology.

Certificate Number: 0186623

Revision Level: 00

Initial Certification Date: 2024-10-23

Certification Effective Date: 2024-10-23

Certification Expiry Date: 2027-10-22



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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 http://www.intertek.com/business-assurance/certificate-validation/

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