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

ISO 13485:2016 CERTIFICATION TM

MDSAP MEDICAL DEVICE SINGLE AUDIT PROGRAM

intertek

Calin Moldovean
President, Business Assurance

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