Certificate

Certificate No.:	MD 2583776-1-1 rev. 1
Manufacturer:	**Veracyte Inc.** 6000 Shoreline Cour, Suite 300. South San Franciso 94080 California USA
REPs Facility ID:	F007570
Certification criteria:	ISO 13485:2016 Australia Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure Canada Medical Devices Regulations – Part 1 – SOR 98/282 United States 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807
Scope:	The Design and development, manufacturing and distribution of in vitro diagnostic medical devices intended to aid in the diagnosis, prognosis, treatment options, disease follow-up and therapeutic efficacy in cancer and other lung diseases.

TUV Rheinland of North America, Inc., an MDSAP recognized Auditing Organization, certifies that the quality management system of the Manufacturer has been audited against and found to conform the Certification criteria for the Scope contained in this certificate. The quality management system is subject to annual surveillance audit(s).

Project No.:	73087732-100
Issue Date:	2025-03-20
Effective Date:	2025-03-20
Expiry Date:	2026-12-08

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

Certification officer: Daniel Świątko
TUV Rheinland of North America, Inc.

The validity of the certificate can be verified on https://www.certipedia.com or calling 1-888-743-4652.

TUV Rheinland of North America, Inc., 400 Beaver Brook Road, Boxborough, MA 01719, USA
Tel: (925) 249-9123, Fax: (925) 249-9124

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Manufacturer:

Veracyte Inc.

6000 Shoreline Cour, Suite 300.
South San Franciso
94080 California
USA

The scope of certification also covers the following sites:

No.	Location	Scope
/01	Veracyte Inc. 6000 Shoreline Cour, Suite 300. South San Franciso 94080 California USA	The design and development. REPs Facility ID: F007570
/02	Veracyte SAS Luminy Biotech Enterprises 163 Avenue de Luminy 13288 Cedex 9, Marseille France	The design and development, manufacturing and distribution. REPs Facility ID: F007571

MDSAP
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