

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Leica Biosystems Newcastle Ltd
Balliol Business Park West
Benton Lane
Newcastle upon Tyne
NE12 8EW
United Kingdom

Facility ID Number: F000261

Holds Certificate No:

MDSAP 696855

The company listed on this certificate has been audited to and found to conform with ISO 13485:2016 including the following country specific requirements:

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure

Brazil: RDC ANVISA n. 67/2009, RDC ANVISA n. 665/2022 - Good Manufacturing Practices, RDC ANVISA n. 551/2021

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

Japan: MHLW MO No 169 (2004), as amended by MHLW MO No 60 (2021), PMD Act

USA: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

The design, development and manufacture of in-vitro diagnostic reagents and their associated ancillaries, consumables, and test kits used in the diagnosis of cancer, disease status, endocrine disorders, genetic testing, protein metabolism, transmissible agents, and immunological typing.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2019-03-04

Effective Date: 2025-02-26

Expiry Date: 2028-02-25



BSI Group America Inc. is an MDSAP recognised auditing organization

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Location	Registered Activities
Leica Biosystems Newcastle Ltd Balliol Business Park West Benton Lane Newcastle upon Tyne NE12 8EW United Kingdom Facility ID Number: F000261	The design, development and manufacture of in-vitro diagnostic reagents and their associated ancillaries, consumables, and test kits used in the diagnosis of cancer, disease status, endocrine disorders, genetic testing, protein metabolism, transmissible agents, and immunological typing.
Leica Biosystems Newcastle Ltd 2 Gosforth Park Way Newcastle NE12 8ET United Kingdom Facility ID Number: F000261	The design and development of in-vitro diagnostic reagents, and their associated ancillaries, consumables, and test kits used in the diagnosis of cancer, disease status, endocrine disorders, genetic testing, protein metabolism, transmissible agents and immunological typing.



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This certificate remains the property of BSI and shall be returned immediately upon request.
An electronic certificate can be authenticated [online](https://www.bsigroup.com/ClientDirectory). Printed copies can be validated at www.bsigroup.com/ClientDirectory
To be read in conjunction with the scope above or the attached appendix.

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