

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016 & EN ISO 13485:2016

This is to certify that:

Thermo Fisher Scientific Oy
Ratastie 2
FI-01620 Vantaa
Finland

Holds Certificate Number:

MD 623249

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 & EN ISO 13485:2016 for the following scope:

The design and development, manufacturing, service, installation and distribution of in vitro diagnostic medical devices including clinical chemistry analysing systems, laboratory automation and in vitro diagnostic assays and reagents.

The design and development, manufacture, distribution and service of in vitro diagnostic medical instruments, liquid handling products and consumables.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2012-02-22

Latest Revision Date: 2024-03-13

Effective Date: 2024-04-01

Expiry Date: 2027-03-31



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Certificate No: **MD 623249**

Location

Thermo Fisher Scientific Oy
Ratastie 2
FI-01620 Vantaa
Finland

Registered Activities

The design and development, manufacturing, service, installation and distribution of in vitro diagnostic medical devices including clinical chemistry analysing systems, laboratory automation and in vitro diagnostic assays and reagents.
The design and development, manufacture, distribution and service of in vitro diagnostic medical instruments, liquid handling products and consumables.

Thermo Fisher Scientific Oy
Rahtikatu 2
FI-80100 Joensuu
Finland

The manufacture and service of in vitro diagnostic liquid handling products and consumable.



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This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract.
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Printed copies can be validated at www.bsigroup.com/ClientDirectory

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000
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