



CERTIFICATE

No. QS1 14 11 71067 003

Certificate Holder:

Liofilchem S.r.l.

Via Scozia

64026 Roseto degli Abruzzi (TE)

ITALY

Certification Mark:



Scope of Certificate:

Design and Development, Production and Sale of In-Vitro Diagnostic Medical Devices: Culture Media for Bacteriology, Identification and Susceptibility Testing Systems, Kits for Plasma Protein Determination. Distribution of other In Vitro Diagnostic Medical Devices

Standard(s):

ISO 13485:2003

The Certification Body of TÜV SÜD America Inc. certifies that the company mentioned above has established and is maintaining a quality management system that meets the requirements of the listed standards.

TÜV SÜD America Inc. is a Health Canada CMDCAS Recognized Registrar.

Report No.:

M5553

Effective Date:

2015-01-20

Expiry Date:

2018-01-19

Gary Minks

Vice President, Regulatory Affairs

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TÜV SÜD America Inc. 10 Centennial Drive Peabody, MA 01960 USA







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LIOFILCHEM S.r.I. Via Scozia 64026 Roseto degli Abruzzi (TE) ITALY

Production of In-Vitro Diagnostic Medical Devices: Culture Media for Bacteriology, Identification and Susceptibility Testing Systems, Kits for Plasma Protein Determination

LIOFILCHEM S.r.I. Contrada Piane Vomano, Traversa di Via Grecia 64026 Roseto degli Abruzzi (TE) ITALY

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