Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 18-1637-Q

TUV USA, Inc. hereby certifies that the quality management system of the company mentioned below is in conformance with ISO 13485:2016 for Medical Devices-Quality Management Systems-Requirements for Regulatory Purposes.

Rovers® Medical Devices

Rovers Medical Devices B.V.
Lekstraat 10, 5347 KV Oss,
The Netherlands

Additional sites covered by QM System: N/A

Scope:

Design, Manufacturing and Distribution of Sterile and Non-Sterile Cell Sampling Devices

The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.

TUV USA, Inc. (a Member of the TÜV NORD Group)
215 Main Street, Suite 1, Salem, NH 03079, USA
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: medical-usa@tuv-nord.com

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Bradley Chen
Vice President – Medical, Americas
Medical Products Division
TUV USA, Inc.