

# 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 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

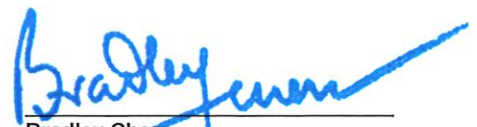
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: [medical-usa@tuv-nord.com](mailto:medical-usa@tuv-nord.com)



Audit Report Reference No.: **20-3848 RC-CA**  
Certificate Initial Issue Date: **2018-12-31**  
Current Cycle Start Date: **2020-10-23**  
Certificate Revised Date: **2020-10-23**

Effective Date:  
**2020-10-23 / ed. 4**

Valid Until:  
**2023-10-19**



Bradley Chen  
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Medical Products Division  
TUV USA, Inc.