Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 19-1610-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.

Stellartech Research Corporation
560 Cottonwood Drive
Milpitas, CA 95035, USA

Additional sites covered by QM System: N/A

Scope:

Design, Manufacture, Contract Manufacture, Service, and Distribution of Electromedical Devices and Accessories, Sterile Surgical Tools, Ablation and Other Catheters

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

Audit Report Reference No.: 21-3976 RC
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Effective Date:
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Valid Until:
2025-08-07

Bradley Chen
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TUV USA, Inc.