

# 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)

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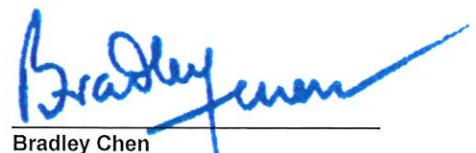
Audit Report Reference No.: **21-3976 RC**

Initial Certification Date: **2019-08-08**

Current Cycle Start Date: **2022-08-08**

Effective Date:  
**2022-08-08 / ed. 6**

Valid Until:  
**2025-08-07**



Bradley Chen  
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Medical Products Division  
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