

EU Quality Assurance Certificate

Regulation (EU) 2017/745, Annex XI Part A

MDR 720081 R001

Manufacturer: STIMULI TECHNOLOGY

Address:

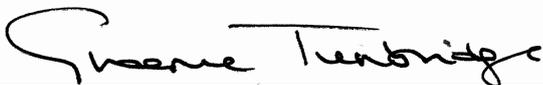
20 bis rue Barthélemy Danjou
Boulogne Billancourt
92100
France

Single Registration Number: FR-MF-000002429

Scope: See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex XI part A, the quality system meets the requirements of the Regulation. For the placing on the market of Class III and Class IIb devices an additional Annex X certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):



Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

First Issue Date: **2021-02-02**

Current Issue Date: **2025-12-04**

Starting Validity Date: **2026-02-02**

Expiry Date: **2031-02-01**

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.
This certificate was issued electronically and is bound by the conditions of the contract.

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Device Schedule: Class IIa, Custom-made and other devices

Device(s)	Risk Classification
TENSI+ (Neuromuscular electro stimulator)	Class IIa



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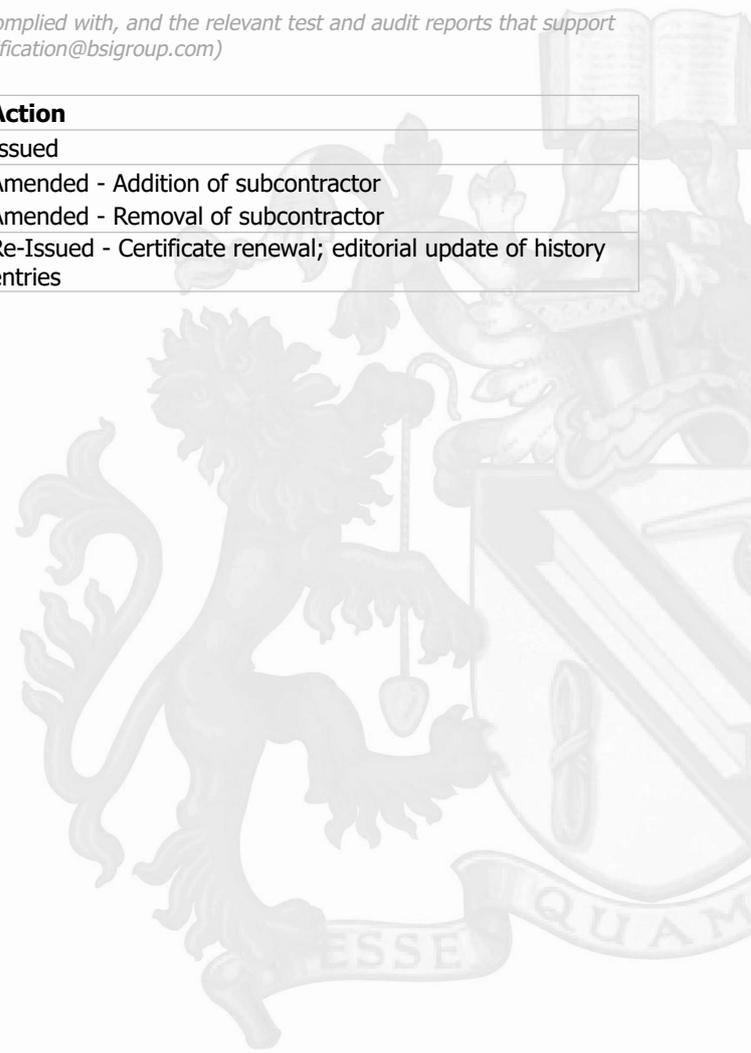
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Certificate History

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)

Date	Reference number	Action
2021-02-02	3095636	Issued
2022-06-01	3643086	Amended - Addition of subcontractor Amended - Removal of subcontractor
Current	30444722	Re-Issued - Certificate renewal; editorial update of history entries



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