Certificate

EC-Certificate

(Production quality assurance system)
according to Annex V of Medical Devices Directive 93/42/EEC

It is herewith confirmed by

BSI Group Deutschland GmbH
Eastgate, Hanauer Landstrasse 115
60314 Frankfurt am Main
Germany

in its function as Notified Body (0535), that the manufacturer:

Neurosoft Ltd.
5, Voronin Str.,
Ivanovo, 153032, Russia

concerning the medical device

portable device for EMG/STIM-guided injections "Neuro-Tox"
UMDNS 16-263, class IIa

fulfils the requirements according to Annex V of the Medical Devices Directive 93/42/EEC. The manufacturer has established a quality assurance system for the production and final inspection of the specified devices.
For the placing on the market of class III products an Annex III certificate is required.

Report No.: SMO7864399
Certificate No.: CE 577334

Notified by
Zentralstelle der Länder für Sicherheitstechnik
ZLS-NB-67/12

First Issue Date:
November 08, 2009.

Based on periodical surveillance this certificate is valid until November 18, 2018.

Current Issue Date: November 19, 2013

Certification Body

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