EC Certificate Full Quality Assurance System: Certificate ES19/86843

The management system of

Xavant Technology (Pty) Ltd

Unit 102 Tannery Industrial Park, 309 Derdepoort Road,
Silverton, Pretoria, 0184, South Africa

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Neuromuscular stimulators for peripheral nerve location. Neuromuscular stimulators for neuromuscular blocking agent monitoring using acceleromyography and/or electromyography. Neuromuscular stimulators for use in somatic and neuropathic pain management

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 11 November 2020 until 26 February 2024
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 11 October 2005
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered ES/MAD 212638

Authorised by

SGS Belgium NV, Notified Body 1639
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