

DECLARATION OF CONFORMITY

Medical Device(s)

Manufacturer: **Viamed Ltd.**
15 Station Road
Cross Hills
Keighley
West Yorkshire BD20 7DT
United Kingdom

Device(s): Supramaximal Nerve Stimulator

Class: IIa

Part number: 2510000

Description: Supramaximal Nerve Stimulator

It is hereby declared that the medical device(s) specified above is (are) in compliance with EC Council Directive 93/42/Eec (Medical Devices Directive) of the 14th June 1993 and considering the amendments by directive 2007/47/EC - as transposed into National Law in the countries where the product is intended to be marketed.

This declaration is supported by:

Technical documentation required by MDD (Annex II) retained by: Viamed Ltd.

EC Quality Assurance Certificate No. CE 01389, first issued by the British Standards Institute (CE2797) on the 23rd August 1996.

Certificate of Registration No. MD 78787 to ISO 13485:2016 of original registration date 26th February 2016, issued by the British Standards Institute (CE0086)

In accordance with 2011/65/EU RoHS Directive

For and on behalf of Viamed Ltd.

Signature:



Name:

Derek Lamb - Managing Director

Date:

07 / 03 / 2019