



Declaration of Conformity.

Class IIa Medical Device(s).

Manufacturer: Viamed Limited.

15 Station Road,
Cross Hills,
Keighley,
West Yorkshire,
BD20 7DT.
United Kingdom.

Device(s): 2510000
Microstim DB III

Description of Device(s): 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 Device Directive) of 14th June 1993 considering the amendments by the 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 the MDD (Annex II) retained by: Viamed Limited.
- EC Quality Assurance Certificate No. CE 01389, first issued by the British Standards Institute (CE0086) on the 23rd August 1996.
- Certificate of Registration No. FS28344 to BS EN ISO 9001:2008 of original registration date 15th June 1994, issued by the British Standards Institute (CE0086).

Signed: 

Date: 30 / 03 / 2011

Position: Managing Director.