VIAMED

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):	Microstim DBS_Mark 3
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 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 9001:2000 of original registration date 15th June 1994, issued by the British Standards Institute (CE0086).

Signed: ……..…… **Date**: ……/…../……

Position: Managing Director.