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

Signed : ..... **Date :** 01/05/03.

Position : Managing Director.