



VIAMED Ltd.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

Microstim DB III

Supramaximal Nerve Stimulator

Product No(s).

2510000

conforms with the essential requirements of Annex I of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices considering the amendments by the directive 2007/47/EC.

In accordance with Annex IX of the Directive 93/42 EEC considering the amendments by the directive 2007/47/EC the product has been classified as

Class IIa

Application of the CE-Marking:

CE0086

Issuer:

Viamed Ltd.

15 Station Road

Cross Hills

Keighley

West Yorkshire, BD20 7DT

United Kingdom

Place, Date:

Keighley , 5 Nov 2010.

Legally binding signature:

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Derek Lamb (Managing Director)