



VIAMED Ltd.

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EC Declaration of Conformity

We hereby declare under sole responsibility that the product

SpO2 Silicone Finger Sensor - Adult. SC4013 GE DATEX Compatible

For use with pulse oximeters in order to determine saturation of haemoglobin non-invasively from light signals of two wavelengths transmitted through tissue or reflected from tissues

Product No(s).

0014725

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 IIb

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)