EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM-2160

hand held pulse oximeter for continuous and spot-check monitoring of functional arterial oxygen saturation (SpO_2) and pulse rate,

Product No.
0012160

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:

CE 0086

Issuer:

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

Place, Date: Keighley, 20 May 2010

Legally binding signature: ……

Derek Lamb (Managing Director)