EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM-2101

Finger Oximeter for monitoring of functional
arterial oxygen saturation (SpO_2) and pulse rate,

Product No.
0012101

conforms with the essential requirements of Annex II of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices.

In accordance with Annex IX of the Directive 93/42 EEC the product has been classified as Class IIa

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, 4 August 2008

Legally binding signature: .....

Steve Nixon (Director)