EC Declaration of Conformity

We hereby declare under our sole responsibility
that the product

VM 2160

handheld 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.

In accordance with Annex IX of the Directive 93/42/EEC 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, 7th November 2007

Legally binding signature: .....

John S Lamb (Managing Director)