EC Declaration of Conformity

We hereby declare under sole responsibility that the product

VM 2500-M and VM 2500-S

hand held capnograph and pulse oximeter for continuous and spot-check monitoring of end-tidal CO_2 concentration (EtCO₂), inspired CO_2 concentration (FiCO₂), functional arterial oxygen saturation (SpO₂) respiration rate and pulse rate,

Product No. **4410500**, **4410501**, **4410520** and **4410521**

conform 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 IIb

Application of the CE-marking:

C€ 0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
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United Kingdom

Place, Date:

Keighley, 4 August 2009

Legally binding signature:

Derek Lamb (Managing Director)