

# 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<sub>2</sub>) 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

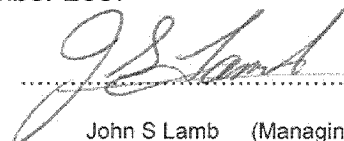
**Issuer:**

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)