

## **EC Declaration of Conformity**

We hereby declare under sole responsibility that the product

**VM-2103**

Finger Oximeter for monitoring of functional  
arterial oxygen saturation (SpO<sub>2</sub>) and pulse rate,

Product No.  
**0012103**

conforms with the essential requirements of Annex II of the Council Directive 93/42/EEC  
of 14 June 1993 considering the amendments by the directive 2007/47/EC as  
transposed into national law in the countries where the medical product is intended to be  
marketed.

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, 26 April 2012

**Legally binding signature:**

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**Derek Lamb** (Managing Director)

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