

EC Declaration of Conformity

We hereby declare under sole responsibility that the

4000 Series SpO₂ Sensors

reusable and disposable pulse oximetry sensors for continuous and spot-check monitoring of functional arterial oxygen saturation (SpO₂) and pulse rate, 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:

CE 0086

Issuer:

Viamed Ltd.
15 Station Road
Cross Hills
Keighley
West Yorkshire, BD20 7DT
United Kingdom

Place, Date:

Keighley, 29 October 2009

Legally binding signature:



Derek Lamb (Managing Director)