

DECLARATION OF CONFORMITY

Medical Device(s)

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

Device(s): 4000 Series SpO2 Sensors,
Pulse Oximeter Extension Cable,

Class: IIb

Part number: Page 2

Description: Multiple Items

It is hereby declared that the medical device(s) specified above is (are) in compliance with EC Council Directive 93/42/Eec (Medical Devices Directive) of the 14th June 1993 and considering the amendments by directive 2007/47/EC - as transposed into National Law in the countries where the product is intended to be marketed.

This declaration is supported by:

Technical documentation required by MDD (Annex II)

EC Quality Assurance Certificate No. CE 540537, first issued by the British Standards Institute (CE0086) on the 20th Aug 2008.

Certificate of Registration No. MD 78787 to ISO 13485:2003 of original registration date 27th January 2004, issued by the British Standards Institute (CE0086)

Certificate of Registration No. FS 28344 to ISO 9001:2008 of original registration date 15th June 1994, issued by the British Standards Institute (CE0086)

In accordance with 2011/65/EU RoHS Directive

For and on behalf of Viamed Ltd. _____

Signature: 

Name: **Derek Lamb - Managing Director**

Date: 21 / 04 / 2015



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Part number: 0014601 0014602 0014605 0014606 0014607 0014608 0014609 0014610
0014612 0014613 0014614 0014615 0014617 0014618 0014619 0014620
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