

DECLARATION OF CONFORMITY

Medical Device(s)

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

Device(s): Oxygen Tents,

Class: IIa

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) retained by: Viamed Ltd.

EC Quality Assurance Certificate No. CE 01389, first issued by the British Standards Institute (CE0086) on the 23rd August 1996.

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: 29 / 07 / 2013

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Part number: 2310000 2310001 2310002 2310003 2310004 2310005 2310006 2310007
2310008 2310009 2310010 2310011 2310012 2310013 2310014 2310020
2310021 2310025 2310026 2310027 2310028 2310029 2310030 2310031
2310032 2310033 2310034 2310035 2310050 2310051 2310052 2310053
2310054 2310055 2310075 2310076 2310077 2310078 2310079 2310080
2310081 2310100 2310101 2310102 2310103 2310104 2310105 2310106
2310107 2310125 2310126 2310127 2310128 2310129 2310130 2310131
2310132 2310133 2310134 2310140 2310141 2310145 2310150 2310151
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Signature: 

Name: **Derek Lamb - Managing Director**

Date: 29 / 07 / 2013