

DECLARATION OF CONFORMITY

Medical Device(s)

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

Device(s): Oxygen Hoods,

Class: IIa

Part number: See page 2

Description: Enclosures to enable oxygen enriched air and humidity to be supplied to the patients head only

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: 23 / 02 / 2016



VIAMED

Viamed Limited - 15 Station Road - Cross Hills
Keighley - West Yorkshire BD20 7DT - United Kingdom
Tel: +44 (0)1535 634542 Fax: +44 (0)1535 635582
Email: info@viamed.co.uk Website: www.viamed.co.uk

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Part number: PP8000 PP8020 PP8050 PP8055 PP8060 PP8065 PP8509 PP8533
PP8540 PP8562 PP8565 PP8589 PP8687 PP8715 PP8729 PP8730
PP8744 PP8746 PP8747 PP8748 PP8752 PP8753 PP8754 PP8755
PP8756 PP8757 PP8758 PP8759 PP8760 PP8771 PP8772 PP8773
PP8774 PP8775 PP8780 PP8781 PP8782 PP8783 PP8784 PP8785

For and on behalf of Viamed Ltd. _____

Signature: 

Name: **Derek Lamb - Managing Director**

Date: 23 / 02 / 2016