

## DECLARATION OF CONFORMITY

### Medical Device(s)

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

Device(s): Sampling Lines

Class: IIa

Part number: See page 2

Description: Sidestream CO2 gas sampling lines

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 14<sup>th</sup> 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

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: 09 Feb 2017



# 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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Class: IIa

Part number:  
4420718 CO2 Nasal Sampling Cannula with O2  
4420738 CO2 Nasal/Oral Sampling Cannula with O2  
4420739 CO2 Nasal/Oral Sampling Cannula with O2  
4420719 CO2 Nasal Sampling Cannula with O2

For and on behalf of Viamed Ltd.

Signature:

Name: **Derek Lamb - Managing Director**

Date: 09 Feb 2017