

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: Gas Sampling Lines for use with sidestream gas exchange monitoring devices

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: 06 / 04 / 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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4420819 4420820 4420854 4420855 4420856 4420860 4420861 4420862
4420865 4420866 4420869 4420870 4420950 4420951 4420952 4420953
4420954 4420955 4420956 4420957 4420958 4420959 4420960 4420961
4420962 4420963 4420964 4420965 4420966 4420967 4420968 4420969
4420970 4420971 4420972 4420973 4420974 4420975 4420976

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