



Declaration of Conformity.

Class IIa Medical Device(s).

Manufacturer: Viamed Limited.

15 Station Road,
Cross Hills,
Keighley,
West Yorkshire,
BD20 7DT.
United Kingdom.

Device(s): Oxygen Sensor R-23V

Description of Device(s): A medical device for monitoring oxygen content in anaesthetic and intensive respiration devices and in incubators Compatible to Draegar 6850645 & 6809777

It is hereby declared that the medical device(s) specified above is (are) in compliance with EC Council Directive 93/42/EEC (Medical Device Directive) of 14th June 1993 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 the MDD (Annex II) retained by: Viamed Limited.
- EC Quality Assurance Certificate No. CE 01389, first issued by the British Standards Institute (CE0086) on the 23rd August 1996.
- Certificate of Registration No. FS28344 to BS EN ISO 9001:2000 of original registration date 15th June 1994, issued by the British Standards Institute (CE0086).

Signed:

Date: 01/06./05.....

Position: Managing Director.