VIAMED

Declaration of Conformity.

Class I Medical Device(s).

Manufacturer:	Viamed Limited. 15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT. United Kingdom.
Device(s):	SpO2 Probes – pulse oximetry extension cables. 0019650 Critikon & Sensormedics P965E10
Description of Device(s):	For use with pulse oximeter sensors in order to extend the reach of the sensor to the patient.

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**: ……/……/……

Position: Managing Director.