## Risk Analysis Report

Date 18 February 1998

Product	
Product Type SpO <sub>2</sub> Probes	MDD Class III
Model Type Various	

**Product Description** 

Pulse Oximeter Compatible Finger Probes

Manufacturer Address

Viamed Ltd.,

15 Station Road,

Crosshills

Keighley,

'Vest Yorkshire BD20 7DT, UK

(In accordance with pr EN 1441:1997)

We hereby declare that the statements made herein are correct and valid

Company: Viamed Ltd. 15, Station Road, Crosshills, Keighley, West Yorkshire, BD20 7DT. UK

Signature

Name J.S.Lamb

Director

Date 18/2/28

Level of Risk Product: CEMARK\ SPOPROBE\RARSPO 18 February 1998

Insignificant 4: Tolerable 3: Significant 2: Catastrophic 1 Page 1

## IR= Initial Risk RR = Residual Risk

Ref		Part of Equipment which pose risks	I R	Design solution Adopted	R R	Document referenced
C	ENERGY					
<u></u> C2.1	Electricity	Instrument & Sensor failure	3		3	
C2.2		Instrument & Sensor failure	3		3	
C2.3	Mechanical Force		4		4	
C2.4	Iononizing Radiation		4	·	4	
C2.5	Non Ionising Radiation		4		4	
C2.6	Electromagnetic Fields		4		4	
C2.7	Moving Parts		4		4	
C2.8	Suspended Masses		4		4	
C2.9	Patient Support Failure		4		4	
C2.10	Pressure (Vessel rupture)		4		4	
C'`1	Acoustic pressure		4		4	
C2.12	Vibration		4		4	
C2.13	Magnetic Fields (MRI)		4		4	
C3	BIOLOGICAL		4			
C3.1	Bio-Burden	·	4			
C3.2	Bio-Contamination		4		4	H
C3.3	Bio-Incompatibility		4		4	
C3.4	Incorrect Output (Substance/energy)		4	<b>,</b>	4	
C3.5	Incorrect Formulation (Chemical Composition)		4		4	
C3.6	Toxicity		4	l .	4	l l
C3.7	Cross Infection	User error	4	l .	4	l I
$c_{\smile}$	Pyrogenicity		4	1	4	1
C3.9	Inability to Maintain Hygienic Standards		4			•
C3.10	Degradation		4	l		
C4.	ENVIRONMENTAL		4	1	4	1
C4.1	Electromagnetic Interference		4			1
C4.2	Inadequate supply of Power or Coolant		4	1	4	
C4.3	Likelihood of Operation outside Prescribed Environmental Conditions		4		4	1

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Ref.	Hazard	Part of Equipment which pose risks	I R	Design solution Adopted	R R	Document referenced
C	Incompatibility with other Devices		4		4	
C4.5	Accidental Mechanical Damage		4		4	
C4.6	Contamination due to Waste Products and or Device Disposal		4		4	
C5	DEVICE USE		4		4	
C5.1	Inadequate Labelling		4		4	
C5.2	Inadequate Operating Instructions		4		4	
C5.3	Inadequate Specification of Accessories		4		4	
C	Inadequate Specification of Pre-Use Checks	·	4		4	
C5.5	Over-complicated Operating Instructions		4		4	
C5.6	Unavailable or Separated Operating Instructions		4		4	
C5.7	Use by Unskilled /untrained Personnel		4		4	
C5.8	Reasonable Foreseeable Abuse		4		4	
C5.9	Insufficient Warning of Side Effects		4		4	
<u>C</u> 0	Inadequate Warnings of Hazards Likely With Re-use of Single Use Devices		4		4	
C5.11	Incorrect Measurement and other Metrological Aspects	Technique limited	3		3	
C5.12	Incorrect Diagnosis		4		4	
C5.13	Erroneous Data Transfer	-	4		4	
C5.14	Misrepresentation of Results		4		4	
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IR= Initial Risk RR = Residual Risk

Ref.	Hazard	Part of Equipment which pose risks	4	Design solution Adopted	4	Document referenced
C 5	Incompatibility with Consumables /accessories / other Devices	Manufacturers use similar connectors	3	Colour spot identification Labelling	4	_
C6	FUNCTIONAL FAILURE MAINTENANCE and AGEING		4		4	
C6.1	Inadequacy of Performance Characteristics for the Intended Use		4		4	
C6.2	Lack of ,or Inadequate Specification for Maintenance including Post Maintenance Functional Tests		4		4	
C6.3	Inadequate maintenance		4		4	
C6.4	Lack of Adequate Determination of End of Device Life	Will fail to function	4		4	
C6.5	Loss of Mechanical Integrity	Will be visible to the user	4		4	
C6.6	Inadequate Packaging ( contaminationand/or Deterioration of the Device)		4		4	
C6.7	Improper Use		4	:	4	