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Wismar, 02.04.2014

Viamed Ltd. 15 Station Road Cross Hills Keighley

## West Yorkshire BD20 7DT United Kingdom

## Mechanical strength tests (drop tests)

Product identification: Pulse oximeter with finger sensor Type/Model/Parameters: VM 2160 with silicone sensor

P/N: 0012160, 0012161, 0012162, 0012163, 0012164

The pulse oximeter equipment VM 2160 with the associated silicone sensors as listed above are tested according IEC 60601-1 and ISO 9919. These are the product standards for pulse oximeter equipment and include mechanical strength requirements as a sub-part of the electrical safety. Please be advised that the electrical safety is not restricted to electrical. It includes thermal requirements, fire hazards, mechanical requirements and all other functional safety requirements related to the safety of the equipment to be tested. The term electrical means other products than electrical equipment will not be certified.

The standard IEC 60601-1:1988+A1:1991+A2:1995 covers in clause 21 mechanical strength requirements. The pulse oximeter was tested according sub-clause 21.5 as a handheld equipment dropping from 1 m high onto hardwood surface, see test report 408.123.2A. With the silicone cover, the equipment including probe remains safe, no mechanical degradation and maintains full functionality as required. In addition ISO 9919:2005 requires additional tests regarding mechanical strength, see the attached excerpt of the test report 408.123.3.

In addition the equipment was tested as with mechanical shocks of 102 g and at least 30 min with mechanical vibration as described in b) of the test report excerpt.

CEcert is a independent test house accredited according national and European law, see link on homepage.

Bernd Schmidt

Head of Test Laboratory

Attachment: Excerpt of Test Report 408.123.3

HRB 6695

KTO 356 138800



Excerpt of Test Report 408.123.3

ISO 9919						
Clause	Requirement + Test	Result - Remark	Verdict			

	1		Р
21.102	Shock and vibration for transport		
	Pulse oximeter equipment or its parts, intended for use during patient transport outside a healthcare facility, shall have adequate mechanical strength when subjected to mechanical stress caused by normal use, pushing, impact, dropping, and rough handling.	see attachment # 7	Р
	After the following tests, pulse oximeter equipment shall not cause a safety hazard and shall function normally.	functional test with functional test equipment, see attachment # 2	Р
	a) Shock test in accordance with IEC 60068-2-27 (102g/ 6ms/ half sine, 3 shocks per direction per axis)	see attachment # 7	Р
	b) Broad-band random vibration test in accordance with IEC 60068-2-64 (30 min per perpendicular axis - 10 Hz to 100 Hz: 5,0 (m/s2)2/Hz;	see attachment # 7	Р
	- 100 Hz to 200 Hz: -7 db per octave;		
	- 200 Hz to 2 000 Hz: 1,0 (m/s2)2/Hz)		
	c) For mobile pulse oximeter equipment, free fall to IEC 60068-2-32, using Procedure 1 (height: 0,1 m; number of falls: one; direction: vertical, (normal operating position)	equipment tested in accordance to sub-clause d) and in accordance to general standard (1 m height free fall)	Р
	d) For portable pulse oximeter equipment, free fall to IEC 60068-2-32, using Procedure 2 (height: 0,25 m; number of falls: one; direction: on each of the six surfaces.)	see attachment # 7	Р
	For portable pulse oximeter equipment that is intended to be used with a carrying case, that case may be applied to the equipment during this test.		Р
	Equipment tested and complying with the requirements in 21.102 in total or part, is considered to comply with the corresponding requirements of 21.101.		Р

Possible	test	Case	verdicts:
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- test case does not apply to the test object...... N/A

- test object does meet the requirement...... P (Pass)

- test object does not meet the requirement...... F (Fail)

Testing:

Date of receipt of test item ...... 2008-05-05