

EC Declaration of Conformity

We hereby declare under our sole responsibility
that the product

VM-2160 with accessories

Handheld Pulse Oximeter with SMARTsat® technology for continuous or spot check monitoring of functional arterial oxygen saturation (SpO₂) and pulse rate (PR) .

is in conformity with the essential requirements of Annex I and Annex II of the Council Directive 93/42/EEC as amended by 2007/47/EC concerning medical devices.

In accordance with Annex IX of the Council Directive 93/42/EEC the products have been classified as Class IIb.

Application of the CE-marking:



MEDCERT GmbH, Pilatuspool 2, 20355 Hamburg

Validity:

Date of expiry: 31 December 2019

Issuer:

bluepoint medical GmbH & Co. KG
An der Trave 15
D-23923 Selmsdorf
Germany

Place, Date:

Selmsdorf, 22 January 2018

Legally binding signature:

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Bernd Lindner
General Manager