EC Declaration of Conformity

We hereby declare under our sole responsibility
that the product

VM2160

Handheld pulse oximeter for continuous and spot check monitoring of functional arterial oxygen saturation (SPO2) and pulse rate

Conforms with the essential requirements of Annex I and Annex II of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices.

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

Application of the CE-marking:

CE 0482 MEDCERT GmbH, Pilatuspool 2, 20355 Hamburg

Validity: **Date of expiry: 31 December 2018**

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

Place, Date: Selmsdorf, 22 December 2017

Legally binding signature: .....

Bernd Lindner
(Managing Director)