ENVITEC
by Honeywell

EC Declaration of Conformity

This declaration of conformity is issued under the sole responsibility of the manufacturer in accordance with the Council Directive 93/42/EEC in its currently valid version (Annex I) concerning medical devices.

Product:	**Pulse Oximeter**
Type:	**MySign®S** **Inkl. Zubehör**
Classification: (RL 93/42/EEC, Annex IX)	**Class IIb**
CE mark:	CE 0123
Notified Body:	TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 Munich, Germany
Conformity Assessment Process:	Annex II, section 3 of the Directive 93/42/EEC
Particular product standard applied:	ISO 80601-2-61
Issued by:	**ENVITEC-WISMAR GMBH** **Alter Holzhafen 18** **D-23966 Wismar** **Germany**
Valid from: Valid until:	**2016-10-26** **2019-10-25**
Place, Date:	Wismar, 2016-10-28
Authorized Signature:	Marcus Ostländer Managing Director