

## EU - Conformity Statement

We declare with sole responsibility that following product comply with the basic requirements according the Council Directive 93/42/EEC in the current valid version (Annex I) concerning medical devices.

Product: **Disposable Pulse Oximetry Sensors**

Typ: DA-2211-1 MySign®  
DP-2211-2 MySign®

Classification: **Class IIb**  
(RL 93/42/EEC, Annex IX)

CE marking:



Notified Body: TÜV SÜD Product Service GmbH, Munich, Germany

Conformity Assessment Process: Annex II, section 3 of the Directive 93/42/EEC

Issued by: **ENVITEC-WISMAR GMBH**  
**Alter Holzhafen 18**  
**D-23966 Wismar**  
**Germany**

Valid from: **2013-04-30**

Place, Date: **Wismar, 2013-04-30**

Authorized Signature:

Marcus Ostländer  
Managing Director