

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: **Oxygen Monitor**

Type: **MySign® O**
incl. Accessories

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

CE mark: 

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 80601-2-55

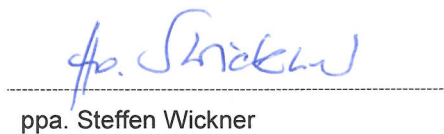
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Valid until: 2022-06-30

Place, Date: Wismar, 2019-05-29

Authorized Signature:


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