

# 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 Sensor**

Typ:

|          |        |
|----------|--------|
| EnviteC  | Viamed |
| OOM102-A | R-22VA |

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

**Class IIa**

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

Issued by:

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

Valid from:

2019-06-07

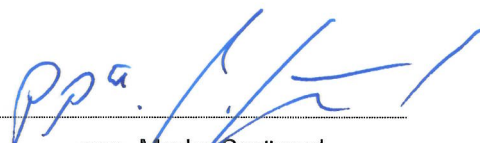
Valid until:

2021-06-06

Place, Date:

Wismar, 2019-06-07

Authorized Signature:

  
ppa. Marko Sprössel

  
ppa. Steffen Wickner