Honeywell
Healthcare Solutions GmbH

# EU Declaration of Conformity

This declaration of conformity is issued under the sole responsibility of the manufacturer in accordance with Regulation 2017/745/EU Article 120 and Directive 93/42/EEC in its currently valid version.

Product: **Oxygen Sensors**

Basic UDI-DI: 4036616013680

Type:

| Honeywell | Viamed | Honeywell | Viamed | Honeywell | Viamed |
|---|---|---|---|---|---|
| OOM101 | R15-V | OOM105 | T-7-V | OOM201 | R-23-V |
| OOM102 | R-22Vi | OOM106 | R-45-V | OOM202 | R-49-V |
| OOM102-A | R-22VA | OOM110 | R-30-V | OOM202-1 | R-41-V |
| OOM102-1 | R-14-Vi | OOM111 | R-45-V | OOM202-2 | R-44-V |
| OOM103 | R-22MEDV | OOM112 | R-43-V | OOM202-2S | R-42-V |
| OOM103-1 | R-17-V | OOM113 | R-48-V | OOM204 | R-47-V |
| OOM104 | R-23-Vi | | | | |

Classification: (Annex VIII) **Class IIa**

CE Mark: CE 0123

Notified Body: TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany

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

Particular product standard applied: ISO 80601-2-55

Issued by:

**Honeywell Healthcare Solutions GmbH**
**Eudamed-SRN: DE-MF-000004924**
**Alter Holzhafen 18**
**D-23966 Wismar**
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