

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: **Class IIa**
(Annex VIII)

CE Mark: 

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

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

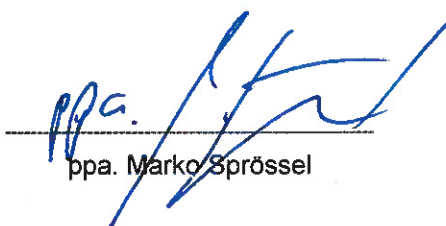
Particular product standard applied: ISO 80601-2-55

Issued by: **Honeywell Healthcare Solutions GmbH**
Eudamed-SRN: DE-MF-000004924
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Germany

Valid from: 20.06.2022
Valid until: 26.05.2024

Place, Date: Wismar, 20.06.2022

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