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 in its currently valid version. Article 120 (3a) case b) applies to the products. The requirements from Article 120 (3c) are met.

Product:	**Oxygen Sensors**		
Basic UDI-DI:	4036616OOMCK		
Type:	OOMLF102 OOMLF102-1 OOMLF102-L OOMLF102-1L OOMLF102-M3	OOMLF103 OOMLF103-1 OOMLF103-1M OOMLF106 OOMLF110	OOMLF111 OOMLF202 OOMLF202-1 OOMLF204
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** **Germany**		
Valid from:	04.03.2024		
Valid until:	31.12.2028		
Place, Date:	Wismar, 04.03.2024		

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