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:	**Flow Sensor**
Basic UDI-DI:	4036616FlowP8L
Type:	SpiroQuant H
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:	
Issued by:	**Honeywell Healthcare Solutions GmbH** **Eudamed-SRN: DE-MF-000004924** **Alter Holzhafen 18** **D-23966 Wismar** **Germany**
Valid from:	13.03.2024
Valid until:	31.12.2028
Place, Date:	Wismar, 13.03.2024

Authorized Signature:

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