

## 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: **Flow Diverter**

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

**Class I**

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

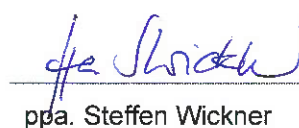
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**Alter Holzhafen 18**  
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**Germany**

Valid from: 2020-11-19  
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Place, Date: Wismar, 2020-11-19

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