Declaration of Conformity acc. to Annex IV of the Regulation 2017/746 on IVD medical devices (IVDR)

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We, company Dr. Müller Gerätebau GmbH, declare that the issuing of this Declaration of Conformity is our sole responsibility.

Product name	REF. No.	Basic UDI-DI	Intended use	Risk class
SensoStar GLH six	920800	42555893500AA2000RB	electrochemical analyser, near-patient testing	A

We hereby declare that listed product complies with all applicable requirements of EU Regulation 2017/746 on IVD medical devices (IVDR). The above-mentioned product is an in vitro diagnostic medical device according to article 2(2) of the IVDR.
It fulfils the basic safety and performance requirements according to Annex I of the IVDR.
Conformity has been established by conformity assessment procedure after drawing up the technical documentation set out in Annexes II and III of the IVDR.
This Declaration of Conformity is valid from serial number 1661.

Freital, September 20, 2022

Matthias Hartwig
Managing Director / CEO
Dr. Müller Gerätebau GmbH