



Declaration of Conformity
Konformitätserklärung

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Produktname: RapidStain

We hereby declare that the *in vitro*
diagnostic devices

Hiermit erklären wir, dass das *In vitro*
Diagnostikum

Product / Produkt: R-SemenStain

Code-No. / Kat.-Nr. 11050/ 11250

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| <p>(i) is classified as a "all other IVD Medical Devices" according to Annex III of the IVDD</p> <p>(ii) conform to the relevant provisions of the EC Council directives 98/79/EC and</p> <p>(iii) is in accordance with the Annex III of the IVDD</p> | <p>(1) Gemäß Anhang III der EG-Richtlinie 98/79/EG als „Sonstiges <i>In vitro</i>-Diagnostikum klassifiziert ist.</p> <p>(2) Allen übrigen relevanten Verpflichtungen der EG-Richtlinie 98/79/EG genügt und</p> <p>(3) Den Verpflichtungen des Anhang III der EG-Richtlinie nachkommt.</p> |
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Harmonized standard applied

Angewandte harmonisierte Normen

EN 13612:2002 EN ISO 13485:2012 + AC:2012 EN 13612:2002

EN ISO 14971:2012 EN ISO 15193:2009 ISO 15223-1:2012

EN ISO 18113-1:2011 EN ISO 18113-2: 2011 EN ISO 19001:2013 EN ISO 23640:2013

Geschäftsführer: Max Müser

Schwelm den 13.12.2019