Declaration of Conformity

Gerätebau GmbH

Dr. Müller Gerätebau GmbH
Burgker Strasse 133
DE-01705 Freital

Herewith we declare on our own responsibility that the medical device meets all applicable requirements of the Directive 98/79/EC and applied standards and guidelines.

We do not guarantee the fulfilment of these norms and guidelines after unauthorized modification of the product.

Name of the product:

CRP ID

Applied norms / guidelines

DIN EN 13485	Medical devices – Quality management systems – Requirements for regulatory purposes
98/79/EG	In vitro diagnostic medical devices

The CE mark was fixed to the product.

Freital, 30.04.2015

Matthias Hartwig
General Manager