

Declaration of Conformity



Dr. Müller Gerätebau GmbH
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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:

Glucose ID, Cartridge Glucose for SUPER ID clinchem

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, 20.03.2013



Ralf Günther
General Manager