HEINZ HERENZ HAMBURG

Export and Production
– since 1949 –

DIN EN ISO 13485

Heinz Herenz Medizinalbedarf GmbH · Rudorffweg 10 · 21031 Hamburg

EC Declaration of Conformity

according to annex V of the Council Directive 93/42/EEC concerning medical devices amended by Directive 2007/47/EC

We Heinz Herenz
Medizinalbedarf GmbH
Rudorffweg 10
21031 Hamburg / Germany

declare under our sole responsibility that following products

Gynobrush®, sterile, item no. 1032930	class I, sterile
Gynobrush® Plus, with soft plastic tip, sterile, item no. 1032934	class I, sterile

fulfill the requirements of the Council Directive 93/42/EEC concerning medical devices.

This EC-Declaration of Conformity is valid till 27th April 2021 and for all products mentioned above which are handled within this period of time. In case of a change of the product this EC-Declaration of Conformity gets unvalid.

Applied standards or other normative documents:
see current listing in the technical documentation

Notified Body:
Medcert GmbH
Pilatuspool 2
20355 Hamburg, Germany

CE-Number:

CE
0482

Hamburg, 20.05.2016

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Dieter Cyll
(Managing Director)

rev. 20.05.2016