HEINE

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ID-Nr. M-0001/ 02

EC-Declaration of Conformity for medical devices
EG-Konformitätserklärung für Medizinprodukte

according to annex VII in combination with annex V of Council Directive 93/42/EEC

We hereby declare that the medical device

HEINE GAMMA Sphygmomanometer

(UMDNS-Code: 16-156)

Non-invasive manual aneroid blood pressure meter and accessory

Nicht-invasives manuelles Aneroid Blutdruckmessgerät und Zubehör

in the following configurations

Name or Type of device	
Aneroid Sphygmomanometer	GAMMA G7
Aneroid Sphygmomanometer	GAMMA G5
Aneroid Sphygmomanometer	GAMMA GP
Aneroid Sphygmomanometer	GAMMA GST
Aneroid Sphygmomanometer	GAMMA XXL LF

complies with

Council Directive 93/42/EEC (Medical Device Directive) of 14th June 1993
Richtlinie 93/42/EWG des Rates vom 14. Juni 1993 über Medizinprodukte

This declaration of conformity refers to the products marketed since 27. Jan. 2012 and is valid until a revised declaration of conformity is issued but not longer than 01. Feb. 2016 (expiry date of the Annex V EC-Certificate certificate).

Notified Body: DQS Medizinprodukte GmbH, August-Schanz-Straße 21, 60433 Frankfurt, Germany

CE 0297

Herrsching, 27. Jan. 2012

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