



ID-Nr. M-0001/04

**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) / Class Im

**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     | M-000.09.232,<br>M-000.09.233,<br>M-000.09.554,<br>M-000.09.560 |
| Aneroid Sphygmomanometer | GAMMA G5     | M-000.09.230,<br>M-000.09.231,<br>M-000.09.555,<br>M-000.09.561 |
| Aneroid Sphygmomanometer | GAMMA GP     | M-000.09.242,<br>M-000.09.243,<br>M-000.09.556,<br>M-000.09.562 |
| Aneroid Sphygmomanometer | GAMMA XXL LF | M-000.09.3**  |

complies with

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

This declaration of conformity refers to the products marketed since 2013-10-28 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, 2013-10-28  
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