



Heidenheim, 2011-03-23

EC-Declaration of Conformity for Medical Devices

We herewith declare,

That the below named products, which are manufactured and/or placed on the market by PAUL HARTMANN AG, meet the applicable provisions, especially the essential requirements of the following EC-Directive:

Council Directive 93/42/EEC for medical devices

The required conformity assessment procedure according to Annex II.3 has been performed and the technical documentation is kept available.

The conformity assessment procedure is under the supervision of the Notified Body: TÜV SÜD Product Service GmbH, DE-80339 München, Ridlerstr. 65, Identification No. 0123.

Product(s): VivanoMed Foam

Medical device class: Class IIb acc. to rule 4(2)

(acc. to Annex IX of the directive)

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Commercial Register of the
District Court of Ulm
file no. HRB 661090

HARTMANN helps healing.