HARTMANN

HARTMANN
GROUP

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)

PAUL HARTMANN AG

ppa.

Dr. W. Neumann

K.-D. Malowaniec

PAUL HARTMANN AG
Paul-Hartmann-Straße 12
89522 Heidenheim

P.O. Box 1420
89504 Heidenheim

Germany

Phone +49-7321-360
Fax +49-7321-36-3636

www.hartmann.info