



# PHILIPS

## Declaration of Conformity

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**Manufacturer:** Philips Medical Systems  
2301 Fifth Avenue, Suite 200  
Seattle, WA 98121  
USA

**European Representative:** Philips Medizinsysteme Boeblingen GmbH  
Hewlett-Packard Str. 2  
71034 Boeblingen  
Germany

**Product:** Heartstart HS1  
Models – M5066A, M5067A

**Classification:** Class IIb, Rule 9 of Annex IX of the MDD

We herewith declare that the above mentioned products meet the provisions of the council Directive 93/42/EEC for Medical Devices. All supporting documentation is retained under the premises of the manufacturer.

**Notified Body:** TÜV Product Service GMBH,  
Zertifizierstelle  
Ridlerstrasse 31  
D-80339 München  
Germany

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**EC Certificate(s):** G1 02 07 46696 001

**Start of CE-marking:** 18 October 2002 – s/n A02I00001

Seattle, WA 18 October 2002

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Teresa Skarr, Regulatory and Medical Affairs Manager