PHILIPS

Declaration of Conformity

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 #0123
EC Certificates(s):	G1 02 07 46696 001
Start of CE-marking:	18 October 2002 – s/n A02I00001

Seattle, WA 18 October 2002

Teresa Skarr, Regulatory and Medical Affairs Manager

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