



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, M5068A

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 65
D-80339 München
Germany

#0123

Start of CE-marking: 31 July 2003 – s/n Y071503-10

Seattle, WA 31 July 2003

Teresa Skarr, Regulatory and Medical Affairs Manager