PHILIPS

Declaration of Conformity

Manufacturer:	Philips Medical Systems 2301 Fifth Avenue, Suite 200 Seattle, WA 98121 USA
European Representative:	Philips Medizin Systeme Boeblingen GmbH Hewlett-Packard Str. 2 71034 Boeblingen Germany
Product:	HeartStart FRx Defibrillators Models – 861304, 861305
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:	2 February 2005, – s/n B04L-00055

Seattle, WA 2 February 2005

Teresa Skarr, Regulatory Affairs Manager

H05023