Declaration of Conformity

PHILIPS

Philips Medical Systems
22100 Bothell Everett Highway
Bothell, WA 98021-8431 USA

Manufacturer:	Philips Medical Systems 22100 Bothell Everett Highway Bothell, WA 98021-8431 USA
European Representative:	Philips Medizin Systeme Boeblingen GmbH Hewlett-Packard Str. 2 71034 Boeblingen Germany
Notified Body:	TÜV SUD Product Service GMBH Zertifizierstelle Ridlerstrasse 65 D-80339 München Germany
Product Name and/or Model:	HeartStart HS1 Infant/Child Pads Cartridge Models – M5072A
Classification:	Class IIb, Rule 9
EU Directive:	Directive 93/42/EEC
GMDN Code and Title:	41587 External defibrillator electrode, pediatric, single-use
Start of CE-marking:	Lot# Y081914-01
Additional Information:	None

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

Place and Date of Issue: Bothell, WA September 22, 2014

Signature: Carlene Comrie, Director Regulatory Affairs