



PHILIPS

Declaration of Conformity

Manufacturer: Philips Medical Systems
2301 Fifth Avenue, Suite 200
Seattle, WA 98121-1825
USA

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

Product: Philips Heartstart FR2
Models – M3860A, M3861A
Laerdal Heartstart FR2
Models – M3840A, M3841A

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

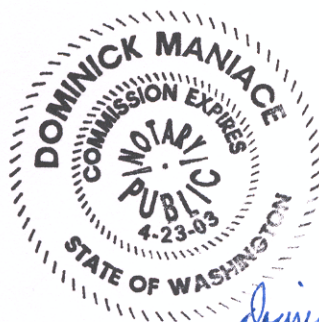
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EC Certificate(s): G1 02 07 46696 001

Start of CE-marking: 20 August 2002, M3860A – s/n 0802071071
M3861A – s/n 0802070901
M3840A – s/n 0802070037
M3841A – s/n 0802069952

Seattle, WA 20 August 2002

Teresa Skarr, Regulatory and Medical Affairs Manager



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Dominick Maniace
my commission expires: 4/23/03