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DECLARATION OF CONFORMITY

We,

ZOLL *Medical Corporation*
269 Mill Road
Chelmsford, MA 01824-4105 USA,

declare under our sole responsibility that the products:

E Series
E Series Pulse Oximetry Option
E Series End Tidal CO_2 Option
E Series 12 Lead Option
E Series NIBP Option
M Series
M Series CCT
M Series Biphasic Option
M Series Pulse Oximetry Option
M Series End Tidal CO_2 Option
M Series 12 Lead Option
M Series NIBP Option
M Series IBP/TEMP Option
AED PLUS
AED PRO
CPR-D padz™
R Series

to which this declaration relates are in conformity with the provisions of Council Directive 93/42/EEC of 14 June 1993 of the Medical Device Directive which apply to them.

The quality system under which these products were designed and manufactured has been certified by TUV Rheinland Product Safety (0197) to be in compliance with Annex II of the Medical Device Directive including European Standard, ISO 13485.

18 OCT 2006

Paul Dias
V.P. Quality Assurance and Regulatory Affairs

ZOLL Medical Corporation

Authorized European Representative
Eric Rozeboom
ZOLL International Holding B.V.
Edisonring 3a
6669 NA Dodewaard
The Netherlands
Tel. +31 488-411-183

CE

Certificate of Conformance

This is to declare under our sole responsibility that the following ZOLL Medical Corporation products:

AED Plus
AED Pro
E Series
M Series
M Series CCT
Smart Battery
Smart XL Battery
R Series
SurePower Battery

are in conformance with the provisions of Council Directive 2002/96/EC of 27 January 2003 on Waste Electrical and Electronic Equipment which apply to them.