



DECLARATION OF CONFORMITY



ZOLL Medical Corporation
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<i>Product Description</i>	<i>Classification</i>	<i>Rule</i>	<i>Annex</i>	<i>Beginning Lot Number</i>
<i>CPR-D padz®</i>	<i>IIb</i>	<i>9</i>	<i>IX</i>	<i>0113</i>
<i>CPR-D padz with Accessory Kit</i>	<i>IIb</i>	<i>9</i>	<i>IX</i>	<i>0113</i>
<i>CPRD-padz without Accessory Kit</i>	<i>IIb</i>	<i>9</i>	<i>IX</i>	<i>0113</i>

ZOLL declares that the above products conform to European Council Directive 93/42/EEC (Medical Device Directive) class IIb per rule 9 of Annex IX, assessed per Annex II.

The quality system under which these products were designed and manufactured has been found to be in compliance with the Medical Device Directive including European Standard EN ISO 13485:2012 certified by the Notified Body TUV SUD Product Service GmbH, Ridlerstr. 65, 80339 München, Germany (Notified Body Number 0123).

23 January 2013

Paul Dias
Vice President, QA & RA

The above products 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.

