

EC Declaration of Conformity

Manufacturer: **HeartSine Technologies Limited**
Canberra House
203 Airport Road West
Belfast, BT3 9ED

Device: **samaritan® PAD 360P**
Model: **SAM 360P**
Description: Automated external defibrillator.
Medical Device Classification: Identified as **Class IIb** under rule 9 of Annex IX of Council Directive 93/42/EEC as amended by 2007/47/EC

HeartSine Technologies declares that the HeartSine samaritan PAD 360P (SAM 360P), a therapeutic medical device in the range of Automated External Defibrillators, and its associated accessories are designed and manufactured in conformity with:

- a) the essential requirements (Annex I) and provisions of the European Medical Device Directive (MDD) **European Council Directive 93/42/EEC** (as amended by 2007/47/EC)
 - o And is subject to the procedure set out in Annex II (excluding section 4), Full Quality Assurance System, of Directive 93/42/EEC, as amended by Directive 2007/47/EC;
 - o Under the supervision of notified body number CE0120, SGS United Kingdom Limited, Worle Parkway, Weston Super Mare, United Kingdom, BS22 6WA.
- Article 4 of ROHS2 Directive (2011/65 EU), with exceptions Annex III (6c – lead in copper alloy) and Annex IV (17 – lead in solder for portable emergency defibrillators).

Certification	SGS Certificate Number
Council Directive 93/42/EEC	GB02/54193
ISO 13485:2003	GB02/54195
EN ISO 13485:2012	GB02/54195
ISO 9001:2008	GB02/54194

Signature 

Date 15 Aug 14

Paul Phillips
VP Quality & Regulatory Affairs
HeartSine Technologies Ltd.

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