



EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 067590 0006 Rev. 01

Manufacturer: **HeartSine Technologies, Ltd.**
207 Airport Road West
Belfast, Northern Ireland, BT3 9ED
UNITED KINGDOM

Product Category(ies): Automatic External Defibrillators for adult and paediatric use. Automated External Defibrillator battery and defibrillator electrode cartridges:

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10675900006Rev.01

Report No.: 75950803_CN

Valid from: 2021-03-30
Valid until: 2024-05-26

Date, 2021-03-30

Christoph Dicks
Head of Certification/Notified Body