



Declaration of Conformity

Manufacturer Cardiac Science Corporation
 N7 W22025 Johnson Road
 Waukesha, Wisconsin 53186-1856
 United States of America

I hereby ensure and declare that the medical device, Powerheart® G5 AED, Automatic, meets the provisions of the Directive 93/42/EEC (M5), and with the Essential Requirements (Annex I), concerning medical devices. This declaration is made on the basis of the EC Certificate number 00357, issued by BSI (ID# 2797) in accordance with Annex II, Section 3 of this directive.

In addition, I hereby ensure and declare under some responsibility of the manufacturer that the Powerheart® G5 AED, Automatic, meets the provisions of the Directive 2011/65/EU of the European Parliament and the Council of 8 June 2011 the restriction of the use of certain hazardous substances in electrical and electronic equipment.

Applicable Standards Refer to Annex A of this Declaration

Product Powerheart® AED G5 Automatic (Model # G5A)

Classification (MDD, Annex IX) Class IIB (Rule 9)

GMDN Code 48047

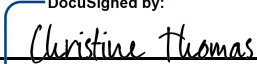
UMDN Code 17116

Start of CE Marking June 8, 2012

Authorized Representative MDSS GmbH
 Schiffgraben 41
 D-30175 Hannover
 Germany

7/29/2019 | 7:23:56 AM CDT

Date

DocuSigned by:

 Signature

Vice President, Regulatory Affairs

Title

Christine Thomas

Printed name