

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. CE 657862
Issued To: **Myocardial Solutions, Inc.**
808 Aviation Parkway
Suite 700
Morrisville
North Carolina
27560
USA

In respect of:

Design, development and manufacture of software for the quantification of cardiac MRI images.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):



Gary E Slack, Senior Vice President Medical Devices

First Issued: **2016-12-20**

Date: **2021-01-29**

Expiry Date: **2024-05-26**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

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Supplementary Information to CE 657862

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| NBOG Code(s) | Device Description | Intended purpose |
|------------------|--|-------------------|
| Class IIa | | |
| MD 1111 | Software for analysis of cardiac MRI images for quantification of cardiac function | N/A for Class IIa |

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Information and Contact: BSI, Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands Tel: + 31 20 346 0780

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