EC Certificate Full Quality Assurance System: Certificate US08/2927

The management system of

LifeLine Software, Inc.

3304 South Broadway Avenue, Suite 200,
Tyler, TX, 75701-7810, United States
has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Radiation Dose Verification software.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 10 January 2017 until 10 January 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 3 January 2020
Issue 5. Certified since 10 January 2008

Certification is based on reports numbered WW/MW 600809

Authorised by

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