EC Certificate Full Quality Assurance System: Certificate US07/844

The management system of

Segami Corporation

8310 Guilford Road, Suite A,
Columbia, MD, 21046, 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

Medical imaging systems for the acquisition, processing, review and archiving of radiological images.

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 2019 until 09 January 2024
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 02 March 2021
Issue 8. Certified since 12 March 2007

Certification is based on reports numbered WW/MW 215666

Authorised by

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