EC Certificate Full Quality Assurance System: Certificate US19/819943515

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 molecular imaging software system for the acquisition, processing and review 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 16 December 2019 until 09 January 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 12 March 2007
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered WW/MC 215666

Authorised by

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