EC Certificate Full Quality Assurance System: Certificate US19/819943496

The management system of

INVIA, LLC

3025 Boardwalk Street, Suite 200,
Ann Arbor, MI, 48108, 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

Software/systems for the analysis of radiographic (nuclear, PET and CT) medical images.
Corridor4DM – Nuclear Cardiac Quantification 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 16 December 2019 until 26 February 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 28 September 2010
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered WW/MW 602506

Authorised by

Pieter Weterings
Certification Manager

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