EC-Declaration of Conformity
according to Directive 93/42/EEC, Annex II excluding section (4)

MiE medical imaging electronics GmbH
represented by the presidents Thomas Kühl and Günther W. Kühl

declares in own responsibility that following described system fulfills the requirements of Annex I of the Directive 93/42/EEC.

Manufacturer: MiE medical imaging electronics GmbH
Hauptstr. 112,
D-23845 Seth, Germany
Tel.: +49 4194 9977-0
Fax: +49 4194 9977-55
mie@miegjermany.de www.mie-scintron.com

Product Name: System ECAM - SCINTRON

Product Description: Anger type of whole-body and SPECT Gamma-Camera with one alt. two high resolution rectangular detectors and Nuclear Medicine Workstation for the whole nuclear medicine diagnostic

Classification: Class IIa

Validity: This Declaration of Conformity is valid until 26.05.2024

Name and Address of Notified Body: DEKRA Certification GmbH
Handwerkstraße 15
D-70565 Stuttgart, Germany

Notified Body number: 0124

Medical Devices – Quality Management Systems – Requirements for regulatory purposes

Seth, 20.05.2021
Date

Signature President