

EC Declaration of Conformity

Date: Feb 27, 2020

Manufacturer:

MIM Software Inc.
25800 Science Park Drive
Suite 180
Cleveland, Ohio 44122 USA

EU Authorized Representative:

Emergo Europe BV
Prinsessegracht 20
2514 AP The Hague
The Netherlands
Phone: (31) (0) 70 345-8570
Fax: (31) (0) 70 346-7299
EmergoVigilance@ul.com

Declares that the medical device described hereafter

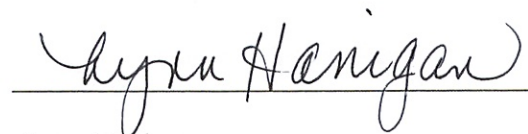
MIM®

*Includes variants: MIMfusion, MIMcardiac, MIMneuro, MIM Maestro, MIM Encore,
MIM Symphony (LDR & Bx), MIM – Thin Client (mobile), MIM
SurePlan Liver Y90, MIM SPECTRA Quant, and MIM SurePlan MRT.*

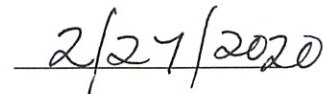
Classification of IIa using Annex IX, Rule 10

is in conformity with the essential requirements and provisions of Council Directive 93/42/EEC as amended by Directive 2007/47/EC and is subject to the procedure set out in Annex II (excluding Section 4) of Directive 93/42/EEC under the supervision of **Notified Body** SGS Belgium NV (1639), Noorderlaan 87, BE-2030 Antwerp, Belgium.

CE Certificate: US19/819943671



Lynn Hanigan
Quality Assurance Director
Management Representative
MIM Software Inc.



Date: