EC Certificate Full Quality Assurance System: Certificate US19/819943671

The management system of

MIM Software Inc. also trading as MIM Software Brussels BVBA, MIM Software Beijing Co., Ltd and MIM Software

25800 Science Park Drive, Suite 180,
Cleveland, OH, 44122, 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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 27 February 2020 until 02 April 2024
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 07 July 2005
and first certified by SGS Belgium NV since 16 December 2019

This is a multi-site certification.
Additional site details are listed on subsequent pages

Certification is based on reports numbered WW/MC 211752

Authorised by

SGS Belgium NV, Notified Body 1639
SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

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Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

Issue 2

Detailed scope

Software/Systems including MIM, MIMviewer and Mobile MIM for medical image viewing applications for diagnosis, treatment evaluation, and treatment planning

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.

Additional facilities

MIM Software Beijing Co., Ltd, Suite 809, Shouxiang Science Building, No.51 Xueyuan Rd., Haidian District, Beijing, 100191, P.R.China

MIM Software Beijing Co., Ltd, , Suite 2315, Shangpu International Office Building, NO.6 Daye RD., Jinjiang District, Chengdu, 610020, P.R.China

MIM Software, Houtparklaan 1 bus 21 , 3600 Genk, Belgium

MIM Software Brussels BVBA, Drukpersstraat 4, 1000 Brussel, Belgium