Number: 3903172CE01

EU Quality Management System Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex IX Chapter I and III

Manufacturer:

iMDsoft® Ltd.

Kiryat Atidim
 # 4
POB 58178
Tel Aviv, 6158101
Israel
SRN ID.: IL-MF-000018368

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EU- Regulation which apply to them:

0344

Supplement to certificate: 2007326CN

Authorized Representative: MDSS GmbH, Schiffgraben 41, 30175 Hannover, Germany

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant requirements of EU Regulation 2017/745, including all subsequent amendments for the above mentioned conformity assessment. The manufacturer/ authorized representative is subject to periodic surveillance as required for the applicable conformity assessment in accordance to Regulation 2017/745.

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.M. McKenzie
Principal Certification Manager

First Issued: **30-11-2023** Date: **30-11-2023** Expiry date**: 01-11-2028**

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 www.dekra.nl Company registration 09085396

2184-5

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This certificate covers the following device(s) / groups of device(s):

MEDICAL DEVICE SOFTWARE - NOT INCLUDED IN OTHER CLASSES (V92, class IIb)	
Device Name: MetaVision® ver. 6.x (beginning from ver. 6.20)	*Intended Purpose: MetaVision is intended for clinical and workflow documentation, interfacing, conversion, presentation, and storage, order and medication management, decision support and analysis in the healthcare environment (e.g. high acuity and acute care). MetaVision may provide the following uses, without controlling or altering the functions or parameters of any other connected medical devices: (i) the electronic transfer of medical device data; (ii) the electronic storage of medical device data; (iii) the electronic conversion of medical device data from one format to another format in accordance with a preset specification; and (iv) the electronic display of medical device data.*

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Conditions for or limitations to the validity of this certificate:

- N/A

Certificate History

Identification of the Common Specifications and Harmonized Standards complied with are documented within the technical documentation and audit assessments carried out. These are traceable through the DEKRA Certification B.V. Certification Notice. The Certification Notice also identifies the necessary information related to the quality management system of the manufacturer, including facilities.

Revision	Date of Issue certificate	Certification Notice Reference	Action
0	30-11-2023	2007326CN18	First issue

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