AI4CMR Software

User Manual



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Manufactured by

Al4MedImaging Medical Solutions S.A.

SRN PT-MF-000002951

Address

Rua do Parque Poente, Lote 35 4705-002 Sequeira, Braga Portugal

Telephone

+351 253 145 430

https://ai4medimaging.com/ai4cmr/

European Union



Al4CMR is qualified as a class IIa medical device. It complies with the requirements of the European Medical Device Regulation MDR 2017/745

MDR Certificate, with CE Mark Number: MDR 740464

UDI (01)05600250059015(11)210729(21)010000

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1. Terms and Conditions

This User Manual shall be accompanied by the specific plugin User Manual.

The plugin User Manual describes in more detail the features and steps necessary to interact with the AI4CMR service. For clarification purposes only, we present an example of a possible sequence of steps to interact with AI4CMR.

1.1 Regulatory Status

Al4CMR bears the MDR Certificate, with CE Mark Number: MDR 740464.

Users are required to investigate the regulatory requirements pertinent to their country or location prior to using AI4CMR.

It is the user's responsibility to obey these rules, statutes, and regulations.

1.2 Intended use

Al4CMR is an image analysis software for post-processing of cardiac magnetic resonance images (CMR), which are used for assessment of cardiac diseases.

Al4CMR uses artificial intelligence and is intended to automize cardiac segmentation allowing instantaneous quantification of ventricular volumes, myocardial mass, ejection fraction of heart ventricles and to automatically assess left ventricle wall motion.

Al4CMR shall be used by qualified medical professionals, experienced in examining and evaluating magnetic resonance images as a support tool that provides relevant clinical information.

AI4CMR can be integrated with DICOM viewers through a standard communication protocol.

The software is not intended to determine or recommend a course of action or treatment for a patient.

AI4CMR software does not preclude the analysis of the reporting physician. The cine sequences analysed by AI4CMR are only a part of the complete CMR exam and all the data should be integrated with other sequences and the clinical background of the patient to conclude the CMR exam.

The reporting physician holds the responsibility for validating the data provided by the AI4CMR software and performing the final interpretation of the CMR exam.

1.3 Intended users

Al4CMR is a medical device and shall be used by qualified medical professionals, experienced in examining and evaluating magnetic resonance images.

1.4 Conditions for use

Al4CMR is a cloud-hosted service. The system shall be used in association with a third-party DICOM Viewer, which provides the interface between Al4CMR and the intended users (Medical professionals), Software Developers, and Others.

Patients will not be in direct contact with AI4CMR. The CMR images acquired for the patient are fed from the MRI Scanner or PACS (picture archiving and communication system) to the third-party DICOM Viewer.

This system context and interfaces to third-party systems are represented in the diagram of Figure 1. The AI4CMR system boundaries, are defined by the box "AI4CMR".

The integration between AI4CMR and the DICOM Viewer is established using plugins that implement a standard communication protocol.

The communication between AI4CMR and the third-party system is initiated by the third-party system which posts a request to the AI4CMR in a networked environment. AI4CMR will validate and process the request and return a response via the same medium.

The patient data is anonymized by the third party before being sent to AI4CMR.

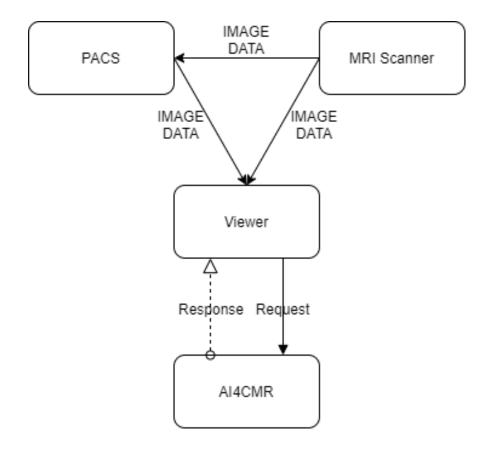


Figure 1. High-level system representation of AI4CMR.

1.5 Contraindications

Al4CMR is contraindicated to be used for post-processing of all image modalities other than cine CMR.

1.6 Limitations

AI4CMR analysis is limited to cine CMR acquisitions.

Other studies such as Late Gadolinium Enhancement or Perfusion are not covered.

Al4CMR's performance has been tested in three MRI scanner vendors: Siemens, GE Healthcare Systems and Phillips Medical Systems. For new MRI scanner vendors, a warning message will be returned to the User stating it has not been tested.

Al4CMR's performance has been validated for a range of image characteristics. Cases with image characteristics that are outside the boundaries of the verified performance of the algorithms will be rejected.

1.7 Warnings and precautions

- The physician must always review the cases even when AI4CMR classifies them as normal.
- Quantitative analysis is dependent on the quality and correctness of the image source data acquisition.
- The patient data must be anonymized by the third-party plugin before sending it to AI4CMR.
- The patient data displayed is initially derived from the DICOM information, if available.
 Editing these values may affect the calculations in all modules. It is the User's responsibility to verify this data before releasing the final results.
- The User must keep a UPS system that provides emergency power when the input power source or mains power fails.
- It is the plugin maintenance team's responsibility to execute the installation procedure to validate the communication between the User and AI4CMR.
- It is Al4MedImaging's responsibility to update/do maintenance periodically to reduce the probability of Al4CMR SW defects.
- At the installation time, a copy of the digital manual must be copied to the User's network, under the domain of the User's IT Manager.
- In the event of loss of access credentials please follow the procedure in section 4.
 User account handling.

1.8 Intended patient population

The software analyses images of patients that were examined by CMR imaging.

Age The use is not recommended for children (age < 18yo), since the clinical evaluation studies targeted a mostly adult population.

Weight No limitations

Height No limitations

1.9 Applicable medical conditions

Al4CMR is an image analysis software for the post-processing of CMR images, used for the assessment of the following cardiac diseases:

- Cardiac failure
- Ischemic heart disease
- Valvular heart disease
- Cardiomyopathy
- Cardiac function and morphology
- Hypertensive cardiac disease
- Other pathologies that need assessment of cardiac function and morphology

1.10 Safety Instructions

Software AI4CMR is hosted in a cloud. The intended users interact remotely with the software.

The equipment communicating with AI4CMR shall:

- Be connected to a secure network;
- Have an access control system, to ensure that only accredited users can access the device and the AI4CMR service.

It is highly recommended the usage of anti-virus software in all computers interacting with the AI4CMR to protect against cyber-attacks. Periodic backups are also recommended to make sure that no patient data is lost in an unforeseen event.

2. Conventions and Abbreviations

2.1 Conventions

EDV – End-Diastolic Volume [ml]

ESV – End-Systolic Volume [ml]

SV – Stroke Volume [ml]

EF – Ejection Fraction [%]

Mass – Myocardial Mass [g]

CO - Cardiac Output [L/min]

EDV/BSA – End-Diastole Volume/Body Surface Area [ml/m²]

ESV/BSA - End-Systole Volume/Body Surface Area [ml/m²]

SV/BSA – Systole Volume/Body Surface Area [ml/m²]

Mass/BSA – Myocardial Mass/Body Surface Area [g/m²]

CI – Cardiac Index [L/(min m²)]

BSA - Body Surface Area [m²]

2.2 Abbreviations

ED - End-Diastole

ES - End-Systole

MRI - Magnetic Resonance Imaging

CMR - Cardiac Magnetic Resonance

DICOM - Digital Imaging and Communications in Medicine

HTTP – Hypertext Transfer Protocol

SA - Short-axis

LA – Long-axis

LV - Left Ventricle

3. Getting Started

3.1 Notice to the user

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

3.2 System requirements

Al4CMR software runs as a Docker image on the cloud. The minimum requirements for the ethernet connection are:

• High bandwidth network (50Mbits/s minimum and 100Mbits/s recommended).

3.3 Setup

For customer registration, the customer environment shall be informed to Al4MedImaging: MRI scanner vendor, PACS vendor and version, Viewer vendor and version and third-party plugin ID and version.

A *client_id* and temporary user authentication credentials will be assigned by Al4MedImaging to the costumer.

The third-party plugin shall be installed and configured as per its User Manual.

To validate the set-up, the following validation protocol shall be followed:

1. Three clinical cases (hereinafter called gold samples GS1; GS2 and GS3) are delivered to the customer by Al4MedImaging. The samples shall be submitted to Al4CMR through the third-party plugin.

GS1 case is a valid case with all the mandatory and optional data;

GS2 case is a valid case with all the mandatory data;

GS3 case is a non-valid case with missing mandatory data.

2. The results displayed by the third-party plugin for each of the golden samples shall be compared with the following expected results:

GS1:

Classification		Confidence		
Normal		0.95		
EDV	EDV index		CI	
111.58	74.	.11	3.18	
ESV	ESV	index	ED frame	
36.84	24.	47	1	
SV	SV ii	ndex	ES frame	
74.74	49.64		12	
Mass	Mass index		BSA	
78.03	51.83		1.51	
EF			CO	
66.98			4.78	
	Final Clas	ssification		
	Nor	mal		

GS2:

A warning shall be displayed stating CO, CI and other indexed volume metrics were not computed since heart rate, height and weight were not given.

word not dempated direct rate, neight and weight word not given.				
Classification	1	Confidence		
Suspect		0.70		
EDV	EDV	index	CI	
209.53		-	-	
ESV	ESV	index	ED frame	
105.66		-	25	
SV	SV ii	ndex	ES frame	
103.87		-	9	
Mass	Mass	index	BSA	
139.71		-	-	
EF			CO	
49.57			-	
Final Classificat				
Suspect				

GS3:

The submission shall be disabled. A warning shall be displayed stating there are missing required fields.

- 3. The customer's IT Manager must send the validation report to Al4MedImaging's Field Support via email. The report must include:
 - A signed statement by the IT Manager confirming if the validation protocol was successful or if it failed.
 - o Screenshots of the results obtained for the 3 golden samples.
- 4. Al4MedImaging's Field Support must confirm reception and sign the report. If the validation protocol is successful Al4MedImaging will provide the final user

credentials to the customer. Otherwise, further support must be offered for problem resolution in articulation with the third-party plugin vendor.

3.4 User Training

After the setup is validated, the users will partake in individual or group training given by Al4MedImaging through the Microsoft Teams platform. The users must be trained before using Al4CMR.

Additional individual/group training or Demos can be requested and scheduled by contacting our Technical Support (Section 5).

3.5 Software overview

The AI4CMR operating sequence of events is summarized in Figure 2. The green coloured boxes include the operations executed on the external DICOM Viewer plugin, to initiate AI4CMR processing (blue boxes).

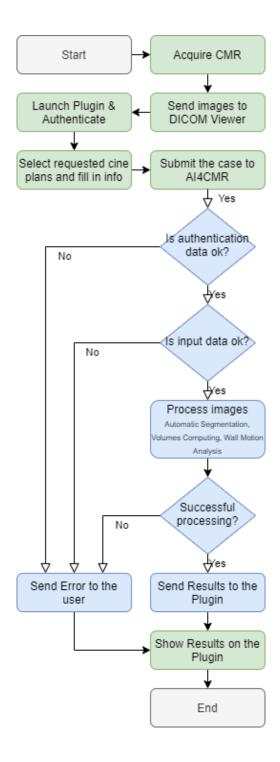


Figure 2. Flowchart of the operating sequence. Green steps are out of Al4CMR's control; blue steps are controlled by Al4CMR.

3.5.1 General operating steps

The general operating steps to use AI4CMR are summarized in the following table:

Operating step	System context
1. The Cardiac Magnetic Resonance is acquired	MRI Scanner
according to standardized protocols	
2. The images are sent to the third-party software	DICOM Viewer
3. The user launches the dedicated plugin on the	Plugin
third-party software and inserts the user credentials	
4. The user selects the correct cine sequence	Plugin
planes and other requested patient data	
5. The user submits the request form to the AI4CMR	Plugin
cloud service	
6. The request is subject to authentication	AI4CMR
7. The data on the request is checked for sanity and	AI4CMR
completion	
8. The patient data is processed by several Al-	AI4CMR
based image processing modules (myocardium	
automatic segmentation, ventricle volumes	
computation, wall motion analysis)	
9. The results and log messages (errors, warnings,	AI4CMR
) are collected and returned to the plugin via the	
communication API	
10. The user sees the output of AI4CMR on the	Plugin
dedicated plugin	

A CMR case to be submitted to AI4CMR shall consist of:

- 1. Required:
 - o DICOM cine SA with metadata:
 - Image Patient Position;

- Image Patient Orientation;
- Pixel Spacing;
- Slice Thickness;
- Rows;
- Columns.
- Patient metadata:
 - Age (on acquisition date);
- o Sex.

2. Optional:

- DICOM cine 2ch, 3ch and 4ch (recommended);
- Height, weight, heart rate.

The outputs of AI4CMR consist of:

- Left Ventricle volumes-derived metrics:
 - EDV End-Diastolic Volume [ml]
 - ESV End-Systolic Volume [ml]
 - SV Stroke Volume [ml]
 - EF Ejection Fraction [%]
 - Mass Myocardial Mass [g]
 - CO Cardiac Output [L/min]
 - EDV/BSA End-Diastole Volume/Body Surface Area [ml/m²]
 - ESV/BSA End-Systole Volume/Body Surface Area [ml/m²]
 - SV/BSA Systole Volume/Body Surface Area [ml/m²]
 - Mass/BSA Myocardial Mass/Body Surface Area [g/m²]
 - CI Cardiac Index [L/(min m²)]
 - BSA Body Surface Area [m²]
- Case classification:
 - Wall motion class normal or suspect
 - Wall motion class confidence
 - Final class normal or suspect

See Section 3.5.2 for details on clinical performance.

Notice:

- The final case class takes both wall motion classification and ejection fraction value into account. A case is only deemed as normal if both criteria are in the normality range.
- 2. The "confidence" is a probability output (0-1) from the machine learning models used to classify each case. The closer the value is to 1, the greater is the confidence in the attributed classification, taking the original training database¹ as a reference.

3.5.2 Summary of outputs validation

The studies conducted under the clinical evaluation phases showed high agreement between AI4CMR measurements and the human observers' in a set of 146 patients:

- Correlation in the measurement of EDV is 0.99, ESV is 0.99, SV is 0.81, EF is 0.96 and Mass is 0.95. The other volume metrics are arithmetically derived from the previous.
- The accuracy and precision on the measurement of the volume is given by the bias (mean difference) and the standard deviation:

		EDV (ml)	ESV (ml)	SV (ml)	Mass (g)	EF (%)
A	AI4CMR vs rater1 (*)	8.6±17.6	-3.7±17.6	11.7±17.1	1.6±19.6	3.9±5.7
1	AI4CMR vs rater2 (*)	8.7±17.6	-2.1±17.9	10.8±16.8	3.3±18.4	3.8±5.8

- (*) The raters (experts) used the certified software QMass® by Medis Medical Imaging Systems, which computes the volumes based on the raters' manual segmentations.
- For case classification, the sensitivity is 100%, the negative predictive value is 100%, the accuracy is 76.7%, the positive predictive value is 70% and the specificity is 50%.

No uncertainty/error information is shown in the software together with the LV measurements.

¹ Training database - Neural networks and other artificial intelligence programs require an initial set of data, called training data, to act as a baseline for further application and utilization. Al4CMR has been trained, validated and tested on a split database of around 1000 patients, labelled by experts.

3.5.3 Plugin

The Plugin User Manual which accompanies the current manual describes in more detail the features and steps necessary to interact with the AI4CMR service. The Plugin consists of a user interface to help the User create the request form. Different viewers can have different user experiences, but the main idea should be the same.

For clarification purposes, this section provides merely indicative (not binding) examples of what the user experience may be like with a third-party Plugin for AI4CMR.

Login

The first step to use the AI4CMR is to successfully authenticate with the service. To do that, the User shall provide a username and password, Figure 3:



Figure 3. Example of a plugin login window.

Form filling

The plugin provides a way for the User to fill in the necessary information to perform a request, as presented in Figure 4.

The User shall identify each sequence (SA, 2ch, 3ch, 4ch) and add it to the Plugin (using drag and drop, for example). The SA sequence is mandatory and should always be

provided. The LA sequences are optional and can be discarded if they do not exist. The inclusion of the LA views is highly recommended if they are available.

Patient metadata shall be filled in by the User if the Plugin does not automatically fill it based on the DICOM information :

- Mandatory:
 - Patient ID
 - Age
 - Sex
- Optional:
 - Heart rate (if not sent to AI4CMR: CO and CI will not be computed)
 - Height (if not sent to AI4CMR: no BSA nor indexed metrics (EDV/BSA, ESV/BSA, SV/BSA, Mass/BSA, CI) will be computed)
 - Weight (if not sent to AI4CMR: no BSA nor indexed metrics (EDV/BSA, ESV/BSA, SV/BSA, Mass/BSA, CI) will be computed)



Figure 4. Example of a plugin form filling window.

Results presentation

After successful processing, the plugin presents the results.

One example of a possible user interface for results reporting is presented in the bottom half of Figure 5 (LV report).

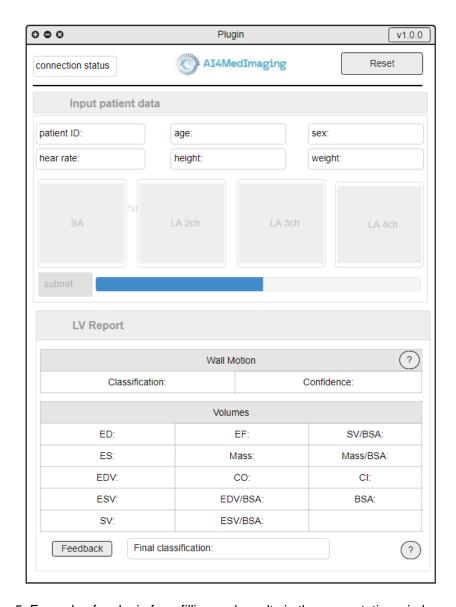


Figure 5. Example of a plugin form filling and results in the presentation window.

After a failed processing, the plugin presents an error message, so that the User can adapt the request or report a possible problem.

Parametrizations

Al4CMR allows for the change of a set of parameters (wall motion threshold and ejection fraction thresholds). These values should only be changed by the customer's IT Manager and after careful consideration.

Additional fields

Connection status

This field is used to indicate that the AI4CMR service is available. If not, the User can force a connection status update by clicking on the status button.



Figure 6. Connection status example.

Reset button

The reset button is used to clean all the info from the current request and make it easier for the User to start a new request without cleaning each field individually.



Figure 7. Reset button example.

Info button

This button presents information regarding the plugin and AI4CMR service, such as email, certifications, versions, and so on.

A suggestion for the presentation of the AI4CMR label can be seen in Figure 9.



Figure 8. Info button example.

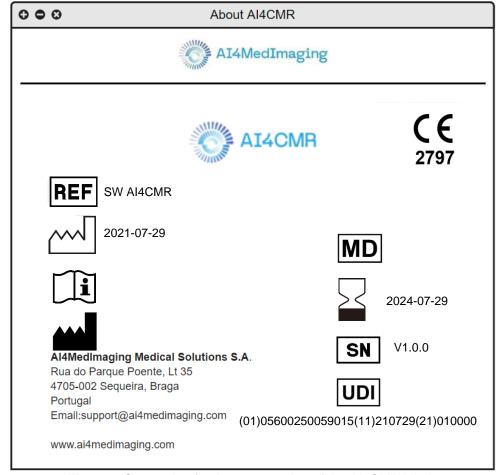


Figure 9. Suggestion for the presentation of the AI4CMR label.

Confidence disclaimer

This button presents information to clarify the wall motion classification results.



Figure 10. Confidence disclaimer button example.

Feedback button

The User can access plugin support by clicking on this button, which redirects him to feedback email software.



Figure 11. Feedback button example.

4. User account handling

The AI4CMR provides a user account handling page, that allows each user to change their personal information and credentials. To access this functionality, the user should use the following link:

https://auth.ai4medimaging.com/auth/realms/srerealm/account

4.1 Authentication

To access the account, the user should provide the assigned username or email and password.



Figure 12. User account login page.

After successful authentication the user handling page will be presented, Figure 13.

4.2 User information handling

On the separator "Account" the user can change the email, first name and last name by filling the respective field and clicking in "Save". If these values already exist, they will be presented in the respective field.

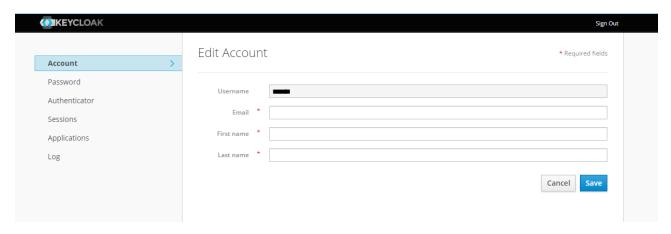


Figure 13. User account personal information edit page.

4.3 Password handling

To change the password, the user should change to the "Password" separator, provide the current password on the "Password" field and the new password on the "New Password" and "Confirmation" fields. After clicking in "Save" the password will be updated.

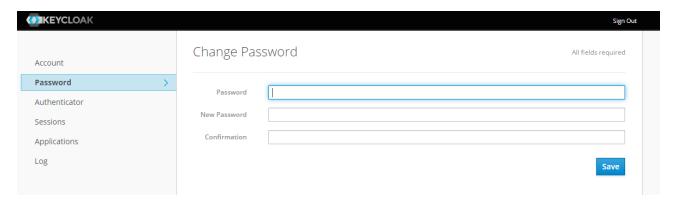


Figure 14. User account password edit page.

In case of password loss or account compromised, the IT Manager of the User's Institution should contact the Al4Med Technical Support (see section 5 for details).

4.4 Open sessions

The user has the option to monitor all open sessions associated with his account. That information is accessible in the "Sessions" separator. If any of the sessions open is considered suspect, the user has the option to close it using the "Log out all sessions" button.

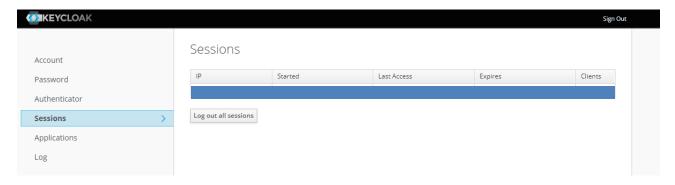


Figure 15. User account open sessions page.

4.5 Logging info

In addition to the open sessions, the user has the option to see all logging events that happen using his account. This information can be accessed on the "Log" separator.

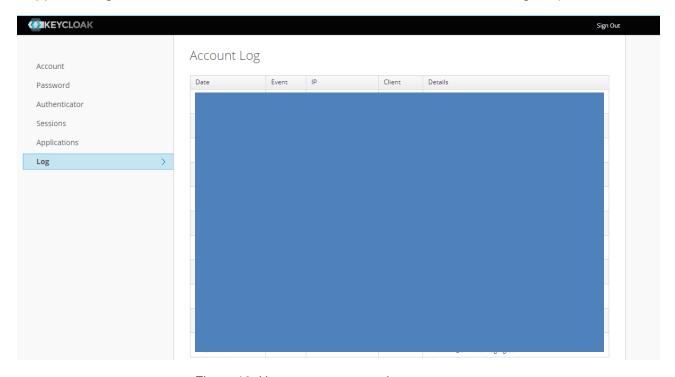


Figure 16. User account access log page.

5. Technical Support

For technical questions, to report a problem, or to request individual/group training or demos, please contact our team by phone, e-mail, or website:

Al4MedImaging Medical Solutions S.A.

Rua do Parque Poente, Lote 35 4705-002 Sequeira, Braga Portugal

Telephone: +351 253 145 430

Website: https://ai4medimaging.com/

Problem reporting e-mail: support@ai4medimaging.com

6. Lifetime

Intended lifetime of 3 years from the release date (2024-07-29)

Approval of the Instructions for Use – User Manual (Record)

Role	Function	Name	Signature	Approval date
Author	Head of Technology and Science	Nelson Martins		
Reviewer	Head of Development	Eva Costa		
Approver	QA/RA Manager	Carla Almeida		

Record Version	Description of changes
01	Initial approval
02	Addition of performance metrics for the measurement of the left ventricle volumes in section 3.5.2 Summary of outputs validation.
03	Change on CE marking of conformity logo
04	Added Notified Body number field to the CE marking of conformity logo.
05	Updated the Regulatory Status, Plugin and Lifetime sections. Added UDI and SRN numbers in the second page of this manual.