

A VARIATIONAL METHOD FOR SCAR SEGMENTATION WITH MYOCARDIAL CONTOUR CORRECTION IN DE-CMR IMAGES

S. Merino-Caviedes ¹, L. Cordero-Grande ², M. T. Pérez Rodríguez ³, M. T. Sevilla-Ruiz ⁴,
A. Revilla-Orodea ⁴, M. Martín-Fernández ¹, C. Alberola-López ¹

¹ Laboratorio de Procesado de Imagen, Universidad de Valladolid, Spain

² Department of Biomedical Engineering, King's College London, London, UK

³ Departamento de Matemática Aplicada, Universidad de Valladolid, Spain

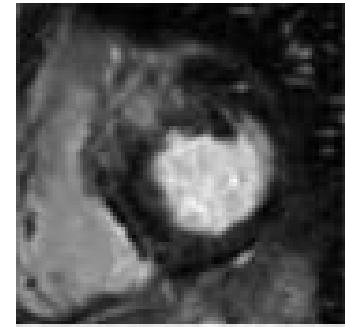
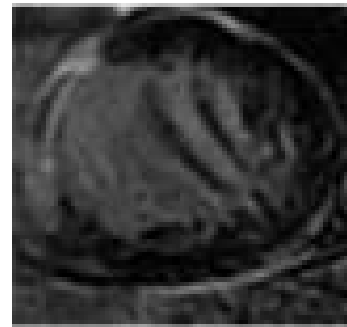
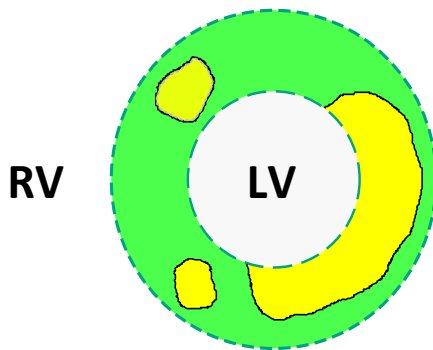
⁴ Instituto de Ciencias del Corazón, Hospital Clínico Universitario, Valladolid, Spain

Index

- Introduction
 - Motivation
 - DE-CMR Segmentation
 - Objectives
- Proposed Algorithm
 - Variational Framework
 - Label Posterior Probability Formulation
 - Contour Regularization
- Experimental Results
- Conclusions

Motivation

- Cardiac MR images:
 - Good contrast between soft tissues
 - The full heart can be imaged
 - There are many modalities: CINE, Tagging, DE-CMR...
- DE-CMR allows the identification of scarred tissue in the myocardium.



Review of DE-CMR Segmentation

- Most current methods restrict the segmentation of the myocardium.
 - It simplifies the problem.
 - What if there are misalignments in the myocardial mask?

Method	Myocard. Contours	Scar identification method	Post Processing included in the method
TFA1 [5]	Fixed	2 step thresholding	FP, FN removal by feature analysis, region growing and hole filling
TFA2 [10]	Fixed	2 step thresholding	FP, FN removal by feature analysis
PWD [3]	Fixed	Mixture Model + Watershed	Noise removal + hole filling

[5] Q. Tao, J. Milles, K. Zeppenfeld, H. J. Lamb, J. J. Bax, J. H. C. Reiber, and R. J. van der Geest, "Automated segmentation of myocardial scar in late enhancement MRI using combined intensity and spatial information", *Magnetic Resonance in Medicine*, vol. 64, no. 2, pp. 586-594, 2010.

Review of DE-CMR Segmentation

- Most current methods restrict the segmentation of the myocardium.
 - It simplifies the problem.
 - What if there are misalignments in the myocardial mask?

Method	Myocard. Contours	Scar identification method	Post Processing included in the method
TFA1 [5]	Fixed	2 step thresholding	FP, FN removal by feature analysis, region growing and hole filling
TFA2 [10]	Fixed	2 step thresholding	FP, FN removal by feature analysis
PWD [3]	Fixed	Mixture Model + Watershed	Noise removal + hole filling

[10] LY Hsu, Natanzon, A, P Kellman, GA Hirsch, AH Aletras, and AE Arai, "Quantitative myocardial infarction on delayed enhancement MRI. Part I: Animal validation of an automated feature analysis and combined thresholding infarct sizing algorithm," *Journal of Magnetic Resonance Imaging*, vol. 23, pp. 298-308, 2006.

Review of DE-CMR Segmentation

- Most current methods restrict the segmentation of the myocardium.
 - It simplifies the problem.
 - What if there are misalignments in the myocardial mask?

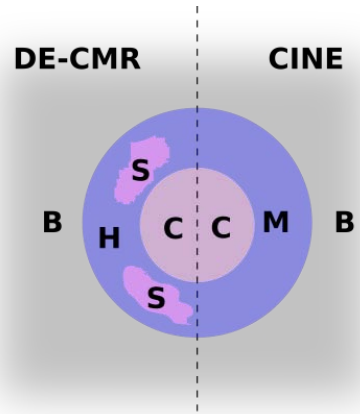
Method	Myocard. Contours	Scar identification method	Post Processing included in the method
TFA1 [5]	Fixed	2 step thresholding	FP, FN removal by feature analysis, region growing and hole filling
TFA2 [10]	Fixed	2 step thresholding	FP, FN removal by feature analysis
PWD [3]	Fixed	Mixture Model + Watershed	Noise removal + hole filling

[3] A. Hennemuth, A. Seeger, O. Friman, S. Miller, B. Klumpp, S. Oeltze, and H.-O. Peitgen, "A Comprehensive Approach to the Analysis of Contrast Enhanced Cardiac MR Images," *IEEE Transactions on Medical Imaging*, vol. 27, no. 11, pp. 1592-1610, 2008.

Objectives

- To propose a segmentation method for DE-CMR that:
 - Is able to modify the myocardial contours if necessary
 - Provides smoothness to the myocardial contours.
 - At the same time, uses the information provided by a CINE segmentation to increase robustness.
- To explore how the state of the art segmentation methods behave:
 - when the myocardial contours have misalignments.
 - with non ischemic cardiopathies .

Variational Framework



$$\mathcal{L} = \{C, H, S, B\}$$

DE-CMR labels

$$\mathcal{A} = \{C, M, B\}$$

CINE labels

- C: Blood cavity
- M: Myocardium
- B: Background

- H: Healthy tissue
- S: Scar

Convex Potts Model (From [7])

$$\min_{\mathbf{u}(\mathbf{x}) \in \Delta_+} \Psi(\mathbf{u}) = \sum_{l \in \mathcal{L}} \int_{\Omega} \left(f_l(\mathbf{x}) u_l(\mathbf{x}) + g_l(\mathbf{x}) |\nabla u_l(\mathbf{x})| \right) d\mathbf{x}$$

Data Fidelity

Contour
Regularization

$$f_{L_i}(\mathbf{x}) = -\ln(P(L_i(\mathbf{x})|I(\mathbf{x})))$$

[7] E. Bae, J. Yuan, and X.-C. Tai, "Global minimization for continuous multiphase partitioning problems using a dual approach," *Int J Comput Vis*, vol. 92, no. 1, pp. 112--129, 2011.

Label Posterior Probability Formulation

$$P(L_i|I) = \frac{\sum_{k=1}^K P(I|A_k, L_i)P(L_i|A_k)P(A_k)}{\sum_{j=1}^L \sum_{k=1}^K P(I|A_k, L_j)P(L_j|A_k)P(A_k)}$$

$P(I|A_k, L_i)$

- Models the probabilistic distribution of $I(x)$.
- Assumption of independence with respect to $\hat{A}(x)$.
- The Rician distribution is chosen for the blood and the myocardial tissues.

$P(A_k)$

- Probability of the CINE label A_k .
- Decays with the distance to the CINE ROI k .
- The binary indicator function for all A_k are smoothed with a Gaussian kernel and normalized.

Label Posterior Probability Formulation

$$P(L_i|I) = \frac{\sum_{k=1}^K P(I|A_k, L_i)P(L_i|A_k)P(A_k)}{\sum_{j=1}^L \sum_{k=1}^K P(I|A_k, L_j)P(L_j|A_k)P(A_k)}$$

$$P(L_i|A_k)$$

- Controls the influence of the CINE segmentation and the image likelihood locally.
- The value at each location is a linear combination of 3 extreme situations:
 - Fully trust the CINE probability
 - Fully trust the a priori tissue probability...
 - ...at the epicardial border
 - ...at the endocardial border
- Weights are computed using the edge information of the DE-CMR image.

Contour Regularization

The regularization local weights $g_i(\mathbf{x})$ depend on the image gradient AND the CINE segmentation:

$$g_C(\mathbf{x}) = \gamma_0 H(1 - a_1(\mathbf{x}), \varepsilon) + \gamma_c r(\mathbf{x}) H(a_1(\mathbf{x}), \varepsilon)$$

$$g_H(\mathbf{x}) = \gamma_0 H(1 - a_2(\mathbf{x}), \varepsilon) + \gamma_t r(\mathbf{x}) H(a_2(\mathbf{x}), \varepsilon)$$

$$g_S(\mathbf{x}) = g_H(\mathbf{x})$$

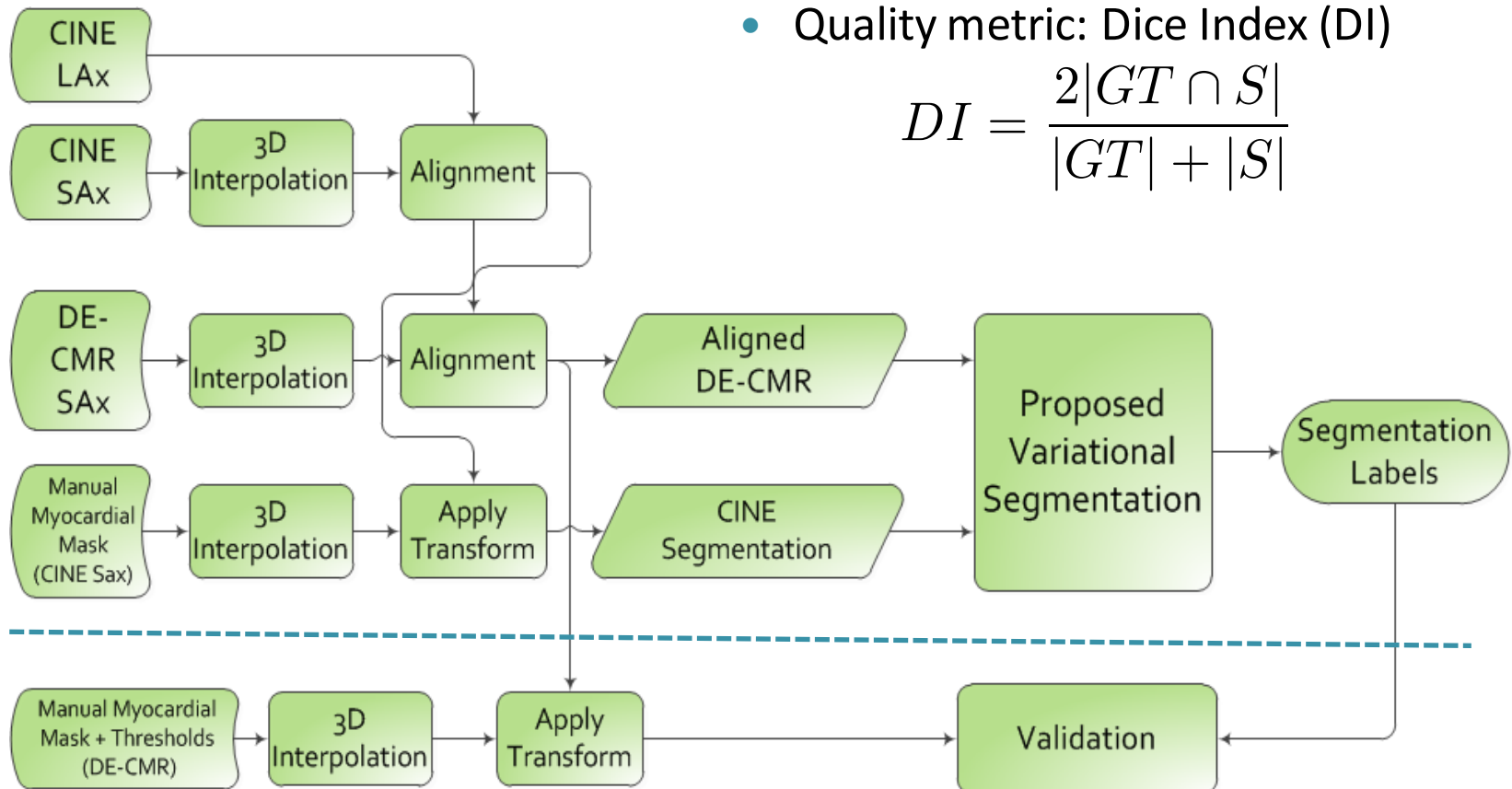
$$g_B(\mathbf{x}) = \gamma_0 H(1 - a_3(\mathbf{x}), \varepsilon) + \gamma_c r(\mathbf{x}) H(a_3(\mathbf{x}), \varepsilon)$$

$$r(\mathbf{x}) = H\left(\left(\frac{3}{2}\right)\sigma_{b(\mathbf{x})} - b(\mathbf{x}), \varepsilon\right)$$

Experimental Setup

- 11 studies from Hypertrophic Cardiomyopathy patients.
- Quality metric: Dice Index (DI)

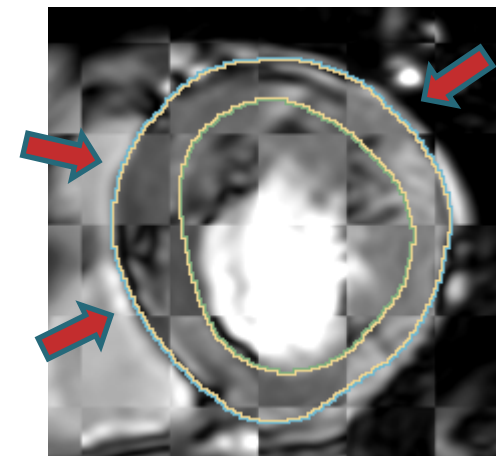
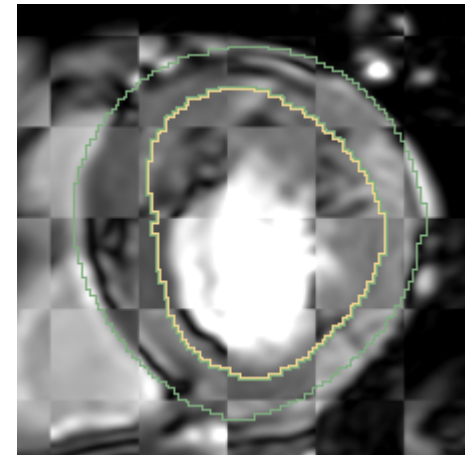
$$DI = \frac{2|GT \cap S|}{|GT| + |S|}$$



Experimental Results

Study	TFA1	TFA2	PWD	PROP
1	0.407	0.638	0.533	0.688
2	0.554	0.595	0.359	0.677
3	0.674	0.569	0.620	0.700
9	0.183	0.213	---	0.207

DI between the scar and healthy tissue ROIs yielded by the considered segmentation methods.



Conclusions

- A variational segmentation method for DE-CMR where:
 - The scar is identified
 - The myocardial contours may be modified.
 - The data fidelity uses a Bayesian approach that takes into account both the image intensity probability distributions and a registered myocardial segmentation coming from CINE.
 - The CINE myocardial segmentation is also used to compute the regularization weights.
- The correction of the myocardial contours improves the scar identification.
- The correction is stronger in the volumes with lower CINE myocardial alignment.



Thank you!