Knee cartilage segmentation using active shape models and contrast enhancement from magnetic resonance images

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ABSTRACT

In this paper, we take advantage from contrast characteristics of our magnetic resonance images improving the performance of Active Shape Models (ASM) applied on knee cartilage segmentation. We perform an image fusion-based contrast enhancement method using time series MRI T2. Then, we apply ASM algorithm and we compare results with ASM without contrast enhancement. The results show that the ASM with contrast enhancement performs better and is consistent. We validate these results using Dice coefficient and Hausdorff distance.

Keywords: Segmentation, Active Shape Models, Contrast Enhancement, Gaussian and Laplacian Pyramids, Image Fusion.

1. INTRODUCTION

Knee Osteoarthritis (OA) [1] is a disease caused by biomechanical stress that affects the articular cartilage and bones of the knee. This condition causes pain and malfunction. OA may be present in any of the medial femoral compartments, either the tibiofemoral or patelofemoral, according to the location of damaged cartilage. Diagnosis of Knee OA is done since the first clinical examination. Pain, morning stiffness and knee swelling in a patient older than 50 years are considered as consequences of OA. However, image analysis of the knee also has an important role since it can confirm the OA diagnosis, determines compartments involved and evaluates the disease stage. Moreover, it can confirm the responsibility of OA in the symptoms and provides information about disease evolution during treatment. Magnetic Resonance Imaging (MRI) [2] provides a non-invasive assessment for evaluating the presence and progression of the Knee OA [3]. Magnetic resonance can show the soft tissue structures and their boundaries with the bones, without significant distortion. Furthermore, MRI does not change the tissue's dimension and there is not superposition between anatomical structures and, more important, it directly visualizes the knee cartilage and its defects [4].

Automatic or semiautomatic knee segmentation has been studied for more than 20 years and many approaches have been reported, for example: growing regions [5-7], Bezier splines [8], active contours [9, 10], Bayesian classifiers [11] and active shape models (ASM) [12, 13]. ASM [14] compute an average shape of the object to segment from a training set and a statistical model of minimal parameters that allow the shape to adjust to different objects within a certain range. ASM have been widely used in medical image analysis because there is sufficient knowledge about the shape of targeted anatomical objects obtained through diverse medical imaging modalities.

The purpose of our work is to take advantage of the contrast characteristics of Resonance Magnetic Images (volume acquisitions in time series) in order to enhance the contrast in the images and therefore, improve the ASM segmentation performance in knee cartilage. We test the contrast enhancement method on real data; this is new because, in the original paper [15], only synthetic images are presented.

2. METHODS AND MATERIALS

In this section, the main characteristics of MRI, segmentation and contrast enhancement algorithms are described. Validation metrics of segmentation and contrast enhancement are also discussed.

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2.1 Medical Images

The images were selected from 16 different volume sets on a 3T MRI system (Siemens Verio). In each set, there are 11 knee volumes and, in each volume, there are 10 images. The images are in T2* in an echo-time series (Multi-echo system). Every 7.54 [ms] a new knee volume is acquired by the sequence is called *Carr-Purcell-Meiboom-Gill* [20]. An important characteristic of this sequence is the gradual contrast decrease along the volume acquisition time (Figure 1). The images were acquired with a 384 x 384 matrix (0.414-mm in-plane resolution and 3.6-mm slice thickness).

2.2 Active Shape Models

The original algorithm of Active Shape Models (ASM) consists of a statistical shape model derived from a Point Distribution Model (PDM) obtained through a set of landmarks on the shape contours. The PDM is used for modeling the shape of an object and its variations.

First, the shapes are aligned using Procrustes Analysis (PA), PA is performed by optimally translating, rotating and uniformly scaling the objects. In other words, both the space location and the size of the objects are freely adjusted. The aim is to obtain a similar location and size, by minimizing a shape difference measure called *Procrustes distance* between two shapes.

The PDM is built applying Principal Component Analysis (PCA) to the aligned set (by PA) of shapes represented by landmarks. The original shapes X_i and their representation model b_i (i = 1, ..., N), N is the number of variation modes (described below), are related by the mean shape \overline{X} and the eigenvectors P:

$$b_i = P^T (X_i - \bar{X}), \qquad X_i = \bar{X} + P b_i \tag{1}$$

We can reduce the representation dimensionality using only the eigenvectors of the major eigenvalues. In this case, equation (1) turns into an approximation, with an error depending on the magnitude of the excluded eigenvalues. Therefore, assuming the data is Gaussian distributed, each component of the vectors b_i is limited to guarantee only plausible shapes are represented:

$$|b_i^m| \le A_{\sqrt{\lambda_m}}, \qquad 1 \le i \le N, \quad 1 \le m \le M,$$
(2)

where A is regularization constant, usually between 1 and 3 (it depends on desired flexibility of model), M is the number of used eigenvectors and λ_m are the eigenvalues of the covariance matrix.

The appearance model is built from a set of Gray Local Profiles (GLP) in the normal direction to the contour at each landmark. The GLPs are built from normalized first order derivatives and describe the local intensity variations at each landmark positions. The profile equation is:

$$y_{si} = \frac{ds_i}{\sum_{0}^{n_s - 2} |ds_{ik}|},$$
(3)

where the profile is $s_i = [s_{i0} \ s_{i1} \dots \ s_{i(n_s-1)}]$ and n_s is the size of the profile.

The next step is to fit the model on interest region. An instance of the model is given by the adaptation of shape X_i with respect to reference shape X_r :

$$\boldsymbol{X}_{i} = \boldsymbol{M}(\boldsymbol{s}_{i}, \boldsymbol{\theta}_{i})[\boldsymbol{X}_{r}] + \boldsymbol{t}_{i}, \tag{5}$$

where $M(s, \theta)$ is a rotation matrix with angle θ and scale s, t_i is the translation and X_r is the combination of \overline{X} and the model parameters:

$$\boldsymbol{X}_r = \boldsymbol{\overline{X}} + \boldsymbol{P}\boldsymbol{b} \tag{6}$$

Combining last equation with (5), the adaptation results in the next equation:

$$\boldsymbol{X}_{i} = \boldsymbol{M}(\boldsymbol{s}_{i}, \boldsymbol{\theta}_{i})[\boldsymbol{\overline{X}} + \boldsymbol{P}\boldsymbol{b}] + \boldsymbol{t}_{i}$$

$$\tag{7}$$

The position parameters are given by t, s and θ and the model parameters are represented by b. The position parameters represent pose modifications and model parameters represent shape modifications.

The adaptation pretends mapping the model X to X + dX. The position parameters are adjusted and model parameters are modified from the deformed version of the shape. The term dX can be written as:

$$dX = (dX_{pos}, dX_{shp}) \tag{8}$$

Where $dX_{pos} = (\mathbf{1} + d_s, d_{\theta}, d_t)$ and $dX_{shp} = P^T d_b$.

The influence of the position parameters on the shape X_i is:

$$\boldsymbol{X}_{align} = M(\boldsymbol{s}_i(1+d_s), \boldsymbol{\theta}_i + d_{\boldsymbol{\theta}})[\boldsymbol{X}_r] + \boldsymbol{t}_i + d_{\boldsymbol{t}}$$
(9)

Where the terms $1 + d_s$, $\theta_i + d_\theta$ and $t_i + d_t$ are obtained minimizing the equation:

$$E_j = (\mathbf{X}_i - \mathbf{M}(s_j, \theta_j) [\mathbf{X}_j] - \mathbf{t}_i)^T \mathbf{W} (\mathbf{X}_i - \mathbf{M}(s_j, \theta_j) [\mathbf{X}_j] - \mathbf{t}_i),$$
(10)

where W is the identity matrix aimed at giving the same weight at each point. The influence of model parameters only modifies X_r in the equation (9), hence $X_r \to X_r + dX_{shp}$.

The distance is dX_{shp} is:

$$d\boldsymbol{X}_{shp} = P^T d_b = \boldsymbol{M} \left(\left(s_i (1+d_s) \right)^{-1}, -\theta_i - d_\theta \right) \left[\boldsymbol{M}(s_i, \theta_i) \boldsymbol{X}_r + d\boldsymbol{X} - d_t \right] - \boldsymbol{X}_r$$
(11)

From the last equation, the new value of $b \rightarrow b + db$ is verified in equation (2). And, if the condition is valid, the next parameters are updated:

$$t_{x} + dt_{x} \rightarrow t_{x}$$

$$t_{y} + dt_{y} \rightarrow t_{y}$$

$$\theta + d_{\theta} \rightarrow \theta$$

$$s(1 + d_{s}) \rightarrow s$$

$$b + db \rightarrow b$$
(12)

The process is repeated by *n* iterations, either predefined or controlled by a stop criterion.

2.3 Image Fusion-based Contrast Enhancement

We enhance contrast using a method that exploits the image characteristics, i.e.: contrast and luminance in time series. The method is called *Image fusion-based Contrast Enhancement* [15], this method takes advantage of these two characteristics and, in a multiresolution scheme, fuses both image characteristics getting the contrast enhancement. The algorithm can be summarized in the next steps:

Step 1: Calculate the image quality measures for each of the input images:

Contrast:

$$C_{i,j,k} = |\nabla I| = \sqrt{(I(x,y) - I(x+1,y))^2 + (I(x,y) - I(x,y+1))^2},$$
(13)

given an input image I(x, y) where x and y are the row and column coordinates, respectively.

Luminance:

$$L_{i,j,k} = L(I(x,y); m_o, \sigma) = exp\left(-\frac{(I(x,y)-m_0)^2}{2\sigma^2}\right),$$
(14)

where σ proposed by authors is 0.2 (giving more weight to pixels which value is closer to the mean) and m_0 is the mean intensity of the original image.

Step 2: For each image obtain the scalar weight map (15) and the normalized scalar weight map using (16).

$$P_{i,j,k} = C_{i,j,k}{}^{\alpha} L_{i,j,k}{}^{\beta} \tag{15}$$

$$P_{i,j,k} = \left[\sum_{k=1}^{N} P_{i,j,k2}\right]^{-1}$$
(16)

where k = 1, 2, ..., N and N is the number of input images. The weight of contrast and luminance of the metric P uses a power function, where α and β are the corresponding weighting exponents (α , $\beta > 0$), that increase or decrease the influence of quality measures.

Step 3: Decompose the input images using a Laplacian pyramid.

Step 4: Obtain the fused pyramid as a weighted average of the original Laplacian decompositions for each level l, with the *l*-th level of Gaussian pyramid of the weight map (16) serving as the weights. The fused pyramid is:

$$L\{F\}_{i,j}^{l} = \sum_{k=1}^{N} G\{\widehat{W}\}_{i,j,k}^{l} L\{I\}_{i,j,k}^{l}$$
(17)

Step 5: Reconstruct the image from the fused Laplacian pyramid.

2.4 Validation

Validation (quantitative evaluation) aims at measuring the algorithm performance. In our particular case, we wish to measure the accuracy of the segmentation algorithm and the fusion-based contrast enhancement performance. An output image has been enhanced over the input image if it enables a better perception of image details. A contrast enhancement is a difficult task because contrast quantification is hard. Thus, it is necessary to establish measures for defining good enhancement. We use *Absolute Mean Brightness Error* (AMBE) [16], *Discrete Entropy* (DE) [17], and *Edge-Based Contrast Measure* (EBCM) [18] as quantitative contrast measures, and *Dice Coefficient* and *Hausdorff Distance* for evaluating segmentation.

2.4.1 Dice Coefficient

The DICE coefficient is a similarity measurement between sets. In this case, it is used as a global measure of the existing differences between the calculated shape and the reference shape. The coefficient is given by:

$$DICE = \frac{2(A \cap B)}{|A| + |B|},\tag{18}$$

where A is the area inside calculated shape and B is the area inside reference shape. DICE coefficient values are between 0 and 1; the closer the coefficient to 1, the more similar the shapes are.

2.4.2 Hausdorff Distance

The Hausdorff distance determines the similarity between two shapes [19]. The Hausdorff distance is defined as the maximum distance from a point set to the nearest point of other set points. If $A = \{a_1, ..., a_m\}$ and $B = \{b_1, ..., b_n\}$ are two point sets, the Hausdorff distance between A and B is:

$$H(A,B) = \max(h(A,B), h(B,A)),$$
 (19)

where

$$h(A,B) = \max_{a \in A} \min_{b \in B} ||a - b||$$
(20)

Equation (20) is properly called the direct Hausdorff distance from the set A to set B, using the underlying norm $\|\cdot\|$ on the point sets A and B, and

$$h(B,A) = \max_{b \in B} \min_{a \in A} ||b - a||$$
(21)

is called inverse Hausdorff distance. The Hausdorff distance is not symmetric. Then, equation (19) considers both cases giving a more general definition.

2.4.3 Absolute Mean Brightness Error

AMBE is the absolute difference between the Mean Brightness (MB) values of an input image *X* and an output image, for example:

$$AMBE(X,Y) = |MB(X) - MB(Y)|.$$
⁽²²⁾

Where MB(X) and MB(Y) are the MB values of X and Y, respectively. The lower AMBE value, the better brightness is preserved.

2.4.4 Discrete Entropy

The DE of an image measures its content, where a higher value indicates an image with richer details. The DE is defined as:

$$DE(X) = \sum_{\forall i} p(x_i) \log(p(x_i))$$
(23)

Where $p(x_i)$ is the probability of the pixel intensity, and it is calculated from the normalized histogram.

2.4.5 Edge-Based Contrast Measure

The EBCM is based on the human perception mechanism; it is very sensitive to contours (or edges). The gray level in the object edges is calculated by computing the mean value of the pixel gray levels weighted by their edge values. The contrast c(i, j) for a pixel of image X located at (i, j) is thus defined as:

$$c(i,j) = \frac{|x(i,j)-e(i,j)|}{|x(i,j)+e(i,j)|}.$$
(24)

Where the mean edge gray level is:

$$e(i,j) = \sum_{(k,l)\in\mathcal{N}(i,j)} g(k,l) x(k,l) / \sum_{(k,l)\in\mathcal{N}(i,j)} g(k,l)$$

$$\tag{25}$$

 $\mathcal{N}(i, j)$ is the set of all neighboring pixels of pixel (i, j), and g(k, l) is the edge value at pixel (k, l). The magnitude of the image gradient is calculated using the Laplacian operator. The EBCM for image X is thus computed as the average contrast value, for example:

$$EBCM(X) = \sum_{i=1}^{H} \sum_{j=1}^{W} c(i,j)/HW$$
(26)

Where *HW* is the total number of pixels. Then, for an output image *Y* of an input image *X*, the contrast is improved when $EBCM(Y) \ge EBCM(X)$.

3. RESULTS

This section shows the results of contrast enhancement and the goal is to find the best fusion option on images. Results validation is based on contrast measures described on previous section. Later, taking the images with the best contrast, we test segmentation algorithm on femur cartilage. Medical images are in DICOM format.

3.1 Contrast enhancement results

As mentioned above, we have 11 images of the same slice in each of the 16 selected sets. We selected the slice of the fourth volume because it presents the best contrast between bone and cartilage perceptually and, with this image, we compare the contrast enhancement results. In Figure 1, slices from different volumes are showed and it is noticeable the contrast characteristics we want to use.



Figure 1(a) First-volume slice (bright). Figure 1(b) Fourth-volume slice (good contrast). Figure 1(c) Sixth-volume slice (dark).

The contrast enhancement algorithm is based on image fusion, and then we fuse the fourth volume slice with the rest of corresponding volumes slices (brighter and darker slices). But, in the results graphs, we only show fusion results with slices of the fifth, sixth, seventh, eighth and ninth volumes because they are more representative. Fused images are evaluated by the following measures: AMBE, DE, EBCM and results are shown in Figure 2. Table 1 shows average results of each fusion.

3.2 Segmentation results

We chose fusing fourth and sixth volume slices (justified on Discussion section). Then, we compare the results with ASM segmentation on slices of the fourth volume (Original) and the fused images. Since the ASM require prior training, we used the "leave-one-out" method to maximize data usage. The Dice coefficient and Hausdorff distance are the metrics used for validating segmentation results (Table 2). Moreover, the best and the worst segmentation result obtained on the enhancement contrast images are shown in Figure 3, Image 10 and Image 1 respectively.

The average of Dice coefficients calculated on the original image is (SD) 0.7586 ± 0.1169 and, on the fusion image is 0.8155 ± 0.052 , and the average of Hausdorff distance calculated on in the original image is 3.3803 ± 1.2336 and, on the fusion images is 2.9158 ± 0.6735 . Each segmentation is achieved in 20 seconds approximately.



(c) Figure 2(a) AMBE Results. Figure 2(b) DE Results. Figure 2(c) EBCM Results. Results on the original image are shown in blue and the best fusion (4-6 fusion) results are in red.

Image	AMBE avg.	DE avg.	EBCM avg.
Original	-	6.2427	0.1956
4-5 Fusion	600.4813	6.0476	0.2347
4-6 Fusion	689.0589	6.0174	0.2626
4-7 Fusion	731.8271	5.0883	0.2776
4-8 Fusion	764.6651	5.9662	0.2851
4-9 Fusion	785.4851	5.9491	0.2899

Table 1. Average results of AMBE, DE and EBCM for each fusion

Image	Dice_Original	Dice_Fusion	Haus_Original	Haus_Fus
Image 1	0.6760	0.7077	2.4755	2.9222
Image 2	0.7345	0.7814	3.5323	3.3891
Image 3	0.8063	0.8207	2.7598	2.0835
Image 4	0.7623	0.7797	3.1228	2.9954
Image 5	0.8220	0.8908	3.1798	2.6852
Image 6	0.7442	0.8468	2.2151	1.6756
Image 7	0.4994	0.7778	4.9496	2.5941
Image 8	0.7643	0.7639	2.9259	2.7121
Image 9	0.4848	0.8452	6.7318	3.0975
Image 10	0.8621	0.8983	3.1684	2.9625
Image 11	0.8245	0.8241	2.9469	2.8717
Image 12	0.7750	0.8256	5.2034	5.2466
Image 13	0.8748	0.7915	2.0863	2.2911
Image 14	0.8407	0.7724	3.4308	3.5774
Image 15	0.8006	0.8367	2.7630	2.9529
Image 16	0.8671	0.8863	2.5940	2.5955

Table 2. For each image, Dice coefficients of ASM with original image and ASM with fusioned image and Hausdorff distances of ASM with original image and ASM with fusioned image.





Figure 3(a) the best segmentation result is on Image 10 and Figure 3(b) the worst segmentation result is on Image 1

4. **DISCUSSION**

Based on the segmentation results, we note that contrast enhancement improves, in most cases, the segmentation results of knee cartilage. Furthermore, we proved ASM segmentation performs better with contrast enhancement images, since it depends mainly on gray levels and their distribution profiles. The average results of Dice coefficient and Hausdorff distance are improved with the contrast enhancement and their variance also decreases in comparison with segmentation on original images. However, in some patients with advanced Knee OA and, who have no cartilage left, is not substantial since there is little to segment. This is a serious problem for the ASM algorithm because there are not gray level variations, so the algorithm. Fortunately, ASM always provides plausible shapes and landmarks are correlated between themselves. Therefore, the failures are not so serious but.

From a purely perceptual opinion, we chose the 4-6 fusion. However, using the results of AMBE, DE and ECBM, we conclude the following: The average ECBM of fused image is greater than the average ECBM of the original images and, although subsequent fusions shown better ECBM, the difference between 4-5 fusion and 4-6 fusion is high and the difference between 4-6 fusion and 4-7 fusion has a significant decay. Furthermore, the greater is the number of the

fusion image, the worse are the AMBE and Entropy results. If AMBE increases and Entropy decreases means quality image is affected. In the particular case of entropy, if it decreases, the image loses details and sharpness decays. For these reasons, we chose 4-6 fused images, because they have equilibrium on AMBE, DE and ECBM metrics. The contrast enhancement technique fits perfectly with the characteristics of our images.

5. CONCLUSION AND FUTURE WORK

In this paper, we have used known methods: ASM and Fusion-based Contrast Enhancement, but they had not been tested together before.

The contrast enhancement method presents good results in [15] but, this work did not evaluate its test results on a real application or on real data. Our images are very well suited to test the proposed contrast enhancement method: they belong to the same data set but each one has a different contrast. This characteristic totally depends on the pulse sequence used during image acquisition. Commonly, knee cartilage is acquired in different sequences, mainly T1, but, using the former sequence, we pretend to find something additional to just the cartilage size or thickness. Additionally, we also carried out a relaxometry study.

Relaxometry characterizes a tissue depending on the relaxation time T2, by fitting an exponential curve. This time is very important for diagnosis OA because an elevated T2 time suggests biomechanical changes on cartilage (deterioration) suggesting that the patient is likely to develop this disease [21].

The segmentation results show that contrast enhancement improves segmentation performance, proving our hypothesis. A better contrast implies a gradient increment, thus a better region differentiation. The knee cartilage segmentation is necessary because we want to estimate T2 time over this specific zone. Our results are good and useful, however, there are a lot of methods that improve ASM performance, and some of them use descriptors and transforms. We could compare results in order to select the best in a future work.

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