

ORIGINAL ARTICLE

The use of an active appearance model for automated prostate segmentation in magnetic resonance

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Abstract

Background. The prostate gland is delineated as the clinical target volume (CTV) in treatment planning of prostate cancer. Therefore, an accurate delineation is a prerequisite for efficient treatment. Accurate automated prostate segmentation methods facilitate the delineation of the CTV without inter-observer variation. The purpose of this study is to present an automated three-dimensional (3D) segmentation of the prostate using an active appearance model. **Material and methods.** Axial T2-weighted magnetic resonance (MR) scans were used to build the active appearance model. The model was based on a principal component analysis of shape and texture features with a level-set representation of the prostate shape instead of the selection of landmarks in the traditional active appearance model. To achieve a better fit of the model to the target image, prior knowledge to predict how to correct the model and pose parameters was incorporated. The segmentation was performed as an iterative algorithm to minimize the squared difference between the target and the model image. **Results.** The model was trained using manual delineations from 30 patients and was validated using leave-one-out cross validation where the automated segmentations were compared with the manual reference delineations. The mean and median dice similarity coefficient was 0.84 and 0.86, respectively. **Conclusion.** This study demonstrated the feasibility for an automated prostate segmentation using an active appearance with results comparable to other studies.

Magnetic resonance (MR) is increasingly used for target delineation in treatment of prostate cancer using radiotherapy because of the superior soft-tissue visualization compared with computed tomography (CT). Studies by Hentschel et al. [1] and Rasch et al. [2] have investigated target delineation in CT and MR for treatment of prostate cancer. They found that using MR instead of CT can lead to a volume reduction of 35–40%, respectively, with a lower inter- and intra-observer variance [1,2]. The overestimation of the prostate volume can cause higher normal tissue toxicity and limit the radiation dose to the prostate.

The clinical practice for target delineation for planning radiotherapy of prostate cancer is today widely performed using a manual contouring of the prostate slice-by-slice using either the axial, coronal, sagittal view or a combination of the views. The manual work is a labor-intensive task and is prone to inter- and intra-observer variance. Therefore, several authors have described semi- or automated methods

for prostate segmentation in MR. Some studies, e.g. [3,4] have used voxel-based segmentations yet while incorporating knowledge about the general prostate shape. However, most previous studies have focused on either atlas- or model-based methods. Atlas-based approaches in three dimensions (3D) have been studied [5–7] while a model-based approach was presented by Pasquier et al. [8] using a landmark-based shape model followed by a contour optimization to deform the model to the prostate. Statistical shape models were presented in [9,10]. Chandra et al. [11] used a deformable model consisting of triangulated surface and image feature model trained specific to the patient's MR scan. Chowdhury et al. [12] used linked statistical shape models using information from both MR and CT in a level-set shape representation.

A few methods use a combination of prostate shape and intensity information to model and segment the prostate. Ghose et al. [13] used active

appearance models (AAM) propagated by the approximation coefficients of Haar wavelet transform in 2D. Recently, Toth and Madabhushi [14] used image-derived features in an AAM where the selection of landmarks were replaced with a level-set representation of the shape. The original AAM suggested by Cootes et al. [15,16] requires that landmarks at the border of the object are selected, which is prone to error and time-consuming. Therefore, recent work has focused on a re-formulation of the shape modeling in a level-set framework of multi-modality MR images of the lateral ventricles in the brain [17] and segmentation of the vertebrae in CT [18].

In this paper we propose a method that combines a level-set shape model and a texture model for prostate segmentation. The level-set representation eliminates the need for selecting corresponding landmarks in all MR scans. In addition prior knowledge learned in the training of the model is incorporated to adjust the model parameter and guide the 3D segmentation.

Material and methods

Materials

The data consisted of T2-weighted 1.5 Tesla MR scans ($0.5469 \times 0.5469 \times 3 \text{ mm}^3$) from 30 patients. The patients had all been diagnosed with local or locally advanced prostate cancer and referred to the Department of Oncology for radiotherapy and enrolled in a prior study investigating the feasibility using a Ni-Ti prostate stent described in [19]. Each patient had a MR scan with a manual delineation of the prostate performed for the planning of the radiotherapy. The prostate size for the 30 patients calculated from the manual delineations was found to range from 17.42 cm^3 to 92.94 cm^3 .

Active Appearance Model

The assumption behind AAM is that changes in the shape and the texture of a given object are correlated. The AAM presented here is a principal component analysis (PCA)-based statistical model where a shape and a texture model are combined. In this way, the statistical model describes both the object shape and the object texture with the same set of parameters. The elimination of the need to select landmarks on the prostate border was achieved using level-set in which the prostate shape was represented as a signed distance map. The methodology is derived from Hu and Collins [17], Stephansen [18], and Cootes and Taylor [20]. For further details see the Supplementary Appendix (available online at <http://informahealthcare.com/doi/abs/10.3109/0284186X.2013.822099>). New images and shapes were synthesized

using the AAM for the shape ϕ and the image intensities g by varying the appearance parameters c :

$$\begin{bmatrix} \phi \\ g \end{bmatrix} = \begin{bmatrix} \hat{\phi} \\ \hat{g} \end{bmatrix} + \begin{bmatrix} P_s W_s^{-1} P_{cs} \\ P_g P_{cg} \end{bmatrix} c \quad (1)$$

Where $\hat{\phi}$ and $\hat{g}$ are the mean shape and mean texture, respectively. P_s is the principal modes of shape variation, W_s is the weight matrix to account for difference in units between shape and texture. The shape and texture parameters were combined into the appearance parameters c . P_{cs} and P_{cg} represent the shape and texture part of the principal modes of appearance variation, respectively. The appearance parameters c values were constrained to the interval of ± 3 standard deviations as in [16].

Appearance model-based segmentation

The aim of the segmentation is to find the model parameters of the AAM that generate a model image that closely corresponds to the target. The matching of the model image to the target image is an optimization problem where the appearance parameters were varied to minimize the squared difference between the images. The strategy was suggested by Cootes and Taylor [20] in which the spatial pattern of the difference encodes knowledge about how to adjust the model parameters to estimate the prior knowledge before segmentation of the target image. For each step size tried in the segmentation, new parameter values were computed from the estimated adjustments and each of the AAM parameters were constrained to be within ± 3 standard deviations to ensure that all possible shapes were generated.

Validation method

The validation of AAM was performed as a leave-one-out cross-validation of all 30 MR data sets. The automated segmentations were compared with the manual reference segmentations both by a visual comparison and by the similarity measures Dice similarity coefficient (DSC) [21], the mean surface distance and the largest distance between the manual and the automated segmentation. The sensitivity and specificity were determined to provide an indication of a possible trend in under- or over-segmentation of the automated method compared with the manual reference delineations. The sensitivity and specificity were also determined based on the overlap (false-positive and false-negative) between the segmentations and a high value was therefore desirable. All measures were calculated for each image and then averaged to get one value and standard deviation for each similarity measure.

Table I. Results from the comparison of the automated segmentation and the manual reference delineation.

Similarity measure	Mean $\pm$ std	Median $\pm$ std	Range
DSC	0.84 ± 0.07	0.86 ± 0.07	0.59–0.94
Sensitivity	0.99 ± 0.02	0.99 ± 0.02	0.88–1.00
Specificity	0.93 ± 0.08	0.96 ± 0.08	0.67–1.00
Mean surface distance [mm]	1.39 ± 1.00	1.12 ± 1.00	0.43–5.38
Largest distance [mm]	7.01 ± 2.31	6.62 ± 2.31	3.59–11.59

Results

The appearance model was trained and tested on 30 MR images of the prostate in a leave-one-out cross validation. Table I shows the similarity measures from the automated segmentation compared with the manual reference delineation.

Figure 1 shows the automated prostate segmentation compared with the manual delineations and the average model. It can be seen that the automated segmentation appears more as a smoother contour but also tend to be smaller than the manual delineation.

Figure 2 shows the 3D rendering of the manual (left) and the automated (right) segmentation. It can be seen that a segmentation based on slice-based delineation appeared layered while the 3D segmentation appeared smooth over the volume.

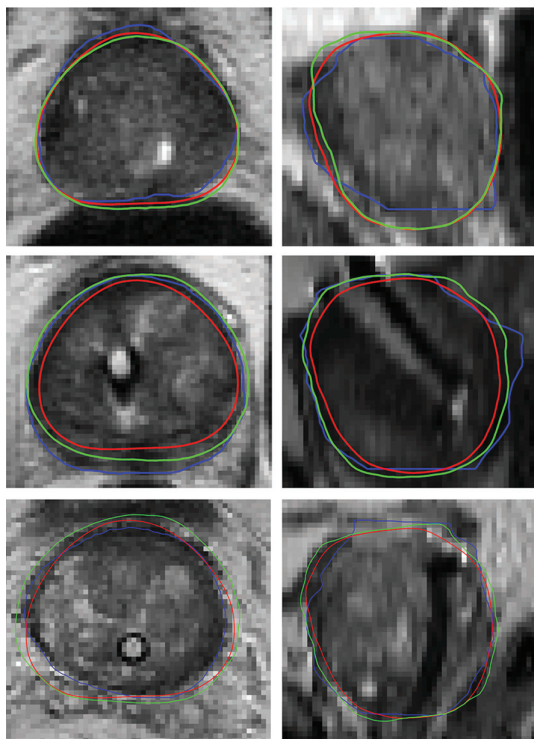


Figure 1. Examples of the prostate segmentation in the axial plane (left) and the sagittal plane (right) from three patients. Blue: Manual reference segmentation, Green: Automated segmentation, Red: Average model. The DSC for the three segmentations was 0.94 (top), 0.79 (middle), 0.89 (bottom).

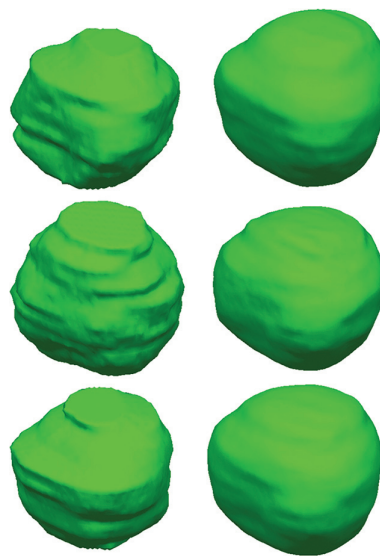


Figure 2. Examples of the prostate segmentation. Left: The reference manual segmentation. Right: Automated 3D segmentation.

Both the similarity measures and the visual comparison indicated that the outliers in the DSC occurred when the model failed to find the correct size of the prostate with the largest volume. This was supported by the sensitivity and specificity of the method indicating cases of under-segmentation.

Discussion

An automated prostate segmentation method has been presented for planning prostate cancer radiotherapy to potentially reduce the inter-observer variation in the manual delineation. The prostate segmentation was performed using MR as it is increasingly used for prostate delineation because of the superior soft-tissue visualization compared with CT. An accurate and fast automated segmentation of the prostate could be used to track the prostate before treatment and could decrease the time used for clinical target delineation in the planning of radiotherapy. This would require that a clinical validation of the automated method was performed.

The method was based on the AAM originally suggested by Cootes et al. [15] and reformulated by Hu and Collins [17] with a level-set representation of the shape by which the need for selecting landmarks is eliminated. Prior knowledge about how to guide the parameter optimization in the segmentation algorithm was incorporated in the training of the model. The iterative segmentation proposed by Cootes et al. [15] to allow for pose changes during segmentation was extended to a 3D segmentation.

Other studies report similar results as obtained here with a mean and median DSC of 0.84 and 0.86, respectively. The range of mean DSC from the previous studies is 0.84–0.88 [8–10,13,14] and the range

for the median DSC is 0.85–0.86 [5,9]. The sensitivity of the study by Martin et al. [10] was 0.86 compared with ours of 0.99. However, the numbers are difficult to compare as a different imaging protocol have been used. For example, Toth and Madabhushi [14] used MR images acquired using an endorectal coil, which makes the prostate boundary easier to detect.

Outliers in the results are cases in which the algorithm fails to determine the correct scale, which is the case for the segmentation with the lowest DSC but the largest volume. Here the model fails to represent the structure, because it is not sufficiently represented in the training set. Visually, the method performed best in the central part of the prostate, which could be explained by the generally better image support here.

There was a trend in the present study that using the automated segmentation method compared with the manual segmentation resulted in an under-segmentation of the prostate. The manual prostate delineations originally used as the clinical target volume were used for training the AAM and the validation in the lack of a true gold standard, which can explain some of the errors in the automated segmentation. The inter-observer variation of the manual delineations was not validated. The 3D rendering of the manual and the automated segmentation illustrates that a segmentation based on slice contours will appear layered in the sagittal and coronal plane. The 3D segmentation appears in contrast as a smooth segmentation in all image planes.

A fully automated 3D prostate segmentation using an AAM was developed and validated for 30 patients to reduce inter-observer variation in the target delineation. The performance of the method was comparable to other studies with a dice similarity coefficient of 0.84 and showed the feasibility for automated prostate segmentation in MR using active appearance models.

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Supplementary material available online

Supplementary Appendix and Figure A1.