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Medical Image Segmentation by Combining Graph Cut and Oriented Active Appearance Models

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Abstract

In this paper, we propose a novel 3D segmentation method based on the effective combination of the active appearance model (*AAM*), live wire (*LW*), and graph cut (*GC*). The proposed method consists of three main parts: model building, initialization, and segmentation. In the model building part, we construct the *AAM* and train the *LW* cost function and *GC* parameters. In the initialization part, a novel algorithm is proposed for improving the conventional *AAM* matching method, which effectively combines the *AAM* and *LW* method, resulting in Oriented *AAM* (*OAAM*). A multi-object strategy is utilized to help in object initialization. We employ a pseudo-3D initialization strategy, and segment the organs slice by slice via multi-object *OAAM* method. For the segmentation part, a 3D shape constrained *GC* method is proposed. The object shape generated from the initialization step is integrated into the *GC* cost computation, and an iterative *GC-OAAM* method is used for object delineation. The proposed method was tested in segmenting the liver, kidneys, and spleen on a clinical *CT* dataset and also tested on the MICCAI 2007 grand challenge for liver segmentation training dataset. The results show the following: (a) An overall segmentation accuracy of true positive volume fraction (*TPVF*) > 94.3%, false positive volume fraction (*FPVF*) < 0.2% can be achieved. (b) The initialization performance can be improved by combining *AAM* and *LW*. (c) The multi-object strategy greatly facilitates the initialization. (d) Compared to the traditional 3D *AAM* method, the pseudo 3D *OAAM* method achieves comparable performance while running 12 times faster. (e) The performance of proposed method is comparable to the state of the art liver segmentation algorithm. The executable version of 3D shape constrained *GC* with user interface can be downloaded from website <http://xinjianchen.wordpress.com/research/>.

Keywords

Object Segmentation; Active Appearance Models; Live Wire; Graph Cut

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I. INTRODUCTION

Image segmentation is a fundamental and challenging problem in computer vision and medical image analysis. In spite of several decades of research and many key advances, a few challenges still remain in this area. Efficient, robust, and automatic segmentation of anatomy on radiological images is one of these challenges.

The image segmentation methods could be classified into several types: image based [1–12], model based [13–30], and hybrid methods [31–39]. Purely image based methods perform segmentation based only on image information; these include thresholding, region growing [1], morphological operations [2], active contours [3, 4, 22], level sets [5], live wire (*LW*) [6], watershed [7], fuzzy connectedness [8, 9] and graph cut (*GC*) [10, 11, 47]. These methods perform well on high quality images. However, the results are not as good when the image quality is inferior or boundary information is missing. In recent years, there has been an increasing interest in model-based segmentation methods. One advantage of these methods is that, even when some boundary information is missing, such gaps can be filled due to the closure and connectedness properties of the model. The model-based methods employ object population shape and appearance prior such as atlases [13–17, 23–25], statistical active shape model [18–20, 26], and statistical active appearance models (*AAMs*) [21, 27, 28]. *MICCAI2007* “Grand Challenge” workshop [29] organized a competition for liver segmentation which attracted a lot of attention. In that competition, the three best-rated approaches [29, 30] were all based on statistical shape models with some form of additional deformation. Such hybrid approaches are rightfully attracting a great deal of attention at present. The relative merits of the synergy that exists between these two approaches – purely image-based and model-based strategies – are clearly emerging in the segmentation field. As such hybrid methods that form a combination of two or more approaches are emerging as powerful segmentation tools [31–39] where their superior performance and robustness over each of the component methods have been well demonstrated.

Many of the above mentioned image-based [1, 4], model-based [14–18, 20–25, 30], as well as hybrid [39] techniques were tailored for specific body regions and image modalities. However, it is desirable to have a general approach that is applicable to any (or most) body regions and image modalities and protocols and not heavily dependent on the characteristics of fixed shape families and image modalities. While perhaps some of the above techniques can be generalized in this spirit, few methods have demonstrated to work in this general setting.

In this paper, we propose a general method which can be used to segment most organs and which effectively combines the *AAM*, *LW*, and *GC* methods, leading to the *GC-OAAM* approach, and arrive at an automatic, efficient, and accurate segmentation method. *LW* is a user-steered 2-dimensional segmentation method in which the user provides recognition help and the algorithm does the delineation precisely. The major limitation of live wire is that the recognition process (selecting anchor points on the boundary) is done by a human operator; hence it is far less efficient. *AAM* methods use landmarks to represent shape and appearance, and use principal component analysis to capture the major modes of variation in shape and appearance observed in the training data sets. However, the specific shape and

appearance information are also lost during the model building. *GC* methods have the ability to compute globally optimal solutions and have proven to be a useful multidimensional optimization tool which can enforce piecewise smoothness while preserving relevant sharp discontinuities. However, some *GC* methods need human operator to label the source and sink seeds. In this paper, our aim is to combine the complementary strengths of the individual methods to arrive at a more powerful hybrid strategy that can overcome the weakness of the component methods.

Several existing approaches embody hybrid integration in the above spirit. Besbes et al. [12] proposed a discrete *MRF*-based segmentation method which combined shape priors and regional statistics. However, this method did not perform segmentation at the pixel level. Freedman and Zhang [34] incorporated a shape template into the graph-cut formulation as a distance function. However, it relied crucially on user input. Based on the latter method, Ayvaci and Freedman [35] proposed a joint registration-segmentation method which removed the user interaction requirement and resolved the problem of template registration. However, this method required proper registration of the shape template for an accurate segmentation. Kumar et al. [36] used a *MRF*-representation where the latent shape model variables were integrated via expectation maximization. While shape information was utilized in a principled Bayesian manner, this approach was computationally intensive where a separate energy minimization was required. Malcolm et al. [37] imposed the shape prior model on the terminal edges and performed graph cut optimization iteratively starting from an initial contour. Their method constructed a statistical shape space using kernel principal component analysis. This method also relied on user input. Vu and Manjunath [38] proposed a shape prior integrated segmentation method using graph cuts suitable for multiple objects. The shape prior energy was based on a shape distance popular in level set approaches. However, the shape used was a simple fixed shape. Most of the above mentioned methods operated on 2D images.

Compared to these methods, the strategy proposed in this paper is a 3D anatomy segmentation method. More importantly, different from all the above shape prior-integrated methods, our technique does not need shape registration. The proposed *GC-OAAM* effectively combines the rich statistical shape and appearance information embodied in *AAM*, effective boundary oriented delineation in *LW*, with the globally optimal delineation capability of the *GC* method.

The remaining part of this paper is organized as follows. In Section 2, we elaborate the complete methodology of the delineation algorithm. In Section 3, we describe an evaluation of this method in terms of its accuracy and efficiency. In Section 4, we summarize our contribution.

II. THE *GC-OAAM* APPROACH

2.1 Overview of the approach

The proposed method consists of two phases: training phase and segmentation phase. Fig. 1 shows the flowchart of the proposed method. In the training phase, an *AAM* is constructed and the *LW* boundary cost function and *GC* parameters are trained. The segmentation phase

consists of two main steps: initialization and delineation. In the initialization step, we employ a pseudo 3D initialization strategy in which the organs are initialized slice by slice via a multi-object *OAAM* method, and a refinement method is applied to further adjust the initialization results. The employment of pseudo 3D initialization strategy is motivated by two reasons: (1) Compared to a full 3D initialization method, the proposed method is much faster. (2) It is difficult to combine *AAM* with *LW* in 3D. The experimental results demonstrate that the proposed method has performance comparable to the full 3D *AAM* initialization method, which may be due to the effective combination of *AAM* with the *LW* method. Finally, for the delineation part, the object shape information generated from the initialization step is integrated into the *GC* cost computation. An iterative *GC-OAAM* method is proposed for object delineation. The details of each step are given in the following sub-sections.

2.2 Model Building and Parameter Training

Before building the model, the top and bottom slices of each organ are first manually identified. Then linear interpolation is applied to generate the same number of slices for the organ in every training image. 2D *OAAM* models are then constructed for each slice level from the training images. The *LW* cost function and *GC* parameters are also estimated in this stage.

2.2.1 Landmark Specification—Due to the nature of the proposed method (slice by slice), we represent a 3D shape as a stack of 2D contours, and manually label the 3D shape slice by slice, although semi-automatic or automatic methods are also available for this purpose. For each slice, operators locate the shape visually, and then identify prominent landmarks on that shape.

2.2.2 AAM Construction—The standard *AAM* method [27, 28] is used to construct the model. The model includes both shape and texture information.

The generative model can be described by

$$\mathbf{x} = Q\mathbf{b}, \quad (1)$$

where Q is a matrix of the selected eigenvectors of the covariance matrix over the training samples for shape and texture, \mathbf{b} is the model parameter vector and \mathbf{x} is a sample generated by the model.

Suppose M_j represents the *AAM* for slice level j and the number of slice levels is n , then the overall model M can be represented as

$$M = (M_1, M_2, \dots, M_n). \quad (2)$$

Although we employ the pseudo 3D initialization strategy, we also build the real 3D *AAM* M_{3D} using the method in [43]. However, this 3D model M_{3D} is used only for providing the delineation constraints as to be explained later.

2.2.3 LW Cost function and GC Parameter Training—Similar to the *OASM* method [31], an oriented boundary cost function is devised for each organ in the model M as per live wire method [6].

A *boundary element*, *bel* for short, of a given image slice I is an ordered pair (a, b) of 4-adjacent pixels a and b . It represents the oriented edge between pixels a and b , (a, b) and (b, a) representing its two possible orientations. To every *bel* of I , we assign a set of features. The features are intended to express the likelihood of the *bel* belonging to the boundary (of a particular object) that we are seeking in I . The cost $c(b)$ associated with each *bel* b of I is a linear combination of the costs assigned to its features

$$c(b) = \frac{\sum_i w_i c_f(f_i(b))}{\sum_i w_i}, \quad (3)$$

where w_i is a positive constant indicating the emphasis given to feature function f_i , and c_f is the function to convert feature values $f_i(b)$ at b to cost values $c_f(f_i(b))$. In live wire technique [6], f_i represents features such as intensity on the immediate interior of the boundary, intensity on the immediate exterior of the boundary, and gradient magnitude at the center of the *bel*. c_f is an inverted Gaussian function, and here, uniform weights w_i are used for all selected features.

For the purpose of *OAAM*, we shall utilize the feature of live wire to define the best oriented path between any two points as a sequence of *bel*s with minimum total cost. The only deviation in this case is that the two points will be taken to be any two successive landmarks employed in the *AAM*, and the landmarks are assumed to correspond to pixel vertices. With this facility, we assign a cost to every pair of successive landmarks of any shape instance \mathbf{x} associated with M_j , which represents the total cost of the *bel*s in the best

oriented path $\langle b_1, b_2, \dots, b_h \rangle$ from landmark x_k to landmark x_{k+1} . That is,

$$\kappa(x_k, x_{k+1}) = \sum_{i=1}^h c(b_i). \quad (4)$$

For any shape instance $\mathbf{x} = (x_1, x_2, \dots, x_n)$ of M_j , the cost structure $\mathbf{K}(\mathbf{x})$ associated with M_j may now be defined as

$$K(\mathbf{x}) = \sum_{k=1}^m \kappa(\mathbf{x}_k, \mathbf{x}_{k+1}), \quad (5)$$

where m is the number of landmarks for this slice and we assume that $\mathbf{x}_{m+1} = \mathbf{x}_1$. That is, $K(\mathbf{x})$ is the weighted sum of the costs associated with the best oriented paths between all m pairs of successive landmarks of shape instance \mathbf{x} . The parameters of GC are also trained during the training stage; more details on this are given at section 2.4.1.

2.3 Initialization

The initialization step plays a key role in our method, which provides the shape constraints to the later GC segmentation step and makes it fully automatic. The proposed initialization method includes three main steps. First, a slice localization method is applied to detect the top and bottom slices of the organ. Second, a linear interpolation is applied to generate the same number of slices for the subject as in the model. And third, the organ is recognized slice by slice via the $OAAM$ method. A multi-object strategy [44] is utilized to help with object initialization. We found from experiments that the initialization performance with multiple organs in the model is much better than with a single organ due to the constraints among multiple organs. It means that, even if just one organ is to be segmented, other organs can be employed in the segmentation to provide context and constraints. Finally, a refinement method is applied to the initialization result. These three steps are described below.

2.3.1 Top and Bottom Slices Localization—There are several recent works related to slice localization. Haas et al. [39] introduced an approach for creating a navigation table using eight landmarks which were detected in various fashions. Seifert et al. [19] proposed a method to detect invariant slices and single point landmarks in full body scans by using probabilistic boosting tree (PBT) and Haar features. Emrich et al. [40] proposed a CT slice localization method via k - NN instance based regression. The aim of slice localization in our approach is to locate the top and bottom slices of the organ. Since we already trained the model for each organ slice, we could use this model for slice localization. The proposed method is based on the similarity to the $OAAM$ model of the slice.

For top slice localization in a given image, the top slice model is applied to each slice in the image using the recognition method detailed in 2.3.2 and evaluating the respective similarity metric (Eqn. (6)). Then the slice corresponding to the maximal similarity (minimal distance) is taken as the top slice of the organ. Fig. 2 shows the distance value computed from Eqn. (6) for the top slice in a patient abdominal CT image. The minimum corresponds to the top slice of the left kidney. A similar method is used for the bottom slice detection.

2.3.2 Object Recognition—The proposed object recognition method is based on the AAM . The conventional AAM matching method for object recognition is based on the root-mean-square difference between the appearance model instance and the target image. Such a strategy is better suited for matching appearances than for the detailed segmentation of target

images (see Fig. 3(b)). This is because the *AAM* is optimized on global appearance, and is thus, less sensitive to local structures and boundary information. Conversely, the *LW* delineates the boundary very well [6], however it needs good initialization of landmarks and is an interactive method. Here, we integrate the *AAM* with the *LW* method (termed *OAAM*) to combine their complementary strengths. That is, the *AAM* provides the landmarks to the *LW*, and as a return, *LW* improves the shape model of the *AAM*. The *LW* is fully integrated with *AAM* in two aspects: (1) *LW* is used to refine the shape model in *AAM*; (2) the *LW* boundary cost is integrated into cost computation during the *AAM* optimization method. Fig. 3(c) shows the proposed *OAAM* segmentation result; compared to conventional *AAM* method (Fig. 3(b)), the boundary delineation is much improved.

2.3.2.1 Refinement of the Shape Model in AAM by LW: First, the conventional *AAM* searching method is performed once to obtain a rough placement of the model. Then the following method is applied to refine the shape model in *AAM*. The shape is extracted from the shape model of the *AAM*, and then the landmarks are updated based on *LW* using only the shape model and the pose parameters (translation, rotation and scale). Subsequently, the refined shape model is transformed back into the *AAM*. At the same time, *AAM* refinement is applied to the image yielding its own set of coefficients for shape and pose.

Suppose P_1 , P_2 and P_3 are three successive landmarks in shape instance \mathbf{x} .

Algorithm: Refine AAM shape model based on LW: begin

1. Extract the shape instance \mathbf{x} from the shape model.
2. Update the landmarks' position in \mathbf{x} based on *LW* as follows (see Fig. 4).
 First, perform *LW* delineation from P_1 to P_2 , and P_2 to P_3 .
 Next, find the middle point Q_1 and Q_2 in the *LW* segments generated from P_1 to P_2 , and from P_2 to P_3 , respectively. Then perform *LW* delineation from Q_1 to Q_2 .
 Finally, find the closest point P_2' on the *LW* segment from Q_1 to Q_2 . Update P_2 to P_2' .
 Perform the above three steps on all landmarks in \mathbf{x} , and produce the updated shape model \mathbf{x}' .
3. Transform \mathbf{x}' into a new shape model instance \mathbf{x}_a' via an affine transformation so as to align it with the mean shape $\bar{\mathbf{x}}$.
4. Apply the model constraints to the new shape model \mathbf{x}_a' so that the new shape is within the allowed shape-space.

end

2.3.2.2 OAAM Optimization: In the conventional *AAM* matching method, the optimization is based only on the difference between the appearance model instance and the target image. The boundary cost is not taken into consideration. By combining the boundary cost, the performance of *AAM* matching can be considerably improved. In the proposed method, the

LW technique is integrated into the cost computation during the optimization process. Combined with the above shape model refinement method, our optimization method is as follows.

Algorithm OAAM Optimization: begin

Let the current estimate of the model parameter vector be b , the pose t , the texture transformation T_u and the image sample (as a vector) at the current estimate be g_{im} .

1. Extract the shape parameters b_s from the entire model parameter b , and refine b_s and pose t using the shape refinement method described above. This results in a new set of parameters b'_s and t' .
2. Resample the image intensity, resulting in the vector g'_{im} , and project the texture sample into the texture model frame using $g'_s = T_u^{-1}(g'_{im})$;
3. Evaluate the error vector, $r(b) = g'_s - g'_{im}$, and the current error, $E_{aam} = |r|^2$.
4. Compute the live wire cost along the shape boundary, E_{lw} , and compute the total error as

$$E_{total} = \alpha 1 \cdot E_{aam} + \alpha 2 \cdot E_{lw}. \quad (6)$$

5. Compute the predicted displacements, $\delta_b = -Rt(b)$, where $R = \left(\frac{\partial r^T}{\partial b} \frac{\partial r}{\partial b} \right)^{-1} \frac{\partial r^T}{\partial b}$.
6. Update the model parameters $b \leftarrow b + k\delta_b$, where initially $k = 1$.
7. Calculate the new shape points \mathbf{x} , and refine the new shape using the shape refinement method described above, and obtain the refined new shape point \mathbf{x}' .
8. Calculate the model frame texture g''_m , and sample the image at the new points \mathbf{x}' to obtain g''_{im} .
9. Calculate a new error vector, $r' = T_u^{-1}(g''_{im}) - g''_m$, and the error $E'_{aam} = |r'|^2$.
10. Compute the live wire cost along the predicted shape boundary, E'_{lw} . And compute the total error, $E'_{total} = \alpha 1 \cdot E'_{aam} + \alpha 2 \cdot E'_{lw}$.
11. If $E'_{total} < E_{total}$ then accept the new estimate; otherwise, try at $k = 0.5$, $k = 0.25$, etc, until no improvement can be made.

end

In our implementation, we set $\alpha 1 = \alpha 2 = 0.5$. During the initialization, we employ a multi-resolution strategy, in which we start at a coarse resolution and iterate to convergence at each level before starting the next level. This strategy is more efficient than searching at a single resolution and can lead to a convergence to the correct solution even when the initial model position is away from the real object(s).

2.3.2.3 Refinement of the 3D Recognized Shapes: After objects are recognized in all slices, the recognized shapes are stacked together to form 3D objects. We observed from experiments that sometimes the initialization result for one slice is far away from the results for its neighboring slices. This signals failure of recognition for this slice. We found that at most two slices failed in recognition in this sense for each subject in all of our experiments (in 80 cases, non-failed: 71; 1 slice failed: 7; 2 slices failed: 2). When failure occurs, we interpolate the new shape from the shapes in neighboring slices. Fig. 5 shows an illustration for the proposed method. The following method is applied to improve the recognized shape results.

Algorithm: Refinement of the 3D recognized shape: begin

for each slice level j , $1 \leq j \leq n$, do

1. Suppose the error in recognition (Eqn. (6)) for the current slice is e , and e_{\max} represents maximum error. Compute the distance d_{j-1} and d_{j+1} between the centroids of the shapes in neighboring slices $j-1$ and $j+1$.
2. Compute the total reliability for slice j as

$$rel_j = \eta_1 \cdot \left(\frac{e_{\max} - e}{e_{\max}} \right) + \eta_2 \cdot \exp\left(-\frac{(d_{j,j-1} - \mu(d_{j,j-1}))^2}{2 \cdot \text{var}(d_{j,j-1})}\right) + \eta_3 \cdot \exp\left(-\frac{(d_{j,j+1} - \mu(d_{j,j+1}))^2}{2 \cdot \text{var}(d_{j,j+1})}\right)$$

(7)

where η_1 , η_2 and η_3 are weights (in our implementation, $\eta_1=0.5$, $\eta_2=0.25$ and $\eta_3=0.25$).

$\mu(d_{j-1})$, $\text{var}(d_{j-1})$, $\mu(d_{j+1})$ and $\text{var}(d_{j+1})$ are the mean and variance of d_{j-1} and d_{j+1} , respectively, which are estimated from training images during the model building process. For the first and last slice, only one neighbor slice is used.

3. If $rel_j > \text{thre}_j$, then the recognized result is considered reliable; otherwise the recognized result is discarded and the new shape is interpolated from the neighboring slices. thre_j is the threshold of reliability, which is also estimated from the training images.

endfor

end

2.4 Segmentation/Delineation

The purpose of this step is to precisely delineate the shapes recognized in the previous step. We propose an iterative *GC-OAAM* (named *IGC-OAAM*) method for the organ's delineation. The *GC-OAAM* algorithm effectively integrates the shape information with the globally optimal 3D delineation capability of the *GC* method.

2.4.1 Shape integrated GC—GC segmentation can be formulated as an energy minimization problem such that for a set of pixels P and a set of labels L , the goal is to find a labeling $f: P \rightarrow L$ that minimizes the energy function $En(f)$.

$$En(f) = \sum_{p \in P} R_p(f_p) + \sum_{p \in P, q \in N_p} B_{p,q}(f_p, f_q), \quad (8)$$

where N_p is the set of pixels in the neighborhood of p , $R_p(f_p)$ is the cost of assigning label $f_p \in L$ to p , and $B_{p,q}(f_p, f_q)$ is the cost of assigning labels $f_p, f_q \in L$ to p and q . In two-class labeling, $L = \{0, 1\}$, the problem can be solved efficiently with graph cuts in polynomial time when $B_{p,q}$ is a submodular function, *i.e.*, $B_{p,q}(0, 0) + B_{p,q}(1, 1) \leq B_{p,q}(0, 1) + B_{p,q}(1, 0)$ [11].

In our framework, the unary cost $R_p(f_p)$ is the sum of a data penalty $D_p(f_p)$ and a shape penalty $S_p(f_p)$ term. The data term is defined based on the image intensity and can be considered as a log likelihood of the image intensity for the target object. The shape prior term is independent of image information, and the boundary term is based on the gradient of the image intensity.

The proposed shape-integrated energy function is defined as follows:

$$En(f) = \sum_{p \in P} (\alpha \cdot D_p(f_p) + \beta \cdot S_p(f_p)) + \sum_{p \in P, q \in N_p} \gamma \cdot B_{p,q}(f_p, f_q), \quad (9)$$

where α, β, γ are the weights for the data term, shape term S_p , and boundary term, respectively, satisfying $\alpha + \beta + \gamma = 1$. These components are defined as follows:

$$D_p(f_p) = \begin{cases} -\ln P(I_p|O), & \text{if } f_p = \text{object label} \\ -\ln P(I_p|B), & \text{if } f_p = \text{background label} \end{cases} \quad (10)$$

$$B_{p,q}(f_p, f_q) = \exp\left(-\frac{(I_p - I_q)^2}{2\sigma^2}\right) \cdot \frac{1}{d(p, q)} \cdot \delta(f_p, f_q), \quad (11)$$

and

$$\delta(f_p, f_q) = \begin{cases} 1, & \text{if } f_p \neq f_q \\ 0, & \text{otherwise,} \end{cases} \quad (12)$$

where I_p is the intensity of pixel p , *object label* is the label of the object (foreground). $P(I_p | O)$ and $P(I_p | B)$ are the probability of intensity of pixel p belonging to object and background, respectively, which are estimated from object and background intensity histograms during the training phase (details given below). $d(p, q)$ is the Euclidian distance between pixels p and q , and σ is the standard deviation of the intensity differences of neighboring voxels along the boundary.

$$S_p(f_p) = 1 - \exp\left(-\frac{d(p, \mathbf{x}_O)}{r_O}\right), \quad (13)$$

where $d(p, \mathbf{x}_O)$ is the distance from pixel p to the set of pixels which constitute the interior of the current shape \mathbf{x}_O of object O . (Note that if p is in the interior of \mathbf{x}_O , then $d(p, \mathbf{x}_O) = 0$.) r_O is the radius of a circle that just encloses \mathbf{x}_O . The linear time method in reference [42] was used in this paper for computing this distance.

During the training stage, the histograms of intensity for each object are estimated from the training images. Based on this, $P(I_p | O)$ and $P(I_p | B)$ can be computed. As for parameters α, β and γ in Eqn. (9), since $\alpha + \beta + \gamma = 1$, we estimate only α and β by optimizing accuracy as a function of α and β and set $\gamma = 1 - \alpha - \beta$. We use the gradient descent method for the optimization. Let $Accu(\alpha, \beta)$ represent the algorithm's accuracy (here we use the true positive volume fraction [45]), α and β are initialized to 0.35 each, then $Accu(\alpha, \beta)$ is optimized over the training data set to determine the best α and β .

2.4.2 Minimizing En with Graph Cuts—Let G be a weighted graph (V, A) , where V is a set of nodes, and A is a set of weighted arcs. Given a set $T \subseteq V$ of k terminal nodes, a cut is a subset of edges $C \subseteq A$ such that no path exists between any two nodes of T in the residue graph $(V, A \setminus C)$. In our implementation, we segment the object using the α -expansion method in [46].

The graph is designed as follows. We take $V = PU L$, i.e., V contains all the pixel nodes and terminals corresponding to the labels in L which represent objects of interest plus the background. $A = A_N \cup A_T$, where A_N is the n -links which connect pixels p and q ($p \in P, q \in N_p$) and with a weight of w_{pq} . A_T is the set of t -links which connect pixel p and terminals $\ell \in L$ and with a weight of $w_{p\ell}$. The desired graph with cut cost $|C|$ equaling $En(f)$ is constructed using the following weight assignments:

$$w_{pq} = \gamma \cdot B_{p,q}; \quad (14)$$

$$w_{p\ell} = K - (\alpha \cdot D_p(\ell) + \beta \cdot S_p(\ell)); \quad (15)$$

where K is a constant large enough to make the weights $w_{p\ell}$ positive.

2.4.3 IGC-OAAM—We assume that the recognized shapes are sufficiently close to the actual boundaries in the given image to be segmented. The *IGC-OAAM* algorithm then determines what the new position of the landmarks of the objects represented in the initialized shape \mathbf{x}_{in} should be such that the minimum graph cut cost is achieved, as presented below.

Algorithm IGC-OAAM: Input: Initialized shapes \mathbf{x}_{in} .

Output: Resulting shapes \mathbf{x}_{out} and the associated object boundaries.

begin

while number of iterations $< nIteration$ *do*

1. Perform *GC* segmentation using Eqn. (9) based on the *OAAM* initialized shapes \mathbf{x}_{in} ;
2. Compute the new position of the landmarks by moving each landmark in \mathbf{x}_{in} to the point closest on the *GC* boundary; call the resulting shapes \mathbf{x}_{new} ;
3. If no landmarks move, then, set \mathbf{x}_{new} as \mathbf{x}_{out} and stop;
Else, subject \mathbf{x}_{new} to the constraints of model M_{3D} , and call the result \mathbf{x}_{in} .

endwhile

Perform one final *GC* segmentation based on \mathbf{x}_{out} , and obtain the associated object boundaries.

end

In our implementation, $nIteration$ is set as 3. Also we limit the distance a landmark can move within any iteration to 6 voxels.

III. EXPERIMENTAL RESULTS

The proposed methods were tested on a clinical *CT* dataset. This dataset contained images pertaining to 20 patients (10 male and 10 female, ages 32 to 68), acquired from pre-contrast phase of two different type of *CT* scanner (GE Medical systems, LightSpeed Ultra, and Philips, Mx8000 IDT 16). The pixel size varied from 0.55 to 1 *mm*, and slice thickness from 1 to 5 *mm*. Four experiments of liver, left kidney, right kidney and spleen segmentation were done to evaluate the proposed method. All objects were manually segmented by two experts to generate the reference segmentations (ground truth). The leave-one-out strategy was used in the evaluation.

3.1 Evaluation of the Localization of the Top and Bottom Slice

The proposed slice localization method was used to detect the top and bottom slices of the liver, left kidney, right kidney and spleen. These organs were manually checked to generate the reference standard of the top and bottom position. Table 1 shows the experimental results. We observe that the localization of the top slice of liver is most accurate which may

be due to the high contrast in the lung region; while the localization of liver bottom has the largest error which may be due to the lack of sufficient contrast in that region. The average localization error is 7.3 mm . Compared to Emrich et al's result [40] of 4.5 cm , the proposed method seems superior.

3.2 Evaluation of Initialization

For helping with the initialization of liver, left and right kidneys, and spleen, the skin object is included in the model in addition to the object of interest. Fig. 7 shows one slice example and its corresponding mean shape and texture model for the four objects. We select only 8 landmarks for the skin object because *LW* works very well for this object even for such a small number of landmarks. Table 2 summarizes the number of interpolated slices and the number of landmarks used in our experiments.

In Figs. 8–11, the left column shows sample initialization results for the four objects. A quantitative evaluation of the initialization approach is presented in Table 3. The accuracy in terms of true positive and false positive volume fractions (TPVF and FPVF) [45] is shown. TPVF indicates the fraction of the total amount of tissue in the true delineation; FPVF denotes the amount of tissue falsely identified, which are defined as follows,

$$\text{TPVF} = \frac{C_{TP}}{C_{td}} \quad (16)$$

$$\text{FPVF} = \frac{C_{FP}}{U_d} \quad (17)$$

Where, U_d is assumed to be a binary scene with all voxels in the scene domain C set to have a value 1, as shown in Fig. 6, more details can be seen in [45].

Experiments were done to compare the performance of pseudo-3D *AAM* (single object, here pseudo-3D means slice-by-slice), pseudo-3D *MAAM* (multi-object), real 3D *MAAM* (multi-object), and the proposed pseudo-3D multi-object *OAAM* using reference images from expert #1 as ground truth. We note that the multi-object strategy improves the accuracy considerably over single object *AAM*. The *MOAAM* method also improves the *MAAM* initialization performance due to the effective combination of *AAM* and *LW*. The pseudo-3D *MOAAM* and the real 3D *MAAM* methods [43] have comparable performance, while the pseudo-3D *MOAAM* method is about 12 times faster (see Table 4). This is one of the reasons that we used the pseudo-3D initialization method.

After object recognition for all the slices, all the recognized shapes are stacked together to form a 3D shape. Then the refinement of shape method proposed in section 2.3.3 is applied if the 3D shape is not transiting smoothly. We found actually our *OAAM* recognition method works very well and the refinement only applied to very few cases. In total 80 (20×4) cases of organ recognition, there are 7 cases with one slice of object recognition failed (liver:3, left

kidney:1, right kidney:2 and spleen:1), 2 cases with two slices of object recognition failed (liver:1 and right kidney 1).

3.3 Evaluation of the *IGC-OAAM* Delineation Method

The accuracy of delineation by *IGC-OAAM* expressed in TPVF, FPVF and average symmetric surface distance [29] using reference images from expert #1 are summarized in Table 3. And the evaluation of the reference image from expert 1 to the reference image from expert 2 is also shown in Table 3 as the second rater. We observe that the average TPVF and FPVF is about 94.3% and 0.15%, respectively. In Figs. 8–11, the right column shows the *IGC-OAAM* segmentation results for the liver, left kidney, right kidney, and spleen, respectively. Additionally, Fig. 12 shows the 3D surface distance between the segmentation result by *IGC-OAAM* and reference image (from expert 1) for the liver, left kidney, right kidney, and spleen segmentation, respectively. The mean distance, over all objects and the whole dataset, between the segmented 3D surface and the reference (true) surface was found to be about 0.78 mm.

In terms of efficiency, Table 4 shows the computation time for the four objects on an Intel Xeon E5440 workstation with 2.83GHz CPU, 8 GB of RAM. The average total time (initialization + segmentation) for segmenting one liver is about 310 seconds. Segmentation of kidney and spleen has similar computational time, about 270 seconds.

The proposed *IGC-OAAM* delineation method was also tested on the MICCAI 2007 grand challenge training dataset using leave-one-out strategy. There are 20 CT volumes of abdomen with contrast agent in the training datasets. All datasets have an in-plane matrix of 512×512 pixels and inter-slice spacing from 0.7 mm to 5.0 mm. Fig. 13 shows the recognition and delineation results for three slice levels of one image on MICCAI grand challenge dataset..

The proposed method was evaluated based on the MICCAI 2007 grand challenge for liver segmentation evaluation criteria [29]: volumetric overlap error (Overlap Error), volume difference, symmetric average surface distance, symmetric RMS surface distance, and maximal surface distance. The results achieved by the proposed method and previous work from literature are summarized in Table 5. Compare to the best performance (Kainmüller et. al [30]), we can see we have comparable performance, but much faster speed (6 min vs 15 min).

IV. CONCLUSIONS AND DISCUSSIONS

In this paper, we proposed a 3D automatic anatomy segmentation method. The method effectively combines the AAM, LW, and GC methods to exploit their complementary strengths. It consists of three main parts: model building, initialization, and segmentation. For the initialization part, we employ a pseudo-3D strategy, and segment the organs slice by slice via multi-object OAAM method which effectively combines the AAM and LW methods. For the segmentation (delineation) part, an iterative GC-OAAM method is proposed which integrates the shape information gathered from initialization with a GC algorithm. The method was tested on a clinical CT dataset with 20 patients for segmenting

the liver, kidneys, and spleen. The experimental results suggested that an overall segmentation accuracy of $TPVF > 94.3\%$, $FPVF < 0.2\%$ can be achieved.

As for the initialization, we employed a pseudo-3D strategy and combined *AAM* and *LW* methods to improve the performance. The multi-object strategy also helped initialization due to increased constraints. Compared to the real 3D *AAM* method, the pseudo 3D *OAAM* approach has comparable accuracy while has roughly a 12-fold speed up. The purpose of initialization is to provide a rough object localization and shape constraints for latter *GC* method which will produce refined delineation. We think it is better to have a fast and robust method than a slow and more accurate technique for initialization.

As for the delineation, shape constrained *GC* method is the core part of the whole system. Several similar ideas were also proposed in the references [34–38]. However, they are mostly tested in the 2D images, and it is difficult to compare with these methods because the testing dataset is different.

From Figs. 8–11 and the testing result in table 5, it seems that the proposed method was a little bit under-segmenting the organ. This may due to two reasons: (1) the shape term designed in our cost function is not symmetric. We didn't put the penalty for the pixel if it is inside the shape. That is because we found there are usually some pixels which don't belong to the target organ inside the shape, such as urine in the kidney (Fig. 10). So our method can separate this kind of pixels out. We think this is the beauty of our method. (2) according to our experiences, the expert is easily tender to over-segmenting the organ during the process of manually segmenting the boundaries.

Although localizing a *CT* slice within a human body can greatly facilitate the workflow of a physician, so far, this area of research has not received much attention. The proposed slice localization method aims to localize the top and bottom slices of organs, which is important part of the whole system. The average localization error over the whole dataset and all organs is only about 7.3 *mm* (much improved compared to the result of 4.5 cm in [40]), good for clinical use. In a similar manner, it can also be used to localize any slice by constructing the corresponding slice model.

In this paper, only one object is segmented at a time. With the shape constraints of multiple organs, the proposed *IGC-OAAM* method can be easily generalized to segment multiple organs simultaneously. However, this brings up an issue for *GC* - of the unavailability of a globally optimal min cut solution for simultaneously segmenting multiple objects. For single object segmentation, global optimality is guaranteed. For multiple objects, the α -expansion method can find segmentations only within a known factor of the global optimum [46].

Current proposed method for segmenting one organ is taking about 5 min. To make it more realistic in clinical application, the parallelization or multi-threads of the algorithm is one of the good solutions. Anderson et al. [48] and Liu et al. [49] proposed the parallelization of *GC* methods, and achieved the good performance. This will also be investigated for the proposed method in near future.

The executable version of 3D shape constrained GC with user interface can be downloaded from website <http://xinjianchen.wordpress.com/research/>. Source codes will be available soon. By opening source, we believe it will benefit to the whole society.

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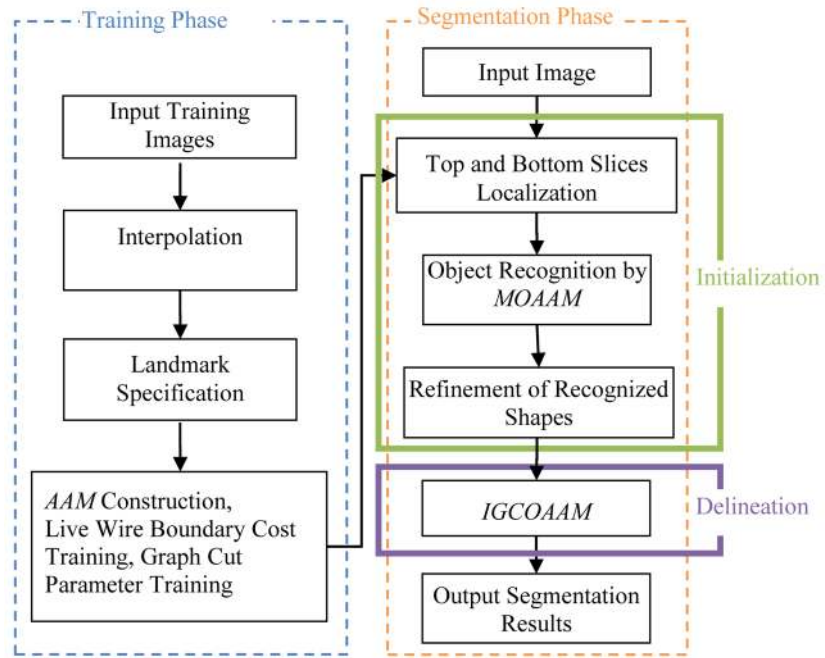


Fig. 1.
The flowchart of the proposed *GC-OAAM* system.

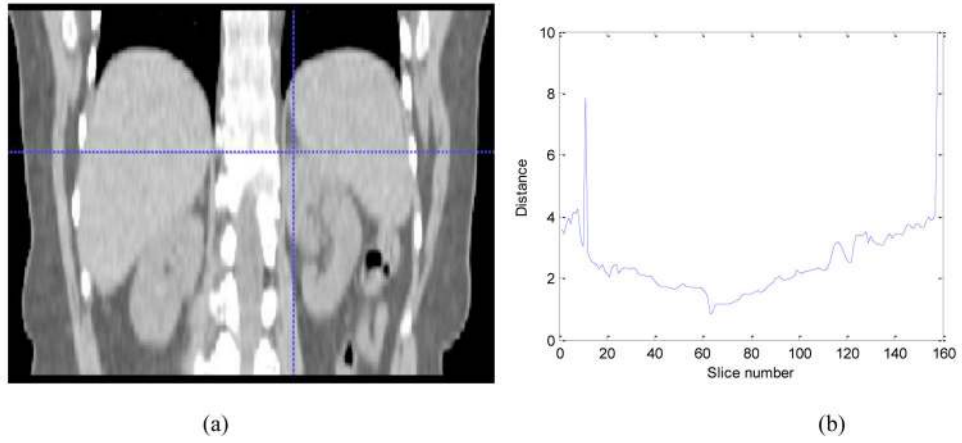


Fig 2. Illustration of top slice recognition. (a) Coronal view of the abdominal region. Cross point represents the top slice of the left kidney. (b) The distance values for the top slice of the left kidney.

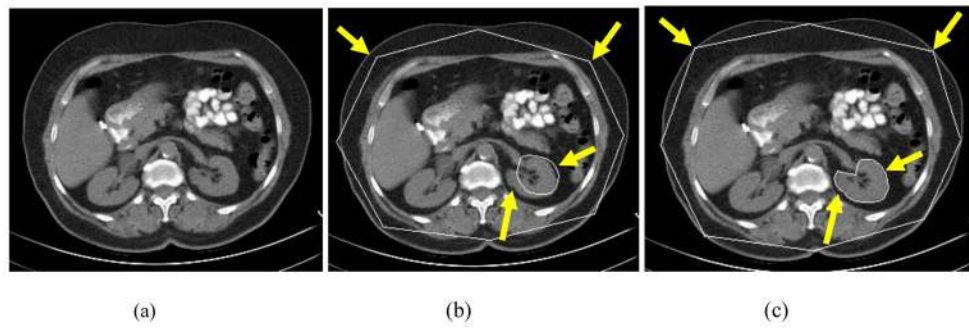


Fig. 3. Comparison of conventional *AAM* and *OAAM* segmentation. (a) Original image. (b) Conventional *AAM* segmentation showing a good appearance fit but poor boundary detection accuracy (arrows). (c) *OAAM* result shows substantial improvement in boundary location (arrows).

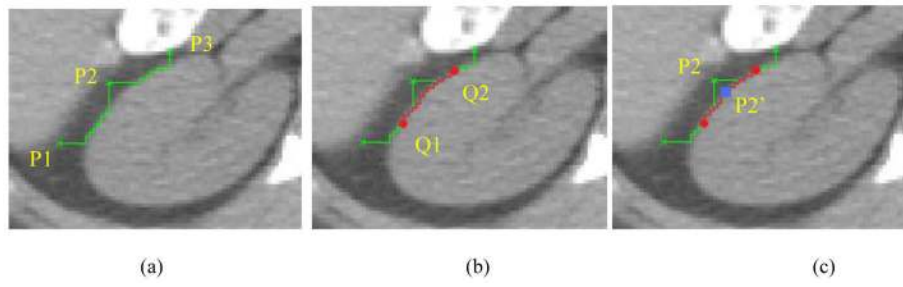


Fig. 4. The illustration of how to update the landmark's position. (a) P1, P2 and P3 are three landmarks from *AAM* shape results. (b) The middle point Q1 of the *LW* segment between P1 and P2, Q2 of P2 and P3 are generated. (c) Landmark P2 is moved to the closest point P2' on the live wire segment from Q1 to Q2.

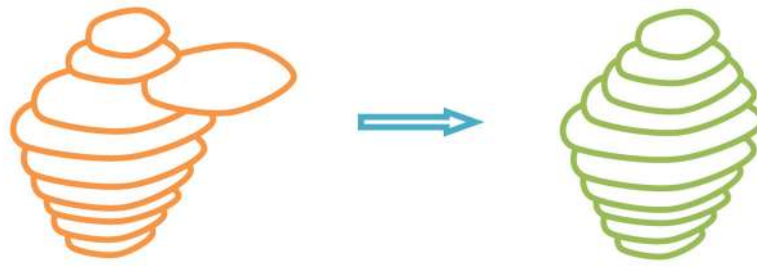


Fig 5.
Illustration of refinement of the 3D recognized shape

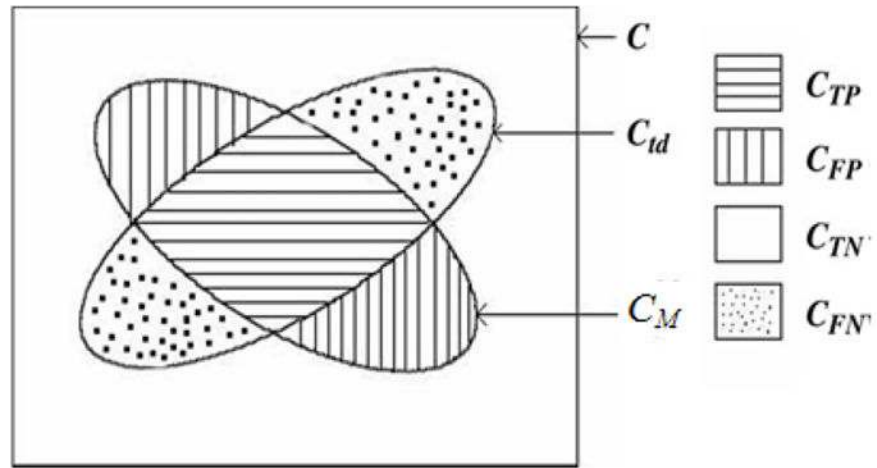


Fig. 6. Illustration of the accuracy factors for delineation for a binary case. Here, C_{id} is the corresponding scene of 'true' delineation, C_M is the delineation result by method M.

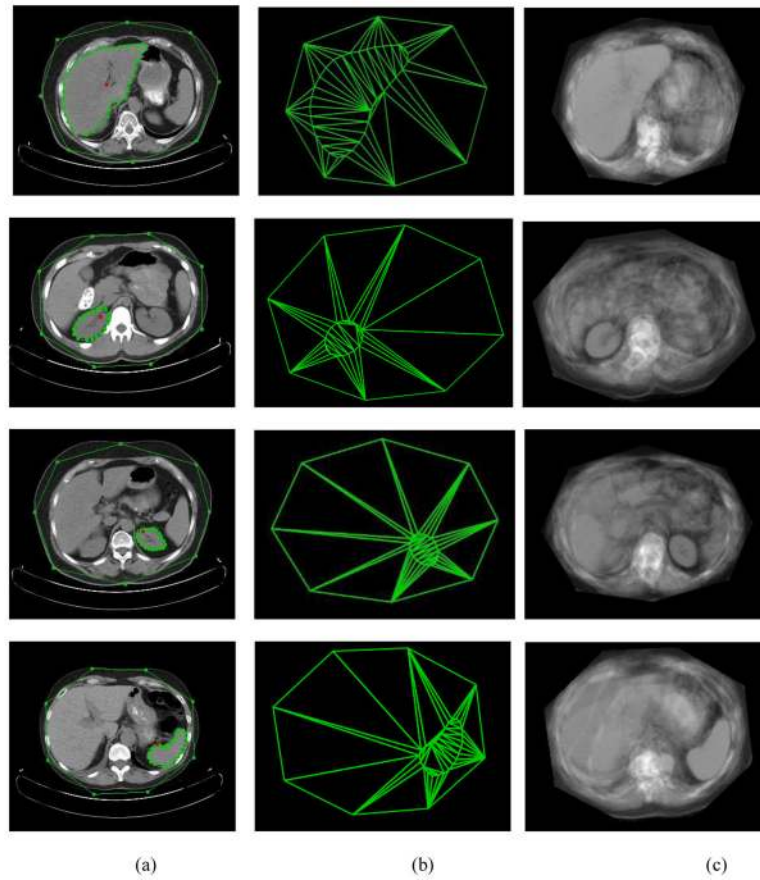


Fig. 7. Illustration of models used in organ initialization. The 1st, 2nd, 3rd and 4th row correspond to liver, right kidney, left kidney, and spleen, respectively. (a) The landmarks of organ and skin on one slice. (b) The corresponding *AAM* shape model for this slice level. (c) The corresponding *AAM* appearance model for this slice level.

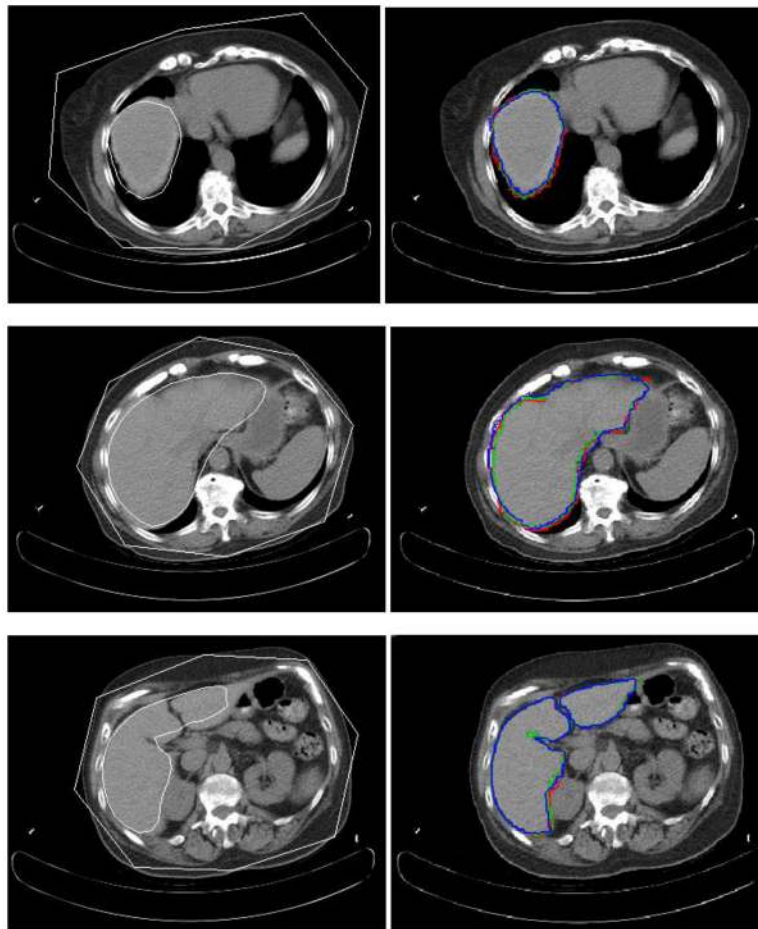


Fig. 8. Experimental results for three slice levels of liver segmentation. The left column is the *MOAAM* initialization result; the right is *IGC-OAAM* result in which red contour represents the reference image 1, green represents the reference image 2, and blue contour represents segmentation by the proposed method.

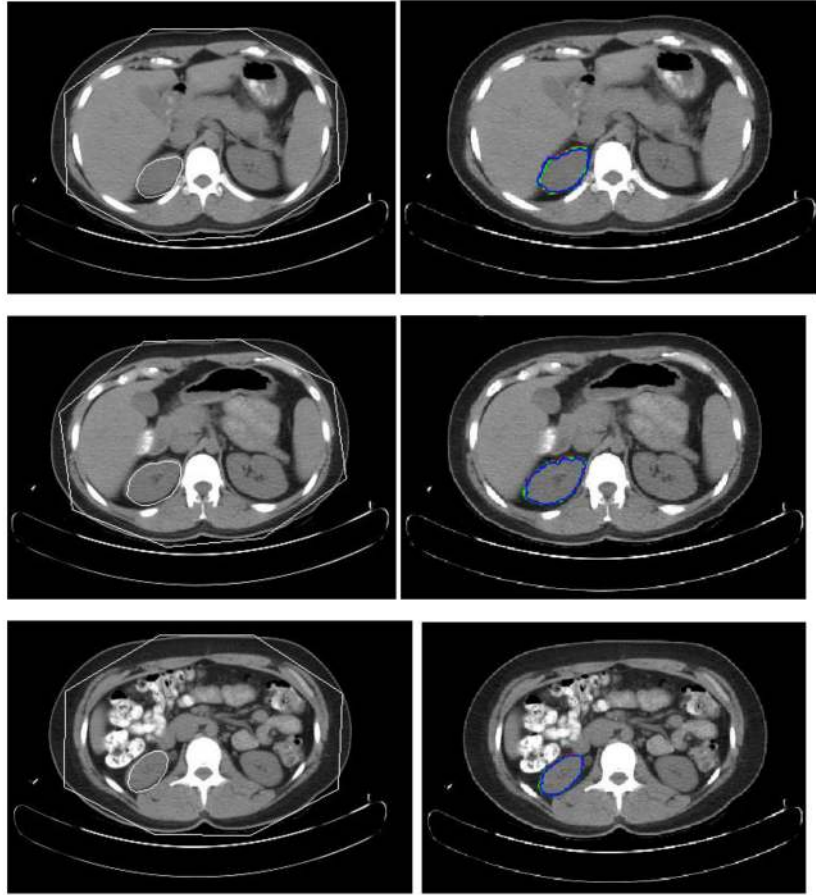


Fig. 9. Experimental results for three slice levels of right kidney segmentation. The left column is the *MOAAM* initialization result; the right is *IGC-OAAM* result in which red contour represents the reference image 1, green represents the reference image 2, and blue contour represents segmentation by the proposed method.

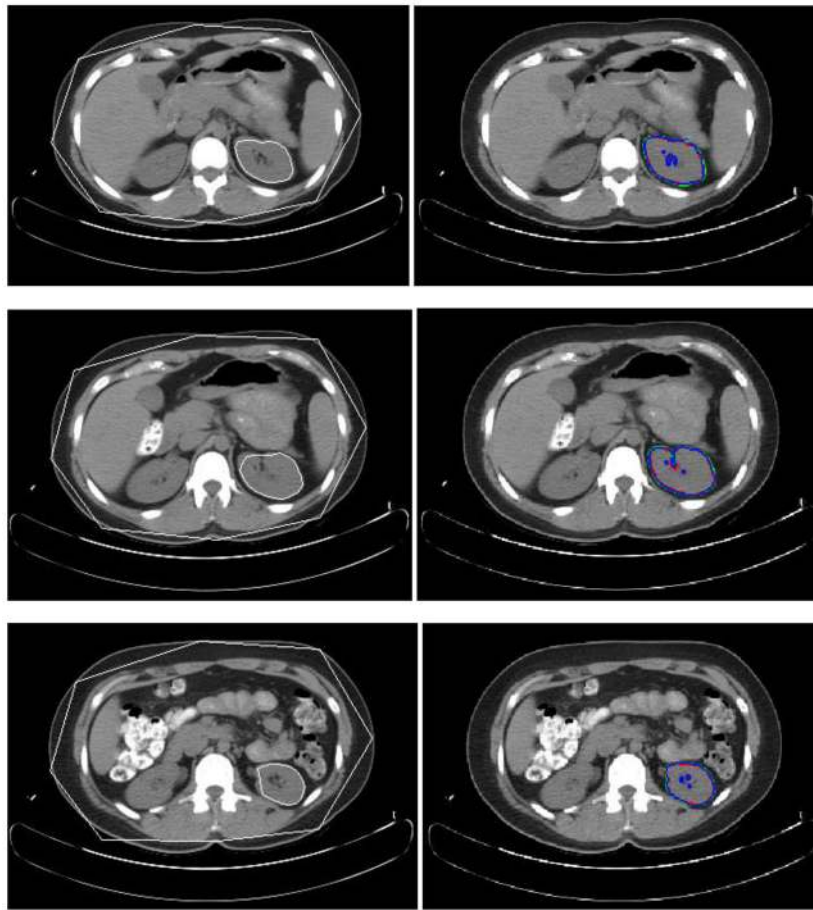


Fig. 10. Experimental results for three slice levels of left kidney segmentation. The left column is the *MOAAM* initialization result; the right is *IGC-OAAM* result in which red contour represents the reference image 1, green represents the reference image 2, and blue contour represents segmentation by the proposed method.

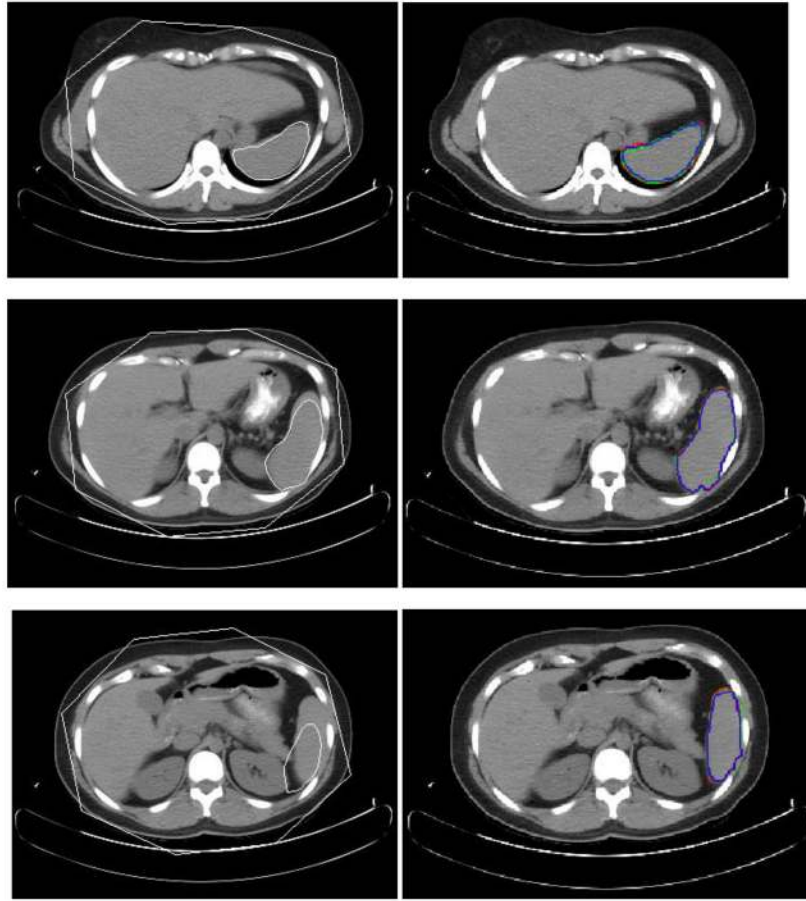


Fig. 11. Experimental results for three slice levels of spleen segmentation. The left column is the *MOAAM* initialization result; the right is *IGC-OAAM* result in which red contour represents the reference image 1, green represents the reference image 2, and blue contour represents segmentation by the proposed method.

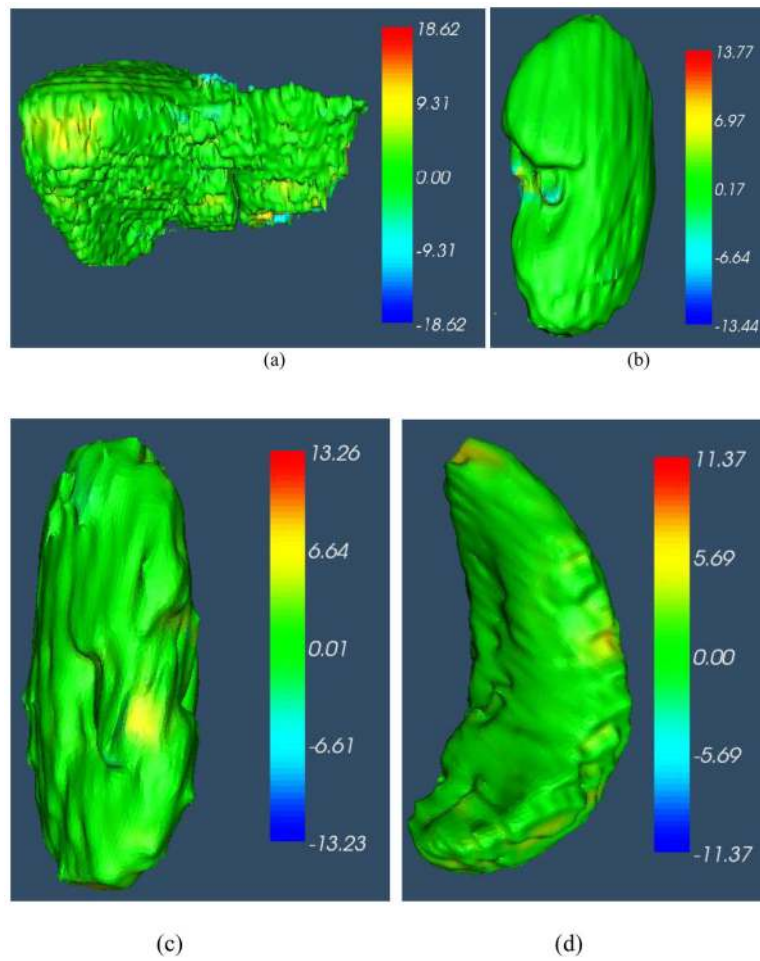


Fig. 12. 3D surface distance (mm) between the segmentation result by *IGC-OAAM* and reference image (ground truth 1). Positive value represents the vertex on the surface of the segmentation is outside of the surface of ground truth, vice versa. (a), (b), (c) and (d) correspond to surface distance for liver, right kidney, left kidney and spleen segmentation, respectively.

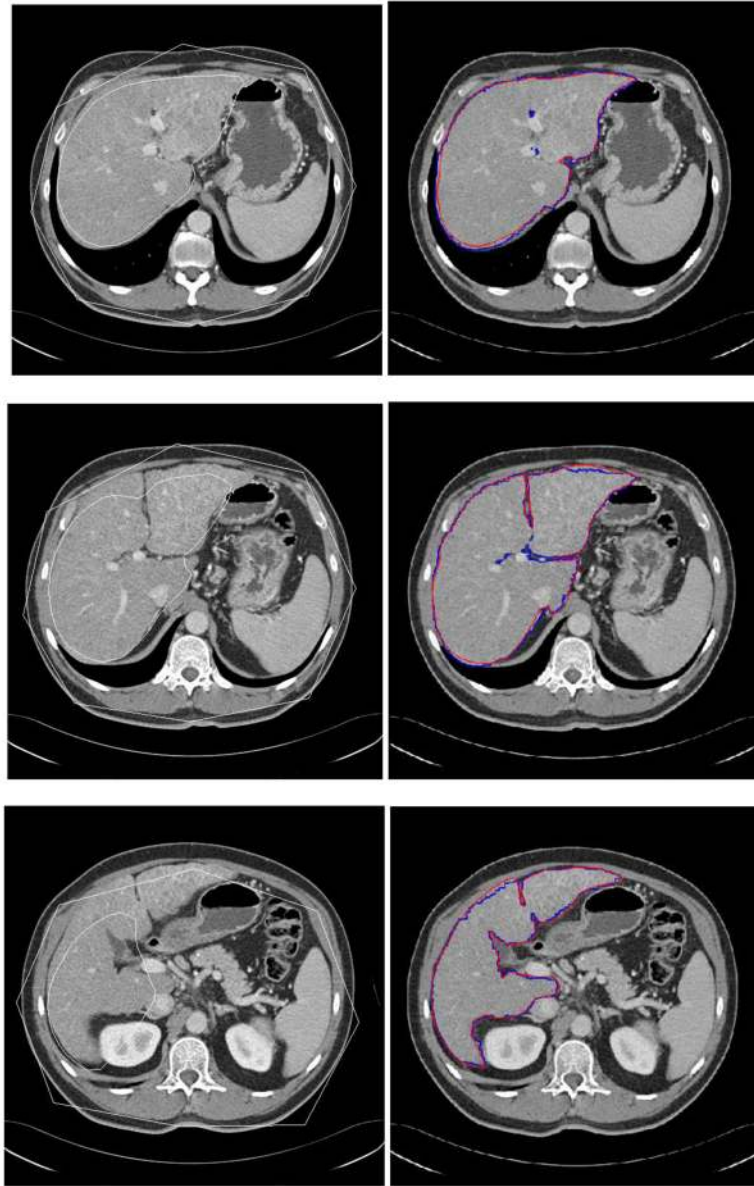


Fig. 13. Experimental results for three slice levels of one image on MICCAI grand challenge dataset. The left column is the *MOAAM* initialization result; the right is *IGC-OAAM* result in which red contour represents the reference (ground truth) and blue contour represents segmentation by the proposed method.

Table 1

Slice localization error (in mm) of the organ top and bottom

Organ		Mean error (in mm) \pm std.dev.
Liver	Top	5.1 \pm 2.5
	Bottom	9.2 \pm 5.1
Left kidney	Top	7.5 \pm 5.2
	Bottom	6.2 \pm 4.6
Right kidney	Top	8.3 \pm 6.5
	Bottom	7.1 \pm 5.8
Spleen	Top	8.1 \pm 5.2
	Bottom	7.3 \pm 6.1
Average		7.3 \pm 5.1

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Table 2

Number of Landmarks and Slices Used in Modeling

	Number of Landmarks in Organ	Number of Landmarks in Skin Object	Number of Interpolated Slices
Liver	35	8	50
Left Kidney	20	8	32
Right Kidney	20	8	32
Spleen	26	8	32

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Table 3

Mean and standard deviation of *TPVF*, *FPVF* for Pseudo-3D *AAM*, Pseudo-3D *MAAM*, 3D *MAAM*, Pseudo-3D *MOAAM*, and *IGC-OAAM* for liver, two kidneys, and spleen.

	TPVF (%)				FPVF (%)				Average Surface Dist (mm)			
	liver	left kidney	right kidney	spleen	liver	left kidney	right kidney	spleen	liver	left kidney	right kidney	spleen
<i>Pseudo-3D AAM</i>	60.01 ±5.12	51.01 ±8.36	53.01 ±6.35	59.31 ±7.25	10.12 ±2.38	15.36 ±3.25	12.15 ±3.01	14.62 ±4.35	12.12 ±5.32	11.59 ±4.39	10.98 ±4.78	11.32 ±4.12
<i>Pseudo-3D MAAM</i>	85.02 ±1.32	84.03 ±1.60	83.18 ±1.68	85.73 ±1.35	1.12 ±0.38	0.93 ±0.45	1.75 ±0.46	1.02 ±0.67	4.59 ±3.81	4.21 ±3.36	4.39 ±3.61	4.12 ±3.28
<i>Pseudo-3D MOAAM</i>	90.52 ±0.93	90.15 ±1.01	90.36 ±1.52	91.32 ±0.88	0.46 ±0.11	0.55 ±0.12	0.59 ±0.13	0.67 ±0.16	2.01 ±1.21	1.82 ±1.06	1.72 ±0.98	1.61 ±0.88
<i>Second rater</i>	90.12 ±1.13	89.32 ±1.22	89.52 ±1.63	90.65 ±1.51	0.53 ±0.05	0.67 ±0.06	0.62 ±0.11	0.71 ±0.13	1.93 ±1.03	1.62 ±0.92	1.81 ±1.25	1.73 ±1.03
<i>IGC-OAAM</i>	94.86 ±0.87	95.32 ±0.96	95.12 ±0.79	95.22 ±1.10	0.12 ±0.05	0.15 ±0.03	0.16 ±0.02	0.12 ±0.02	0.79 ±0.35	0.71 ±0.28	0.75 ±0.29	0.75 ±0.22
	94.03 ±0.92	94.15 ±1.05	94.35 ±0.89	94.68 ±1.39	0.16 ±0.03	0.13 ±0.02	0.21 ±0.03	0.15 ±0.02	0.81 ±0.32	0.75 ±0.25	0.79 ±0.33	0.76 ±0.29

Table 4

Average computational time (in seconds) in all experiments for pseudo-3D *MAAM*, 3D *MAAM*, pseudo-3D *MOAAM*, and *IGC-OAAM*.

Organ	Average computational time (in seconds)			
	<i>Pseudo-3D MAAM</i>	<i>3D MAAM</i>	<i>Pseudo-3D MOAAM (initialization)</i>	<i>IGC-OAAM (delineation)</i>
Liver	50	732	60	310
Left Kidney	33	495	40	275
Right Kidney	32	476	40	260
Spleen	35	556	45	280

Comparison with other methods on MICCAI 2007 liver segmentation grand challenge

Table 5

Method	Runtime [min]	Overlap Error [%]	Volume difference [%]	Avg. distance [mm]	RMS distance	Max distance [mm]
Kainmuller et al.	15	6.1±2.1	-2.9±2.9	0.9±0.3	1.9±0.8	18.7±8.5
Heimann et al.	7	7.7±1.9	1.7±3.2	1.4±0.4	3.2±1.3	30.1±10.2
Proposed IGC-OAAM method	6	6.5±1.8	-2.1±2.3	1.0±0.4	1.8±1.0	20.5±9.3