

Research Article

Automatic Liver Segmentation on Volumetric CT Images Using Supervoxel-Based Graph Cuts

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Accurate segmentation of liver from abdominal CT scans is critical for computer-assisted diagnosis and therapy. Despite many years of research, automatic liver segmentation remains a challenging task. In this paper, a novel method was proposed for automatic delineation of liver on CT volume images using supervoxel-based graph cuts. To extract the liver volume of interest (VOI), the region of abdomen was firstly determined based on maximum intensity projection (MIP) and thresholding methods. Then, the patient-specific liver VOI was extracted from the region of abdomen by using a histogram-based adaptive thresholding method and morphological operations. The supervoxels of the liver VOI were generated using the simple linear iterative clustering (SLIC) method. The foreground/background seeds for graph cuts were generated on the largest liver slice, and the graph cuts algorithm was applied to the VOI supervoxels. Thirty abdominal CT images were used to evaluate the accuracy and efficiency of the proposed algorithm. Experimental results show that the proposed method can detect the liver accurately with significant reduction of processing time, especially when dealing with diseased liver cases.

1. Introduction

Liver cancer is one of the most common cancers worldwide, with increasing morbidity and high mortality [1]. Computed Tomography (CT) has been widely used for clinical diagnosis of hepatic disease because of its high resolution. Accurate liver segmentation from abdominal CT scans is critical for computer-assisted diagnosis and therapy, including patient-specific liver anatomy evaluation, functional assessment, treatment planning, and image-guided surgery [2]. Traditionally, radiologists or physicians have to manually delineate the liver region slice by slice, which is tedious and time-consuming due to the large amount of data [3]. Therefore, accurate and efficient methods for liver segmentation are demanded.

Liver segmentation from CT images remains an open challenge due to the high variability in the shape and size of liver, presence of pathologies like tumor or cirrhosis, and low contrast with adjacent tissues or organs [4]. Recently, a large variety of semiautomatic and automatic methods have been developed to improve the liver segmentation procedure.

These methods are commonly based on region growing [5–7], clustering [8, 9], deformable models or level sets [10–13], statistical shape models (SSMs) [14, 15], probabilistic atlases [16–18], and graph cuts [19–24]. Several comprehensive reviews of liver segmentation techniques have been conducted [25–27].

Table 1 gives an overview of selected liver segmentation methods for CT images. Semiautomatic segmentation methods tend to obtain more precise results compared to automatic methods, while automatic methods can reduce interoperator and intraoperator variability. Region growing or clustering approaches are fast and easy to implement, but they may become ineffective when the liver is inhomogeneous due to large lesions. Deformable models and level set based methods are often used as the final step to refine the segmented contour/surface to improve the accuracy, but they are sensitive to the initial contour/surface and need an iterative process which is time-consuming. Probabilistic atlases and SSMs can offer highly accurate results, but they demand sufficient training data with gold standards to generate the atlases or shape models. Graph cuts methods have been

TABLE 1: Overview of liver segmentation methods for CT images: auto = automatic; semi = semiautomatic; VOE = volumetric overlap error; RVD = relative absolute volume difference; MaxD = maximum symmetric surface distance; DSC = dice similarity coefficient; RG = region growing; DM = deformable model; SSM = statistical shape model; PA = probabilistic atlas; GC = graph cuts; local = from local hospitals; Sliver07 = MICCAI 2007 grand challenge in segmentation of liver datasets.

| Study | Year | Method | Accuracy | Dataset | Auto | Time (s) | CPU (GHz) |
|------------------------------|------|-----------------|--|----------|------|-------------|--------------------------------------|
| Kumar et al. [7] | 2013 | RG | DSC = 98% | local | Auto | 40/slice | — |
| Goryawala et al. [8] | 2014 | Clustering + RG | DSC = 92% RVD = 2.78% | local | Semi | 10.96/slice | — |
| Peng et al. [10] | 2014 | DM | VOE = 6.10% RVD = -0.00% MaxD = 16.80 mm | Sliver07 | Semi | 180 | 3.16 GHz 4 GB RAM |
| Kainmüller et al. [15] | 2007 | SSM + DM | VOE = 6.09% RVD = -2.86% MaxD = 18.69 mm | Sliver07 | Auto | 900 | Intel 3.2 GHz |
| Linguraru et al. [16] | 2010 | PA + DM | DSC = 96.2% VOE = 2.20% ASD = 1.20 mm | Sliver07 | Auto | — | — |
| Platero and Tobar [18] | 2014 | PA + GC | VOE = 7.60% RVD = -0.50% MaxD = 24.70 mm | Sliver07 | Auto | 261.35 | Intel Xeon E5520 2.27 GHz |
| Massoptier and Casciaro [19] | 2007 | GC | DSC = 95% | local | Auto | — | — |
| Li et al. [20] | 2015 | SSM + GC | VOE = 6.24% RVD = 1.18% MaxD = 18.82 mm | Sliver07 | Auto | 284.95 | Core(TM) i5 3.1 GHz 4 GB RAM |
| Chen et al. [23] | 2012 | GC | VOE = 4.16% RVD = 3.53% MaxD = 16.70 mm | Sliver07 | Semi | 60–180 | Intel Core 2 2.66 GHz 3.25 RAM |

widely used in medical image segmentation because they can achieve global optimum solution. For liver segmentation, graph cuts methods based on adaptive thresholding [19], SSMs [20], probabilistic atlas [16], and deformable models [24] have been developed. However, directly constructing the graph over the voxels of CT volume data will lead to a high computational cost.

In this paper, we proposed a new method for automatic liver delineation on CT volume images using supervoxel-based graph cuts. It was highly efficient compared to other methods and provided accurate segmentation results. Both the liver volume of interest (VOI) and the foreground/background seed points for graph cuts were extracted automatically. Supervoxels of the liver VOI were generated, and the graph used for three-dimensional (3D) graph cuts segmentation was defined over the obtained VOI supervoxels.

The rest of this paper is organized as follows. Section 2 introduces the details of the proposed method. Section 3 describes the experiments. Section 4 shows the experimental results, and discussion is given in Section 5. Conclusions are summarized in Section 6.

2. Methods

The flow chart of the proposed approach is illustrated in Figure 1. The proposed segmentation framework consisted of five steps. (1) In the preprocessing step, smoothing and resampling were conducted by using recursive Gaussian filtering and linear interpolation, respectively. (2) The patient-specific liver VOI extraction procedure was performed firstly

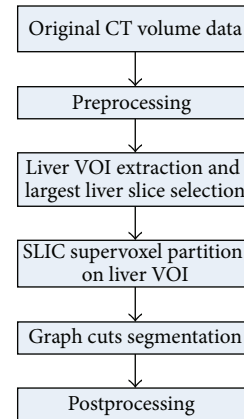


FIGURE 1: Flow chart of the proposed 3D liver segmentation method.

by determining the region of abdomen using maximum intensity projection (MIP) [28] and thresholding methods, followed by applying a histogram-based adaptive thresholding method and morphological operations. The largest liver slice, which was the axial slice containing the maximal amount of liver tissue, was automatically selected. (3) The generation of supervoxels was performed on the liver VOI using the simple linear iterative clustering (SLIC) method [29]. (4) In the segmentation step, foreground/background seed points for graph cuts were selected on the largest liver slice, and additional background seed points on regions of heart and kidney were extracted automatically on selected

heart and kidney slices. Gaussian mixture models (GMMs) were used to estimate the intensity distributions of foreground/background seeds [30]. Subsequently, the graph cuts algorithm [31, 32] was applied to the VOI supervoxels. (5) Finally, morphological opening, cavity filling, and median filtering were used to refine the segmented liver. After being resampled back to the original spatial resolution and size, the segmentation result was output for evaluation.

2.1. Preprocessing. The preprocessing step included smoothing and resampling. Recursive Gaussian filtering was first applied to smooth the input CT volume image I . Then, the smoothed CT volume image was resampled from anisotropic to isotropic voxel size ($1.5 \times 1.5 \times 1.5$ mm) by linear interpolation to obtain the preprocessed volume image I_{pre} .

2.2. Liver VOI Extraction and Largest Liver Slice Selection. For computational efficiency, the region of abdomen I_{ROA} was firstly extracted from I_{pre} to remove several nonabdominal slices and voxels. Then, the liver VOI image I_{VOI} was determined in I_{ROA} .

2.2.1. 3D Abdominal Region Extraction. To extract the region of abdomen I_{ROA} , a 3D abdominal bounding box (ABB) was measured using MIP and thresholding. Coronal and axial MIP images were obtained by applying the MIP algorithm to I_{pre} . Let $[X_{min}, X_{max}, Y_{min}, Y_{max}, Z_{min}, Z_{max}]$ be the coordinates of ABB, where Z_{min} and Z_{max} were the lower and upper bounding coordinates along z -axis and $X_{min}, X_{max}, Y_{min}$, and Y_{max} were the in-plane bounding coordinates. The dimensions of I_{pre} were denoted by u_x, u_y , and u_z in x, y , and z directions, respectively.

(1) Calculation of the Lower Bounding Coordinate Z_{min} . For the coronal MIP image $M_{coronal}$ (Figure 2(a)), which had a size of $u_x \times u_z$, segmentation of bones was performed using the Otsu algorithm [33] to obtain the binary bone mask M_{bone} (Figure 2(b)). Let n_{total} denote the total number of pixels in M_{bone} . The number of bone pixels n_b (in white color) was counted.

If $n_b/n_{total} < 3/4$, M_{bone} was regarded as successful bone segmentation. Otherwise, if $n_b/n_{total} \geq 3/4$, which meant the extraction of bones failed, Otsu algorithm with two thresholds was applied to $M_{coronal}$ to generate thresholds T_1 and T_2 ($T_1 < T_2$); then, M_{bone} was obtained by applying binary thresholding $[T_2, T_{max}]$ to $M_{coronal}$, where T_{max} was the maximum CT value in Hounsfield unit (HU).

For the bone mask M_{bone} , morphological opening with round structuring elements (radius $r = 3$), cavity filling, and median filtering were conducted to generate the processed binary bone mask M_{bone}^* (Figure 2(c)).

For M_{bone}^* , the number of bone pixels was calculated in each column: $i = 0, 1, \dots, u_x - 1$. Among multiple columns with the maximum number of connected bone pixels, the left-most column was regarded as the position of spine i_{spine} (shown as the blue line in Figure 2(c)). Then, among columns $i = 0, 1, \dots, i_{spine} - 30$ and $i = i_{spine} + 30, \dots, u_x - 1$, the local maximums of the number of connected bone pixels, c_{left} and c_{right} , were computed on both sides of the spine.

Assign $Z_{min} = \min(c_{left}, c_{right}) - 5$, shown as the red line in Figure 2(c).

(2) Calculation of the In-Plane Bounding Coordinates $X_{min}, X_{max}, Y_{min}$, and Y_{max} . For the axial MIP image M_{axial} (Figure 2(d)), Otsu algorithm with two thresholds was used to generate thresholds T_3 and T_4 ($T_3 < T_4$), and then binary thresholding $[T_3, T_{max}]$ was applied to M_{axial} to obtain the binary abdomen mask $M_{abdomen}$ (Figure 2(e)).

For $M_{abdomen}$, morphological closing ($r = 1$), cavity filling, and selection of the largest connected region were performed to generate the processed binary abdomen mask $M_{abdomen}^*$ (Figure 2(f)).

A two-dimensional (2D) bounding box was determined based on the abdominal region (in white color) in $M_{abdomen}^*$. The coordinates of the 2D bounding box were then taken as $[X_{min}, X_{max}, Y_{min}, Y_{max}]$.

Assign $[X_{min}, X_{max}, Y_{min}, Y_{max}] \leftarrow [X_{min} - 5, X_{max} - 5, Y_{min} - 10, Y_{max}]$, as the liver was located in the right upper quadrant of the abdomen, shown as the yellow rectangle in Figure 2(f).

(3) Calculation of the Upper Bounding Coordinate Z_{max} . Lungs are filled with air and have very low intensity values in CT images. By applying binary thresholding $[T_{min}, -300]$ to I_{pre} , where T_{min} was the minimum CT value in HU, candidate air-filled regions were extracted. Then, the binary lung mask I_{lung} was obtained by selecting lung regions among the candidate air-filled regions. Figure 2(g) shows the 3D visualization of a lung mask.

For I_{lung} , the number of lung pixels was calculated in each axial slice $j = 0, 1, \dots, u_z - 1$. The slice j_{max} with the maximum number of lung pixels was regarded as the upper bounding of ABB (Figure 2(h)). Assign $Z_{max} = j_{max}$, shown as the green line in Figure 2(c).

Based on the extracted 3D abdominal bounding box ABB, I_{pre} was cropped to obtain the region of abdomen I_{ROA} .

2.2.2. Liver VOI Extraction. In the region of abdomen, liver is the largest organ and located in the right upper quadrant of the abdomen. Also, liver is the largest object in middle axial slices of I_{ROA} . According to this prior knowledge, the liver VOI was extracted from I_{ROA} by applying a histogram-based adaptive thresholding method and morphological operations.

For adaptive thresholding, a rough estimation of the liver intensity range $[T_{lower}, T_{upper}]$ was calculated from the histogram of I_{ROA} , where T_{lower} and T_{upper} denoted the minimal and maximal intensity values of liver voxels, respectively.

By analyzing the volumetric histogram of I_{ROA} , I_{ROA} could be classified as high contrast or low contrast [34]. The image with two high peaks in the histogram was regarded as high contrast (Figure 3(a)). If only one high peak was detected in the histogram, the image was regarded as low contrast (Figure 3(b)).

In Figure 3, V denoted the intensity value and $h(V)$ was the histogram value corresponding to V . If I_{ROA} was a high contrast image, V_1 and V_2 were taken as the intensity values of

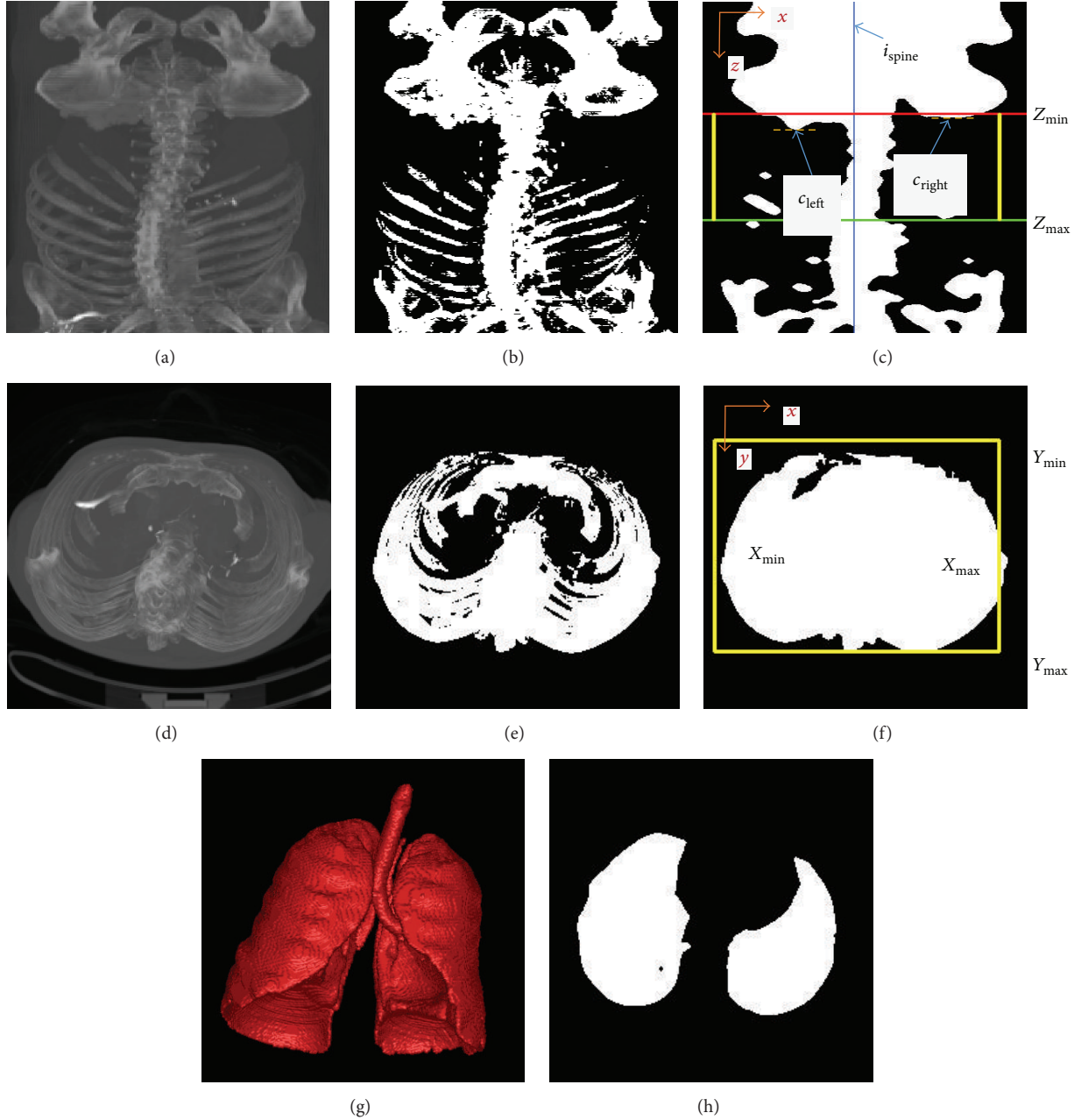


FIGURE 2: An example of the 3D abdominal region extraction. (a) Coronal MIP image. (b) The binary bone mask of (a). (c) The processed binary mask. Blue line shows the position of spine. Red and green lines show the lower and upper bounding along z -axis, respectively. Yellow lines are corresponding to the in-plane bounding box in (f). (d) Axial MIP image. (e) The binary abdomen mask of (d). (f) The processed binary mask. Yellow rectangle shows the in-plane bounding box. (g) The extraction of lungs. (h) The binary lung mask with the largest lung region area.

the valley and the second peak, respectively; V_3 was calculated by $h(V_3) = h(V_2)/30$, $V_3 > V_2$; then, assign $T_{\text{lower}} = V_1$ and $T_{\text{upper}} = V_3$. If I_{ROA} was a low contrast image, V_5 was taken as the intensity value of the peak; then, V_4 and V_6 were calculated by $h(V_4) = h(V_5)/2$, $V_4 < V_5$ and $h(V_6) = h(V_5)/30$, $V_6 > V_5$, respectively; assign $T_{\text{lower}} = V_4$ and $T_{\text{upper}} = V_6$.

By applying binary thresholding $[T_{\text{lower}}, T_{\text{upper}}]$ to I_{ROA} , the initial binary liver mask I_{liver}^0 was obtained (Figure 4(a)).

For I_{liver}^0 , selection of the largest connected component on the right side of abdomen was conducted on each axial slice to obtain the binary mask I_{liver}^1 . Then, morphological opening ($r = 1$) and selection of the largest connected region were applied to each coronal slice of I_{liver}^1 to obtain the binary liver mask I_{liver}^2 (Figure 4(b)).

To determine the liver VOI, the bounding box of the largest 3D connected object in I_{liver}^2 was calculated. The liver

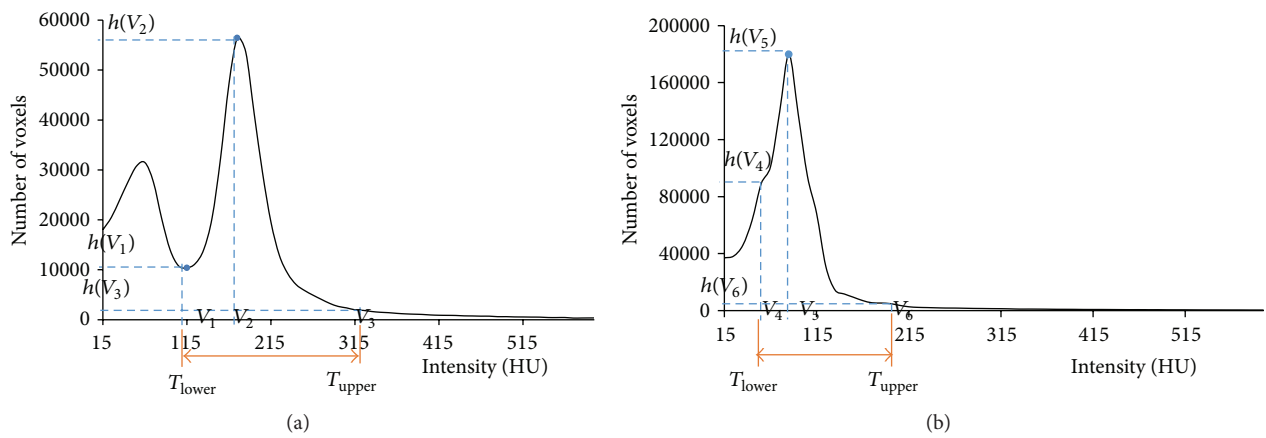


FIGURE 3: Examples of volumetric histograms in range [15, 600] HU. (a) A high contrast image with two high peaks. (b) A low contrast image with only one high peak.

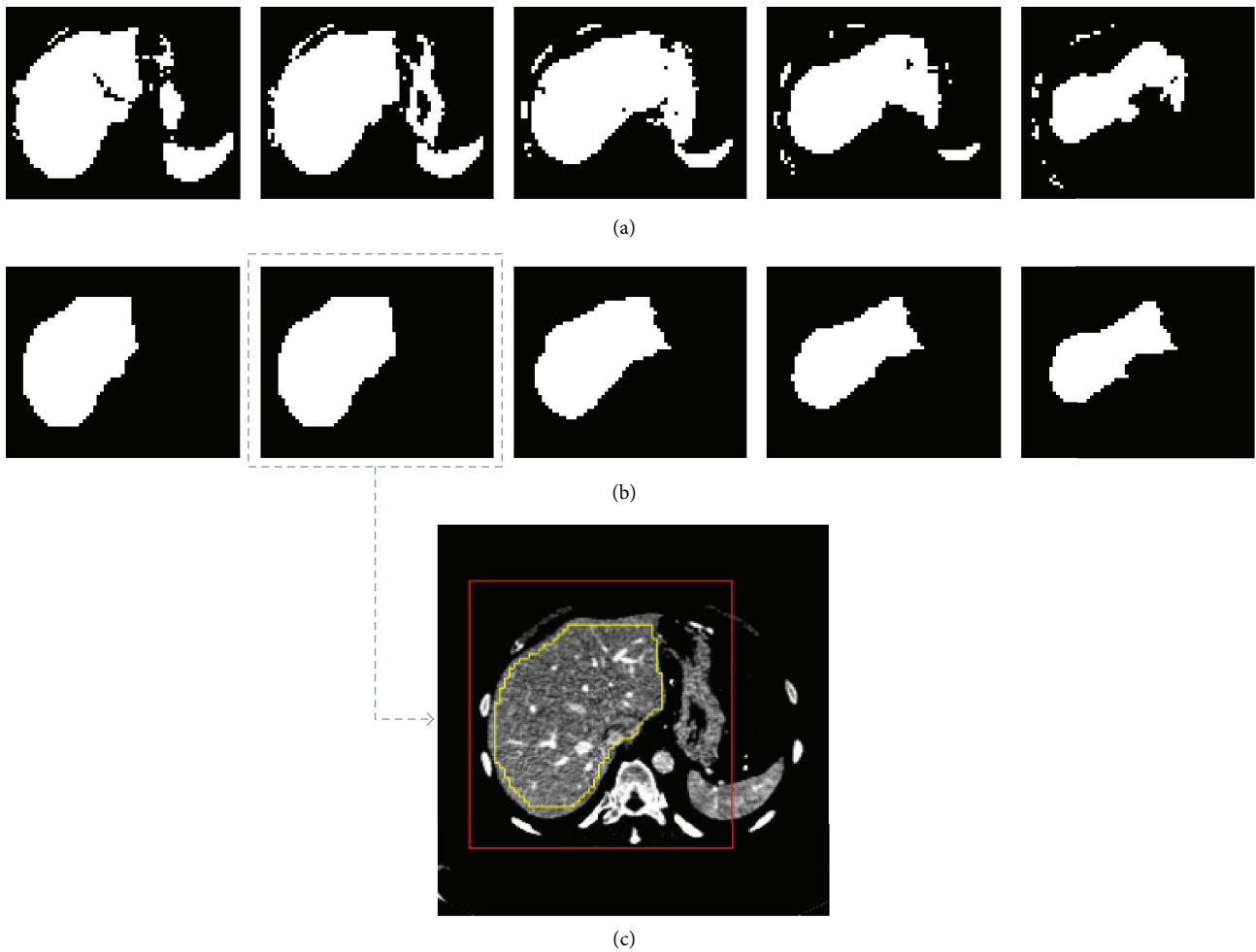


FIGURE 4: An example of liver VOI extraction and the largest liver slice selection. (a) Axial slices of the initial binary liver mask by using adaptive thresholding. (b) Axial slices of the processed binary liver mask. (c) The selected largest liver slice with the initial liver region in yellow and the liver VOI in red.

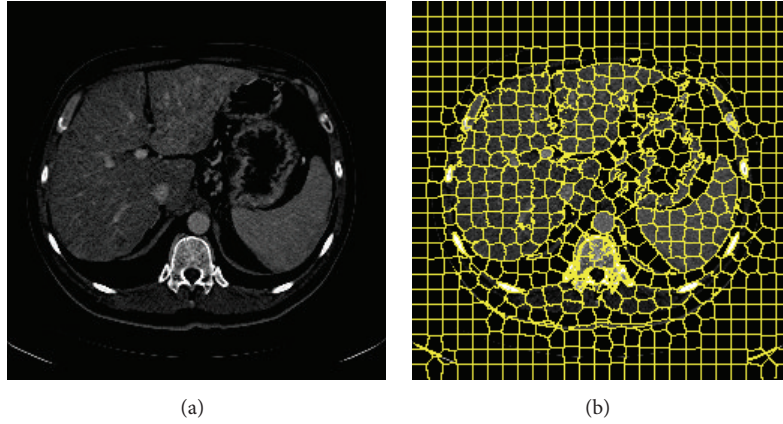


FIGURE 5: Illustration of superpixel generation. (a) Original image. (b) Supervoxels generated by the 2D simple linear iterative clustering (SLIC) algorithm. Yellow contours show the boundaries between the superpixels.

VOI image I_{VOI} was taken as the obtained bounding box with a margin of ten voxels around it.

2.2.3. Largest Liver Slice Selection. In the largest liver slice corresponding to the largest cross section of the liver, the liver was a whole object. Therefore, the seed points could be extracted on the largest liver slice without missing any separate parts of the liver.

To select the largest liver slice, the number of liver pixels was calculated in each axial slice of I_{liver}^2 (Figure 4(b)). The slice with the maximum number of liver pixels was regarded as the largest liver slice. As shown in Figure 4(c), the largest liver slice M_{liver} was selected with the corresponding binary liver mask M_{liver}^* , where the red rectangle represented the liver VOI and the yellow contour denoted the liver region in M_{liver}^* .

2.3. SLIC Supervoxel Partition on Liver VOI. SLIC is a novel k -means based clustering algorithm which can generate supervoxels quickly and efficiently [29, 35]. The SLIC supervoxels have nearly uniform size, while their boundaries closely match true image boundaries. Figure 5 shows the result of superpixel partition on one CT slice using 2D SLIC algorithm.

Let N_v denote the number of voxels in the liver VOI image I_{VOI} :

$$N_v = N_p S^3, \quad (1)$$

where S denoted the supervoxel spacing in each dimension and N_p was the desired number of supervoxels. The intensity of each supervoxel was computed as the average intensity of all voxels within the supervoxel.

2.4. Graph Cuts Segmentation. The liver segmentation problem can be posed as a binary labeling problem and formulated in terms of energy minimization. Graph cuts segmentation achieves an optimal solution by minimizing the energy function via the max-flow/min-cut algorithm [31, 32]. Necessary hard constraints (seed points) and intensity distributions were required for the graph cuts segmentation.

2.4.1. Necessary Hard Constraints. Necessary foreground/background seed points should be selected for graph cuts. On the largest liver slice M_{liver} , foreground seed points were sampled automatically in the liver region corresponding to the mask M_{liver}^* . By applying morphological dilating ($r = 20$), negative operation, and binary thresholding $[-50, T_{\text{max}}]$ to M_{liver}^* , the background mask M_{bkg}^* was obtained. Then, background seed points were sampled automatically on M_{bkg}^* around the liver region.

Additional background seeds on regions of heart and kidney were also selected to prevent oversegmentation. In our experiments, the upper bounding of the liver VOI in z direction was taken as the heart slice, and one or two kidney slices were selected among slices between the lower bounding of I_{VOI} and the largest liver slice.

In liver CT images, regions of heart and kidney always have higher intensities compared to the liver region. Based on the histogram mentioned in Section 2.2, binary thresholding $[V_m, T_{\text{max}}]$ was applied to the heart and kidney slices to obtain the binary background masks M_{heart}^* and M_{kidney}^* , where V_m denoted the intensity value of the highest peak in the histogram.

For M_{heart}^* and M_{kidney}^* , morphological opening ($r = 1$) and eroding ($r = 2$) were applied. Then, the connected regions with average intensity values less than $V_m + 10$ and $V_m + 50$ were removed on M_{heart}^* and M_{kidney}^* , respectively. Additional background seed points were sampled in the remaining regions of M_{heart}^* and M_{kidney}^* .

In our method, the graph cuts algorithm was applied to the liver VOI supervoxels. Therefore, the seed points were all converted to the corresponding supervoxels in the liver VOI.

2.4.2. Gaussian Mixture Models. For liver segmentation on CT images, there were foreground tissues including liver parenchyma, vessels, and tumors as well as background tissues including intercostal muscles, bones, heart, and kidneys. The intensity distribution of each tissue was assumed to follow Gaussian distribution [36].

Let V_p denote the intensity of a supervoxel p , and let N be the number of supervoxels in the foreground or background samples. The intensity distributions of foreground and background were represented as GMMs:

$$\begin{aligned}
P_{\text{GMM}}(V_p) &= \sum_{k=1}^K \omega_k P_N(V_p | \mu_k, \sigma_k^2), \\
P_N(V_p | \mu_k, \sigma_k^2) &= \frac{1}{\sqrt{2\pi\sigma_k^2}} \exp\left[-\frac{1}{2\sigma_k^2} (V_p - \mu_k)^2\right], \\
\mu_k &= \frac{1}{N_k} \sum_p V_p, \\
\sigma_k^2 &= \frac{1}{N_k} \sum_p V_p^2 - \mu_k^2, \\
\omega_k &= \frac{1}{N} N_k,
\end{aligned} \tag{2}$$

where ω_k , μ_k , and σ_k^2 were the weight, mean, and variance of the k th Gaussian components, respectively; K denoted the number of Gaussian components. One GMM was used for the foreground $P_{\text{fg}}(V_p)$, and another one was used for the background $P_{\text{bkg}}(V_p)$. The expectation maximization (EM) algorithm [37] was applied to estimate the parameters of GMMs.

2.4.3. Graph Cuts. The graph $G = (Q, W)$ was created, where Q represented the set of nodes. The nodes comprised the liver VOI supervoxels p , a source terminal Q_S , and a sink terminal Q_T . The set of edges W consisted of two types of undirected edges: n -links (neighborhood links) connecting all unordered pairs (p, q) of neighboring supervoxels in the liver VOI and t -links (terminal links) connecting each supervoxel to the two terminals.

Let $L = (l_1, \dots, l_p, \dots, l_{N_p})$ be a binary vector whose components l_p specified assignments to supervoxels p ; $l_p \in \{0, 1\}$, with l_p equaling 0 for background and 1 for foreground (i.e., liver). Vector L defined a segmentation, and the goal was to assign a unique label to each p by minimizing the following energy function $E(L)$:

$$E(L) = \alpha \sum_{p \in P} R(l_p) + \beta \sum_{(p,q) \in W} B(l_p, l_q), \tag{3}$$

where $R(l_p)$ and $B(l_p, l_q)$ were the regional term and boundary term, respectively; α and β were weighting factors. The minimum cost cut could be computed in polynomial time for the two terminal graph cuts.

Table 2 gives the weights of edges in W . We specified that Q_S and Q_T corresponded to label 0 (background) and label 1 (liver), respectively. Two sets of the seed points were defined as foreground nodes Q_{fg} and background nodes Q_{bkg} , respectively.

The regional term $R(l_p)$ specified the cost of assigning a label l_p to p based on its intensity V_p and the intensity

TABLE 2: Graph edge weights.

| Edge | Weight | For |
|------------|---------------------------|------------------------|
| (p, q) | $\beta \cdot B(l_p, l_q)$ | $(p, q) \in W$ |
| (p, Q_S) | 0 | $p \in Q_{\text{fg}}$ |
| | ∞ | $p \in Q_{\text{bkg}}$ |
| (p, Q_T) | $\alpha \cdot R(l_p = 0)$ | Others |
| | ∞ | $p \in Q_{\text{fg}}$ |
| (p, Q_T) | 0 | $p \in Q_{\text{bkg}}$ |
| | $\alpha \cdot R(l_p = 1)$ | Others |

probabilistic model. With the foreground and background GMMs, we defined $R(l_p)$ as negative log-likelihood:

$$\begin{aligned}
R(l_p = 0) &= -\log(P_{\text{fg}}(V_p)), \\
R(l_p = 1) &= -\log(P_{\text{bkg}}(V_p)).
\end{aligned} \tag{4}$$

The boundary term $B(l_p, l_q)$ represented the penalty of the discontinuity between two adjacent supervoxels p and q [38]. The penalty became large when p and q had similar intensity values. Consider

$$\begin{aligned}
B(l_p, l_q) &= \delta(l_p, l_q) \cdot \frac{1}{(V_p - V_q)^2 + 1}, \\
\delta(l_p, l_q) &= \begin{cases} 1, & \text{if } l_p \neq l_q, \\ 0, & \text{if } l_p = l_q. \end{cases}
\end{aligned} \tag{5}$$

2.5. Postprocessing. After finishing the 3D graph cuts segmentation procedure, the resulting binary liver mask I_{seg} was obtained. Morphological opening, cavity filling, largest region selection, and median filtering methods were applied to I_{seg} to remove spicules and smooth the liver surface. Then, the binary image was resampled back to the original spatial resolution and size for further evaluation.

3. Experiments

3.1. Datasets. The MICCAI Sliver07 datasets [4] used in this study contained 30 clinical contrast-enhanced (portal venous phase) abdominal CT volume images acquired using a variety of scanners, including 20 images with expert segmentations (Sliver07-train) and 10 testing images (Sliver07-test). Most datasets were pathologic, and Sliver07-test datasets involved relatively more extreme cases. The ground truth of Sliver07-test was not publicly available, and the evaluations on Sliver07-test were performed by the organizer of the MICCAI Sliver07.

For Sliver07 datasets, the number of slices, in-plane resolution, and interslice resolution varied between 64 and 394, 0.58 and 0.81 mm, and 0.7 and 5.0 mm, respectively. The segmented liver region was defined to include the entire liver tissue and all internal structures, including vessels, tumors, and cirrhosis.

3.2. *Evaluation Measures.* To quantitatively evaluate the performance of the proposed method, five different error measures were used to measure the volumetric overlap or surface distances of the segmentation result compared to the ground truth [4]. These were volumetric overlap error

$$\begin{aligned}
 \text{VOE} &= 100 \left(1 - \frac{|A \cap B|}{|A \cup B|} \right), \\
 \text{RVD} &= 100 \left(\frac{|A| - |B|}{|B|} \right), \\
 \text{ASD} &= \frac{1}{|S(A)| + |S(B)|} \left(\sum_{S_A \in S(A)} d(S_A, S(B)) + \sum_{S_B \in S(B)} d(S_B, S(A)) \right), \\
 \text{RMSD} &= \sqrt{\frac{1}{|S(A)| + |S(B)|} \left(\sum_{S_A \in S(A)} d^2(S_A, S(B)) + \sum_{S_B \in S(B)} d^2(S_B, S(A)) \right)}, \\
 \text{MaxD} &= \max \left\{ \max_{S_A \in S(A)} d(S_A, S(B)), \max_{S_B \in S(B)} d(S_B, S(A)) \right\},
 \end{aligned} \tag{6}$$

where A denoted the segmented volume produced by the proposed method; B was the segmented volume by experts; $S(A)$ denoted the set of surface voxels of A . The shortest distance of a voxel v to $S(A)$ was defined as $d(v, S(A)) = \min_{s_A \in S(A)} \|v - s_A\|$, where $\|\cdot\|$ denoted the Euclidean distance. A positive value of RVD meant oversegmentation, while a negative value meant undersegmentation.

For MICCAI Sliver07 datasets, the score of 100 indicated a perfect segmentation when all the five measures were zero [4]. The manual segmentation of the average quality (VOE = 6.4%, RVD = 4.7%, ASD = 1.0 mm, RMSD = 1.8 mm, and MaxD = 19.0) was worth the score of 75. The total running time for segmentation was also recorded.

3.3. *Implementation.* To validate the proposed method, experiments were conducted on HP EliteBook 8470w (Intel Core 2.40 GHz CPU and 4 GB RAM). The proposed algorithm was implemented in C++ under Visual Studio 2008, with the use of SLIC code by Achanta et al. [29] (<http://ivrl.epfl.ch/research/superpixels>) and max-flow/min-cut code by Boykov and Kolmogorov [31] (<http://vision.csd.uwo.ca/code/>). The GMM algorithm was implemented using the open source computer vision toolkit OpenCV (<http://www.opencv.org/>), with seeds initialization performed by the k -means++ algorithm [39]. The insight segmentation and registration toolkit ITK (<http://www.itk.org/>) and the visualization toolkit VTK (<http://www.vtk.org/>) were used for basic 3D image processing and 3D visualization of segmentation results, respectively.

The key parameters of the proposed algorithm were determined experimentally on five datasets, which were selected randomly from Sliver07-train. SLIC supervoxels were generated with step $S = 3$ and compactness $m = 20$. The number of components K in foreground/background GMM models was set to $K = 5$. In graph cuts segmentation,

(VOE) [%], relative volume difference (RVD) [%], average symmetric surface distance (ASD) [mm], root mean square symmetric surface distance (RMSD) [mm], and maximum symmetric surface distance (MaxD) [mm]:

the n -links of the graph were specified to 6-neighborhood connectivity. The weighting factors of the energy function were set to $\alpha = 0.01$ and $\beta = 100$.

In the experiments, a larger value of K had no obvious improvement on the segmentation result. A smaller value of K led to imprecise segmentations in some cases, due to the presence of pathologies like tumors, similar intensity with adjacent tissues, and low CT image quality.

In the supervoxel partition step, when S was set to a larger step size, fewer number of supervoxels would be generated, resulting in a lower segmentation accuracy. However, as shown in Table 3, when $S = 3$, the number of the liver VOI supervoxels was suitable, on an average of $46 \times 53 \times 51$ supervoxels. A smaller value of S led to more supervoxels, but it would take a longer time for supervoxel generation.

4. Results

Figure 6 illustrates an example of liver segmentation using the proposed method. Figures 6(a) and 6(b) show the 3D abdominal region extraction using MIP and thresholding methods. In Figures 6(c)–6(f), the orange rectangles show the extracted liver VOI, which was obtained by analyzing the histogram and by using adaptive thresholding and morphological methods. Figure 6(c) shows the largest liver slice with initial liver region in blue and background region in red. Foreground and background seed points were sampled in the liver and background region, respectively. To avoid oversegmentations of heart and kidney, additional background seeds were extracted on the heart and kidney slices, as shown in Figures 6(e) and 6(f). The segmented liver after postprocessing is shown as the yellow contour in Figure 6(f). Figure 6(g) shows the 3D liver volume reconstructed by using surface rendering algorithms in VTK.

TABLE 3: Examples of reduction of the volume size: train = Sliver07-train datasets; test = Sliver07-test datasets.

| Data | Original CT [voxels] | Resampled CT [voxels] | Abdominal region [voxels] | Liver VOI [voxels] | VOI supervoxels [supervoxels] |
|-----------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|-------------------------------|
| Train number 5 | $512 \times 512 \times 319$ | $197 \times 197 \times 212$ | $163 \times 175 \times 200$ | $121 \times 148 \times 192$ | $40 \times 49 \times 64$ |
| Train number 11 | $512 \times 512 \times 388$ | $200 \times 200 \times 258$ | $172 \times 136 \times 148$ | $172 \times 136 \times 148$ | $57 \times 45 \times 49$ |
| Train number 14 | $512 \times 512 \times 129$ | $245 \times 245 \times 427$ | $217 \times 162 \times 138$ | $137 \times 160 \times 138$ | $45 \times 53 \times 46$ |
| Test number 1 | $512 \times 512 \times 502$ | $253 \times 253 \times 267$ | $216 \times 219 \times 122$ | $129 \times 183 \times 122$ | $43 \times 61 \times 40$ |
| Test number 4 | $512 \times 512 \times 165$ | $253 \times 253 \times 328$ | $234 \times 202 \times 176$ | $143 \times 178 \times 169$ | $47 \times 59 \times 56$ |

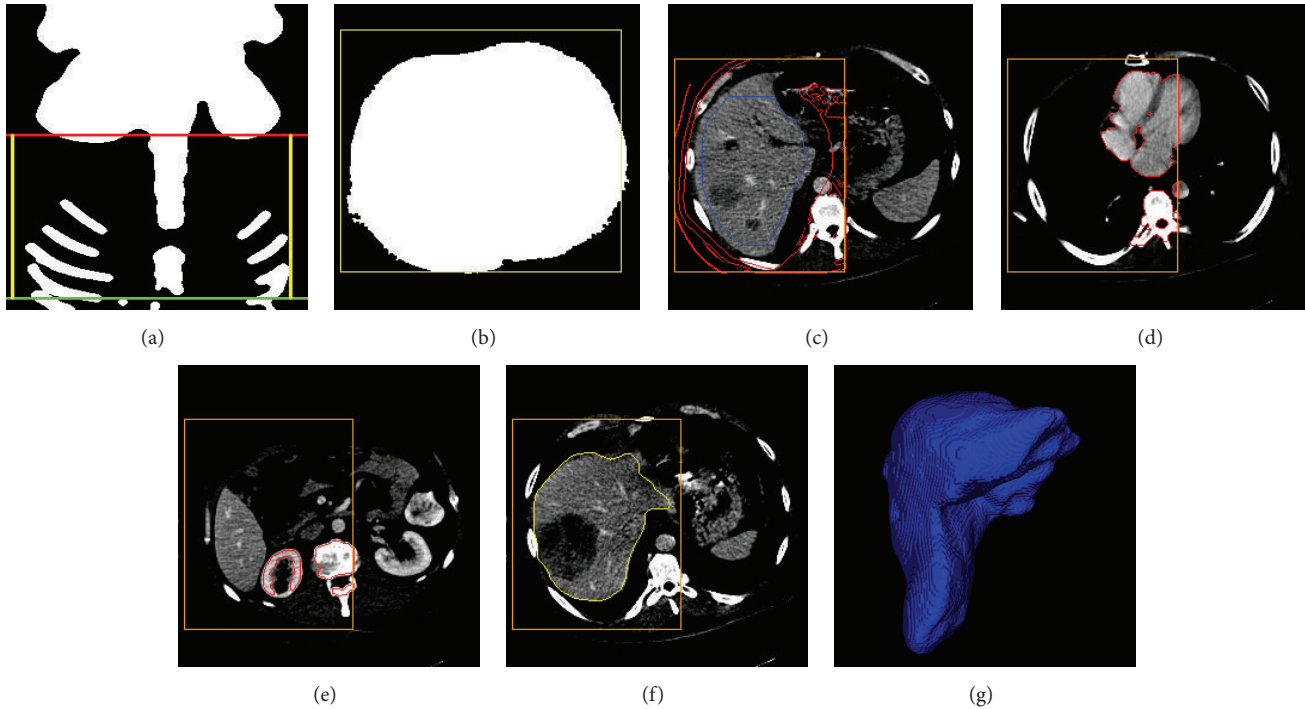


FIGURE 6: An example of liver segmentation using the proposed method. (a) and (b) show the 3D abdominal region extraction. (c) The selected largest liver slice with initial liver region in blue contour and background regions in red contours. From (c) to (f), orange rectangles show the liver VOI. (d) and (e) show the heart and kidney slices, respectively, with additional background regions in red contours. (f) The segmented result after postprocessing in yellow contour. (g) The reconstructed 3D liver volume.

Figure 7 illustrates some representative slices of the segmentation results from three Sliver07-train datasets compared with expert segmentations, where the red contours indicate the expert segmentations and the yellow contours are the segmentation results by the proposed method. Each column shows slices of one specific case in axial, coronal, and sagittal directions, respectively. It can be seen that the livers extracted by the proposed method are comparable to the expert segmentations.

In Figures 7(g) and 7(i), our method succeeded in separating the liver from heart and kidney to avoid oversegmentations. In the third column of Figure 7, the presence of tumors in the liver also could be handled. However, there were still some small oversegmentation or undersegmentation errors occurring near the liver boundaries, mainly due to the low contrast near the liver boundaries. Undersegmentation errors occurred at the tip of the liver (Figure 7(c)) and high intensity regions near the liver edges (Figures 7(b) and 7(e)).

Oversegmentation error was mainly at the place of vena cava (Figure 7(h)).

Table 3 shows examples of the reduction of volume size after each step before graph cuts segmentation. It can be seen that the extraction of liver VOI and generation of supervoxels can significantly reduce the volume size, therefore reducing computational cost.

For Sliver07-train datasets, Table 4 shows the comparative results of the proposed method with the traditional graph cuts method (TGC) [32] and the semiautomatic graph cuts method on SLIC supervoxels (SGC). SGC was similar to our method except that there was no step of liver VOI extraction and largest liver slice selection, and the extraction of seed points was conducted by manual interaction on one or two axial slices. In addition, for SGC, the original CT image was firstly converted to the 8-bit image in the intensity range $[0, 255]$ by using linear contrast stretching with a fixed intensity window. Then, the 8-bit image was used in the

TABLE 4: Comparative results for the Sliver07-train datasets: auto = automatic; semi = semiautomatic; TGC = traditional graph cuts method; SGC = semiautomatic graph cuts method on supervoxels.

| Method | Auto | Runtime [s] | VOE [%] | RVD [%] | ASD [mm] | RMSD [mm] | MaxD [mm] | Score |
|------------|------|-------------|---------|---------|----------|-----------|-----------|-------|
| TGC | Semi | 180–240 | 11.52 | 6.02 | 1.72 | 3.74 | 31.92 | 57.5 |
| SGC | Semi | 30 | 9.73 | 2.00 | 1.64 | 3.26 | 27.82 | 66.1 |
| Our method | Auto | 21 | 7.54 | 4.16 | 0.95 | 1.94 | 18.48 | 75.2 |

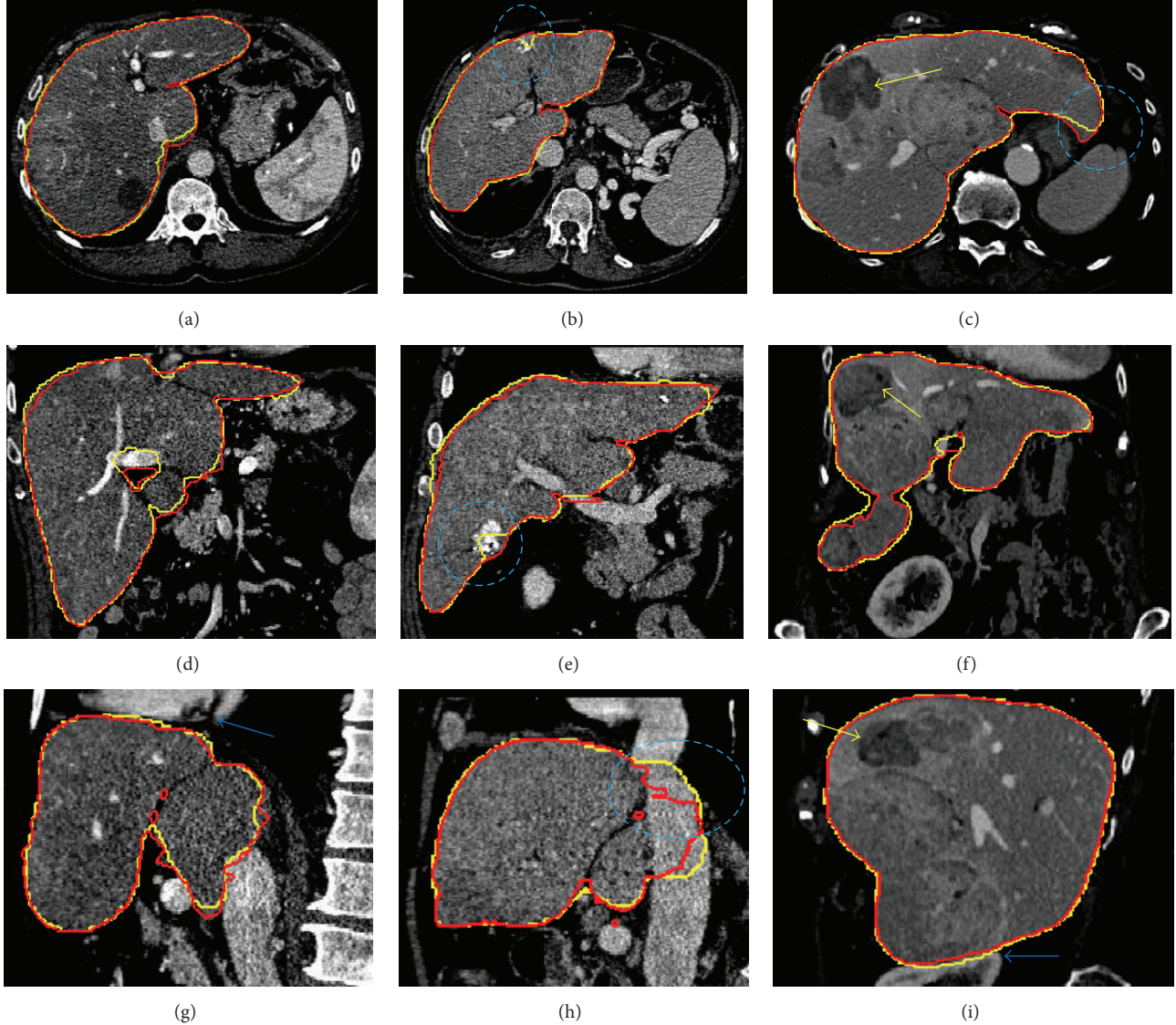


FIGURE 7: Illustrations of the segmentation results. Each column shows slices of one case in the axial, coronal, and sagittal directions, respectively. The contour of the ground truth is in red. The contour of the segmented liver by the proposed method is in yellow.

procedures of supervoxel generation and GMM modeling. For Sliver07-train datasets, the proposed method had a VOE and RVD of 7.54% and 4.16% and an ASD, RMSD, and MaxD of 0.95 mm, 1.94 mm, and 18.48 mm, respectively. The total runtime was 21.21 s on average. It can be seen that, compared with TGC, our method had a great improvement on the time efficiency, which was achieved mainly by using the supervoxels. Compared with SGC, the proposed automatic

segmentation method took less runtime and could obtain more robust segmentation without user interaction.

The evaluation results on the Sliver07-test datasets are presented in Table 5, comparing results of the proposed algorithm and human experts. The average VOE of 7.87%, RVD of 1.31%, ASD of 1.286 mm, RMSD of 2.498 mm, and MaxD of 23.563 mm were obtained, which were comparable to the performance of human experts. The total runtime, including

TABLE 5: Quantitative evaluation of segmentation result for Sliver07-test datasets.

| Data number | Runtime [s] | VOE [%] | RVD [%] | ASD [mm] | RMSD [mm] | MaxD [mm] |
|-------------|-------------|---------|---------|----------|-----------|-----------|
| 1 | 27.04 | 9.37 | 6.68 | 1.54 | 2.77 | 26.84 |
| 2 | 30.76 | 9.22 | 5.41 | 1.37 | 2.23 | 21.66 |
| 3 | 31.47 | 5.98 | -1.68 | 1.14 | 2.02 | 19.73 |
| 4 | 36.88 | 7.16 | 2.97 | 1.19 | 2.38 | 18.02 |
| 5 | 23.93 | 7.60 | -2.21 | 1.37 | 2.85 | 30.51 |
| 6 | 27.13 | 8.94 | 1.34 | 1.75 | 4.29 | 41.46 |
| 7 | 10.31 | 7.08 | 1.16 | 1.07 | 2.12 | 25.46 |
| 8 | 16.91 | 6.90 | 2.46 | 1.10 | 1.80 | 15.19 |
| 9 | 7.52 | 9.57 | -1.00 | 1.31 | 2.58 | 22.10 |
| 10 | 12.03 | 6.89 | -2.03 | 1.02 | 1.94 | 14.66 |
| Average | 27.04 | 7.87 | 1.31 | 1.29 | 2.50 | 23.56 |

TABLE 6: Comparative results with previous methods for the Sliver07-test datasets.

| Method | Runtime [s] | Auto | VOE [%] | RVD [%] | ASD [mm] | RMSD [mm] | MaxD [mm] | Score |
|--------------|-------------|------|---------|---------|----------|-----------|-----------|-------|
| Peng [10] | 180 | Semi | 6.10 | -0.00 | 0.90 | 1.60 | 16.80 | 81.8 |
| Chen [23] | 60-180 | Semi | 4.16 | 3.53 | 0.72 | 1.26 | 16.70 | 81.5 |
| Peng [24] | 120-180 | Semi | 4.58 | 1.08 | 0.68 | 1.45 | 16.88 | 83.4 |
| Li [20] | 285 | Auto | 6.24 | 1.18 | 1.03 | 2.11 | 18.82 | 77.9 |
| Platero [18] | 261 | Auto | 7.60 | -0.50 | 1.30 | 2.90 | 24.70 | 70.5 |
| Our method | 27 | Auto | 7.87 | 1.31 | 1.29 | 2.50 | 23.56 | 71.4 |

the time taken for VOI extraction, largest liver slice selection, supervoxel generation, and graph cuts segmentation, was 27.04 s on average.

Table 6 shows the quantitative comparative results of the proposed method with previous methods for the Sliver07-test datasets. It can be seen that, compared with automatic segmentation methods, semiautomatic methods might have a higher accuracy. In automatic graph cuts based segmentation methods, Li [20] obtained the highest accuracy by using deformable graph cuts with SSM based initialization. The proposed method was comparable to the performance of human experts and the method of Platero [18]. Compared with other methods, the proposed method significantly improved the time efficiency of liver segmentation.

5. Discussion

5.1. Contributions of This Study. In this paper, an efficient supervoxel-based graph cuts method was proposed for automatic liver segmentation from CT images.

To reduce computational time and memory requirement, the proposed method effectively incorporated the graph cuts method with supervoxels partition. The SLIC method was applied to generate supervoxels in the liver VOI. By constructing the graph over supervoxels, the number of nodes in the graph was reduced significantly. Therefore, the computational complexity of graph cuts was decreased.

An automatic liver VOI extraction method was introduced, which contained two steps. First, the region of abdomen was extracted by using MIP and thresholding methods. By analyzing the region of bones in the coronal MIP image and finding the largest lung slice in the extracted

lung region, unnecessary nonliver slices were removed. By measuring the in-plane abdominal bounding box in the axial MIP image, more nonliver voxels were excluded. Second, the liver VOI was extracted from the region of abdomen by applying adaptive thresholding and morphological methods. The adaptive threshold values were calculated by estimating the liver intensity range in the volumetric histogram. According to the prior knowledge of the liver's location and size, an initial binary liver mask was generated and the liver VOI was determined based on the binary liver mask. Voxels outside the liver VOI were removed.

For the purpose of extracting foreground and background seed points automatically, the largest liver slice and heart/kidney slices were extracted to generate foreground and background regions by using adaptive thresholding methods. On the largest liver slice, foreground seeds were sampled inside the liver region without missing any separate parts of liver. To tackle the problem of separating the liver from heart and kidney with similar intensities, additional background seeds were selected on the heart and kidney slices. Then, sufficient seed points were used to estimate the intensity distributions of foreground and background.

5.2. Comparison with Previous Work. Compared with the traditional graph cuts method (Table 4), the proposed approach can realize 3D liver segmentation in an automatic and fast manner. The main improvements come from (1) constructing the graph over supervoxels instead of the voxels, which can greatly reduce the number of nodes in the graph, therefore improving the efficiency of the energy minimization procedure, and (2) generating seed points automatically instead of manual interaction. The foreground and background seed

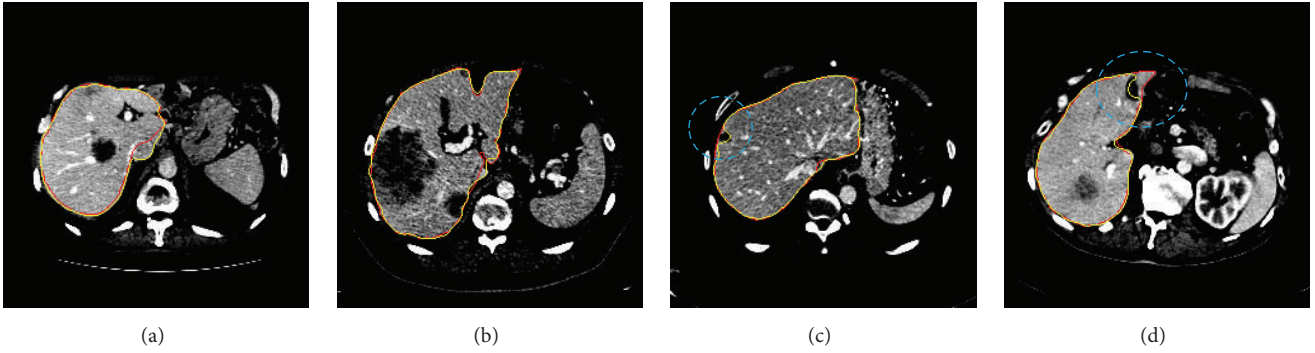


FIGURE 8: Illustrations of the segmentation results compared with expert segmentations from Sliver07-train datasets. The contour of the ground truth is in red. The contour of the segmented liver by the proposed method is in yellow.

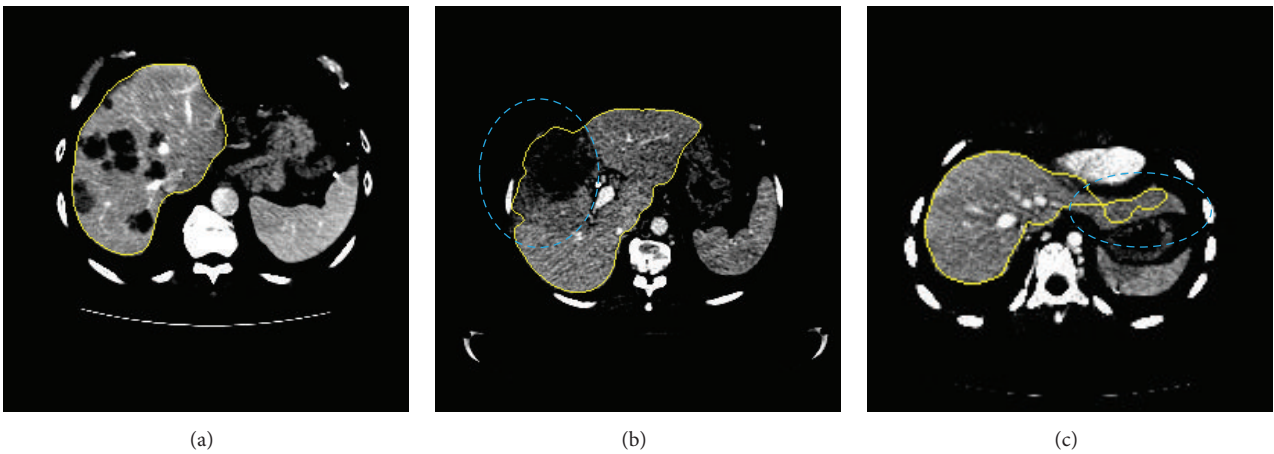


FIGURE 9: Illustrations of the segmentation results from Sliver07-test datasets. The contour of the segmentation result of the proposed method is in yellow.

points were sampled in the initial liver region and background regions, respectively. This can avoid interoperator or intraoperator variability and improve the robustness of the proposed method.

As shown in Table 6, in the semiautomatic graph cuts based methods, Chen [23] incorporated domain knowledge of the intensity, location, and spatial connectivity into the optimization framework, and Peng [24] incorporated appearance and intensity constraints for both the liver and tumor. Unlike Chen's and Peng's methods, both the liver VOI and foreground/background seed points were extracted automatically in our method. Prior knowledge of liver and histogram information were introduced such that the extraction procedures of VOI and seed points were conducted in an adaptive way.

In the automatic graph cuts methods, Platero [18] combined probabilistic atlases with graph cuts, and Li [20] incorporated SSM with graph cuts. Compared to the initialization procedure based on atlases or shape models in Platero's and Li's methods, the proposed method generated hard constraints for graph cuts segmentation based on histogram analysis and simple morphological operations. Also, the nodes of the graph corresponded to supervoxels instead of the voxels in the liver VOI. Although the segmentation

accuracy of our method was lower than that of Li's SMM initialized graph cuts method [20], it was comparable to the performance of human experts. Moreover, the proposed method was faster than other graph cuts based methods. Such a fast segmentation speed may be required in practical applications like computer-assisted ablation needle trajectory planning in image-guided hepatic ablation therapy [40]. Needle trajectory planning involves segmentation of all relevant structures including liver, tumor, hepatic vessels, and other surrounding organs [41, 42]. In cases like CT-guided hepatic ablation procedures, the time efficiency of needle trajectory planning is highly demanded. Therefore, it would be beneficial if the liver segmentation procedure was performed in a fast way with a low computational cost.

Compared with other previous methods based on SSMs, probabilistic atlases, and deformable models, no training process of atlases or shape models was needed for initialization in the proposed method. As shown in Figure 7, although there was a great variation in the intensity, shape, and position of liver among different datasets, the proposed algorithm correctly segmented liver from surrounding organs or tissues.

5.3. Limitations. Figure 9(c) shows an inadequate segmentation at the long and thin boundaries of the liver. This is mainly

due to the similarity in intensity and the shrinking bias of graph cuts minimization. Surface refinement work could be applied to get a better result. Also, the energy function of graph cuts may be improved to obtain better performance on the tiny parts of liver.

On most pathological cases, our approach can handle the presence of tumors in liver, as shown in Figures 8(a), 8(b), and 9(a). However, in Figures 8(c), 8(d), and 9(b), tumors near the liver boundaries still might be misclassified. For such cases, algorithms for generating automatically additional foreground seeds on both the liver and tumor regions could be incorporated in future work.

Another limitation of this study is that the number of datasets for evaluation is small. More datasets should be used to evaluate the performance of the proposed method in the future.

6. Conclusions

This paper presented a new method for automatic liver CT image segmentation using SLIC supervoxels-based graph cuts. The liver VOI was extracted to reduce computational cost. Foreground and background seed points were generated automatically on the largest liver slice and on additional heart and kidney slices to avoid undersegmentations or oversegmentations. By supervoxel partition and liver VOI extraction, the computation time of the proposed method was reduced to less than one minute, while accurate segmentation result can be obtained. In the future, methods for reducing undersegmentation or oversegmentation errors need to be designed to improve the accuracy of the proposed method, especially for pathological cases with tumors. Also, more datasets for evaluation are needed.

Competing Interests

The authors declare that there are no competing interests regarding the publication of this paper.

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