

TEL AVIV UNIVERSITY

The Iby and Aladar Fleischman Faculty of Engineering

The Zandman-Slaner School of Graduate Studies

**Kidney Segmentation and Renal Lesion
Detection in 3D CT**

A thesis submitted toward a degree of
Master of Science in Electrical and Electronic Engineering

by

Neta Blau

October 2017

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This research was carried out in The
School of Electrical and Electronic Engineering

This work was carried out under the supervision of
Prof. Nahum Kiryati

October 2017

Abstract

Here comes the 1-2 pages long abstract of the thesis in English.

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Acknowledgments

I would like to thank my ...

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1 Introduction

In a radiologist's line of work time is of the essence. The flux of imaging cases that require human attention is often overwhelming, forcing radiologists to increase their rate of diagnosis. This, in turn, inevitably reduces the quality of their work, causing some pathologies to go undetected or misrepresented. The consequences may vary between suboptimal treatment and considerable damage to patients' health. Automated tools that aid radiologists with their tasks are therefore highly desirable. Automation of medical image analysis have been the focus of much attention in recent years, supported by the rapid development of computer vision technologies.

1.1 Renal lesions

Among the overlooked pathologies are renal lesions. Their asymptomatic nature makes their detection largely incidental, but not less important. Renal masses can be cystic or solid, benign or malignant, and they range from simple renal cysts to cancerous tumors.

Simple renal cysts are benign and asymptomatic fluid collections in the kidney, commonly occurring in aging kidneys. They are usually found incidentally in patients undergoing abdominal imaging for other reasons [24]. In most cases, no treatment is required. However, cysts must be reported by the radiologist and sometimes require follow-up as they may evolve in time. At the other end of the scale, renal cancer is a lethal disease that manifests as malignant renal lesions. Renal cancer causes approximately 14,000 deaths in the United States every year [11][23]. The majority of renal cell carcinomas, which account for 90% of kidney cancers, are asymptomatic [14], and they as well are usually found incidentally on

noninvasive imaging performed for unrelated problems [12][19][9]. Early diagnosis of kidney cancer significantly improves survival rates, which fall dramatically once the cancer has spread beyond the kidney [15]. In this work, we address the detection of both types of renal lesions as incidental findings in a real clinical setting.

1.2 Computed tomography (CT)

Among the most frequent imaging procedures carried out in hospitals is the three-dimensional (3D) computerized tomography (CT). CT scans are widely used for preventive medicine and screening for diseases in different parts of the human body. Almost 79 million CT scans were performed in the U.S. during the year of 2015, 245 scans per 1000 population [20].

To produce a CT scan, an X-ray generator rotates around the bed where the patient is lying. The x-ray beams shot by the generator are picked up by X-ray detectors positioned on the opposite side of the circle. When a full rotation is completed, a 2D image is constructed. By moving the bed along the rotation axis, a series of cross-sectional images is obtained, composing the 3D scan. The voxels in the scan are displayed according to the mean attenuation of the tissues that they correspond to on a scale from +3071 (most attenuating) to -1024 (least attenuating). This scale is known as the Hounsfield unit (HU) scale. Water, for example, has an attenuation of 0 HU, while air is approximately -1000 HU. Depending on the diagnostic task, the scan can be viewed as images in each of the anatomical planes - axial, sagittal or coronal. The scans are often anisotropic, and the resolution of each slice (also referred to as the pixel spacing), and the spacing between slices vary from scan to scan.

In order to allow higher visibility, an intravenous (IV) contrast agent is often administered. The contrast material can help enhancing the contrast between a lesion and its surrounding tissue. The time interval between the contrast administration and the image acquisition is determined according to the purpose of the imaging and the investigated organs. To see renal lesions better and detect them more easily, the images will usually be acquired at the portal phase of enhancement, 70-80 seconds after the initiation of contrast injection.

1.3 Kidney segmentation

When a radiologist goes over a CT scan and processes it, he first identifies the patient's organs and then looks for any irregularities or abnormalities in them. Adopting the radiologist's strategy, our first step towards detecting renal lesions is to identify and segment the kidneys. Automatic kidney segmentation is a challenging task due to the high inter-patient variability of the kidney shape and location. This task is made harder still by the gray-levels inhomogeneity of the kidney, its similarity to surrounding organs such as the liver, and the possible presence of pathologies.

Numerous methods for kidney segmentation in CT scans have been investigated in the literature. Several atlas-based approaches were proposed in which the segmentation is obtained by registering one or more labeled atlases to the unlabelled target volume [26][4][27]. However, volume registration is very time consuming, and the computation time increases when a multi-atlas approach is used to improve the segmentation accuracy. The increasing availability of medical images enabled the utilization of supervised-learning approaches for organ detection and segmentation. Several proposed methods employ regression forests for organ localization [6][5], classification forests for initial segmentation [5] and joint classification-regression forests for spatially-structured segmentation [8]. These methods are much more efficient than the atlas-based approaches, but their drawback is their dependency on the selected input features.

In recent years, deep convolutional neural networks gained great success in various computer vision tasks, including object detection and segmentation. Given sufficient labeled data, deep networks can learn complex features and representations of the objects to be segmented, thus eliminating the need to manually define and select features. With modern GPUs, the convolution computations have been accelerated significantly, making deep networks a viable option for real-life segmentation tasks. Several studies utilized fully convolutional networks (FCN) for end-to-end segmentation in medical images, and specifically in CT scans. Zhou et al. [28] trained an FCN to segment 19 different organs in 2D slices from all three anatomical planes of a CT scan. The resulting segmentations were later fused by majority voting into a 3D segmentation. Hu et. al [10] employed a 3D

FCN for liver, spleen and kidney segmentation. This step was followed by a time-implicit multi-phase level-set algorithm for segmentation refinement. Cascaded FCNs were also applied to segment the liver and its lesions [3][7]. Thong et. al [25] applied convolutional networks for kidney segmentation in contrast-enhanced CT scans and reported promising results.

1.4 Cyst detection

Provided with a high quality segmentation of the kidneys, we can address the problem of detecting renal lesions, starting with simple renal cysts. Due to their random origins, lesions vary significantly in size, location and quantity. Owing to the homogenous fluid consistency of simple cysts, one may think that simple thresholding of the kidney area can suffice for cyst detection. However, being part of the urinary system, other fluid collections are normally present in the kidneys, making it difficult to distinguish renal cysts from healthy functioning parts of the kidneys such as the renal pelvis and renal calyces.

Only few studies were published on automatic tools for detection or segmentation of renal cysts. Battiato et al. [2] proposed a method to refine a cyst segmentation given an initial boundary marked by the radiologist. The refinement is done in the annotated slice using a bank of filters and is then propagated to the other slices. However, in this method, the cysts must be detected by the users, and their intervention is required for the segmentation. Piao et al. [21] proposed to segment cysts using fuzzy C-mean clustering, but did not report any qualitative results. An algorithm proposed by Badura et. al [1] employs thresholding and morphological operations for the detection of cyst candidates. The candidates are then classified into cysts and non-cysts based on 3D shape-related features. This approach produced promising results, but the algorithm was evaluated on a fairly small data set of 16 patients with 25 cysts.

1.5 Tumor detection

Unlike renal cysts, cancerous tumors are inhomogeneous in their consistency and are more similar to the kidneys in their HU values, thus making it harder to detect

them. Summers et al. [22] employed thresholding and morphological operations to segment renal cysts and solid tumors and estimate their volume. However, the work focused on the quantitation of the lesions and detection rates were not reported. Kim and Park [13] detected renal lesions using texture analysis and homogeneous region growing, but their method was evaluated on 12 tumors only. Linguraru et al. [16] proposed a semi-automated algorithm for segmentation and quantification of renal tumors, based on fast marching and geodesic level sets. This algorithm was later extended to also classify the lesion type using support vector machine [17]. This method generated accurate lesion segmentations but user intervention was still required for their detection. Another solution, proposed by Liu et al. [18], relies on geometric changes on the kidney surface for cyst candidate detection. Shape features are then utilized for the classification of the candidates. This algorithm was validated on a large data-set but is limited to detecting only exophytic lesions, and the reported false-positive rate (15 per patient) was high.

1.6 Outline of Thesis

So far, no fully automatic algorithms have been shown to detect simple renal cysts reliably on a large data set. Here, for the first time, we report an algorithm that achieves this goal and demonstrate its performance. The algorithm is constructed from two main modules, a kidney segmentation module and a cyst detection module. In section «<sec...>» the kidney segmentation module is described, followed by a quantitative evaluation of its performance. Using its output as a starting point, the cyst detection module is then described in section «<sec...>», followed by the overall performance of the algorithm. Additionally, in section «<sec...>» we relate our efforts to automatically detect cancerous renal tumors, based on the kidney segmentation results. Finally, in section «<sec>» we conclude our work and propose some next steps...

2 Kidney segmentation

2.1 Methods

Describe the kidney segmentation method in details: the input/output, the data preprocessing steps and the FCN architecture.

2.2 Data-set

Describe the data set used: randomly selected, number of patients/scans, different resolutions, ground truth annotations, heterogeneous data (including sick kidneys with different pathologies).

2.3 Experiments and results

Describe the training process and validation. Show qualitative and quantitative results.

2.4 Discussion

Discuss the results of the kidney segmentation: can they be improved by gathering more data / better quality of ground truth / 3D networks instead of 2D / handling the cyst presence in the ground truth differently.

3 Cyst detection

3.1 Identification of cyst candidates

Describe the candidate selection algorithm: the input/output, the use of distance maps, and the chosen parameters (thresholding range, minimal radius).

3.2 Classification of cyst candidates

Describe the candidate classification method in detail: the 2.5D patch construction, the data preprocessing (thresholding + Gaussian filter), the network architecture, and the removal of redundant cysts.

3.3 Data-set

Describe the data-set used: randomly selected, number of patients/scans, number of detected cysts, different resolutions and ground truth annotations.

3.4 Experiments and results

Describe the training process and validation. Show qualitative and quantitative results. Elaborate on the different false-positives that were detected and the false-negatives that were missed. Show that the minimal radius parameter controls the trade-off between TP and TP.

3.5 Discussion

Discuss the results of the cyst detection: can they be improved by gathering more data / better quality of ground truth / 3D networks instead of 2D / other approaches (e.g. shape priors).

4 Tumor detection

4.1 Data-set

Describe the data-set used: number of patients/scans with number of detected tumors, different resolutions and ground truth annotations. Show the high variability of the tumors.

4.2 Experiments and results

Describe different methods and approaches we tried and the unsatisfying results...

4.3 Discussion

Discuss the unsolved problem of tumors segmentation - what are we missing? Is it only a matter of gathering more data?

5 Conclusions

What we did, what we achieved, possible next steps.

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תקציר

במסגרת מחקר זה, בוצעו מדידות מהירות של גלי זרימה שונים בעלי פאזה חוזרת במגוון תדרים ומהירויות ממוצעות בעזרת שיטת ה PIV המאפשרת מדידת שדה מהירות במישור בפאזות שונות לאורך מחזור הזרימה ושיטת ה 3D-PTV המאפשרת עקיבה אחר חלקיקים במרחב. מאמצי הגזירה על דופן

אוניברסיטת תל אביב

הפקולטה להנדסה ע"ש איבי ואלדר פליישמן

בית הספר לתארים מתקדמים ע"ש זנדמן סליינר

תכונות שדה הזרימה ומאפייני מאמצי גזירה

בזרימות פועמות בעלות פאזה חוזרת

חיבור זה הוגש כעבודת מחקר לקראת התואר "מוסמך אוניברסיטה" בהנדסה מכנית

על ידי

ישראל ישראלי

העבודה נעשתה בבית הספר להנדסה מכנית

בהנחיית ד"ר ישראלה ישראליוביץ

אייר תשע"ב

אוניברסיטת תל אביב

הפקולטה להנדסה ע"ש איבי ואלדר פליישמן

בית הספר לתארים מתקדמים ע"ש זנדמן סליינר

כותרת של התיזה

חיבור זה הוגש כעבודת מחקר לקראת התואר "מוסמך אוניברסיטה" בהנדסה מכנית

על ידי

ישראל ישראלי

אייר תשע"ב