
NESTED 3D NEURAL NETWORKS FOR KIDNEY AND TUMOR SEGMENTATION

A PREPRINT

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ABSTRACT

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1 Introduction

Kidney cancer has an high number of diagnosed cases each year, being invasive techniques the most common treatment[5]. Machine learning techniques have been proven to provide valuable tools in medical imaging, increasing the accuracy in diagnosis for several diseases[2]. In recent years, deep learning techniques have increased the performance of classical machine learning approaches in several imaging problems, including medical image analysis[4]. In this work, we propose a deep learning approach for kidney cancer detection and segmentation, as participants of the Kits19 challenge.

The code will be freely available for others to use on their own data at <https://github.com/naclet/Kits19> after the publication of this work.

2 Methodology**2.1 Data**

The data used in this study belongs to the Kits19 challenge[3], which consists of 544 patients who underwent radical nephrectomy or partial nephrectomy, reviewed at the University of Minnesota, with a selection of 300 patients that met the inclusion criteria. These patients have annotated CT abdominal images, with two annotations: kidneys and tumors. The data was separated in training and testing data, and the annotations were only available in the 210 patients training set to the challenge participants.

2.2 Preprocessing

The proposed network architecture, described in section 2.3, receives an image of size 168^3 . In order to obtain an image with this dimensions for all our training cases, they had to be preprocessed to a region of interest (ROI) with these dimensions. The preprocessing steps include: a resampling of the image to an isotropic pixel size of $2 \times 2 \times 2$ mm and cropping the CT image to a resolution of 168^3 from the center of the image, if the CT image was smaller, the ROI is padded with zeros. Finally the ROI is normalized so that the intensity values of all the images range from 0 to 1.

2.3 Neural Network Architecture

The proposed deep learning architecture is 3D V-Net [REF] with fourteen convolutional layers. This network is first trained to learn how to segment the kidneys, after that the weights are saved and the same architecture is used to learn how to segment the tumors. The architecture of this network is shown in figure 1. Each convolutional layer has relu as the activation function and filters size $3 \times 3 \times 3$, with the exception of the last convolutional layer which uses a sigmoid activation function. Each convolutional layer in the decoding phase has batch normalization to improve the generalization of the network. Finally, the dimensions of the input and output images is 168^3 .

The network uses stochastic gradient descent as the optimization method, the number of steps per epoch is 100 and the number of steps when computing the metrics on the validation set is 20. The training is stopped after 1000 iterations or when the loss function on the validation set does not improve after 90 iterations.

The difference between training the network to learn how to segment kidneys vs tumors, lays on the loss function being used. In both cases, the metric that is minimized is the negative Sorensen-Dice coefficient (DSC) [1]. But, in the case of the training for tumors, the DSC only takes into account the pixels that are inside the kidneys. This helps the network to avoid learning only zero-valued weights.

$$loss = -\frac{2 \sum_{i=1}^N p_i t_i}{\sum_{i=1}^N p_i + \sum_{i=1}^N t_i + \varepsilon} \quad (1)$$

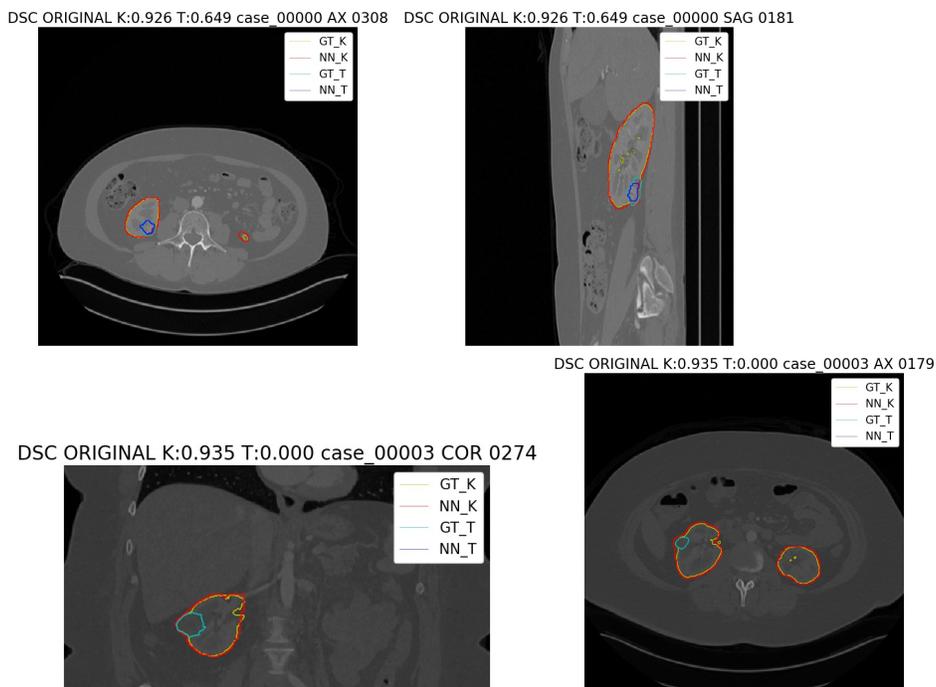


Figure 2: NN architecture

- [3] Nicholas Heller et al. “The KiTS19 Challenge Data: 300 Kidney Tumor Cases with Clinical Context, CT Semantic Segmentations, and Surgical Outcomes”. en. In: *arXiv:1904.00445 [cs, q-bio, stat]* (Mar. 2019). (Visited on 05/23/2019).
- [4] Geert Litjens et al. “A survey on deep learning in medical image analysis”. In: *Medical Image Analysis* 42 (Dec. 2017), pp. 60–88.
- [5] Maxine Sun et al. “Treatment management of small renal masses in the 21st century: a paradigm shift”. eng. In: *Annals of Surgical Oncology* 19.7 (July 2012), pp. 2380–2387. ISSN: 1534-4681. DOI: 10.1245/s10434-012-2247-0.