

# Maximal Contrast Adaptive Region Growing for CT Airway Tree Segmentation

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**Abstract.** In this paper we propose a fully self-assessed adaptive region growing airway segmentation algorithm. We rely on a standardized and self-assessed region-based approach to deal with varying imaging conditions. Initialization of the algorithm requires prior knowledge of trachea location. This can be provided either by manual seeding or by automatic trachea detection in upper airway tree image slices. The detection of the optimal parameters is managed internally using a measure of the varying contrast of the growing region. Extensive validation is provided for a set of 20 chest CT scans. Our method exhibits very low leakage into the lung parenchyma, so even though the smaller airways are not obtained from the region growing, our fully automatic technique can provide robust and accurate initialization for other methods.

**Key words:** CT, airways, segmentation, multi-tolerance, adaptive region growing, median filter

## 1 Introduction

CT scans are frequently used for pulmonary disorder assessment [1]. Pathologies that could affect sufficient lung function include tumors, pulmonary embolism, atelectasis, pneumonia, emphysema, asthma, bronchiectasis, and many others. Certain lung diseases can be diagnosed based on airway wall thickness measurements, diameter, branching geometry and rate of tapering. CT is currently the only readily accessible, relatively noninvasive technique that is capable of providing airway tree quantitative structural data in vivo [2].

Traditionally, analysis of CT chest scans was performed manually by skilled radiologist who recognized areas of abnormal airway properties in consecutive slices of the examined scan. However, analyzing about 400 slices covering the

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chest area is very tedious and too cumbersome for everyday clinical use. Moreover, manual analysis performed by the radiologist was only qualitative estimation of airway abnormalities without accurate quantification of pathological changes.

Even though airway tree abnormalities can be detected based on 2D slices, the ability to extract a full 3D model of the airway tree from a 3D image has several key advantages. For example, slice based measurements can be inaccurate if the airway is not perpendicular to the slice. Also, information available from the slices is deprived of useful context, making it harder for a radiologist to keep track of the generation number of an airway, the structure of nearby airways or the overall shape of a segment [3].

There has been a number of efforts to try to delineate the airway tree in chest CT scans. Airway tree segmentation is a complex task, mainly due to inhomogeneous grey level of the voxels located inside the bronchial lumen, artefacts caused by blood vessels adjacent to airway walls and changes of intensity levels along airway walls.

Many airway segmentation methods use region-growing algorithms, which attempt to separate air and soft tissue voxels using an HU (Hounsfield Unit) threshold [4–8]. Region growing is fast and assumes no prior knowledge of the shape or size of the airways. Choosing an appropriate global HU threshold is difficult, however, as the lungs are filled with air and misclassifying a single wall voxel can allow the segmentation to leak into the lung parenchyma.

Other methods make use of grey-level morphological operators [9–12], or front wave propagation schemes [13–15], to impose structural properties derived from a priori anatomical knowledge.

Region growing is the preferred method for initializing several of the aforementioned algorithms, being a fast and intuitive technique [3,10,11]. Some recent segmentation techniques perform region growing stages on images derived from the intensity of the scan, like the morphological gradient [16], or the posterior of a classification stage [17]. Thus, fully automatic region growing approaches become mandatory, specially those that can provide as much information as possible for the following algorithmic stage, with very low leakage.

Few fully automatic region growing methods have proven successful in these tasks over a significant variety of imaging devices. One of the most common techniques for threshold determination was proposed by Mori et al. [6]. Their approach, widely known as explosion-controlled region growing, tries to determine optimal thresholds by detection of sudden volume increase in the segmented region. The difficulty lies in defining how much volume increase is considered explosion, as compared to normal region growth. In our work, we also propose a self-assessed region growing algorithm. Our assessment strategy is founded on a previous normalization stage, and makes use of an iteratively computed contrast measure. In our method there is no need to define any confidence margins, as the segmented region is determined precisely by the evolution of the proposed contrast measure.

Apart from manual trial-and-error adaptive threshold selection [18], some self-assessed adaptive region growing strategies have been proposed in the past, in a variety of application contexts.

In their work [19], Hojjatoleslami and Kittler proposed a method based on finding the global maxima for two different contrast measures which they computed iteratively, as intensity-decreasing pixels were added to the segmented region. The success of the assessment was founded on the assumption that maximal contrast occurred on region boundaries, which is a reformulation of approaches assuming that the variation of the gray values within regions is smaller than across regions, an inherent assumption in all region growing techniques [20]. Unfortunately, the exhaustivity of their per-pixel approach entailed very low computational efficiency, aggravated by their multiple complex peripheral measures. Revol-Muller et al. [21] used morphological measures to assess the multiplier of the adaptive range in region growing. Instead of computing their assessment function for every pixel addition to the region, they sampled the function for an evenly-spaced set of values.

In our method we propose an assessment function based on a simple measure of the evolving contrast for the region growing sequence. To make this approach computationally feasible in 3D, we produce only evenly-spaced samples of this function, along the values of the assessed parameter defined on the normalized dynamic range of the image. This sampling strategy dramatically reduces computational complexity while preserving most critical values.

Globally, our goal is to provide fast automatic segmentation based only on the location of the upper part of the trachea. Our automatic segmentation algorithm extracts a large fraction of the visible airways with few false positive branches. For some applications, however, even this accuracy rate can be insufficient. Our method can then be considered as an initialization stage for other refining strategies, in a wide variety of imaging devices, and with no significant leakage, as suggested by our experimental results.

## 2 Method

### 2.1 Airway Lumen Intensity Model

Since our goal is providing a mechanism for airway segmentation with minimal user intervention, we have established a model that takes into account their intensity distribution in CT images.

We model then our object of interest as a connected region whose pixel intensities are sampled from a Gaussian distribution with unknown mean and standard deviation. We know that our region of interest is surrounded by other tissues derived from other, sometimes adjacent, intensity distributions. Although common in the literature, this assumption for the intensities is rarely met in practice, in the sense that the intensity distributions of tissues are only approximately Gaussian, as can be inferred from direct observation of histograms. Besides, partial overlap between adjacent distributions often occurs. For dealing with these inconveniences we propose the use of an assessment function, that

is to be evaluated along a sequence of region growing stages (region growing sequence).

## 2.2 Segmentation Algorithm

**Normalization and Denoising** Since our method was conceived for images from a wide range of scanners and acquisition protocols, we have developed a normalizing stage that accounts for such variability. As we will introduce later on, for the self-assessed region growing stage of the algorithm, we require the input intensity dynamic range to be normalized with respect to some parameter estimates of the objective intensity distribution.

In the following equations in which we describe the normalization process,  $N$  is a cubic neighborhood of radius  $R$  around the seed,  $\mathbf{x}$  is a voxel position,  $f(\mathbf{x})$  is the intensity for voxel at  $\mathbf{x}$ ,  $\bar{f}_N$  is the mean intensity estimate in  $N$  and  $|N|$  is the cardinality of  $N$ . Moreover,  $\sigma_{f_N}$  is the estimated standard deviation for intensities in  $N$ ,  $K$  is a constant parameter, and  $f(\mathbf{x})$ ,  $f'(\mathbf{x})$  are the input and output intensities for the non-linear mapping described below.

$$\bar{f}_N = \frac{1}{|N|} \sum_{\mathbf{x}_k \in N} f(\mathbf{x}_k) \quad , \quad (1)$$

$$\sigma_{f_N} = \sqrt{\frac{1}{|N|} \sum_{\mathbf{x}_k \in N} (f(\mathbf{x}_k) - \bar{f}_N)^2} \quad , \quad (2)$$

$$f'(\mathbf{x}) = \left( 1 + \exp \left( -\frac{f(\mathbf{x}) - \bar{f}_N}{\left(\frac{K\sigma_{f_N}}{3}\right)} \right) \right)^{-1} \quad . \quad (3)$$

In a first step we proceed by Gaussian distribution maximum-likelihood (ML) estimation of the mean and standard deviation as in (1-2), and then perform a non-linear normalization using a sigmoidal transfer function centered on the estimated mean as in (3). The width of the sigmoidal window extends  $K\sigma$  around the center  $\bar{f}_N$  of the mapping. For  $K = 3$  the width of the window would be enough to map 99.7% of the samples, of a Gaussian distribution with similar mean and standard deviation. Greater values of  $K$  ensures robust mapping for the estimated distribution (that of the tissues of interest). The sigmoidal mapping has been chosen because of its smoothness, and its ability to focus the output dynamic range on a given input intensity range of interest.

Finally, we perform non-linear denoising using an in-slice bidimensional median filter with kernel radius  $L$ . Other denoising schemes would be valid, always keeping in mind that stronger smoothing usually involves loss of smaller airways.

**Self-Assessed Region Growing** Departing from a normalized and filtered version of the image under study, whose intensities lie in the range  $[0, 1]$ , we apply our self-assessed contrast-maximizing algorithm. The initial region that

needs to be provided (the upper trachea) can be obtained by means of manual seeding, or by automatic detection using one of the many methods available in the literature [22, 23]. For this initial implementation we decided to simply provide the seeds manually, as described below.

Considering an initial region  $R_0$  defined by several seeds along the upper trachea, we provide the analytical description of the  $i$ -th iteration of the algorithm:

1. Update multiplier  $k_i = k_0 + i\Delta k$
2. Compute, in last iteration grown region  $R_{i-1}$ , ML estimates for the mean (available from last iteration) and standard deviation ( $\bar{f}'_{R_{i-1}}, \sigma_{f'_{R_{i-1}}}$ )
3. For every candidate voxel  $\mathbf{x}_{c_{i-1}}$  being 26-connected to  $R_{i-1}$ ,  $\mathbf{x}_{c_{i-1}} \in R_i$  if

$$f'(\mathbf{x}_{c_{i-1}}) \in \left[ \bar{f}'_{R_{i-1}} \pm k_i \sigma_{f'_{R_{i-1}}} \right] \quad (4)$$

4. Compute the assessment function  $O_i(\bar{f}'_{R_i}, \bar{f}'_{P_i})$  using the intensity average  $\bar{f}'_{R_i}$  in  $R_i$  and the intensity average  $\bar{f}'_{P_i}$  in the external perimeter  $P_i$  of  $R_i$  according to (1) and the following eqs.:

$$P_i = \{x_{c_i}\} \cap R_i^C, \quad (5)$$

$$O_i(\bar{f}'_{R_i}, \bar{f}'_{P_i}) = \left| \frac{\bar{f}'_{P_i} - \bar{f}'_{R_i}}{\bar{f}'_{P_i} + \bar{f}'_{R_i}} \right| \quad (6)$$

5. If  $O_{i-1}$  was a local maximum, when compared to  $O_{i-2}$  and  $O_i$  (only when  $i \geq 2$ ), then the algorithm stops and the output is  $R_{i-1}$ . Otherwise another iteration takes place

Of all aforementioned parameters only  $k_0$  and  $\Delta k$  are critical for the performance of the algorithm.  $k_0$  affects computational efficiency requiring a greater number of iterations before a local maximum of  $O(\bar{f}_{R_i}, \bar{f}_{P_i})$  is actually found. Therefore, its fine tuning for a specific scanner, could save some computational time. From observation of the region growing sequence, we conclude that these first iterations are typically very fast, so the improvement is frequently negligible. In what concerns  $\Delta k$ , the choice must guarantee that the assessment function is being sampled adequately in order to detect its local variations. Since the estimates for the mean and standard deviation are continually updated as the region grows, the estimates become increasingly close to the theoretical values. We argue that setting  $\Delta k$  below one tenth of 3 (which is the theoretical value multiplying the standard deviation of a Gaussian distribution for 99.7% of its samples to be included in a range of that width around the mean) is enough for the segmentation process to be able not to miss the available local maxima of the assessment function. This claim is supported by our experimental results.

### 3 Results

We have implemented our algorithm using open source medical image processing libraries, more precisely the Insight Toolkit [24] for algorithm development, and

the command line executable module infrastructure provided by 3DSlicer for fast prototyping, calibration, evaluation, and manual segmentation on real images for further validation [25]. The algorithm that we will validate, uses the following parameter values:  $R = 2$ ,  $K = 12$ ,  $\Gamma = 1$ ,  $k_0 = 1$  and  $\Delta k = 0.1$ . The values for these parameters were determined from other non-thoracic CT images, and are intended to suit any imaging conditions. No thoracic CT scans were used for the tuning of these parameters, as the algorithm was initially conceived for general purpose segmentation. The algorithm was implemented and executed in a 2 GHz Intel Core 2 Duo Windows PC with 2 GB RAM, and the average running time was  $129 \pm 27$  s.

For generation of the presented airway segmentations, the initial region was provided using three manually selected seeds along the upper trachea. The segmentation process took between 1 and 2 minutes for each dataset, including the reading/writing of the images.

The evaluation of this algorithm has taken place in comparison with other 14 algorithms using a set of 40 CT scans, with different acquisition parameters. First 20 datasets were used for training (unnecessary in our approach), and last 20 for testing. The produced segmentations were centrally evaluated by a team of trained observers. The objective was to compare performance. For this purpose, a ground truth was constructed from all available segmentations from the different algorithms, and all results were subsequently evaluated with respect to this ground truth.

Evaluation of each individual segmentation is performed in the following steps:

1. Airway segments or branches were extracted from submitted airway tree segmentations using a fast marching based algorithm [14].
2. Airway segments were evaluated visually on a set of extracted slices from both a reoriented view and a reformatted view with straightened airway centerlines.
3. Each segment was scored as "correct" or "wrong", by at least two observers. The criterion used is whether the extracted airway segment indeed belongs to the airway tree; the exact airway shape and dimensions are not taken into account.

The ground truth is then defined as the union of all valid airway segments from all submitted segmentations.

The following measurements are computed and used for comparing the submitted results:

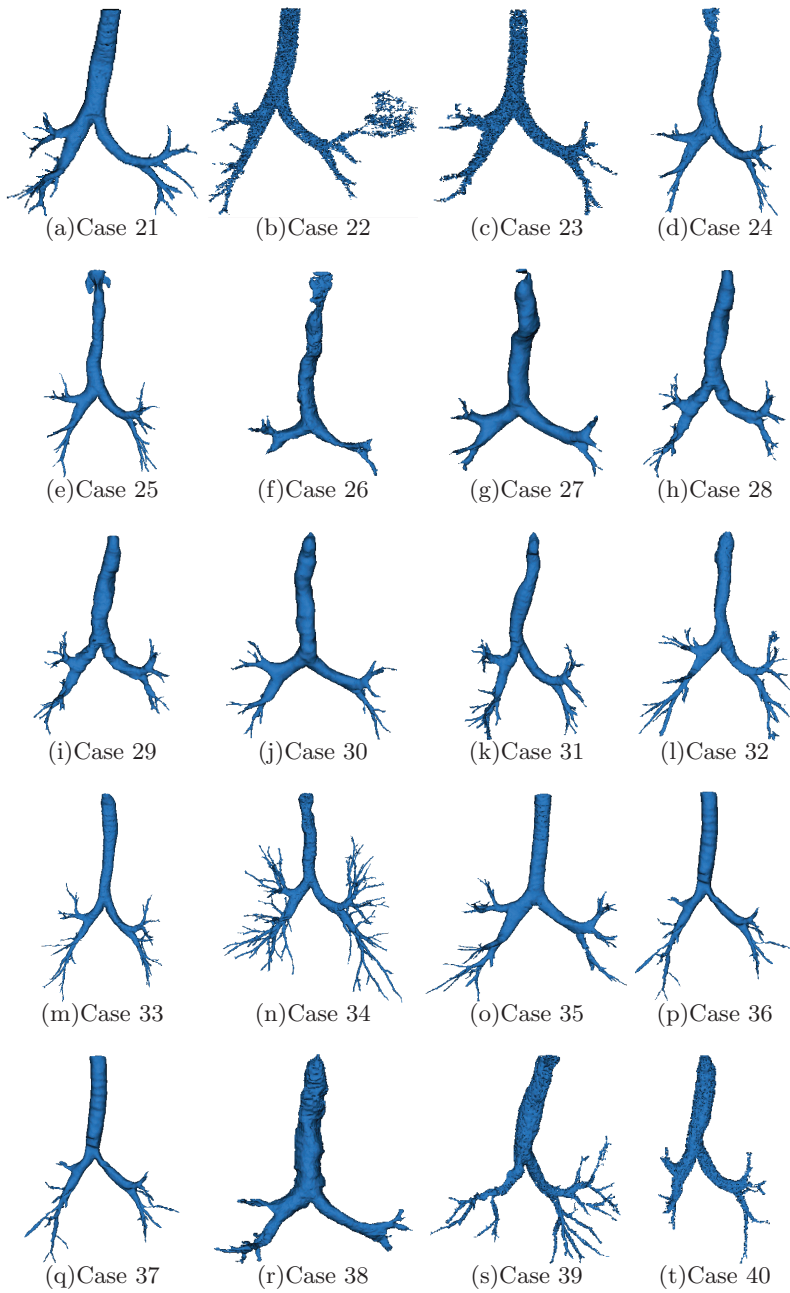
1. Branch count: The number of branches that were detected correctly. A branch is considered detected as long as the length of the centerlines is more than 1 mm.
2. Branch detected: The fraction of branches that were detected, with respect to the branches present in the ground truth.
3. Tree length: The sum of the length of the centerlines of all correctly detected branches.

**Table 1.** Evaluation measures for the twenty cases in the test set.

	Branch count	Branch detected (%)	Tree length (cm)	Tree length detected (%)	Leakage count	Leakage volume (mm <sup>3</sup> )	False positive rate (%)
CASE21	89	44.7	46.0	41.6	0	0.0	0.00
CASE22	54	14.0	37.0	11.2	8	2085.7	30.67
CASE23	33	11.6	27.3	10.5	0	0.0	0.00
CASE24	49	26.3	43.2	26.6	0	0.0	0.00
CASE25	83	35.5	63.5	25.2	0	0.0	0.00
CASE26	22	27.5	15.8	24.0	0	0.0	0.00
CASE27	35	34.7	26.0	32.1	0	0.0	0.00
CASE28	56	45.5	40.5	37.0	0	0.0	0.00
CASE29	74	40.2	44.4	32.2	0	0.0	0.00
CASE30	44	22.6	30.2	19.8	0	0.0	0.00
CASE31	77	36.0	53.7	30.6	3	31.1	0.35
CASE32	80	34.3	62.4	28.6	2	314.3	2.81
CASE33	83	49.4	56.9	38.7	0	0.0	0.00
CASE34	266	58.1	189.8	53.1	2	39.3	0.18
CASE35	112	32.6	78.4	25.3	0	0.0	0.00
CASE36	59	16.2	54.1	13.1	0	0.0	0.00
CASE37	46	24.9	39.2	22.0	0	0.0	0.00
CASE38	35	35.7	26.9	40.5	0	0.0	0.00
CASE39	93	17.9	73.8	18.0	4	65.8	0.95
CASE40	40	10.3	30.9	8.0	0	0.0	0.00
Mean	71.5	30.9	52.0	26.9	0.9	126.8	1.75
Std. dev.	51.7	13.1	36.5	11.8	2.0	466.5	6.84
Min	22	10.3	15.8	8.0	0	0.0	0.00
1st quartile	40	17.9	30.2	18.0	0	0.0	0.00
Median	58	33.4	43.8	25.9	0	0.0	0.00
3rd quartile	89	44.7	63.5	38.7	2	39.3	0.35
Max	266	58.1	189.8	53.1	8	2085.7	30.67

4. Tree length detected: The fraction of tree length in the ground truth that was detected correctly.
5. Leakage count: The number of unconnected groups of "correct" regions that were neighboring with a "wrong" region. Indicates how easy/difficult it is to manually separate leakages from the correctly detected branches.
6. Leakage volume: The volume of regions that were wrongly detected.
7. False positive rate: The fraction of the volume of regions that were detected wrongly over the volume of all detected regions.

Trachea was excluded from the branch length and branch count related measurements. For the leakage based measures, both trachea and main bronchi were excluded. In Table 1 we provide all the computed experimental measures. We also present 3D renderings for the 20 datasets used for validation, in Fig. 1. These results prove that our algorithm is able to produce reasonably complete segmen-



**Fig. 1.** (a)-(t). Segmented airways for test cases 21 through 40



tations, with very limited leakage. In most of the available test sets, the maximal contrast condition has produced fully automatic segmentations which compare fairly with those obtained using more sophisticated methods. The results were obtained with the same parameters for all different acquisition varieties, in a reduced time frame.

## 4 Concluding Remarks and Future Work

Our experimental results suggest that our modified region growing strategy, when used as an initialization stage, might benefit many airway segmentation algorithms currently available. Also, in those techniques which exploit region growing approaches applied over images obtained through processing of the image intensities, the use of our maximal-contrast stopping condition could be useful for further automatization.

Our immediate future line of work will include further study of our novel region-growing stopping criterion. We intend to develop an exhaustive comparison between previously available explosion detection and our newer approach. Further along the way, we would like to develop a hybrid method, using our contrast-assessed region growing for initialization, and adding some refining stages using morphological operations and surface interpolation. This will provide for further comparison between the different initialization strategies.

## References

1. Sluimer, I., Schilham, A., Prokop, M., Ginneken, B.V.: Computer analysis of computed tomography scans of the lung: A survey. *IEEE Transactions on Medical Imaging* **25**(4) (2006) 385–405
2. Coxson, H.O., Rogers, R.M.: Quantitative computed tomography of chronic obstructive pulmonary disease. *Academic Radiology* **12**(11) (2005) 1457–1463
3. Szymczak, A., Vanderhyde, J.: Airway segmentation by topology-driven local thresholding. In: *Progress in Biomedical Optics and Imaging - Proceedings of SPIE*. Volume 6914. (2008)
4. Mori, K., Hasegawa, J.I., Suenaga, Y., Toriwaki, J.I.: Automated anatomical labeling of the bronchial branch and its application to the virtual bronchoscopy system. *IEEE Transactions on Medical Imaging* **19**(2) (2000) 103–114
5. Summers, R.M., Feng, D.H., Holland, S.M., Sneller, M.C., Shelhamer, J.H.: Virtual bronchoscopy: Segmentation method for real-time display. *Radiology* **200**(3) (1996) 857–862
6. Mori, K., Hasegawa, J., Toriwaki, J., Anno, H., Katada, K.: Recognition of bronchus in three-dimensional X-ray CT images with application to virtualized bronchoscopy system. *Proceedings of the 13th International Conference on Pattern Recognition* **3** (1996) 528–532
7. Kiraly, A.P., Higgins, W.E., Hoffman, E.A., McLennan, G., Reinhardt, J.M.: 3D human airway segmentation for virtual bronchoscopy. In: *Proceedings of SPIE - The International Society for Optical Engineering*. Volume 4683. (2002) 16–29

8. Singh, H., Crawford, M., Curtin, J.P., Zwiggelaar, R.: Automated 3D segmentation of the lung airway tree using gain-based region growing approach. *MICCAI* **2** (2004) 975–982
9. Fetita, C.I., Prteux, F., Beigelman-Aubry, C., Grenier, P.: Pulmonary airways: 3-D reconstruction from multislice CT and clinical investigation. *IEEE Transactions on Medical Imaging* **23**(11) (2004) 1353–1364
10. Graham, M.W., Gibbs, J.D., Higgins, W.E.: Robust system for human airway-tree segmentation. In: *Progress in Biomedical Optics and Imaging - Proceedings of SPIE*. Volume 6914. (2008)
11. Sonka, M., Park, W., Huffman, E.A.: Rule-based detection of intrathoracic airway trees. *IEEE Transactions on Medical Imaging* **15**(3) (1996) 314–326
12. Aykac, D., Huffman, E.A., McLennan, G., Reinhardt, J.M.: Segmentation and analysis of the human airway tree from three-dimensional X-ray CT images. *IEEE Transactions on Medical Imaging* **22**(8) (2003) 940–950
13. Tschirren, J., Huffman, E.A., McLennan, G., Sonka, M.: Intrathoracic airway trees: Segmentation and airway morphology analysis from low-dose CT scans. *IEEE Transactions on Medical Imaging* **24**(12) (2005) 1529–1539
14. Schlathölter, T., Lorenz, C., Carlsen, I.C., Renisch, S., Deschamps, T.: Simultaneous segmentation and tree reconstruction of the airways for virtual bronchoscopy. In: *Proceedings of SPIE - The International Society for Optical Engineering*. Volume 4684 I. (2002) 103–113
15. Ginneken, B.V., Baggerman, W., Rikxoort, E.M.V.: Robust segmentation and anatomical labeling of the airway tree from thoracic CT scans. *Lecture Notes in Computer Science (including subseries Lecture Notes in Artificial Intelligence and Lecture Notes in Bioinformatics)* **5241 LNCS**(PART 1) (2008) 219–226
16. Fabijanska, A.: Two-pass region growing algorithm for segmenting airway tree from MDCT chest scans. *Computerized Medical Imaging and Graphics* (2009) Article in Press.
17. Lo, P., Bruijne, M.D.: Voxel classification based airway tree segmentation. In: *Progress in Biomedical Optics and Imaging - Proceedings of SPIE*. Volume 6914. (2008)
18. Adams, R., Bischof, L.: Seeded region growing. *IEEE Transactions on Pattern Analysis and Machine Intelligence* **16**(6) (1994) 641–647
19. Hojjatoleslami, S.A., Kittler, J.: Region growing: A new approach. *IEEE Transactions on Image Processing* **7**(7) (1998) 1079–1084
20. Haralick, R.M., Shapiro, L.G.: Image segmentation techniques. *Computer Vision, Graphics, & Image Processing* **29**(1) (1985) 100–132
21. Revol-Muller, C., Peyrin, F., Carrillon, Y., Odet, C.: Automated 3D region growing algorithm based on an assessment function. *Pattern Recognition Letters* **23**(1-3) (2002) 137–150
22. Hu, S., Hoffman, E.A., Reinhardt, J.M.: Automatic lung segmentation for accurate quantitation of volumetric X-ray CT images. *IEEE Transactions on Medical Imaging* **20**(6) (2001) 490–498
23. Sluimer, I., Prokop, M., Ginneken, B.V.: Toward automated segmentation of the pathological lung in CT. *IEEE Transactions on Medical Imaging* **24**(8) (2005) 1025–1038
24. Yoo, T.S., Ackerman, M.J., Lorensen, W.E., Schroeder, W., Chalana, V., Aylward, S., Metaxas, D., Whitaker, R.: Engineering and algorithm design for an image processing API: a technical report on ITK—the Insight Toolkit. *Studies in health technology and informatics* **85** (2002) 586–592

25. Pieper, S., Lorensen, B., Schroeder, W., Kikinis, R.: The NA-MIC Kit: ITK, VTK, pipelines, grids and 3D Slicer as an open platform for the medical image computing community. In: 2006 3rd IEEE International Symposium on Biomedical Imaging: From Nano to Macro - Proceedings. Volume 2006. (2006) 698–701