

Segmentation of Tubular Structures in Presence of Other Objects

(Research Report)

Matúš Straka



AUSTRIAN ACADEMY OF SCIENCES
ÖSTERREICHISCHE AKADEMIE DER WISSENSCHAFTEN

Overview

Motivation: Peripheral CT-A Data Processing

Challenge: Vessel Segmentation

Approach: Modified Tubular Segmentation

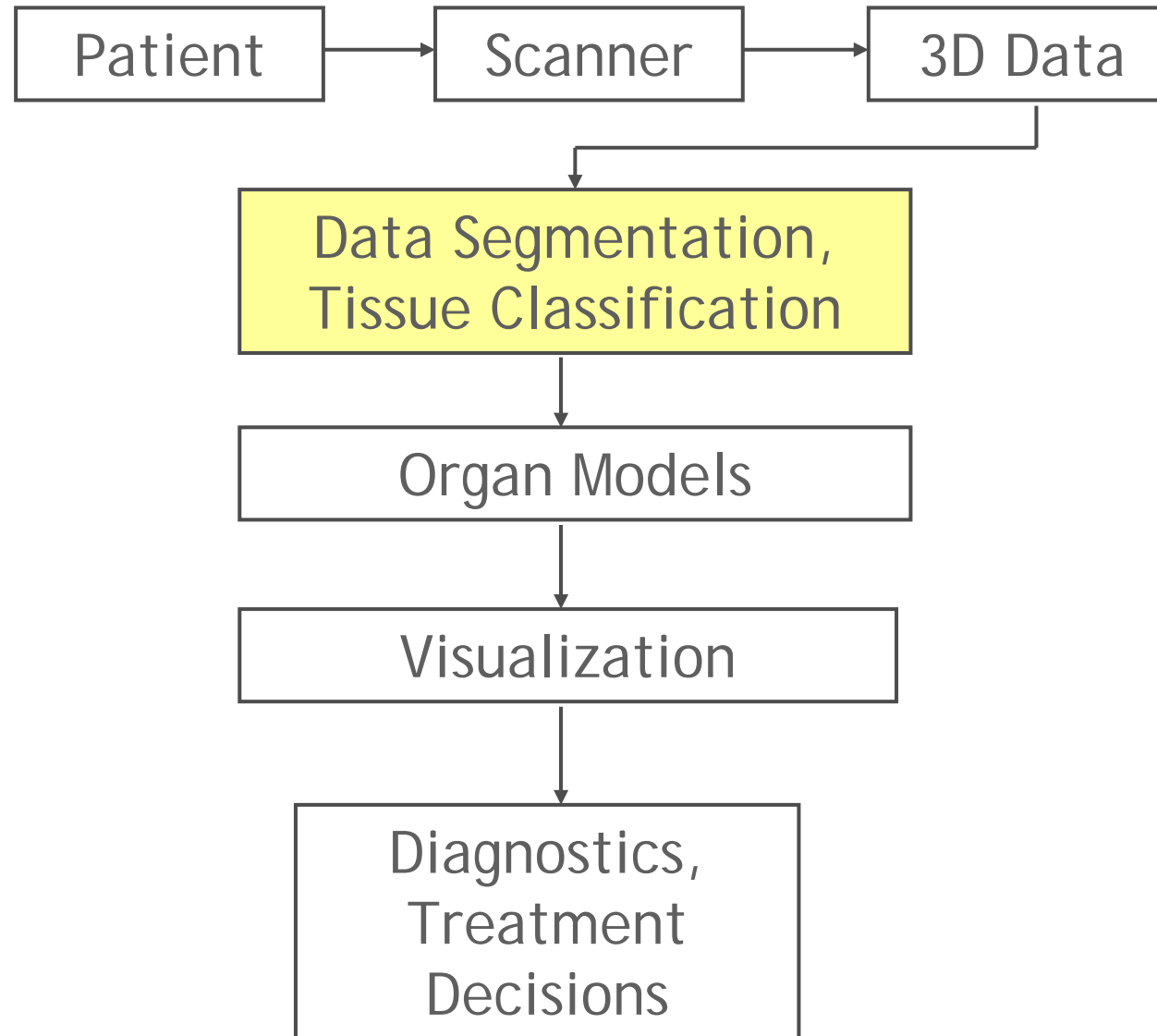
Results & Discussion

Motivation

- Angiography: vessel visualization
- Diagnostics & treatment purposes
- Imaging: helical CT scanner
- Contrast agent enhancement
- Fast, minimally invasive, cheap
- Data:
 - acquisition in 40 secs
 - 3D modality
 - ~1400 axial slices (512x512 each)
 - Element size 0.5x0.5x1.0mm



Medical Data Processing



Overview

Motivation: Peripheral CT-A Data Processing

Challenge: Vessel Segmentation

Approach: Modified Tubular Segmentation


Results & Discussion

CT-A Data Properties

- Vessels and bone denser as soft tissue
- Cortical bone denser as vessels
- Trabecular bone and vessels have similar densities
- Vessel density varies significantly
- Pathologies (calcifications, occlusions, ...)



Goal of CT-A Data Processing

- Visualization:
 - Of thick or isolated vessels - solved
 - Of thin vessels (near bones) - this talk
- Visualization with current techniques = get a model
- Model of a vessel:
 - Presence
 - Centerline & Orientation
 - Diameter/Radius

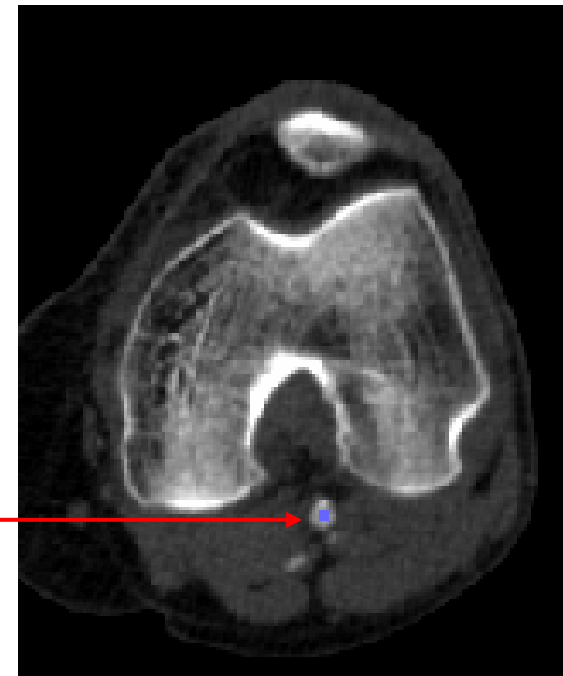
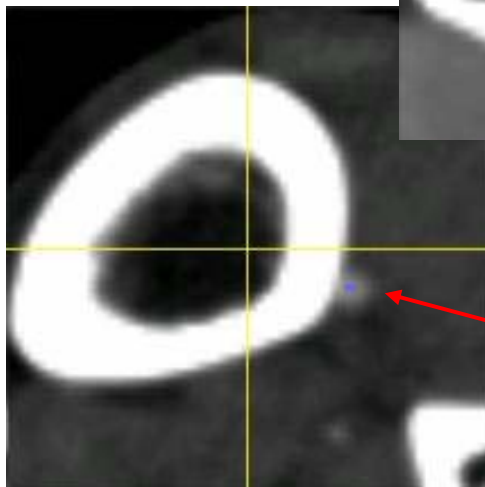
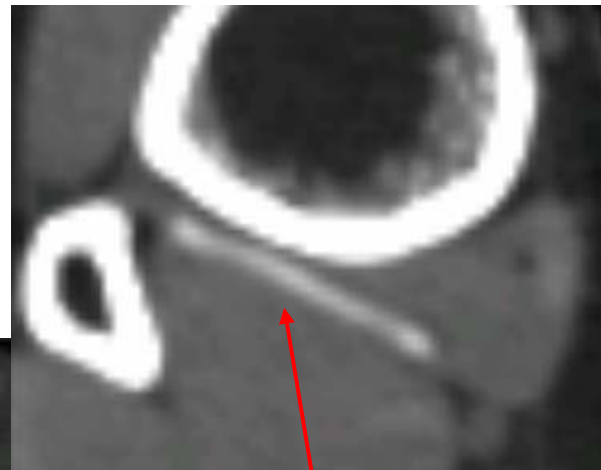
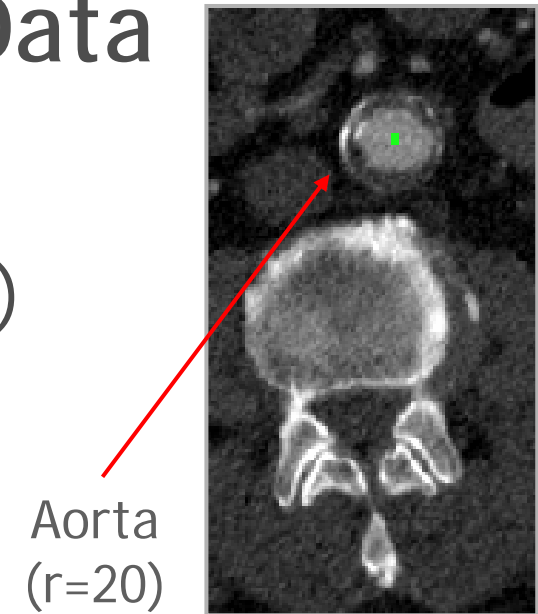
Tubular Model
- Cooperation: Model \leftrightarrow Segmentation
- Naive segmentation approach: match a tubular template

Segmentation of Tubes in 3D data

- Well-studied problem in literature
- Often easy, then
 - Thresholding, Region growing
 - Snakes, Active contours
 - Skeletonization
- Modeling based on tubular shape, tubularity defined as
 - High density change in two directions
 - Low density change in third third direction
 - Eigenvalue analysis

Vessels in Peripheral CT-A Data

- Bone and vessels lie very close
- Data blurred (partial volume effect)
- Vessel size vary ($r = 1-30$ voxels)



Problems in PeriphCTA Processing

- **Thresholding/region growing:** density uncertainty, noise
- **Snakes:** too many false edges, noise
- **Watersheds:** region merging/classification problematic
- **Hessian matrix+eigenvalues:** fails if both darker and brighter objects are close, response depends on contrast, no radius info
- **Anatomic atlases:** inter-patient anatomic variability too high
- **DoG filter:** only for larger objects, orientation info requires PCA computation
- **Distance field based skeletonization:** requires good pre_segmentation
- **Too many structures with quasi-tubular shape**

Overview

Motivation: Peripheral CT-A Data Processing

Challenge: Vessel Segmentation

Approach: Modified Tubular Segmentation

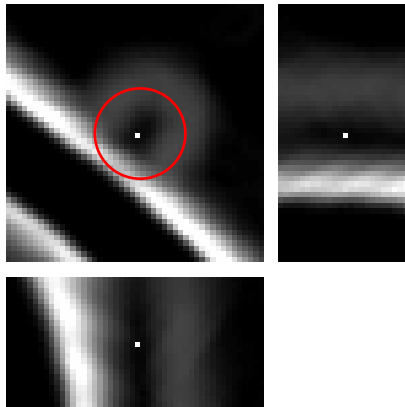
Results & Discussion

Eigenvalue analysis

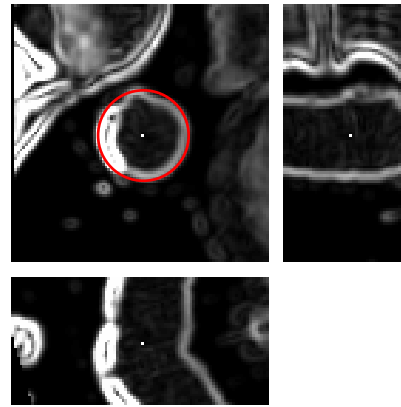
- Derives orientation of object
- Eigen decomposition:
$$\mathbf{A}\bar{x} = \lambda\bar{x}$$
- Returns 3 independent vectors (orientation) and 3 scalars (variation in given direction)
- Used in:
 - PCA (Hotelling Transform)
 - Hessian-Matrix-based analysis
 - Structure Tensor, ...
- Matrix \mathbf{A} : "2nd order element matrix" (2nd partial derivations, covariations, ...)
- No precise radius information

Radius Estimation

- Maximum of GradMag averaged on cylindrical surface (3D)
- In along-axis-resampled data
- Requires precise centre and orientation
- Works for blurred or non-ideal contours



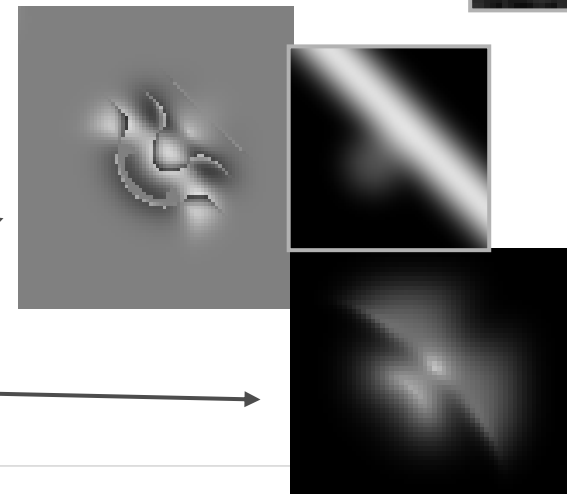
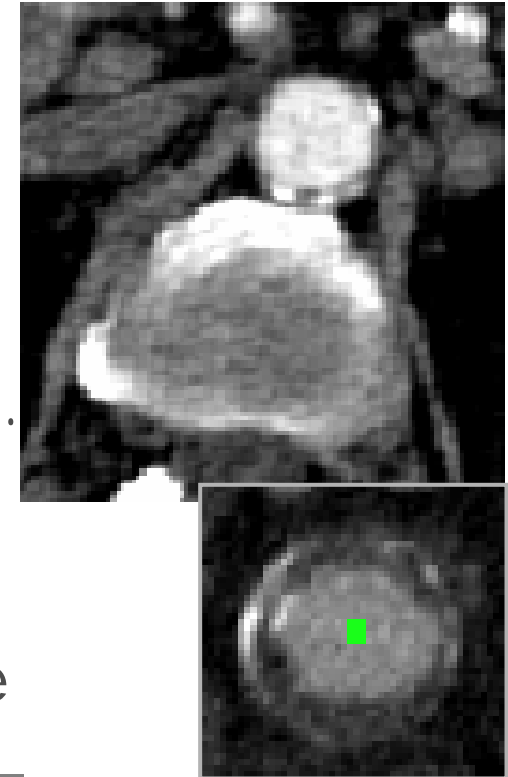
GradMag image,
 $r=2$, trilinear int.



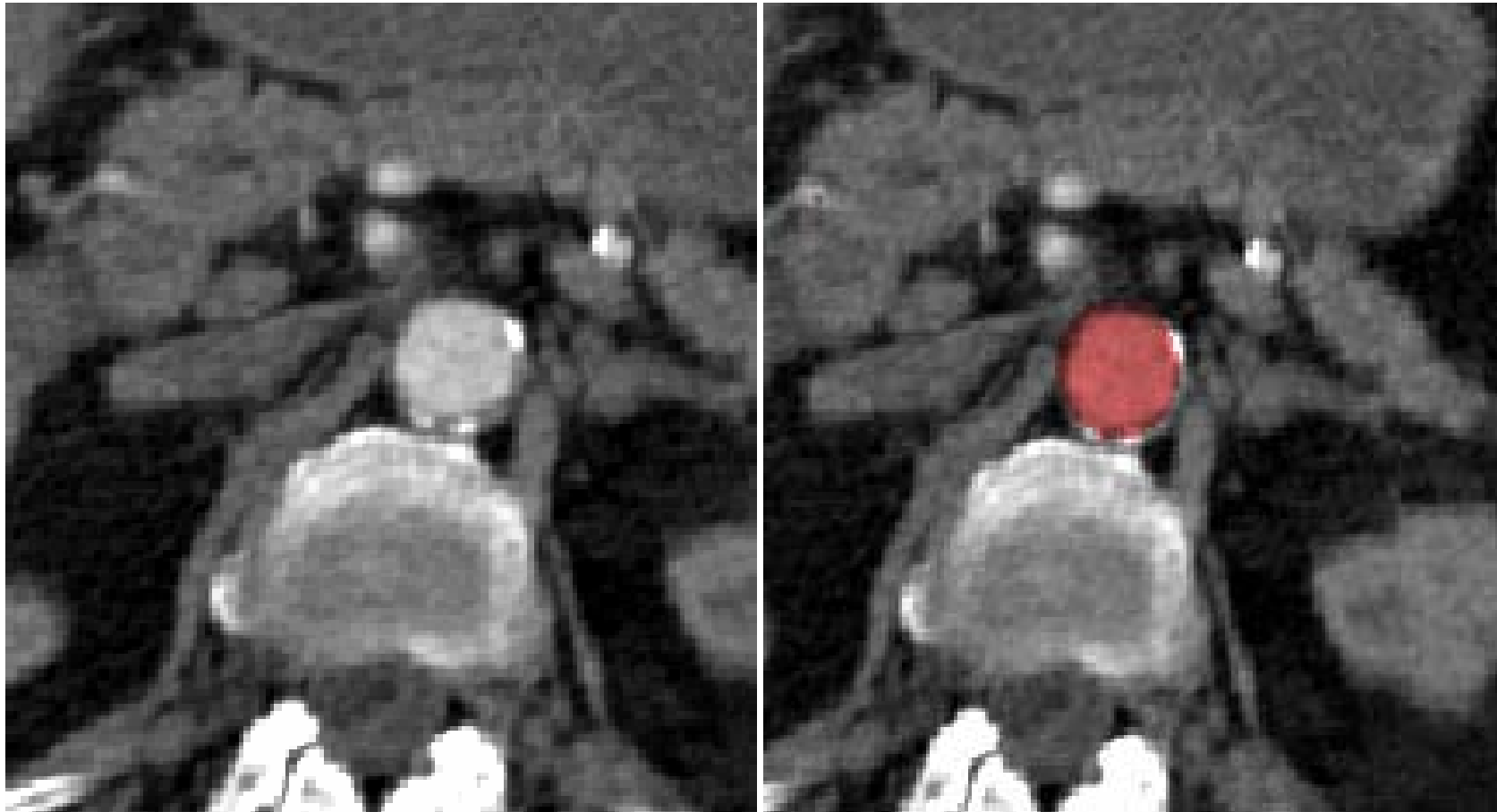
GradMag image,
 $r=22$

Segmentation of Large Vessels

- Large: a lot of samples (>50 per slice)
 - Tubular shape nicely recognizable
 - Quite easy:
 - Orientation: DoG + PCA, Hessian Matrix, ...
 - Center: PCA, CoG, maximum
 - Radius estimation: GradMag maximum
 - Tracking: "Shift along axis and recompute parameters"
 - Center estimation fails for:
 - thin vessels, close to bone
- Hessian-Matrix-Eigenvalues
- Structure-Tensor-Eigenvalues



Segmentation of Large Vessels



Segmentation of Small Vessels

- Precise center computation needed
 - For radius computation
 - For precise modeling
 - For “shift along axis and recompute” algorithm
- Orientation info and approximate position available
⇒ Template Matching feasible
- Algorithm: evaluate best position for vertical cylinder template (compute difference between template and data densities in given position)



Template Matching

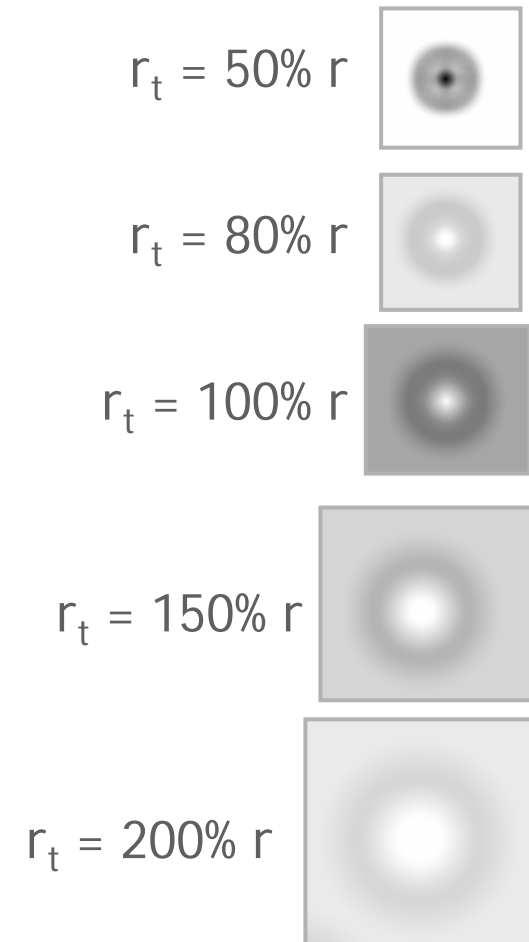
- Parameters to estimate:
 - Vessel density
 - Surrounding tissue density (bones, soft tissue)
 - Vessel template radius
- Standard template matching - two rectangular volumes - axially not symmetric
- Modified template matching:
 - Vessel density similar “inside the object”
 - Enough difference in density “outside the object”
 - Radius



Template Match. - Parameter Estimation

- Template vessel density D_V - use data from previous match
- Radius: use primary user-defined estimation
- Density for surrounding tissue not needed:
 - Check if "outside" is the density different enough
 - D_d - density diff (fore-/background)

$$f(\bar{x}_0) = \frac{1}{xyz} \sum_{x,y,z} \left(\frac{D_d - D_V - D(x,y,z)}{D_d} \right)^2 \quad \begin{array}{l} r(x,y,z) < r_t \\ r_{t2} > r(x,y,z) > r_t \end{array}$$



Results



Old system, would require ~25min
of manual segmentation



Proposed solution, would require
<5min of manual interaction

Conclusions

- Global filters not usable \Rightarrow local user interaction always needed
- Always compute result from multiple samples
- After precise segmentation, higher level models can be computed or applied

Limitations

- Requires user interaction (start point)
- Requires approximate estimation of the radius for:
 - σ in filters
 - size of search area in template matching
- No “native” stopping criteria
 - Wrong segmentation if underlying data change too much (radius)
 - Always computes results, even if nonsense
- Speed / response time (seconds)

Approach

- Eigenvector analysis gives good results for orientation
- Structure Tensor:

$$J(\bar{x}) = \int_V W() \left(\begin{array}{cc|cc|c} \frac{dG(\bar{x})}{dx} & \frac{dG(\bar{x})}{dx} & \frac{dG(\bar{x})}{dx} & \frac{dG(\bar{x})}{dy} & \dots \\ \dots & \dots & \frac{dG(\bar{x})}{dy} & \frac{dG(\bar{x})}{dy} & \dots \\ \dots & \dots & \dots & \dots & \frac{dG(\bar{x})}{dz} & \frac{dG(\bar{x})}{dz} \end{array} \right) dV$$

Thank you for your attention

matus.straka@oeaw.ac.at

<http://www.viskom.oeaw.ac.at/~straka/AngioVis>

