Optimized Strict Multiscale Frangi Prefiltering for Segmentation

Towards an automated PCSVN extraction

Luke Wukmer

Advisor: Dr. Jen-Mei Chang
Department of Mathematics and Statistics
California State University, Long Beach
lwukmer@gmail.com

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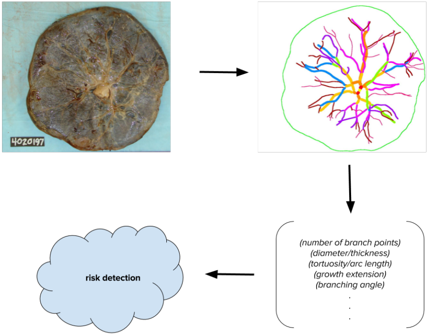
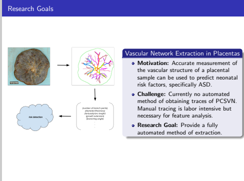
Research Goals



Vascular Network Extraction in Placentas

- **Motivation:** Accurate measurement of the vascular structure of a placental sample can be used to predict neonatal risk factors, specifically ASD.
- **Challenge:** Currently no automated method of obtaining traces of PCSVN. Manual tracing is labor intensive but necessary for feature analysis.
- **Research Goal:** Provide a fully automated method of extraction.



1. In the figure, a manual trace of the placental chorionic vascular surface network (PCSVN) is performed. This trace is measured in multiple ways. Those measurements are turned into a feature vector, which can be used to predict a risk. Refer to Boruta paper.
2. Manual tracing requires like 5 hours or something and requires training. There is some guesswork that's done in it too and some limitations in the ground truth itself (will cover later)

The Image Processing Problem

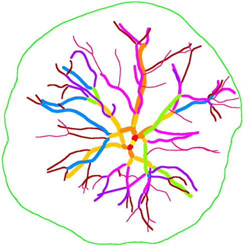
Our image domain

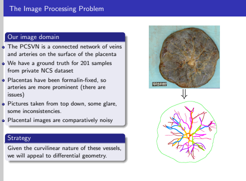
- The PCSVN is a connected network of veins and arteries on the surface of the placenta
- We have a ground truth for 201 samples from private NCS dataset
- Placentas have been formalin-fixed, so arteries are more prominent (there are issues)
- Pictures taken from top down, some glare, some inconsistencies.
- Placental images are comparatively noisy

Strategy

Given the curvilinear nature of these vessels, we will appeal to differential geometry.

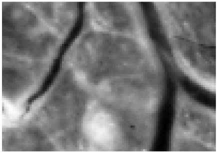
$\Downarrow$

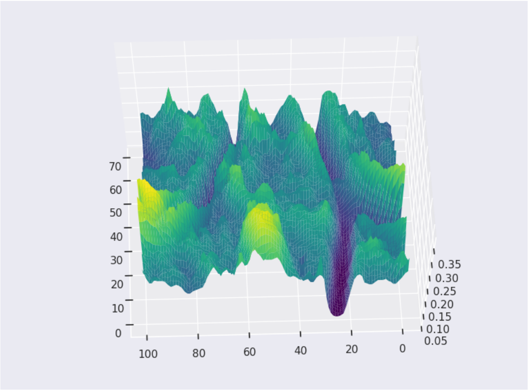


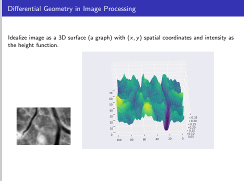


1. The surface of the placenta has a lot of changes in color/topology apart from the PCSVN so a lot of techniques that work elsewhere for vascular segmentation seem to fail here. Thus segmentation is more complicated that than say, an eyeball MRI (like original Frangi paper)
2. Mention colors are simply vessel widths (3 to 19 odds) are part of the tracing protocol. that's really outside of the scope of this thesis, but kept anytime we show a ground truth because they're pretty
3. redo this page with a placenta from your data set

Idealize image as a 3D surface (a graph) with (x, y) spatial coordinates and intensity as the height function.







1. Point of this slide is show that finding curvilinear surfaces is reasonable
2. This point is a lot clearer when you show multiple vessel widths.
3. Also way clearer later when you show the surface after Gaussian blur, not sure if I should put that here and "lie" or not.
4. Crop graph, center, etc

The Frangi Filter

Implementation Detail: Calculating Discrete Hessian

Principal Curvatures and Principal Directions

$$\mathrm{L} \in \mathbb{R}^{m\times n} \iff L \in \mathcal{C}^2\Big([0, m] \times [0, n]\Big)$$

$$\mathrm{H}(x, y) = \begin{bmatrix} L_{xx} & L_{xy} \\ L_{yx} & L_{yy} \end{bmatrix}$$

$$\kappa_i, u_i \ \text{ for } i = 1, 2$$

such that

$$\mathrm{H}u_i = \kappa_i u_i \ , \ |\kappa_1| < |\kappa_2|$$

Frangi Filter Measure

$$F\{\cdot\} = \begin{cases} 0 & \text{if} \kappa_2 < 0, \\ \exp\left(\frac{-A^2}{2\beta^2}\right)\left(1 - \exp\left(\frac{-S^2}{2c^2}\right)\right) & \text{else} \end{cases} \tag{1}$$

$$S = \sqrt{\kappa_1^2 + \kappa_2^2} \quad \text{(structureness)} \tag{2}$$

$$A = \left|\frac{\kappa_1}{\kappa_2}\right| \quad \text{(anisotropy)} \tag{3}$$

with β, c, parameters

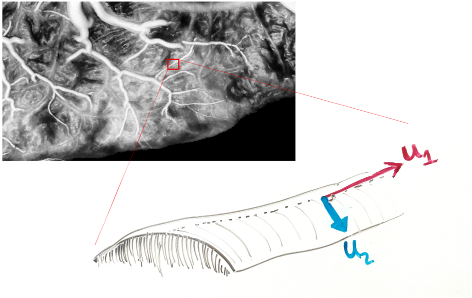
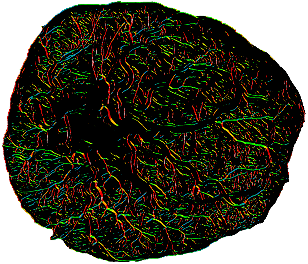


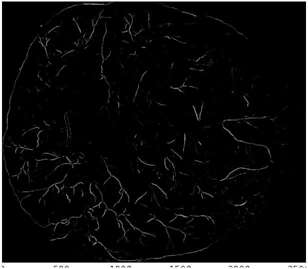
Figure: The principal curvatures (eigenvectors of the Hessian matrix) point in the direction of greatest and least curvature at each pixel

The Frangi filter [2] finds tubular structures on the surface. Corresponds to areas where κ_2 is large and κ_1 is small.

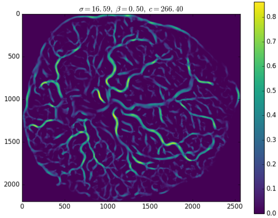
Improved Parameters for Frangi Filter



(a) Placental sample (bad parameters)

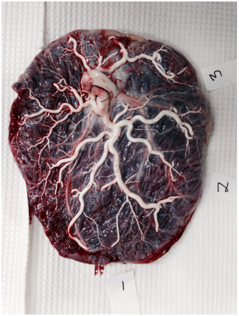


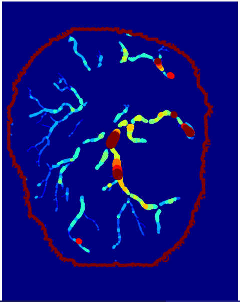
(b) improved parameters



(c) improved parameters (larger scale space)

Future Work

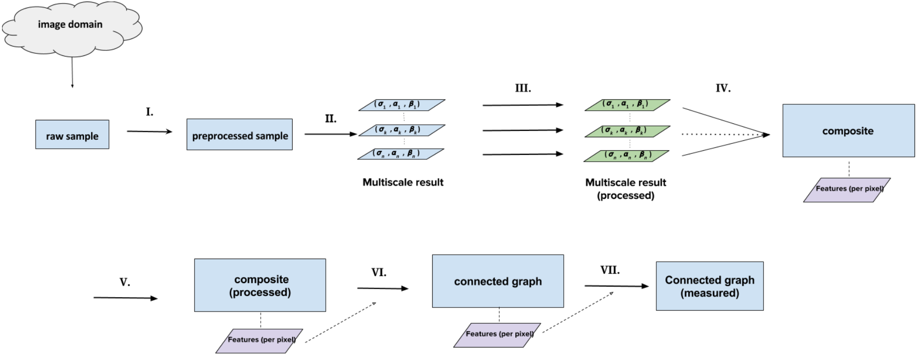




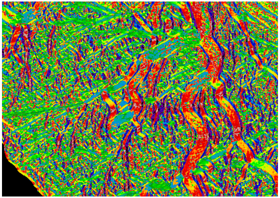
Additional Work Required

- Extend to other samples and image domains.
  - new barium samples
  - other placental samples (NCS, EARLI)
  - STARE, DRIVE retinal database
  - other curvilinear image sets?
- Adapt previous morphological filtering to improved frangi targets
- Speedups with FFT (completed!)
- Find good way to quantify results
- Graph connection problem, etc.

Appendix: Main Extraction Process

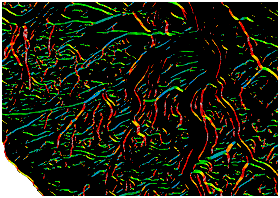


Principal Directions with Frangi Filter (Example with $\sigma = 2$)



(a) all $\theta(u_2, e_1)$

(b) Frangi targets



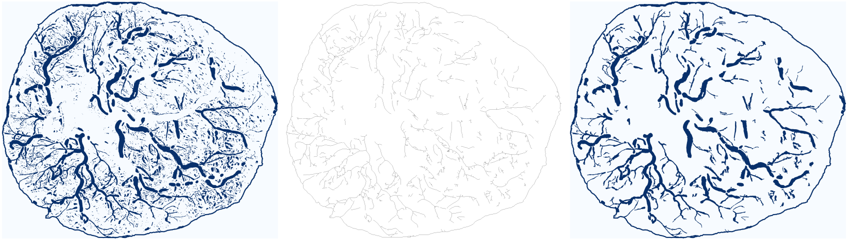
(c) frangi targets with θ

Morphological Filter

- Rectangular filter rotated by $\theta(u_1, e_1)$
- Rectangle size dependent on σ (scale size)
- Binary closing $\rightarrow$ binary opening with disk

Morphological Filtering Algorithm

a. Join all scales.
b. Skeletonize.
c. Iterate over each scale extraction and keep content only if it is connected to the skeleton.



(a) Union of all scales (b) Skeletonization of (a) (c) Sieving (a) through (b)

Figure: Morphological Filtering (smaller scales only)