

Journées du GT Géométrie Discrète et Morphologie Mathématique

# Statistical modeling of pulmonary vasculatures with topological priors in CT volumes

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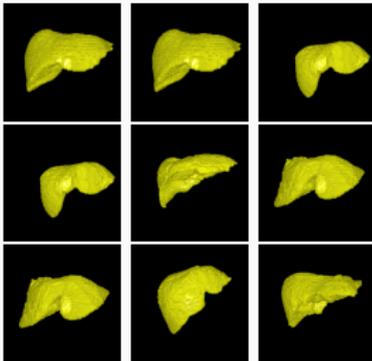
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# Statistical anatomical models

## Statistical models

- represent organ shapes and intensity distributions by a few number of parameters;
- have important roles in medical image analysis, for example,
  - segmentation,
  - super-resolution,
  - modality transform, etc.



Liver shape variations

Model 1

Model 2

# Statistical modeling of blood vessels

- **Less study on modeling blood vessels** has been made, compared to solid organs such as livers, as the shapes and the intensity distributions are highly complex.
- There are **more variations** with respect to the intensities, directions, sizes, etc.



Pulmonary CT image

3D volume patches ( $9 \times 9 \times 9$ ) <sup>2</sup>

# Various studies on modeling of blood vessels

## 1. Geometrical profile model

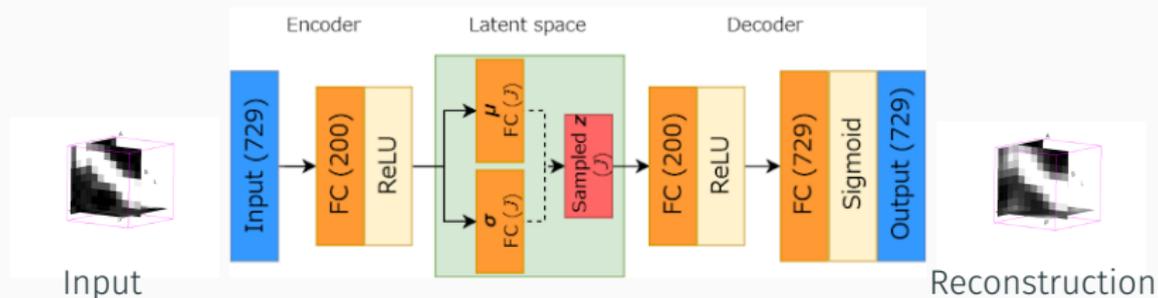
- based on cylinder whose cross sections are ellipses

## 2. Statistical intensity model

- Principal component analysis (PCA)
  - incompatible with non-linear distributions
- Manifold learning
  - incompatible with complex distributions
- Deep learning
  - Stacked AutoEncoder (SAE)
  - Variational AutoEncoders (VAE)

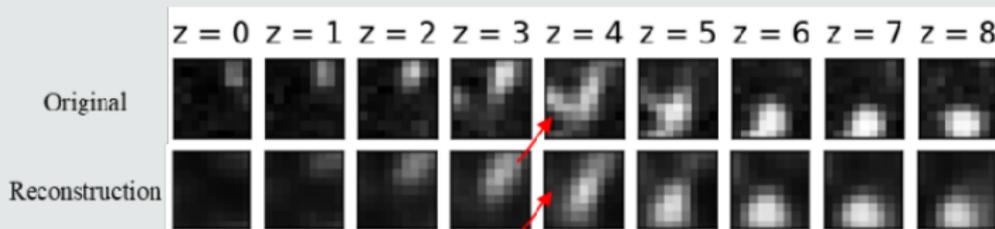
# VAE (previous work) and the problem

Volume patches containing a blood vessel are modeled based on VAEs.



## Problem

The model fails to represent anatomical features of blood vessels: **missing a bifurcation**.



## Loss function of $\beta$ -VAE and its limitation

Given a volume patch  $\mathbf{x}^{(i)}$  and its reconstruction  $\mathbf{y}^{(i)}$ , the loss function of  $\beta$ -VAE is defined by

$$\mathcal{L}_{\beta\text{-VAE}} = -\frac{\beta}{2} \sum_{j=1}^J \left( 1 + \log((\sigma_j^{(i)})^2) - (\mu_j^{(i)})^2 - (\sigma_j^{(i)})^2 \right) + \frac{1}{2} \|\mathbf{x}^{(i)} - \mathbf{y}^{(i)}\|_2^2$$

where  $\mu^{(i)}$  and  $\sigma^{(i)}$  are the mean and variance of the latent space variables, computed from the training data and  $\mathbf{x}^{(i)}$ .

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## Problem

Impossible to consider global topological features!

# Objective

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Building a **statistical intensity model of pulmonary vasculatures** in CT volume patches that incorporates **topological priors**.

More precisely, topologically correct modeling of blood vessels, which allows to represent **bifurcations**.

## Summary:

- Use a conventional network of  $\beta$ -VAE (Saeki et al., 2019), whose loss function is

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- Make use of **differentiability of Persistent Homology** (Zomorodian et al., 2005).
- Design the model such that the reconstructions meet topological priors, whose **loss function** is defined by

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Pre-training

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Fine tuning

# Persistent Homology (PH)

PH enables to quantify topological features of gray images.

- Gray image and its thresholded images



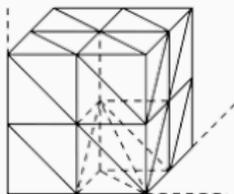
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- Convert a gray image to a simplicial (cubical) complex.



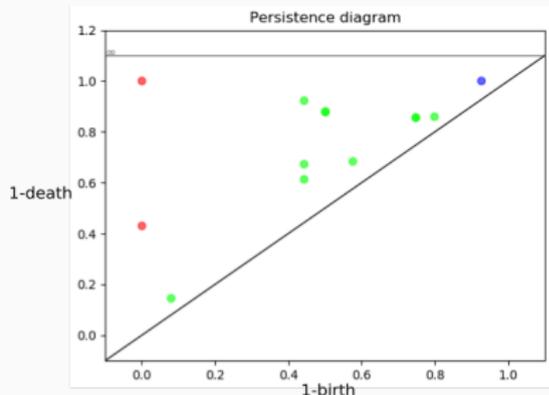
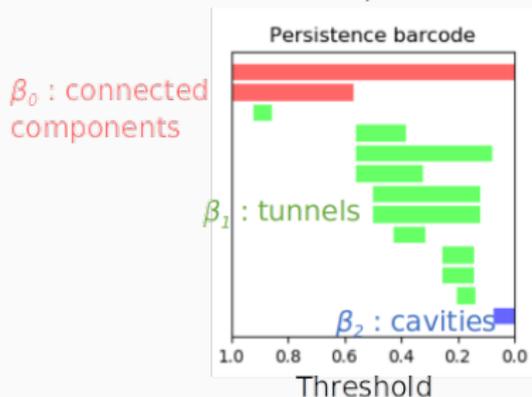
# Persistent Homology (PH)

PH enables to quantify topological features of gray images.

- Gray image and its thresholded images



- Convert a gray image to a simplicial (cubical) complex.
- Calculate its PH (persistent barcode and diagram).



## Topological loss function (Clough et al., 2020)

Assuming that the first  $\beta_k^*$  long bars are the “correct” ones (topological prior), the topological loss function is defined by

$$\mathcal{L}_{\text{topo}}(\beta_0^*, \beta_1^*, \beta_2^*) = \sum_{k \in \{0,1,2\}} \left\{ \lambda_k^* \sum_{l=1}^{\beta_k^*} (1 - |b_{k,l} - d_{k,l}|^2) + \lambda_k \sum_{l=\beta_k^*+1}^{\infty} |b_{k,l} - d_{k,l}|^2 \right\}$$

where  $\lambda_k^*$  and  $\lambda_k$  are weights.

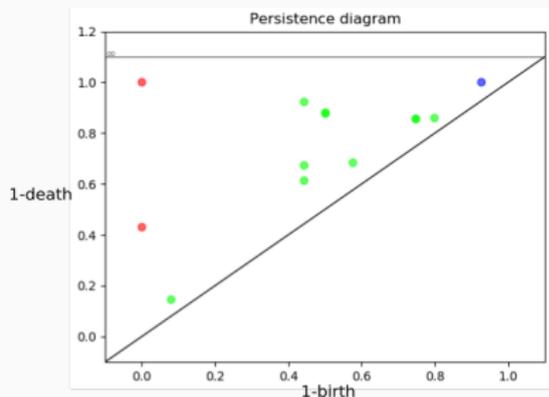
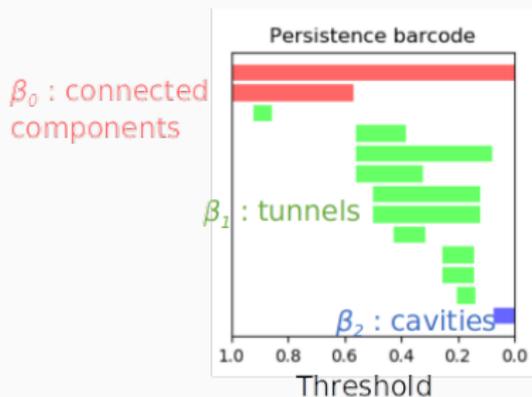
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where  $\lambda_k^*$  and  $\lambda_k$  are weights.

- **Elements to conserve:** move to (0, 1).



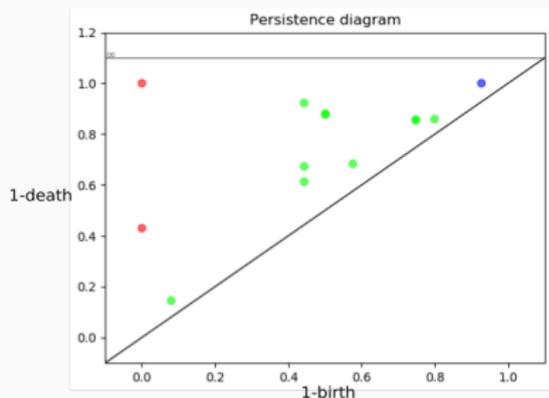
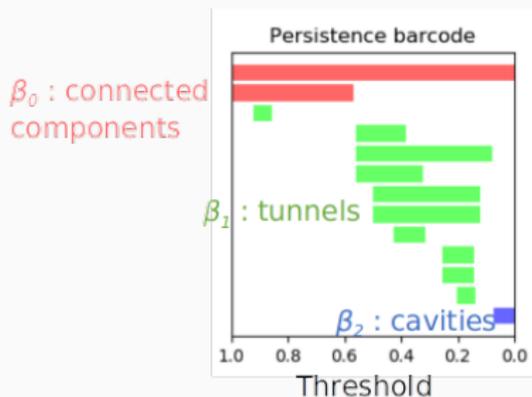
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where  $\lambda_k^*$  and  $\lambda_k$  are weights.

- **Elements to conserve:** move to (0, 1).
- **Elements to eliminate:** move to the diagonal line.



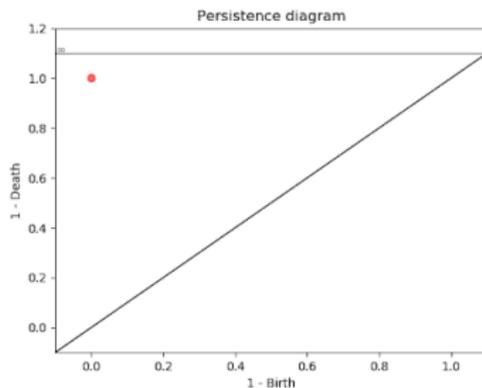
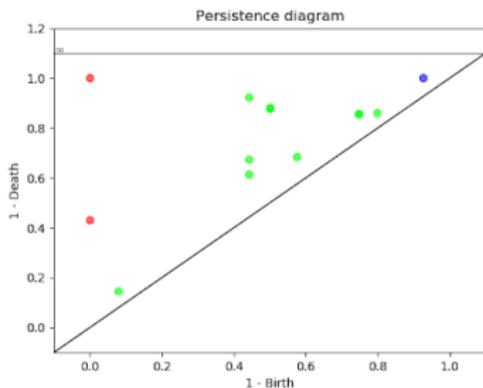
# Topological prior $\mathcal{L}_{\text{topo}}(1, 0, 0)$

Setting  $(\beta_0^*, \beta_1^*, \beta_2^*) = (1, 0, 0)$  favors one connected component.

Input



Ideal reconstruction



## Topological priors $\mathcal{L}_{\text{topo}}^{\square}(1, 1, 1)$ and $\mathcal{L}_{\text{topo}}^{\square}(1, 2, 1)$

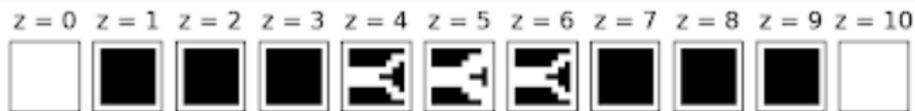
In order to distinguish a single tube and a bifurcation structure, we consider the **padded volume patches with a one-voxel border of value 1** and set

- $(\beta_0^*, \beta_1^*, \beta_2^*) = (1, 1, 1)$  for a single **tube** structure,
- $(\beta_0^*, \beta_1^*, \beta_2^*) = (1, 2, 1)$  for a single **bifurcation** structure.

Without bifurcation: Betti numbers (1, 1, 1)



With bifurcation: Betti numbers (1, 2, 1)



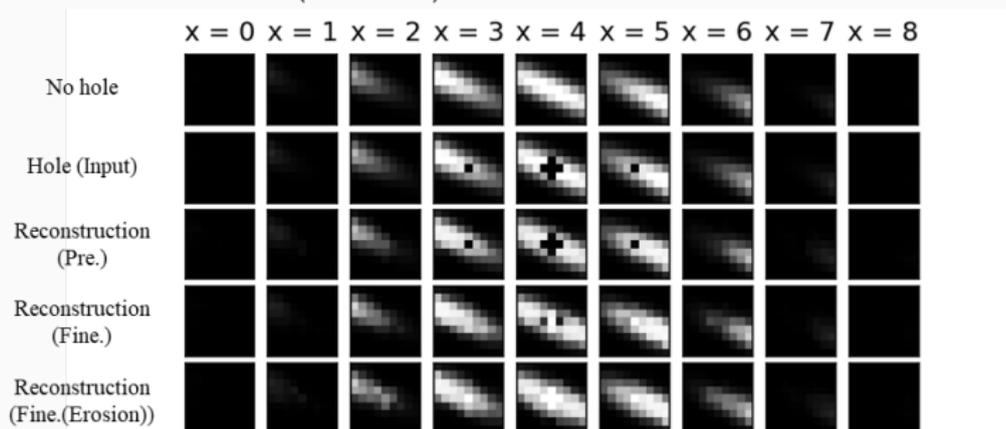
## Experiment: blood vessels with hole artifacts

The network was **pre-trained with only the  $\beta$ -VAE loss** and fine-tuned by adding the **topological prior** of one connected component  $\mathcal{L}_{\text{topo}}(1, 0, 0)$ .

- **Datasets:** artificial volume patches (size  $9 \times 9 \times 9$ ) of blood vessels with hole artifacts are generated; 6000, 2000 and 2000 are used for training, validation and testing.
- **Other setting:** the latent dimensions  $J = 6$ ; the weights of the loss function  $\beta = 0.1$ ,  $\lambda_k^* = \lambda_k = 60000$ .
- **Evaluation strategies:**
  - for intensity, **generalization** and **specificity** (Styner, 2003) were used,
  - for topology, the **topological loss function** with  $\lambda_k^* = \lambda_k = 1$  was used.

## Experiment: blood vessels with hole artifacts (results)

Given original volume patches without/with hole, compare its reconstruction without/with topological prior (Pre/Fine), and with the multiscale version (Erosion).



### Evaluations

	Generalization	Specificity	Topology
Pre	<b>0.00251</b>	<b>0.0134</b>	0.430
Fine	0.00478	0.0143	0.0976
Fine + Erosion	0.00728	0.0156	<b>0.00139</b>

# Experiment: blood vessels with bifurcations

The network was **pre-trained with only the  $\beta$ -VAE loss** and **fine-tuned by adding topological prior of bifurcation case  $\mathcal{L}_{\text{topo}}^{\square}(1, 2, 1)$** .

**Dataset:** volume patches of Y-shaped blood vessel

- CT images taken at Tokushima University Hospital (47 cases)
- thickness is approximately from 1 to 4 mm
- blood vessels are located between hilar and peripheral regions
- patches of size  $9 \times 9 \times 9$  are extracted
  - after applying Hessian filter,
  - around local maximum of voxel values,
  - s.t. the PHs must be close to the topological prior.

1533, 517 and 517 volume patches are used for training, validation and testing.

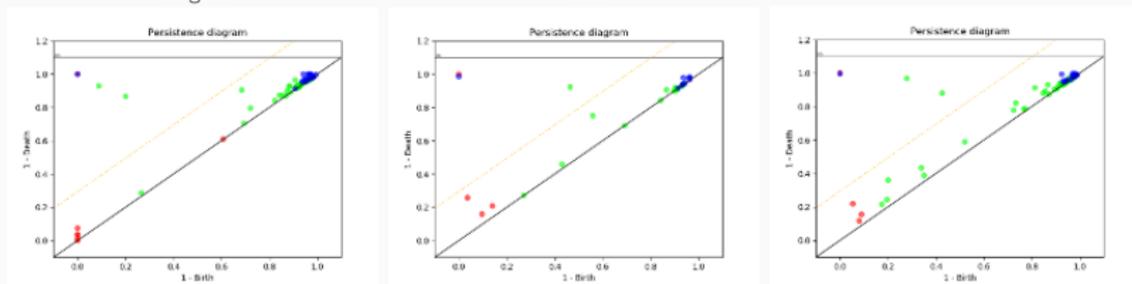
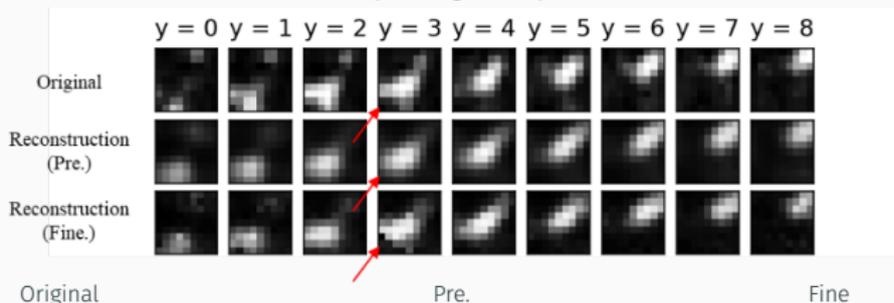
Sagittal sections of  
25 volume patches

## Experiment: blood vessels with bifurcations (cont.)

- **Other setting:** The latent dimensions  $J = 22$ ; The weights of the loss function  $\beta = 0.1$  and  $\lambda_k^* = \lambda_k = 50$ .
- **Evaluation strategies:**
  - for intensity, **generalization** and **specificity** (Styner, 2003) were used,
  - for topology, the **Bottleneck distance** between the PDs of the original and reconstructed images was used.

# Experiment: blood vessels with bifurcations (results)

Given an original volume containing a bifurcation, compare its reconstruction without/with topological prior (Pre/Fine).



## Evaluations

Generalization

Specificity

Bottleneck distance

Pre

**0.00704**

**0.0283**

0.288

Fine

0.0102

0.0410

**0.236**

## Conclusions

- In order to incorporate topological priors of bifurcations into statistical modeling, we integrated topological loss into the deep generative model ( $\beta$ -VAE).
- Topological evaluation of reconstructed images was improved, and the improvement was also qualitatively identified.

# Conclusions and Perspectives

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- Topological evaluation of reconstructed images was improved, and the improvement was also qualitatively identified.

## Perspectives

- How to guarantee topological prior?
- Apply the model to other tasks such as segmentation and super-resolution.