Scanner Variability

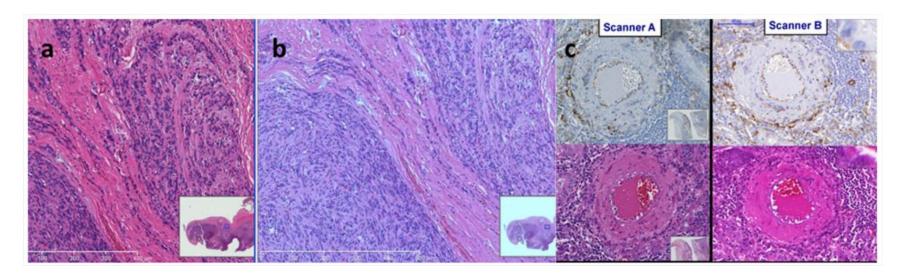


🔬 SeeGene Project Report 🔬



Scanner Variability | What's the issue?

Each time the scanner is changed to digitize tissue images, the properties of the whole slide image change. Although for the pathologists these changes are meaningless, they might have a great effect on the machine learning model.



Scanner Variability | Literature Review

- Conversion to grayscale ignores the important information regarding the color representation.
- **Color normalization** tries to adjust the color distribution of the source image to that of a reference image. However, this method might be expensive because we need to adjust for every reference data.
- Color augmentation is performed by applying random hue, saturation, brightness, and contrast. The advantage of color augmentation lies in the easy implementation regardless of the object being analyzed.

Scanner Variability | Literature Review

Lafarge et al. deals with dataset variabilities caused by differences in pathology labs (e.g., scanners).

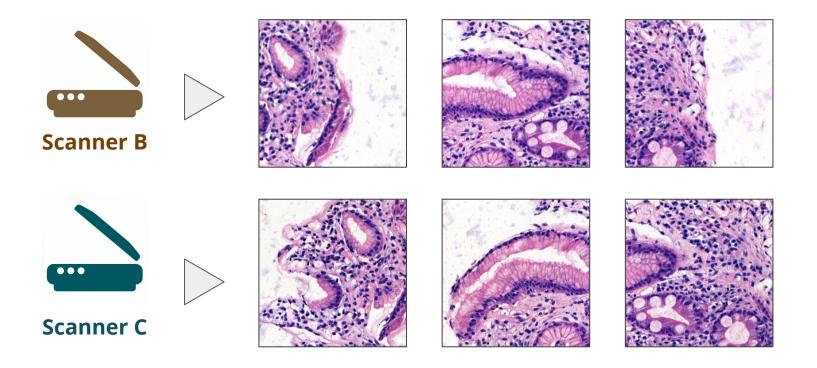
- 1. Color Augmentation
- 2. Staining Normalization (Macenko)
- 3. Domain-Adversarial Neural Network

| CA | | | | | | | | |
|------|---------------|---------------|---------------|---------------|---------------|-------------|---------------|---------------|
| SN | | | | | | | | |
| DANN | | | | | | | | |
| ITS | $.61 \pm .02$ | $.61 \pm .01$ | $.57 \pm .06$ | $.61 \pm .02$ | $.55 \pm .01$ | $.62\pm.02$ | $.61 \pm .01$ | $.57 \pm .01$ |
| ETS | $.33 \pm .08$ | $.58 \pm .03$ | $.46 \pm .02$ | $.55 \pm .05$ | $.48 \pm .08$ | $.62\pm.00$ | $.51 \pm .02$ | $.53 \pm .03$ |

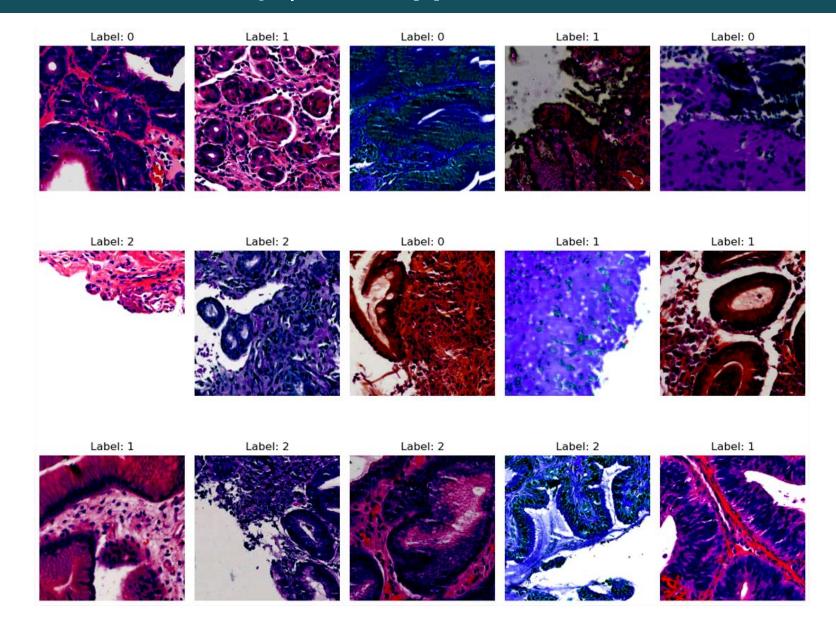
There are some approaches to deal with this kind of variability, which include **adversarial training** and **color augmentation**. Color Augmentation has been shown useful and effective in some domain (according to the literature review), hence:

- Split the dataset (WSI) into:
 - Train → Scanner A
 - Test → Scanner B and Scanner C
- Implement color augmentation during the training process
- If the model performs well on scanner B and scanner C, having been trained with Scanner A, then we can have more confidence that the model can be robust to scanner variability.

Depending on the scanner, the tiling code outputs different patches for the same sample:



First, second, and third patches extracted from scanner 1 and scanner 2



Backbone: ResNet50

Adam Optimizer - lr: 0.001 | Exponential Decay - lrd: 0.85

Epochs: 20 | Early Stopping - Patience: 3 epochs

| | Same Scanner | Scanner B | Scanner C |
|-----------------|--------------|-----------|-----------|
| Baseline | 0.768 | 3 | ? |
| Color Augmented | 0.750 | ; | ? |

Backbone: ResNet50

Adam Optimizer - lr: 0.001 | Exponential Decay - lrd: 0.85

Epochs: 20 | Early Stopping - Patience: 3 epochs

| | Same Scanner | New slides are not "fine-annotated", but weakly-annotated |
|-----------------|--------------|---|
| Baseline | 0.768 | Cannot perform tile-level evaluation |
| Color Augmented | 0.750 | |

Backbone: ResNet50

Adam Optimizer - lr: 0.001 | Exponential Decay - lrd: 0.85

Epochs: 20 | Early Stopping - Patience: 3 epochs

Slide Level Prediction:

D if #_patches_predicted_as_D > #_patches_predicted_as_M else M

| | Same Scanner | Scanner B | Scanner C |
|-----------------|--------------|-----------|-----------|
| Baseline | 0.885 | 0.795 | 0.663 |
| Color Augmented | 0.855 | 0.819 | 0.837 |

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Adam Optimizer - lr: 0.001 | Exponential Decay - lrd: 0.85

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Although this slide-level classification is somehow heuristic, there is some evidence of <u>robustness</u>

~ THANK YOU ~