

# Conditional Prompt Learning for Anomaly Detection in Medical Image Slices

Dohoon Kim<sup>1</sup> and Jae Sung Lee<sup>2</sup>

Department of Data Science, Hanyang University, Seoul, Korea<sup>1</sup> Department of Nuclear Medicine, Seoul National University, Seoul, Korea<sup>2</sup>



#### Introduction

## **♦** Robust Algorithm applicable to diverse domains

- Diverse MRI sequences (T1, T2, T1CE, FLAIR)
- Diverse Modalities (X-ray, OCT...)

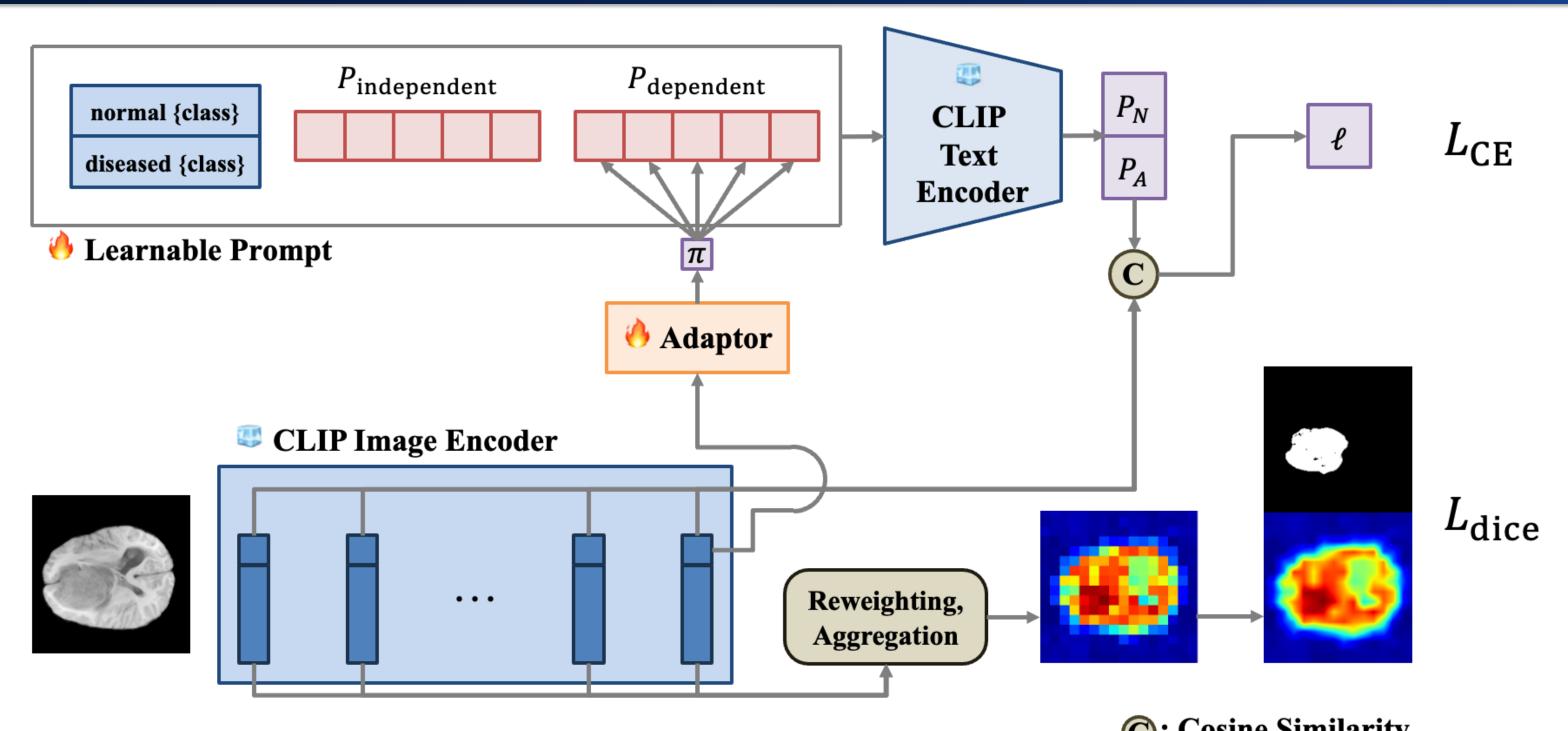
# Giving Condition to Prompt for Adaptation

 Existing works (AnomalyCLIP, AdaCLIP, MediCLIP) use CoOp method solely, limiting model's adaptability

# **◆** A Prompting module to guide CLIP:

- CLIP prompt gUidance for mEdical image
- Guiding CLIP patch embedding to understand medical anomalies in diverse slices

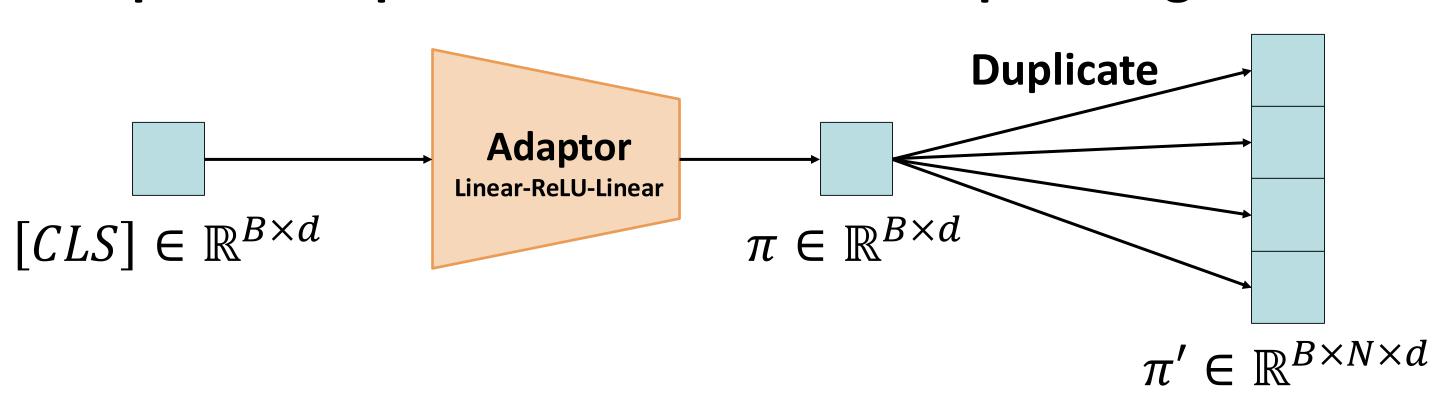
# Methods



**©**: Cosine Similarity P: Prompt Embedding  $\ell$ : Logits

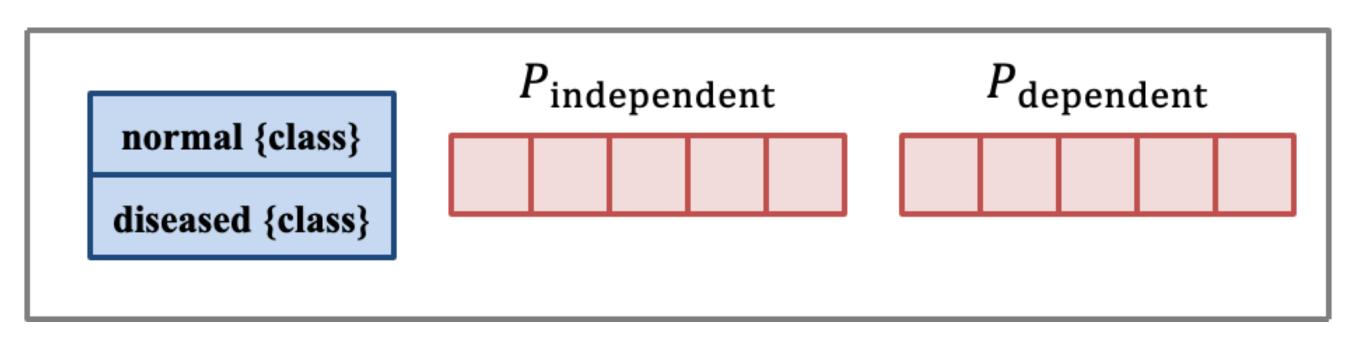
#### ◆ Image Adaptor

- Input: CLS token from final layer of the image encoder
- Output: Prompt bias conditioned on input image



#### Prompt Design

Learnable prompt is composed as below;



- $P_{\text{dependent}} \in \mathbb{R}^{B \times N \times d}$ 
  - $\pi'$  is added to incorporate the information of the image.
- $P_{\text{independent}} \in \mathbb{R}^{B \times N \times d}$
- $P_{\text{class}} \in \mathbb{R}^{B \times 2 \times d}$
- Finally, the prompt fed into CLIP is as follows.

• 
$$P = \text{concat}(P_{\text{class}}, P_{\text{independent}}, P_{\text{dependent}})$$

- In this work, we used ViT-L/14, and N=32
- $v \in \mathbb{R}^{B \times (16 \times 16 + 1) \times d}$  is obtained from last layer

## Reweighting & Upscaling

• This makes patches into 224x224 anomaly maps.

Algorithm 1 Anomaly Map Computation **Require:** Image Feature I, Prompt Feature PInitialize M as an empty list for l=1 to L do Extract patch features  $F_l$  from I[l]Compute anomaly score for patch:  $M_l = \cos(F_l, P_A) - \cos(F_l, P_N)$ end for Compute weighted anomaly scores:  $w = \operatorname{softmax} \left( \frac{1}{HW} \sum_{l} M_{l} \right)$ Compute final anomaly map:  $M_{final} = \sum w_l M_l$ Upscale  $M_{final}$  to size (224, 224) and normalize Return  $M_{final}$ 

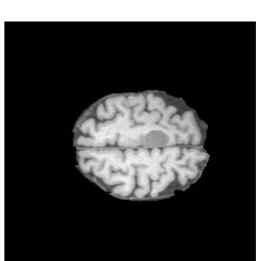
#### Results

- **♦** Experiment on diverse views and sequences
  - Train on T1w MRI with axial Slices
  - Test on the others

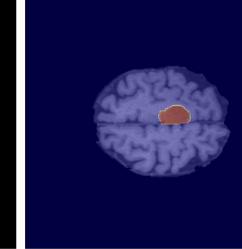
(Pixel-AUROC, %)

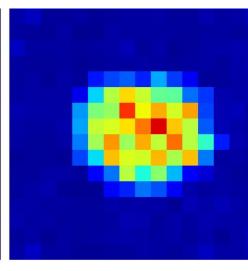
MRI Sequences	Viewpoint		
	Axial	Sagittal	Coronal
T1	93.19	87.43	87.16
T1CE	85.97	83.52	84.74
<b>T2</b>	90.26	85.92	87.42
FLAIR	88.43	84.78	85.20

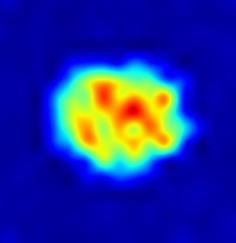
**♦** In-Domain (T1w MRI, axial slice)

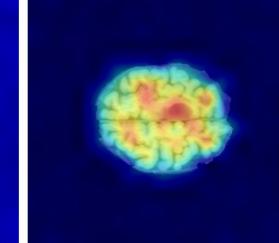




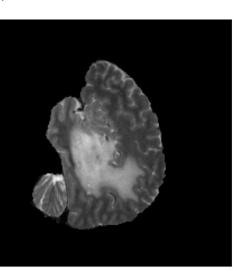


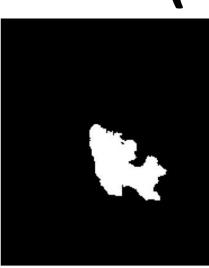


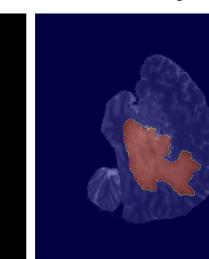


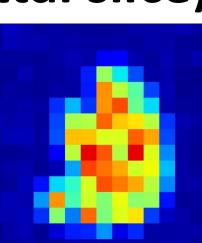


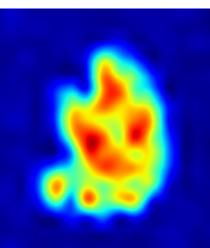
**♦** Domain Shift (T2w MRI, sagittal slice)

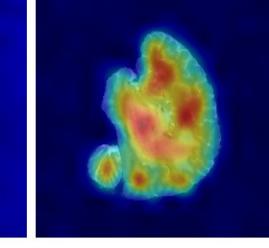








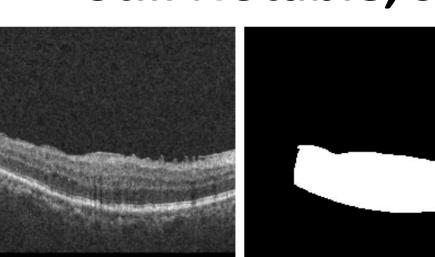


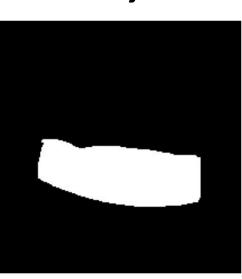


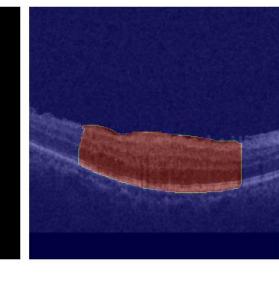
- **♦** Experiment on diverse organs and modalities
  - Zero-Shot Transfer
  - Chest X-ray, Retina OCT

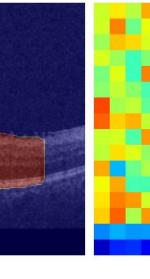
	Image-AUROC	Pixel-AUROC
Chest	55.47	N/A
Retina	55.83	53.15

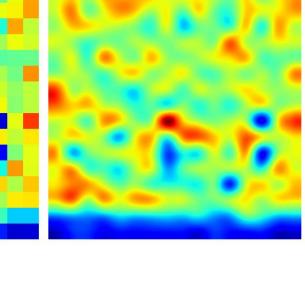
**◆** Actually NOT a successful performance. Rather, FAILED. Still Notable, since better than random prediction

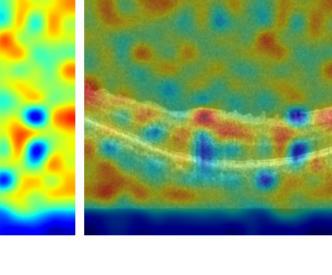












#### Discussion

- ◆ Map generation algorithm might not be optimal
  - Unstable due to non-parametric method, Interpolation
- **♦** Diverse class or domain during training can improve
  - Unstable due to non-parametric method, Interpolation
  - Fine tuning or using auxiliary data might be solution

### Conclusion

- **♦** Without fine-tuning CLIP itself, prompt learning can guide CLIP to understand medical anomalies
- **♦** Without additional decoder, patch embeddings can represent the anomalous location at large