



# Multiview Long-Short Spatial Contrastive Learning for 3D Medical Image Analysis

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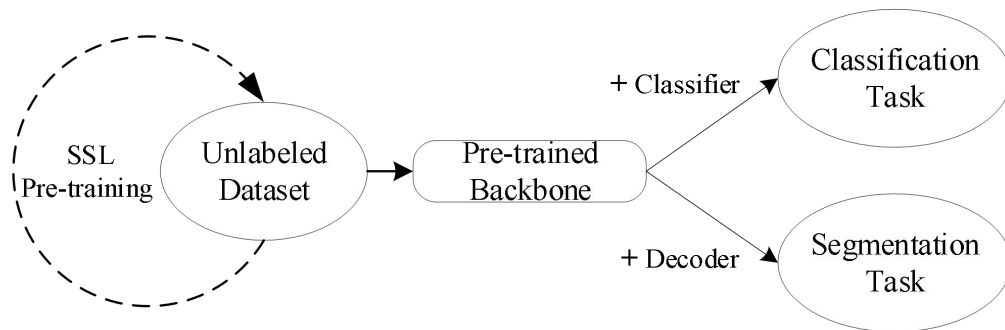
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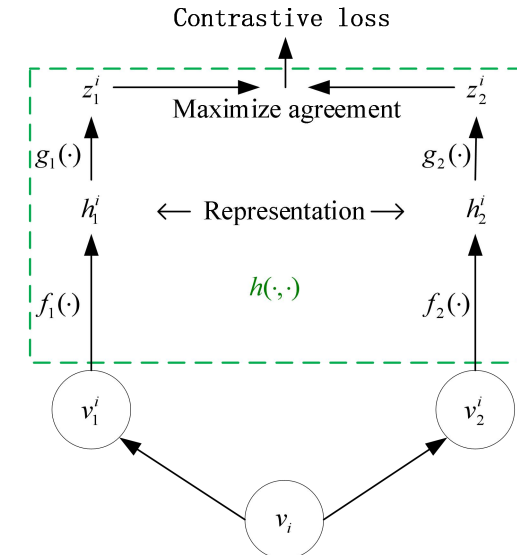
# Introduction

## Background:

- Supervised deep learning requires sufficient labeled data.
  - ❑ Expert knowledge in specific fields.
  - ❑ Time-consuming and Laborious.
- Self-supervised learning (SSL):
  - ❑ Without human-annotations.
  - ❑ Generic representations transferred to downstream tasks.



## Contrastive Learning:



- An effective implementation of the self-supervised learning.
- Contrastive learning trains neural networks to discriminate between “positive” pairs  $(v_1^i, v_2^i)$  and “negative” pairs  $(v_1^i, v_2^j)_{j \neq i}$ .
- Learning is formulated as minimizing a contrastive loss (we use InfoNCE in this paper).

$$L_N(v_1, v_2) = -\mathbb{E} \left[ \log \frac{e^{h(v_1^i, v_2^i)}}{\sum_{j=1}^{K+1} e^{h(v_1^i, v_2^j)}} \right]$$

# Motivation

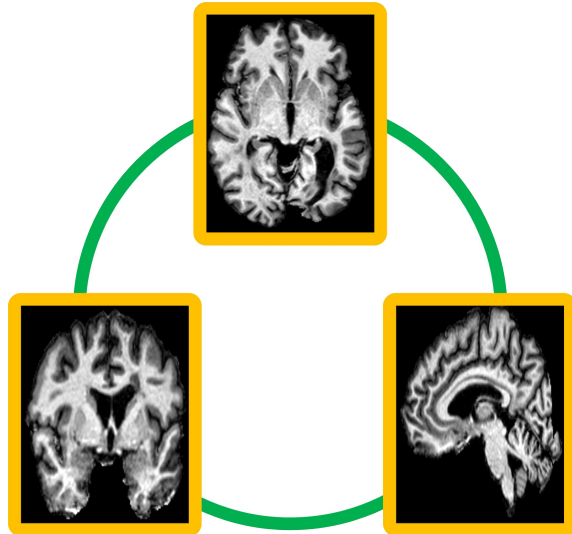
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Two limitations of existing contrasting strategies when applied to 3D medical images:

- Ignore the **intrinsic structural similarity**.
- Ignore **local representation**.

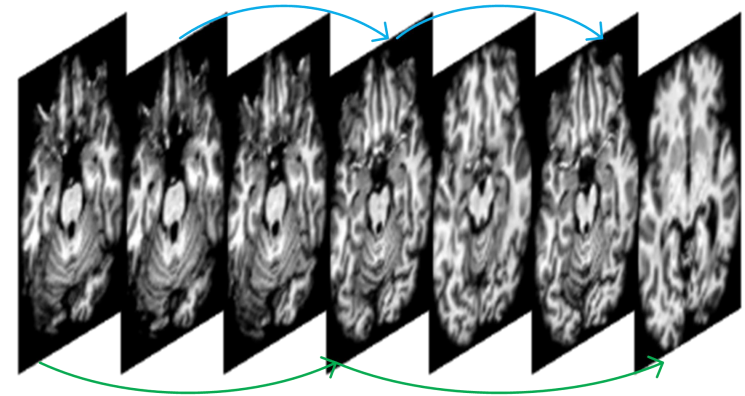
## Observation 1:

The information **shared between three views** (axial, coronal and sagittal views) can capture the **global representation** of volumetric medical image.



## Observation 2:

Matching the short spatial clip to long spatial clip forces the model to extrapolate **local information**.



# Method: Multiview Contrasting Strategy & Long-Short Spatial Contrasting Strategy

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## Multiview Contrasting Strategy:

- To learn global representation, we need to maximize the mutual information between three views ( $v_a, v_c, v_s$ )

$$\max\{I(v_a; v_c) + I(v_a; v_s) + I(v_c; v_s)\}$$

- InfoNCE loss can estimate the lower bound of mutual information. For two views  $v_1, v_2$ :

$$I(v_1; v_2) \geq \log(K) - L_N(v_1, v_2)$$

- Maximizing mutual information between three views → Multiview contrastive learning:

$$L_{multiview} = L_N(v_a, v_c) + L_N(v_a, v_s) + L_N(v_c, v_s)$$

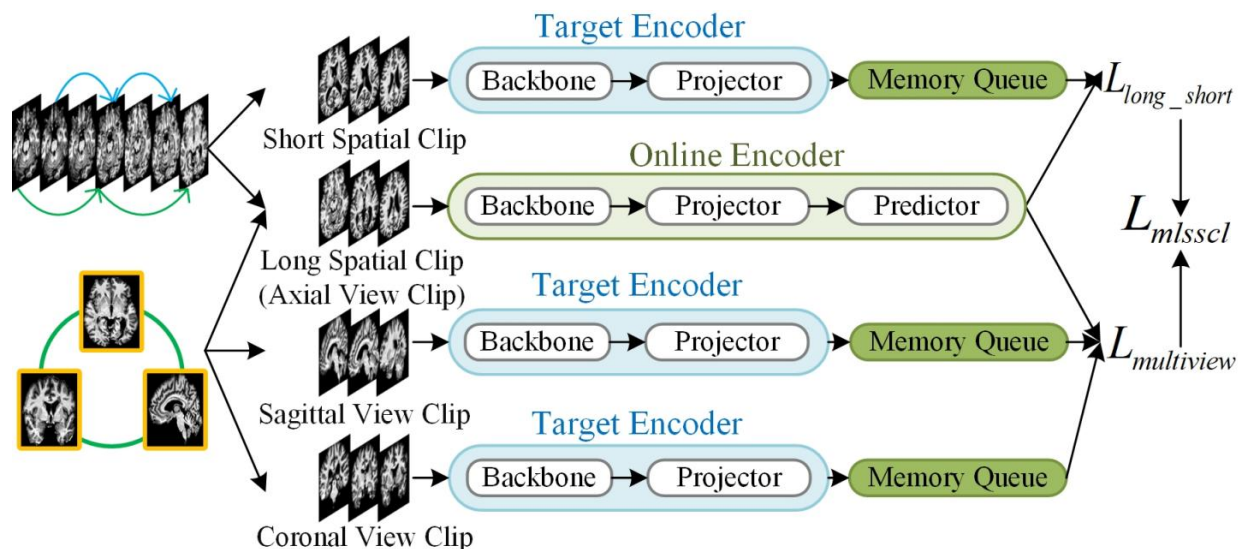
## Long-Short Spatial Contrasting Strategy:

- Maximizing representation similarity between a long spatial clip  $v_L$  and a much shorter spatial clip  $v_S$ :

$$L_{long-short} = L_N(v_S, v_L)$$

Matching the short-clip representation to the long-clip representation forces the model to understand and recognize the **structure and correlation** of **local tissues** in volumetric medical images.

# Method: Multiview Long-Short Spatial Contrastive Learning Framework



## Clip Sampling:

- Sample axial, coronal and sagittal clips from a 3D volumetric medical image with  $C$  slices and a stride of  $\delta_L$ .
- Regard the above axial clip as the long spatial clip and then randomly sample  $C$  axial slices with spatial stride  $\delta_S$  ( $\delta_S < \delta_L$ ) as the short spatial clip.

## Network Architecture:

- One online encoder:
  - ❑ a backbone + a projector head (2-layer MLP) + a prediction head (2-layer MLP).
  - ❑ updated by back-propagation.
  - ❑ the backbone will be transferred to downstream tasks after pre-training.
- Three target encoders: (share weights)
  - ❑ a backbone + a projector head (2-layer MLP)
  - ❑ updated in the manner of momentum.
  - ❑ memory queue to store previous representations.
  - ❑ discarded after pre-training.

## Contrastive Loss:

$$L_{mlsscl} = \alpha L_{multiview} + \beta L_{long\_short}$$

# Experiments: Pre-training on Large-Scale Unlabeled Dataset

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## **Pre-training Dataset:**

- ADNI pre-training set (5953 T1-weighted MRI scans).

## **Instantiation of Network:**

- AD classification task:
  - ▣ 3D ResNet-18 as backbone.
- MS lesion segmentation task:
  - ▣ 3D UNet-based encoder as backbone.

## **Optimization:**

- We pre-train models on ADNI pre-training set for 100 epochs with SGD optimizer.

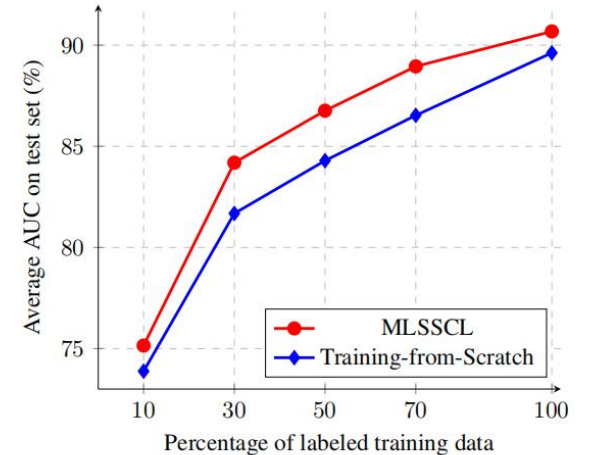
*Other details can be found in paper.*

# Experiments: Transferring Learned Features to AD Classification

**Table 1.** Results(mean $\pm$ std) for AD classification (AD vs. HC) on the ADNI-AD classification test set.

Method	ACC	SEN	SPE	AUC
Training-from-Scratch	0.793 $\pm$ 0.011	0.874 $\pm$ 0.055	0.711 $\pm$ 0.058	0.896 $\pm$ 0.006
BYOL [12]	0.809 $\pm$ 0.004	0.866 $\pm$ 0.032	0.752 $\pm$ 0.037	0.886 $\pm$ 0.016
MoCo [10]	0.825 $\pm$ 0.020	0.886 $\pm$ 0.043	0.764 $\pm$ 0.060	0.895 $\pm$ 0.001
Model Genesis [8]	0.827 $\pm$ 0.004	0.911 $\pm$ 0.061	0.744 $\pm$ 0.061	0.904 $\pm$ 0.009
Age-Aware [13]	0.831 $\pm$ 0.007	0.882 $\pm$ 0.007	0.780 $\pm$ 0.012	0.899 $\pm$ 0.011
<b>MLSSCL</b>	<b>0.858 <math>\pm</math> 0.013</b>	<b>0.911 <math>\pm</math> 0.019</b>	<b>0.805 <math>\pm</math> 0.044</b>	<b>0.907 <math>\pm</math> 0.012</b>

- MLSSCL achieves a **remarkable improvement** over training-from-scratch:  $\uparrow$  6.5%(ACC),  $\uparrow$  3.7%(SEN),  $\uparrow$  9.4%(SPE)
- MLSSCL **outperforms** other SSL methods:  $\uparrow$  2.7%(ACC),  $\uparrow$  2.5%(SPE)
  
- MLSSCL can effectively deal with the situation with **few labeled** training samples. 70% labeled data (MLSSCL)  $\approx$  100% labeled data (training-from-scratch)



**Fig. 2.** The AD classification performance of networks trained with different amounts of labeled data.

# Experiments: Transferring Learned Features to MS Lesion Segmentation

**Table 2.** The segmentation results of different approaches on the ISBI 2015 longitudinal MS lesion segmentation test set.

Method	DSC <sup>†</sup>	PPV <sup>†</sup>	LTPR <sup>†</sup>	LFPR <sup>†</sup>
Training-from-Scratch	0.6176	0.8229	0.4451	0.3485
<b>SSL</b>				
Age-Aware [13]	0.6320	0.8103	0.4586	0.3034
BYOL [12]	0.6337	0.7991	0.4675	0.3442
MoCo [10]	0.6369	0.7972	0.4641	0.3092
Model Genesis [8]	0.6434	0.8200	0.4647	0.3082
<b>MS SOTA</b>				
Aslania et al. [3]	0.6114	<b>0.8992</b>	0.4103	0.1393
Andermatt et al. [18]	0.6298	0.8446	0.4870	0.2013
Valverde et al. [4]	0.6304	0.7866	0.3669	0.1529
Hu et al. [5]	0.6345	0.8682	0.4787	<b>0.1299</b>
<b>MLSSCL</b>	<b>0.6482</b>	0.8007	<b>0.4933</b>	0.2796

- MLSSCL consistently outperforms training-from-scratch and other SSL methods. Compared with training-from-scratch:  
    ↑ 3.06%(DSC), ↑ 4.82%(LTPR), ↑ 6.89%(LFPR)
- MLSSCL still achieves higher DSC and LTPR compared to SOTA segmentation methods.



## Experiments: Ablation to contrasting strategies on AD classification

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**Table 3.** Ablation to contrasting strategies on AD classification task (mean  $\pm$  std).

<b>Contrasting Strategy</b>	<b>ACC</b>	<b>AUC</b>
Long-Short	$0.823 \pm 0.012$	$0.892 \pm 0.014$
Multiview	$0.833 \pm 0.027$	$0.906 \pm 0.024$
<b>Multiview &amp; Long-Short</b>	<b><math>0.858 \pm 0.013</math></b>	<b><math>0.907 \pm 0.012</math></b>

The results demonstrate the **complementarity** of global representation and local representation.

# Conclusion

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- ✓ We introduce multiview contrasting strategy to learn global representations by maximizing the mutual information between three views of the same volumetric medical image.
- ✓ We introduce long-short spatial contrasting strategy to learn local representations by matching a short spatial clip to a long spatial clip in the latent space under the given view.
- ✓ We propose multiview long-short spatial contrastive learning (MLSSCL) framework to combine these two contrasting strategies, which can effectively learn generic 3D representations.
- ✓ Extensive experimental results showed that MLSSCL outperformed training-from-scratch method, especially when fine-tuned on only small amounts of labeled data, and also showed a clear superiority compared with other self-supervised learning methods.



# Thank you!

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