

# Benchmark of U-Net Architectures in Medical Image Segmentation

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0.925

0.875

# METHODOLOGY AND EXPERIMENTATION

### BACKGROUND

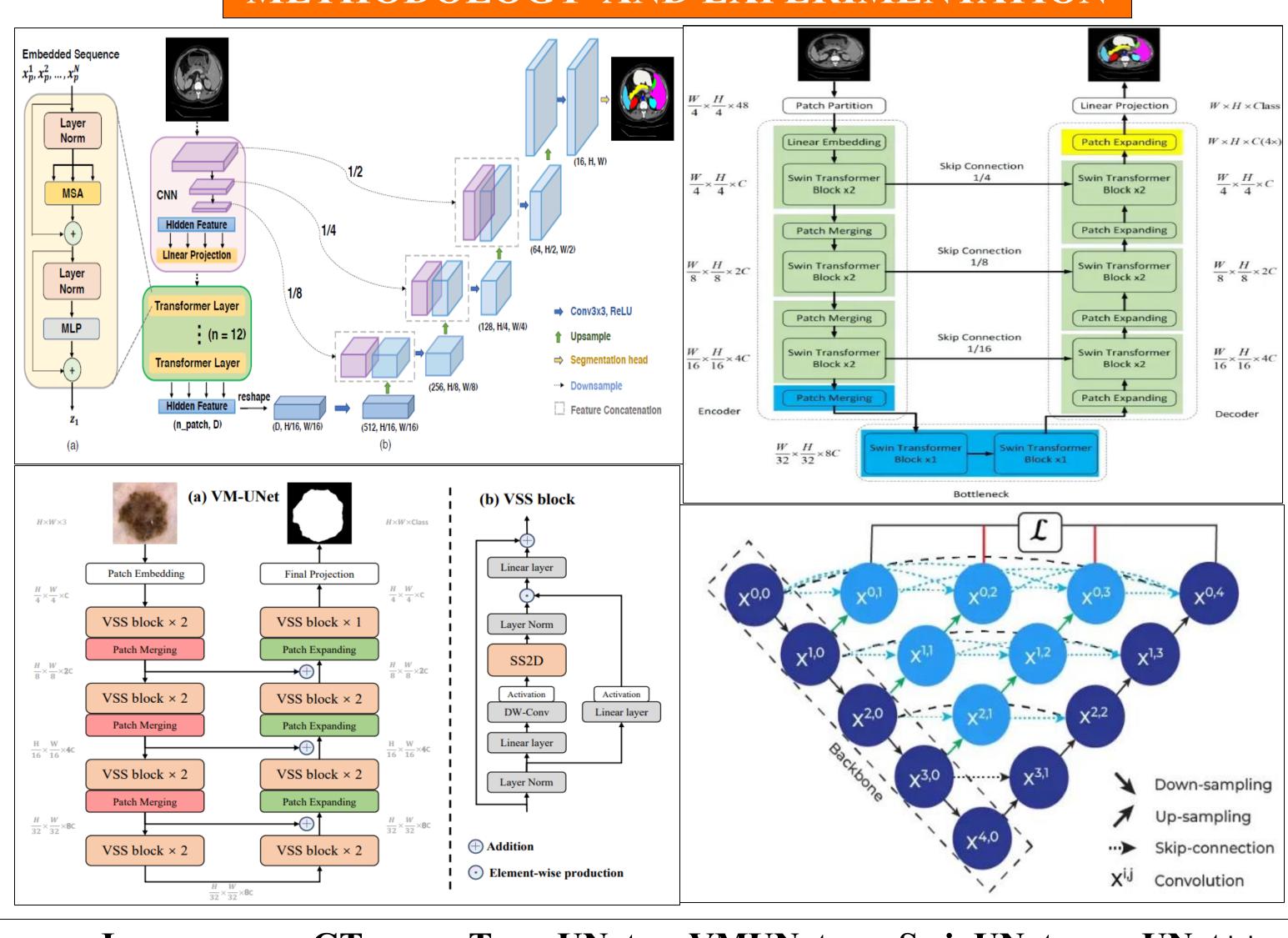
Medical image segmentation is a critical task in healthcare applications, enabling precise localization and identification of anatomical structures. Deep learning models, particularly the U-Net architecture, have become the foundation for many segmentation methods. In this study, we evaluate four U-Netbased architectures—TransUNet, SwinUNet, VMUNet, and UNet++—which integrate different advancements such as transformers, attention mechanisms, and nested architectures. These models were benchmarked on two publicly available datasets representing different anatomical regions. We compare their segmentation performance using standard evaluation metrics such as F1 score, Dice coefficient, recall, and precision.

### **PURPOSE**

The purpose of this study is to benchmark and compare the performance of four different U-Net-based deep learning architectures—TransUNet, SwinUNet, VMUNet, and UNet++ to evaluate how various architectural innovations impact segmentation accuracy in medical image tasks. Each model introduces distinct enhancements to the original U-Net framework: TransUNet incorporates transformers for better nandling of long-range dependencies, SwinUNet leverages a window-based approach for efficient global context aggregation, VMUNet uses state space models for improved feature representation, and UNet++ introduces broader skip connections and deep, nested architectures for more precise segmentation. This study evaluates these architectures across two publicly available datasets: ISIC2018 for skin lesion segmentation and PolyPGen (from MedSegBench) for polyp segmentation.

# Hyperparameters

	TransUNet	SwinUNet	VMUnet	UNet++
Optimizer	SGD	SGD	AdamW	Adam
Scheduler	Polynomial Decay	Polynomial Decay	CosineAnneal ingLR	ReduceLROn Plateau
Batch Size	24	24	64	16
Loss Function	0.5 * CEL + 0.5 * Dice_Loss	0.4 * CEL + 0.6 * DiceLoss	BCEDiceloss (wb=0.4, wd=0.6)	0.5 * DiceLoss + 0.5 * CEL
Epochs	150	150	300	150



$\frac{H}{32} \times \frac{W}{32} \times 8c$ $VSS block \times 2$ $\frac{H}{32} \times \frac{W}{32} \times$	$\frac{H}{32} \times \frac{W}{32} \times 8c$ VSS block × 2	Addition  Element-wise production		X <sup>4,0</sup>	→ Skip-connection  X <sup>i,j</sup> Convolution
Image	GT	TransUNet	VMUNet	SwinUNet	UNet++

### RESULTS

VMU-Net

Unet++

Metrics	Accuracy	Recall	Specificity	F1/ Dice Coefficient	Mean IOU	Precision
TransUNet	0.911	0.837	0.9393	0.838	0.721	0.840
SwinUNet	0.982	0.986	0.981	0.958	0.933	0.931

0.911

0.842

#### PolypGen Dataset

0.969

0.898

0.818

**ISIC 2018 Dataset** 

Metrics	Accuracy	Recall	Specificity	F1/ Dice Coefficient	Mean IOU	Precision
TransUNet	0.973	0.791	0.990	0.832	0.712	0.877
SwinUNet	0.979	0.849	0.992	0.874	0.776	0.900
VMU-Net	0.962	0.859	0.976	0.840	0.724	0.821
Unet++	0.957	0.592	0.985	0.640	0.478	0.723

## CONCLUSIONS AND DISCUSSION

SwinUNet had the best metrics in both the ISIC 2018 dataset and PolyPGen dataset. While all models demonstrate strong performance, a notable limitation observed is their difficulty in accurately segmenting lesions with coloration closely matching the surrounding skin. This suggests a reliance on strong color/contrast cues and highlights an area for future improvement, potentially through advanced feature extraction focusing on texture and boundary details or targeted data augmentation strategies. SwinUNet was the least affected on this limitation.

Additionally, while results are promising, current findings are limited to two datasets, and it remains unclear how each model will perform on other segmentation tasks such as organ segmentation, cell nuclei detection, or multi-class problems.

Future work will involve training and evaluating all four models across the full MedSegBench suite, which contains 35 medical image datasets spanning a wide range of anatomical regions. This broader evaluation will help determine the generalizability, robustness, and scalability of each architecture in diverse clinical scenarios.

### **BIBLIOGRAPHY**

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