



# **BRAIN TUMOR RADIOGENOMIC CLASSIFICATION - AN ADAPTED SWIN TRANSFORMER APPROACH**

By

**M. Satya Sai Teja (Enrolment No. 20/11/EC/011)**

under the guidance of

**Dr. Anupama Namburu, School of Engineering, Jawaharlal Nehru University, Delhi**



# OVERVIEW

- Introduction
- Acknowledgement
- Objective
- Tools and Technologies
- Methodology
- Swin-T Architecture
- Results
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- Conclusion
- References





# INTRODUCTION

- Glioblastoma is a common and deadly type of brain cancer in adults.
- This aggressive cancer has a poor prognosis, making treatment very challenging.
- To understand its genetics, doctors need to perform surgery to get a tissue sample.
- Analyzing the genetic characterization of the tumor through these invasive surgical methods take several weeks.
- Development of an accurate method to predict the genetics of cancer through radiogenomics, refines the type of therapy required.





# ACKNOWLEDGEMENT

I would like to express my sincere gratitude to everyone who supported me this project:

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- RSNA-MICCAI organizations, for their collaborative effort in formulating the problem statement, organizing the challenge, and providing the BraTS 2021 dataset.





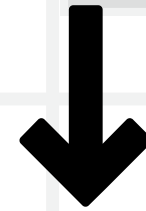
# OBJECTIVE

- Genetic sequence O6-Methylguanine-DNA-Methyltransferase (MGMT) promoter methylation acts favourably for prognosis and responsiveness to chemotherapy.
- Develop a predictive model, using the techniques from Computer Vision, Machine Learning, Deep Learning and Radiogenomics.
- By training and testing this model with MRI (Magnetic Resonance Imaging) scans, we estimate the probability of MGMT promoter methylation.
- Improve treatment decision making and enhance patient outcomes.





# TOOLS AND TECHNOLOGIES



## TOOLS

- Python
- Pandas
- Numpy
- OpenCV
- Pydicom
- Pytorch
- Scikit-learn
- Seaborn



## TECHNOLOGIES

- Radiogenomics
- Computer Vision
- Machine Learning
- Deep Learning
- Swin Transformer





# METHODOLOGY

## PHASE 1

- Visualization
- Analysis
- Preprocessing
- Augmentation

## PHASE 2

- Data Loading
- Dataset Retrieval
- Modelling
- Training

## PHASE 3

- Optimization
- Prediction
- Evaluation
- Submission

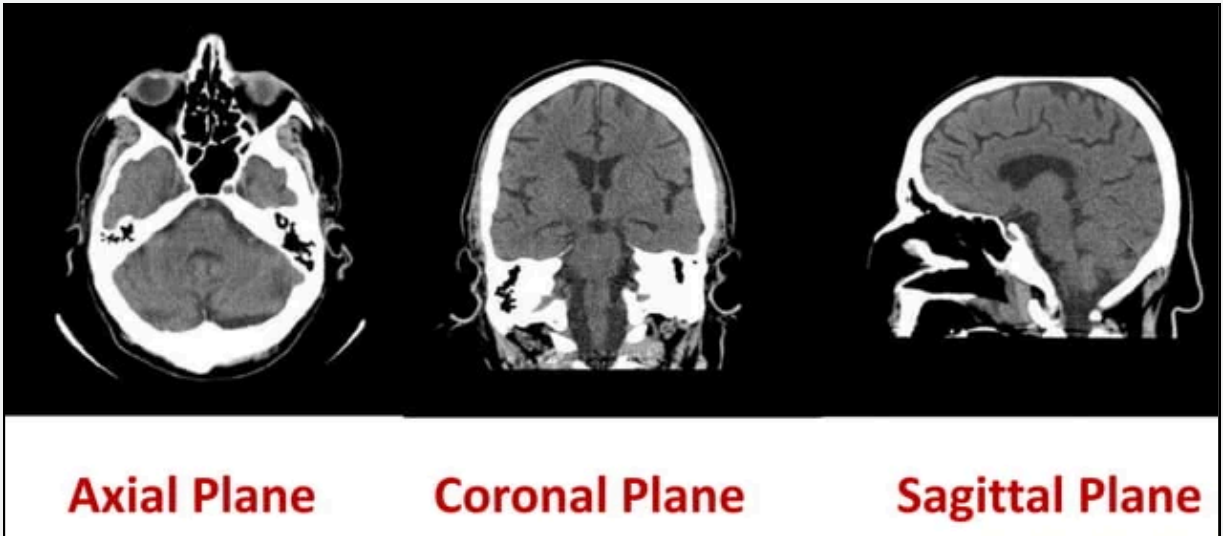




# METHODOLOGY

```
Training/Validation/Testing
├── 00000
│   ├── FLAIR
│   │   ├── Image-1.dcm
│   │   ├── Image-2.dcm
│   │   └── ...
│   ├── T1w
│   │   ├── Image-1.dcm
│   │   ├── Image-2.dcm
│   │   └── ...
│   ├── T1wCE
│   │   ├── Image-1.dcm
│   │   ├── Image-2.dcm
│   │   └── ...
│   ├── T2w
│   │   ├── Image-1.dcm
│   │   ├── Image-2.dcm
│   │   └── ....
│   └── 00001
│       └── ...
└── ...
```

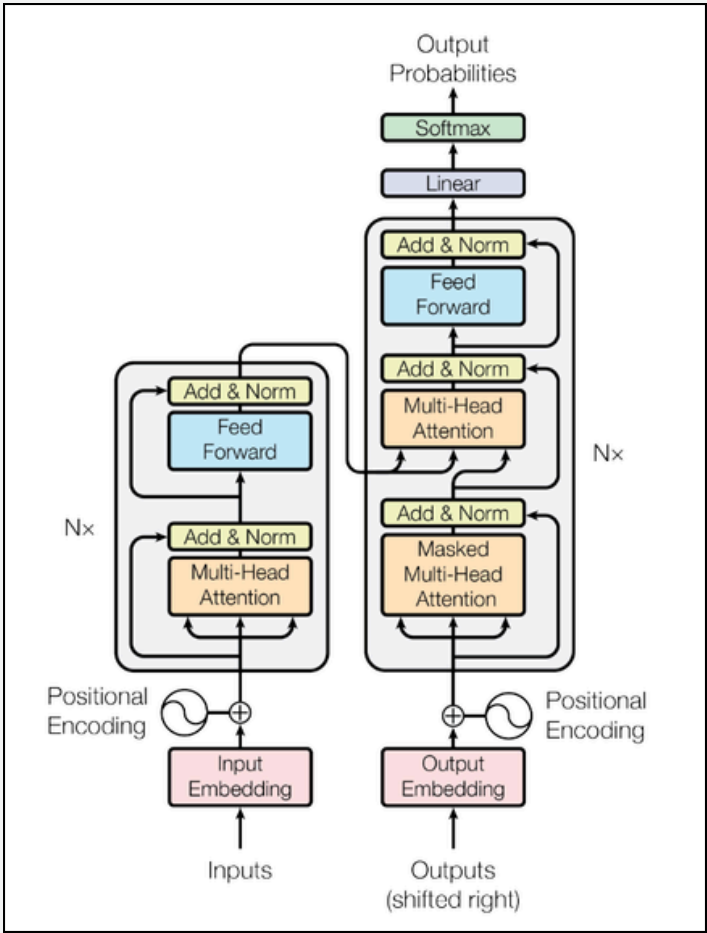
```
BraTS21ID, MGMT_value
00001, 0.5
00013, 0.5
00015, 0.5
etc.
```



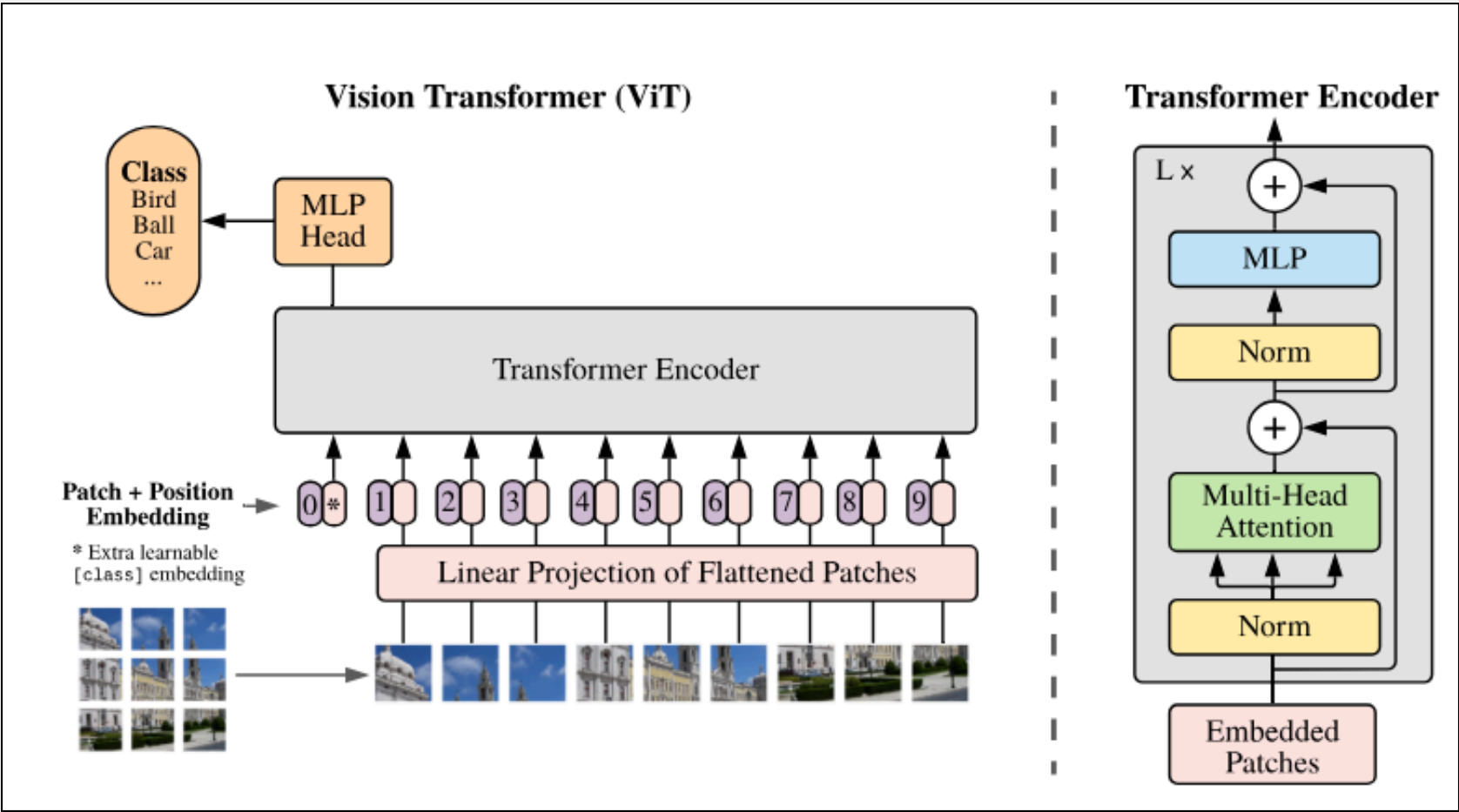




# SWIN-T ARCHITECTURE



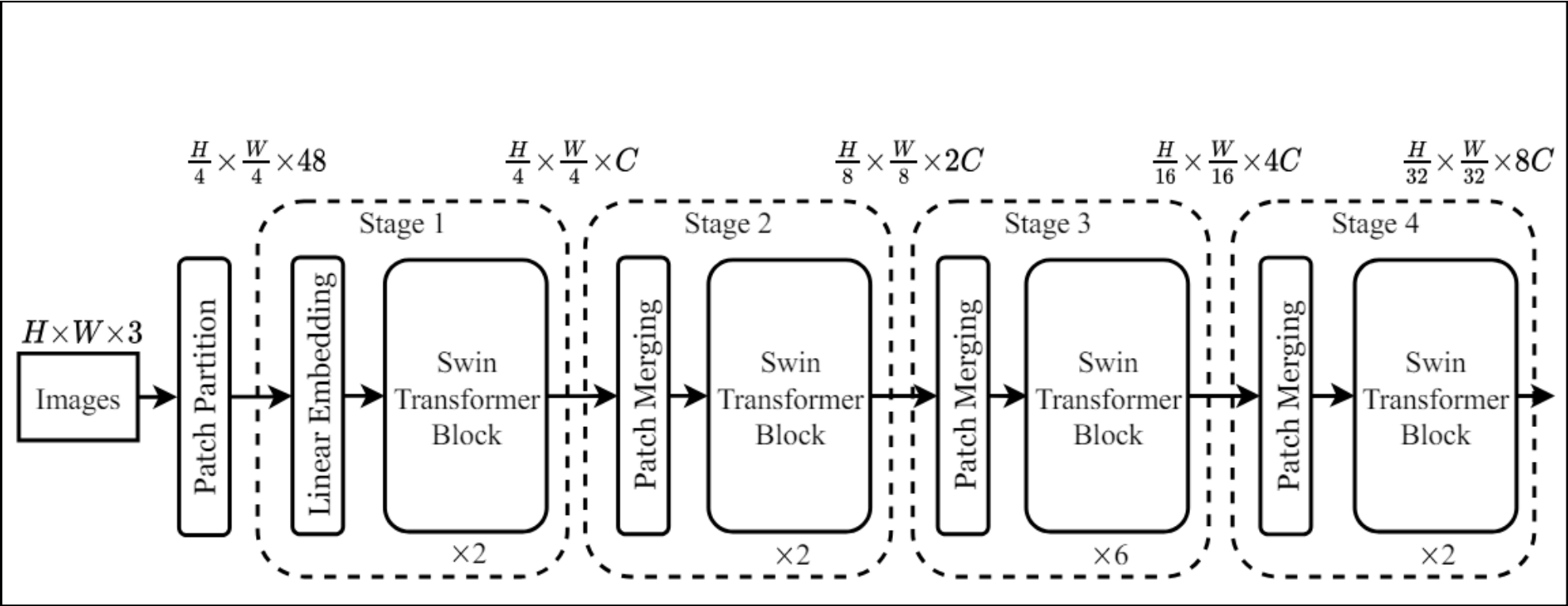
original transformer



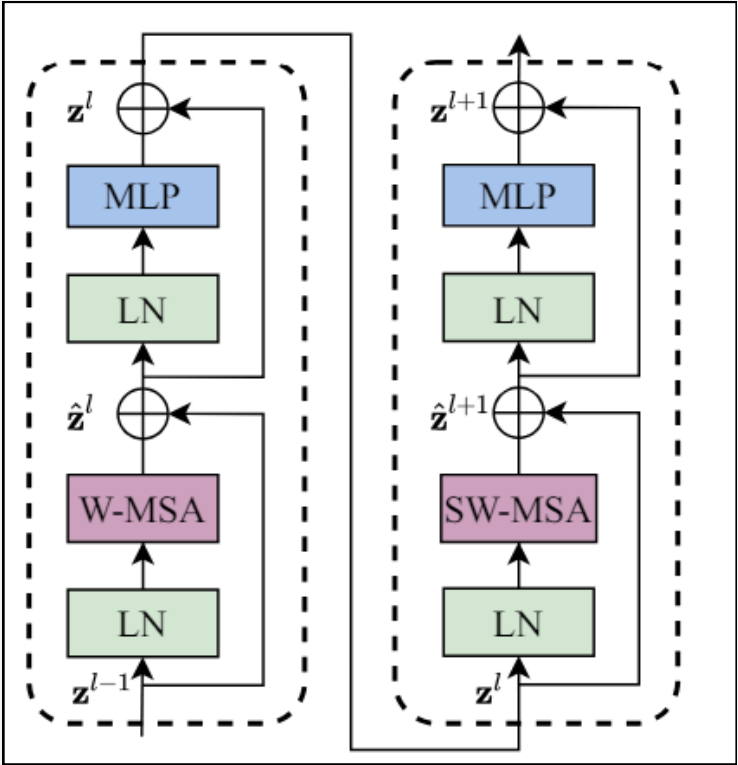
vision transformer



# SWIN-T ARCHITECTURE



Architecture of Swin-T



Transformer Block



# SWIN-T ARCHITECTURE

## IMPLEMENTATION

- Convert the loaded image to sequence of patches
- Patch Embeddings
- Positional Encoding
- Transformer Encoder
- Token Classification
- Training and Fine-Tuning
- Evaluation





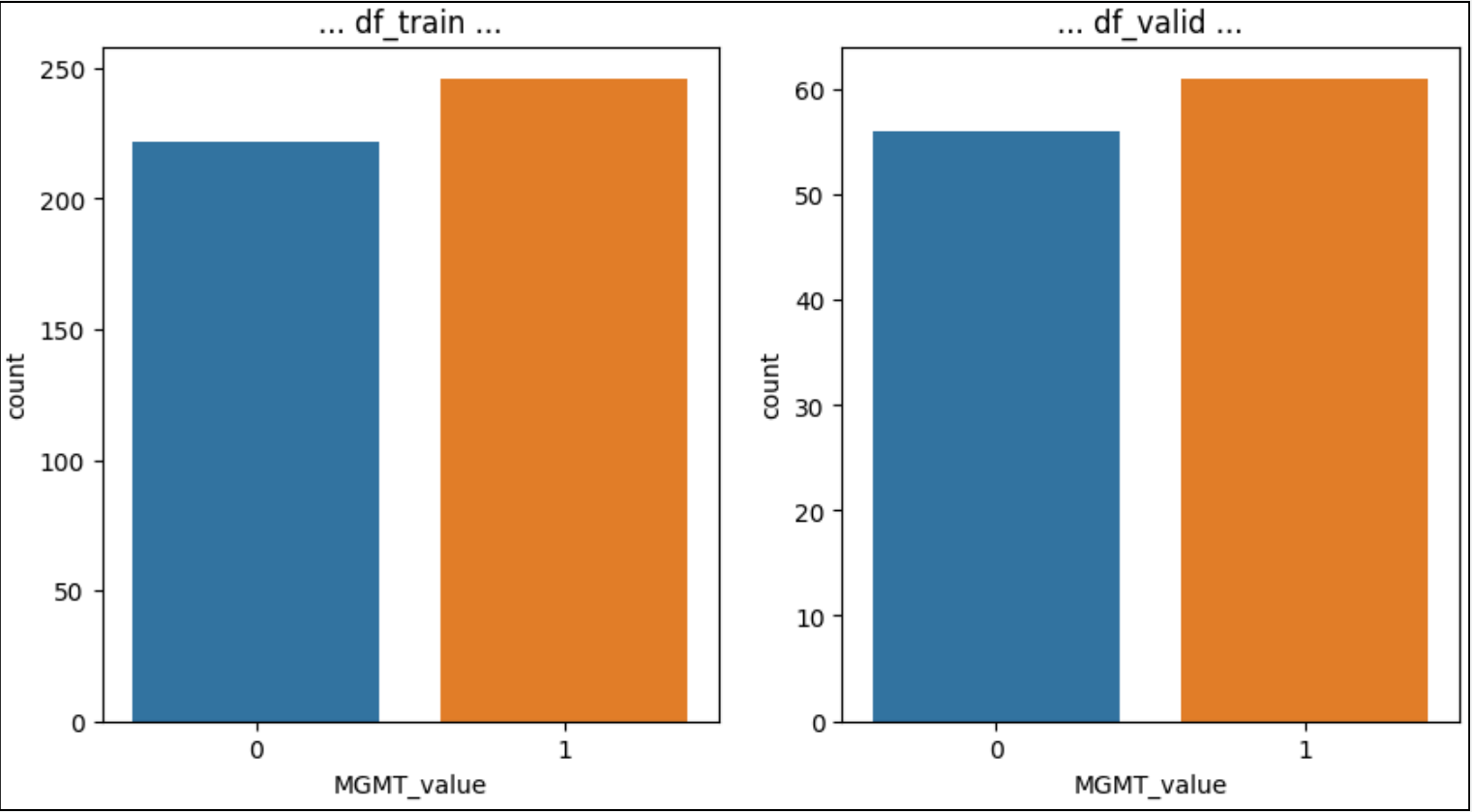
# RESULTS

BraTS21ID	MGMT_value
0	1
1	1
2	0
3	1
4	1
...	...
580	1
581	1
582	1
583	0
584	0

585 rows × 2 columns

df

after splitting df in 80:20 ratio



df\_train

df\_valid





# RESULTS

```
df_train['BraTS21ID_full'][481]

'00481'

df_train['FLAIR'][481]

'../input/rsna-miccai-brain-tumor-radiogenomic-classification/train/00481/FLAIR/'
```

dataframe manipulation for easy access  
of the images from each BraTS21\_ID folder

df\_valid.head()

	MGMT_value	BraTS21ID_full	FLAIR	T1w	T1wCE	T2w
BraTS21ID						
563	0	00563	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...
195	0	00195	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...
28	1	00028	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...
608	1	00608	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...
21	0	00021	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...	../input/rsna-miccai-brain-tumor-radiogenomic-...

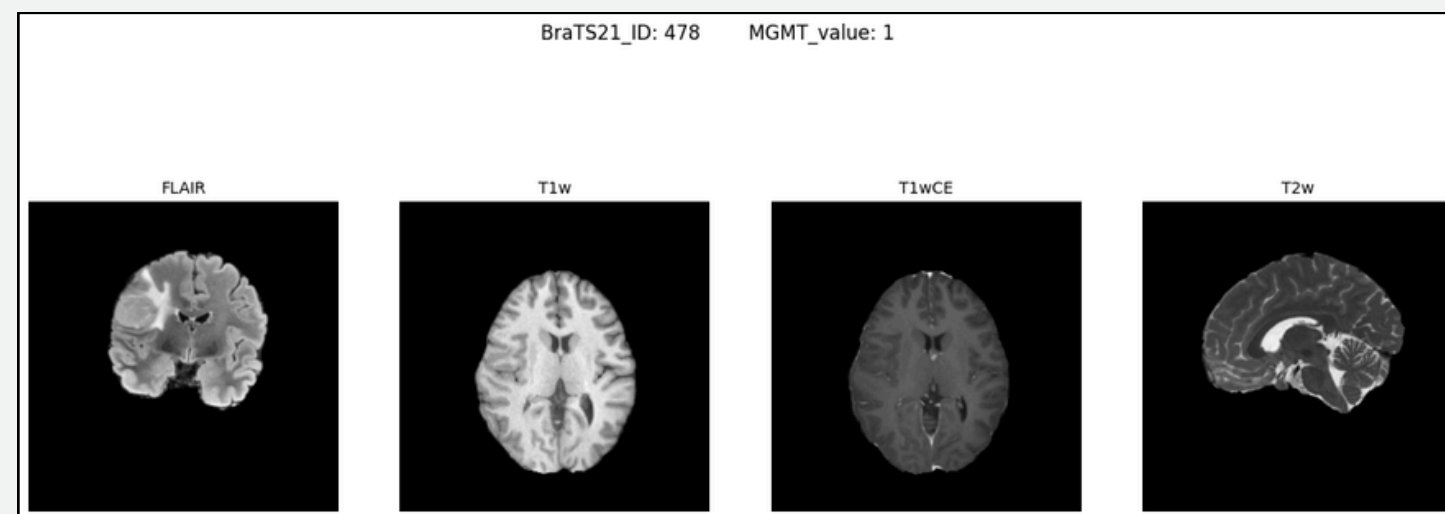
df\_valid



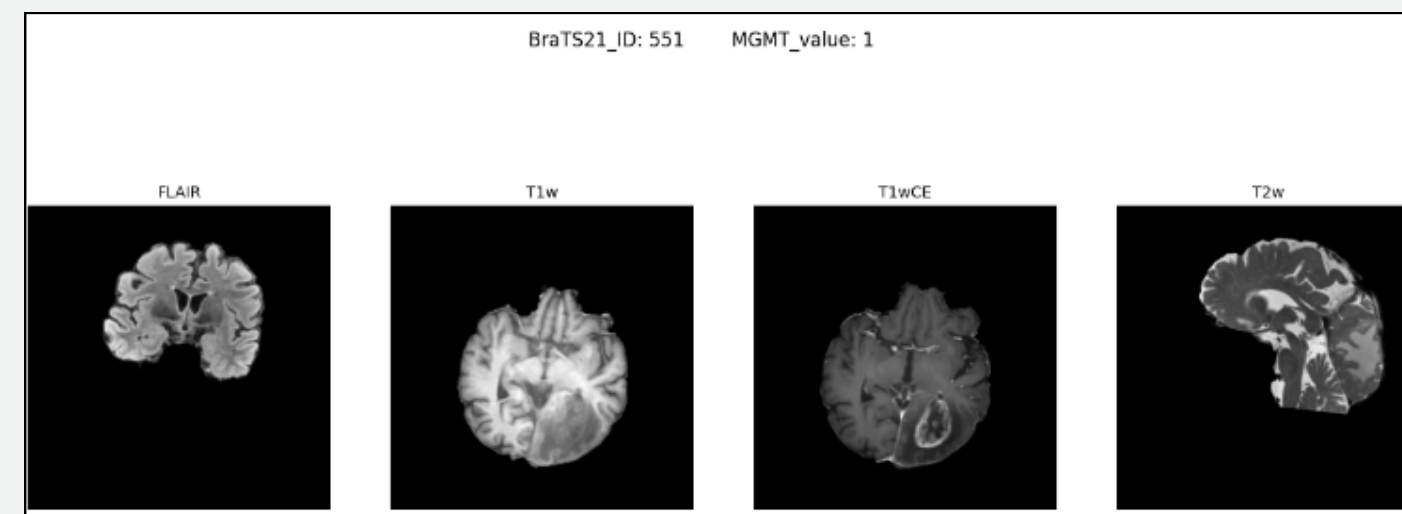


# RESULTS

Visualisation of different MRI types (FLAIR, T1w, T1wCE, T2w)



BraTS21\_ID (478)

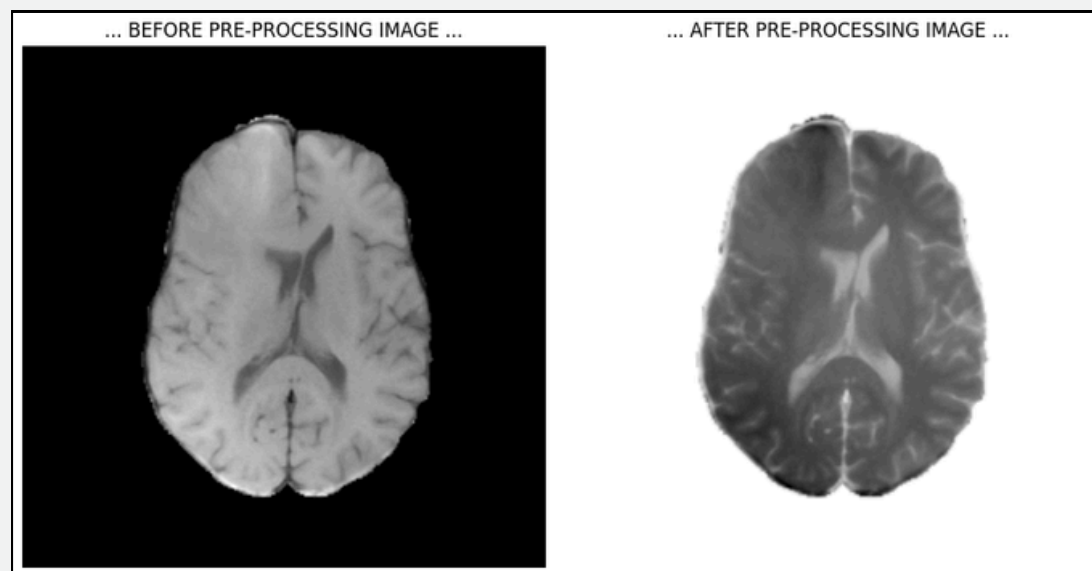


BraTS21\_ID (551)





# RESULTS



apply voit\_lut

```
a = load_dicom_images_3d("00481")  
print(a.shape)  
print(np.min(a), np.max(a), np.mean(a), np.median(a))
```

```
(1, 256, 256, 64)  
0.0 1.0 0.05419003281395742 0.00012306291639827978
```







# RESULTS

## Parametric values

<i>BraTS ipynb version</i>	<i>Optimizer</i>	<i>Learning rate</i>	<i>Epochs</i>	<i>Patience</i>	<i>Batch size</i>	<i>Depth</i>	<i>Patch embedding</i>
<i>(VIT Model) - version 2</i>	<i>Adam</i>	<i>0.001</i>	<i>10</i>	<i>10</i>	<i>4</i>	<i>2</i>	<i>Linear projection</i>
<i>(Swin VIT Model) - version 1</i>	<i>Adam</i>	<i>0.001</i>	<i>3</i>	<i>3</i>	<i>1</i>	<i>12</i>	<i>Linear projection</i>
<i>(Swin VIT Model) - version 4</i>	<i>Adam</i>	<i>0.001</i>	<i>5</i>	<i>5</i>	<i>2</i>	<i>12</i>	<i>Conv2D projection</i>
<i>(Swin VIT Model) - version 5</i>	<i>Adam</i>	<i>0.0005</i>	<i>10</i>	<i>10</i>	<i>4</i>	<i>16</i>	<i>Conv2D projection</i>
<i>(Swin VIT Model) - version 21</i>	<i>Adam</i>	<i>0.0003</i>	<i>15</i>	<i>10</i>	<i>4</i>	<i>16</i>	<i>Conv2D projection</i>
<i>(Swin VIT Model) - version 23</i>	<i>Adam</i>	<i>0.0002</i>	<i>25</i>	<i>10</i>	<i>4</i>	<i>16</i>	<i>Conv2D projection</i>

## Validation ensemble AUC and accuracy after prediction

```
df_valid["MGMT_pred"] = 0
for m, mtype in zip(modelfiles, mri_types):
    pred = predict(m, df_valid, mtype, "train")
    df_valid["MGMT_pred"] += pred["MGMT_value"]
df_valid["MGMT_pred"] /= len(modelfiles)

auc = roc_auc_score(df_valid["MGMT_value"], df_valid["MGMT_pred"])
print(f"Validation Ensemble AUC: {auc:.4f}")

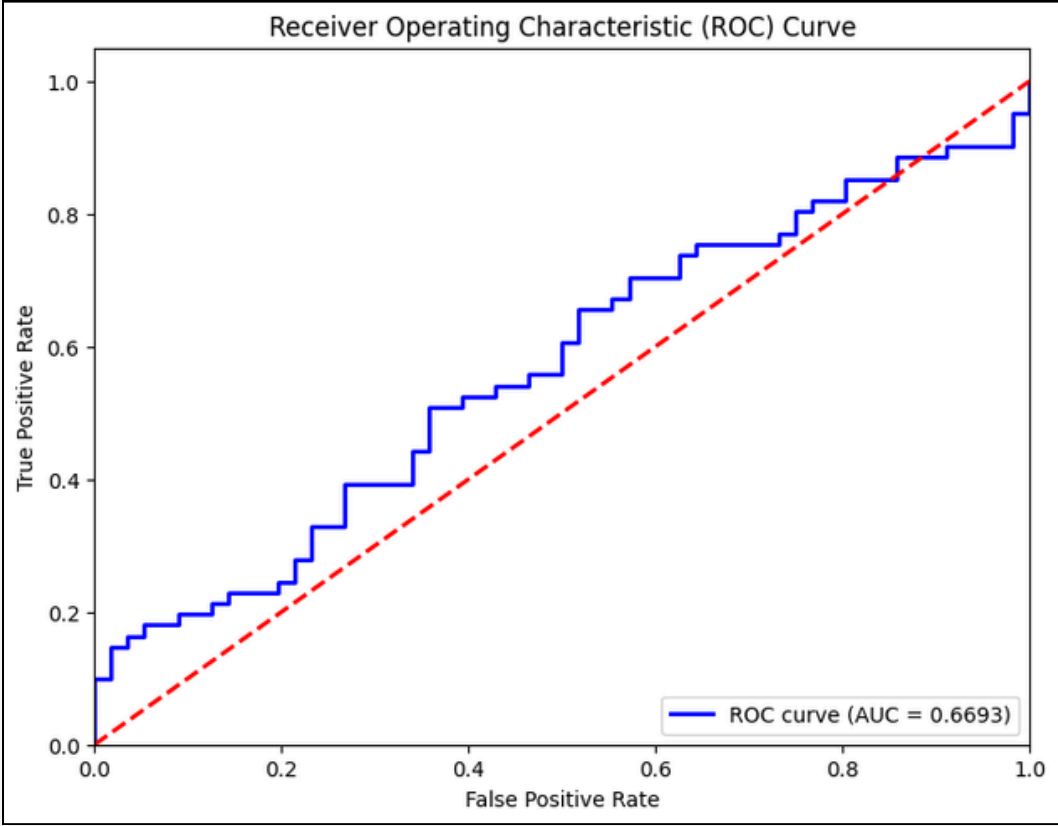
predicted_labels = (df_valid["MGMT_pred"] >= 0.5).astype(int)
actual_labels = df_valid["MGMT_value"]
accuracy = accuracy_score(actual_labels, predicted_labels)
print(f"Validation Ensemble Accuracy: {accuracy:.4f}")
```

Predict: FLAIR-best.pth FLAIR (117, 8)  
Predict: T1w-best.pth T1w (117, 8)  
Predict: T1wCE-best.pth T1wCE (117, 8)  
Predict: T2w-best.pth T2w (117, 8)  
Validation Ensemble AUC: 0.6693  
Validation Ensemble Accuracy: 0.6714

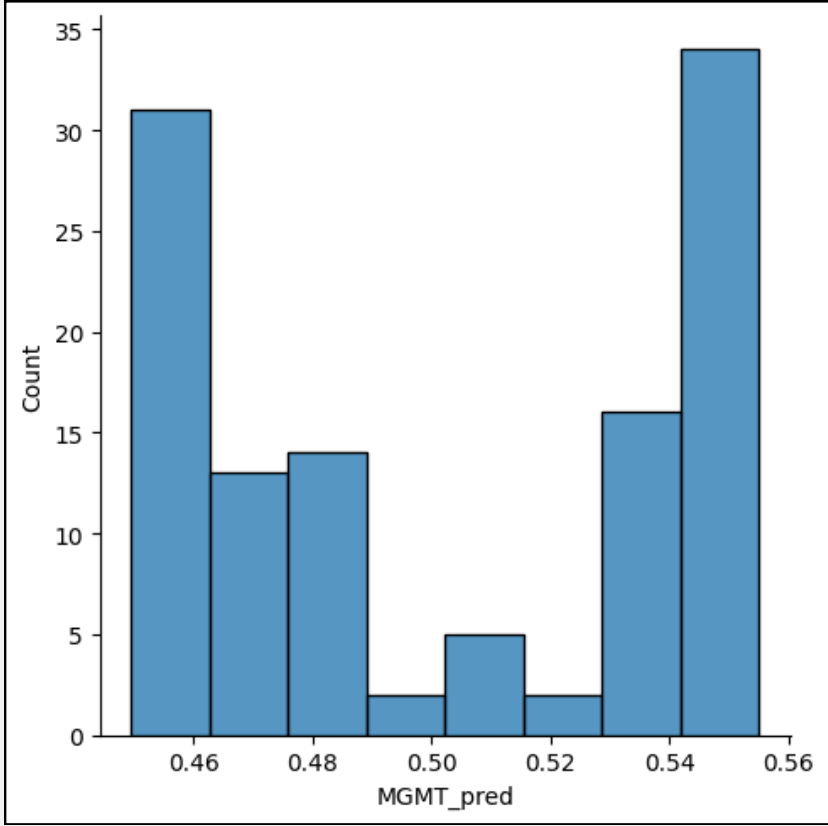




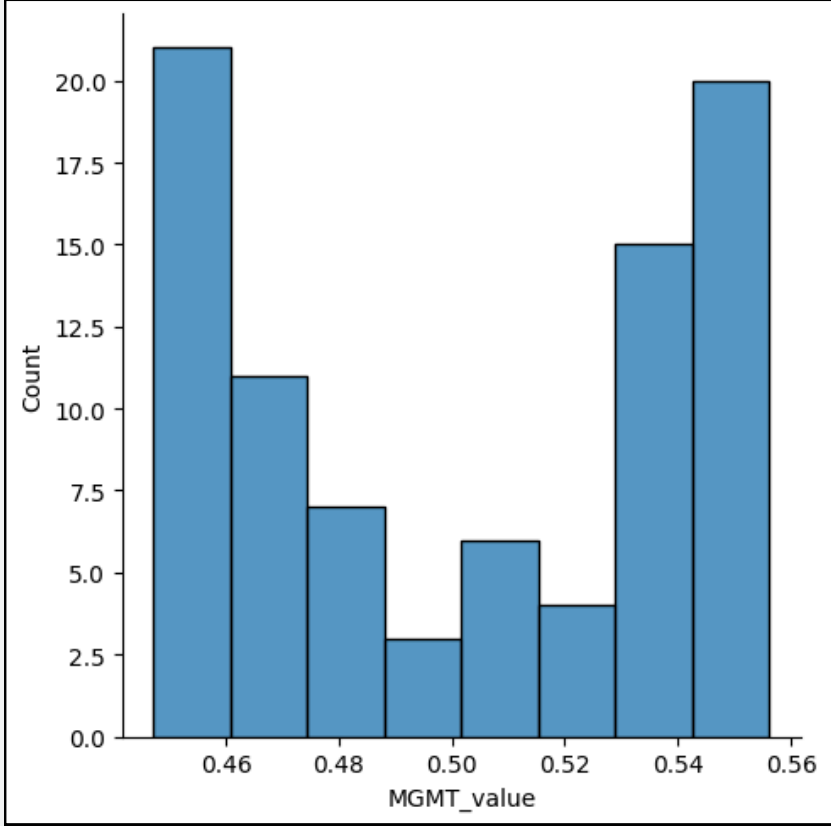
# RESULTS



ROC curve obtained on df\_valid



df\_valid[MGMT\_pred]






submission[MGMT\_value]








# RESULTS

Submission and Description	Private Score ⓘ	Public Score ⓘ
 <b>BraTS (VIT Model) - Version 2</b> Succeeded (after deadline) · 1mo ago · BraTS (VIT Model)   Versio...	<b>0.44390</b>	<b>0.43208</b>
 <b>BraTS (Swin VIT Model) - Version 1</b> Succeeded (after deadline) · 1mo ago · Notebook BraTS (Swin VIT...	<b>0.58305</b>	<b>0.56421</b>
 <b>BraTS (Swin VIT Model) - Version 4</b> Succeeded (after deadline) · 1mo ago · Notebook BraTS (Swin VIT...	<b>0.59593</b>	<b>0.58694</b>

Initial submissions and poor performance






Submission and Description	Private Score ⓘ	Public Score ⓘ
 <b>BraTS (Swin VIT Model) - Version 5</b> Succeeded (after deadline) · 1mo ago · Notebook BraTS (Swin VIT...	<b>0.61586</b>	<b>0.62896</b>
 <b>BraTS (Swin VIT Model) - Version 21</b> Succeeded (after deadline) · 13d ago · Notebook BraTS (Swin VIT ...	<b>0.61503</b>	<b>0.61945</b>
 <b>BraTS (Swin VIT Model) - Version 23</b> Succeeded (after deadline) · 13d ago · Notebook BraTS (Swin VIT ...	<b>0.61125</b>	<b>0.62262</b>

Best submissions sorted based on private scores





# RESULTS

#	△	Team	Members	Score
1	▲ 1039	[Tunisia.ai] I love this competition		0.62174
2	▲ 595	Minh Phan		0.61881
3	▲ 676	Cedric Soares		0.61732
4	▲ 469	Leaky Folds	 	0.61562

Surpassed the fourth best submission by “Leaky Folds” with a private AUC score of 0.61586





# NOVELTY

- Introduction of the latest computer vision techniques, such as Vision Transformers and their variants, in the field of brain tumor radiogenomics.
- Development of a transformer model that incorporates patch embedding and positional embedding layers from the Vision Transformer.
- Incorporation of Shifted Windows and a hierarchical structure from the Swin Transformer, resulting in a hybrid model that combines elements of both the Vision Transformer and Swin Transformer.





# CONCLUSION

- Submission was evaluated based on the area under the ROC curve between the predicted probability and the observed target.
- Obtained an AUC score of 0.61586
- Predict genetic subtypes of glioblastoma from MRI scans and reduce invasive procedures to enhance patient outcomes.
- Represents a significant advancement in precision medicine for neuro-oncology





# REFERENCES

- 🔍 Radiogenomic classification for MGMT promoter methylation status
- 🔍 Structural and Advanced Imaging in predicting MGMT promoter methylation
- 🔍 RSNA - MICCAI Brain Tumor Radiogenomic Classification Challenge
- 🔍 Vision Transformer Model
- 🔍 EfficientNet3D Brain Tumor Radiogenomic Classification





# THANK YOU

Presentation by M. Satya Sai Teja

