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Abstract

- A brain tumor: an **abnormal growth of tissue**, which is found to be unchecked by the normal cell-cycle control mechanism.
- Early detection of brain tumors is imperative for **enhancing patient care, tailoring treatment strategies, and saving lives**.
- Manual segmentation requires a high level of expertise and is time-consuming.
- This study proposes a **graph attention** (GAT)-based model by exploiting multichannel (FLAIR, T1CE, T1, and T2) MRI modalities for brain tumor segmentation.
- On average, the model performs 6% better than a baseline, on the dice score evaluation metric.

Objectives

- Localizing core, enhancing, and whole tumors precisely.
- Introducing a computationally efficient GAT model for unstructured MRI brain tumor segmentation.

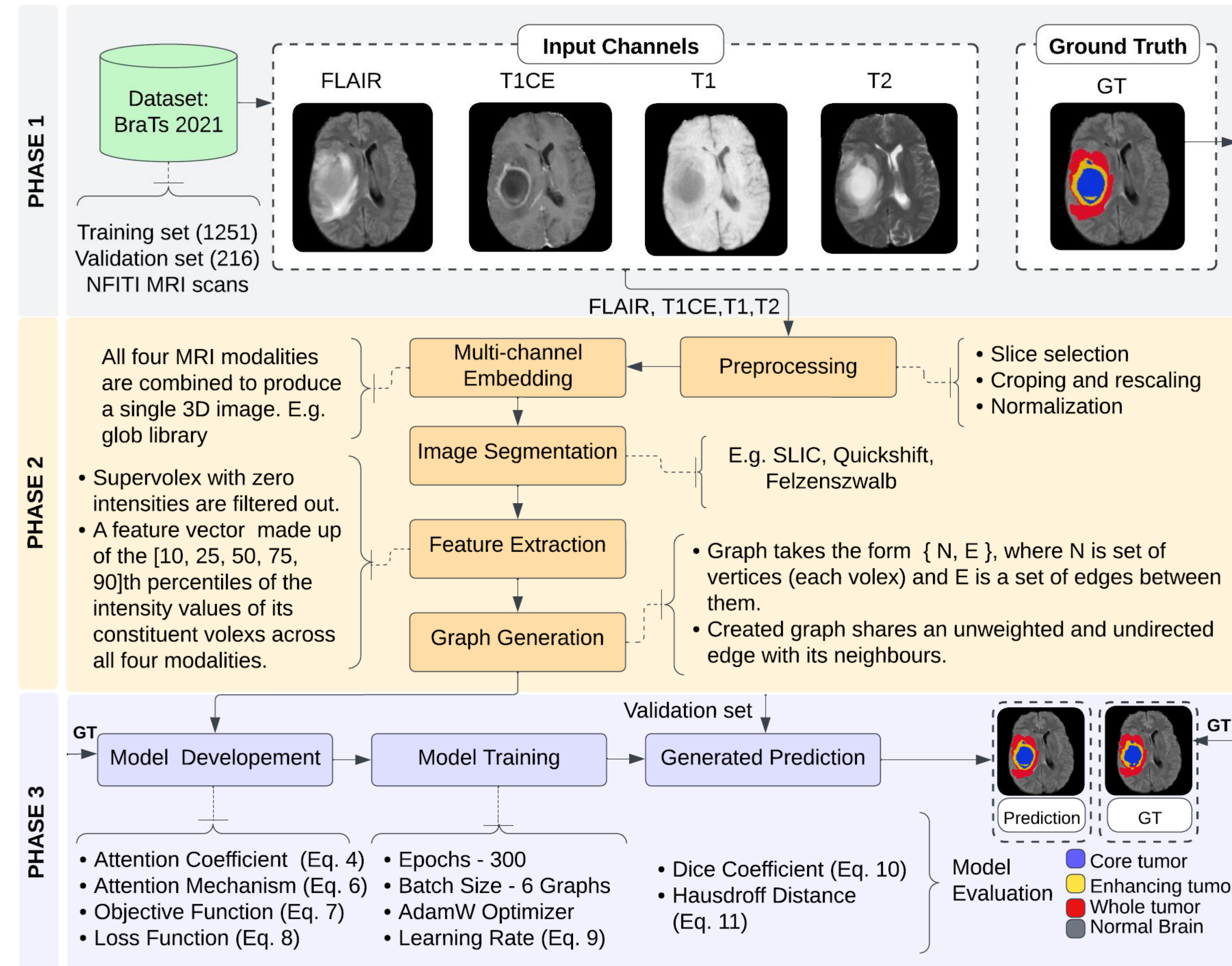
Challenges

- Brain tumors can occur anywhere in the brain and can vary greatly in size, form, and morphology.
- MRI scans are distorted and poorly contrasted due to non-uniformity in the static magnetic field and non-linearity in the gradient magnetic field.
- Training requires high computational power.

Datasets

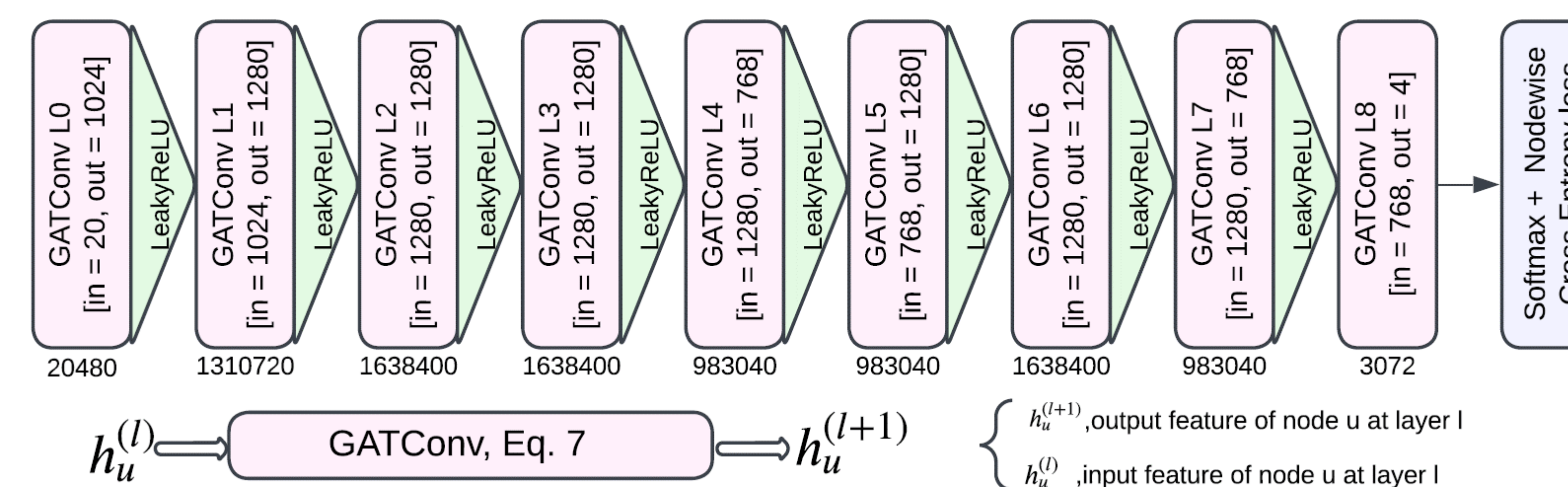
- BRATS 2021.
- Two different sets of data namely Training set (1251 patient's data), and Validation set (216 patient's data).

Proposed Solution



Detailed flow diagram of the proposed solution. It consists of three phases. **Phase 1:** Data Collection, **Phase 2:** Data Curation, and **Phase 3:** Model Development and Evaluation.

Layer-wise Schematic of the Proposed GAT



Detailed layer-wise schematic of the GAT: It subsumes GATConv layers alternated with a non-linear activation. Every GATConv layer updates each node's features by sampling neighboring nodes and aggregating the features by concatenation. LeakyReLU is applied with negative input slope $\alpha = 0.2$ for calculating the. The output of GAT has an associated node-wise multi-label cross-entropy loss. Numbers below each layer represent the trainable parameters for respective layers.

Methodology

Phase 1: Fusion of multimodality MRI data, including FLAIR, T1CE, T1 and T2.

Phase 2:

2.1. Performed image segmentation of 3D MRI using SLIC and generated feature vector based on the intensity values for each Voxel.

2.2. Produced graph in the form of $\{N, E\}$, where N is set of vertices and E is a set of edges between them.

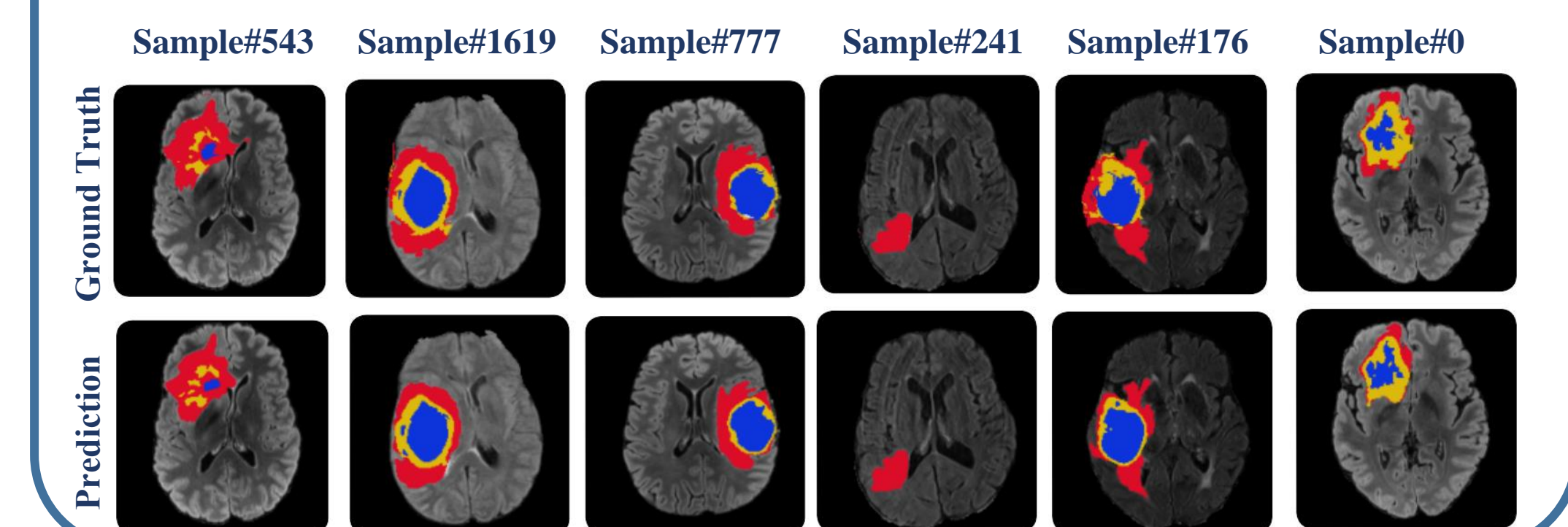
Phase 3:

3.1. Training and validation of GAT model using BRATS datasets.

3.2. Evaluation - segmentation results of all type of tumors are evaluated using dice coefficient.

Experimental Results

Method	Tumor Subregion Dice Score			Avg. Dice Score	% of Improvement
	WT	TC	ET		
GNN [2]	0.87	0.78	0.74	0.80	Baseline
GNN-CNN [2]	0.89	0.81	0.73	0.81	1.5 \uparrow
3D CMM-net [3]	0.84	0.81	0.75	0.80	-
3D-UNet [4]	0.87	0.76	0.73	0.79	1.0 \downarrow
GAT (ours)	0.91	0.86	0.79	0.85	6.0 \uparrow



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