An UNet-based Brain Tumor Segmentation Framework via Optimal Mass Transportation Pre-processing



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BraTS2022 Challenge

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Problems:

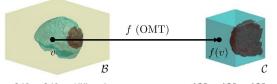
- Due to the large input size for training, we only use small batch sizes.
- Enhancing tumor (ET) is so small and dispersed that models are difficult to detect.

Proposed method:

- Use OMT to deform irregular objects into regular objects, which reduces the shape of model input and speeds up model training.
- Detect the WT roughly in phase 1 and segment the WT, TC, and ET in phase 2. The tumor's proportion can enlarge by the density function in phase 2.
- Generate the data augmentation by using OMT with different density function.

Optimal Mass Transportation (OMT) Problem

OMT Problem [2, 4]:



 $240\times240\times155\times4$

 $128\times 128\times 128\times 4$

$$f^* = \underset{f \in \mathsf{MP}}{\arg\min} \sum_{v \in \mathcal{V}(\mathcal{B})} \|v - f(v)\|_2^2 \ m(v), \tag{1}$$

• the function space of mass-preserving map is

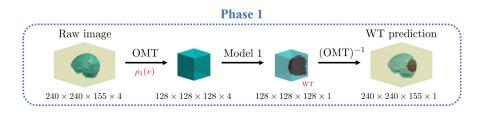
 $\mathsf{MP} = \left\{ f: \mathcal{B} \to \mathcal{C} \mid \rho(\tau) | \tau | = |f(\tau)|, \forall \tau \in \mathcal{T}(\mathcal{B}) \right\},$

• the local mass with the density function is

$$m(v) = \frac{1}{4} \sum_{\tau \in N(v)} \rho(\tau) |\tau|.$$

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Two-Phase OMT (2P-OMT) Framework: Phase 1

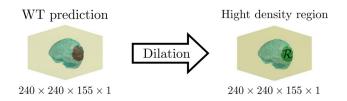


In phase 1, the density function is defined as

$$\rho_1(v) = \exp(\gamma \cdot \operatorname{HE}(g(v))), \tag{2}$$

where g is the grayscale of flair mode, HE is the histogram equalization, and $\gamma > 0$ is a hyper-parameter.

Two-Phase OMT (2P-OMT) Framework: Phase 2



In phase 2, the density function is defined as

$$\rho_{2}(v) = \begin{cases} \exp(\gamma \cdot \operatorname{HE}(g(v))), & \text{if } v \in \mathcal{R} \\ 1.0, & \text{if } v \in \mathcal{B} \setminus \mathcal{R} \end{cases}$$

$$Phase 2$$
Raw image
$$\overbrace{Q_{40} \times 240 \times 155 \times 4}^{\operatorname{OMT}} \overbrace{Q_{2}(v)}^{\operatorname{OMT}} \overbrace{Q_{2}(v)}^{\operatorname{Model 2}} \overbrace{Q_{40} \times 128 \times 128 \times 4}^{\operatorname{Model 2}} \overbrace{Q_{40} \times 128 \times 128 \times 3}^{\operatorname{Prediction}} 240 \times 240 \times 155 \times 3 \end{cases}$$

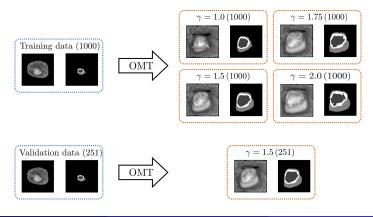
$$(3)$$

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Data Pre-processing and Augmentation (I)

- Split the dataset into 1000 for training and 251 for validation.
- Generate the OMT data with different densities (γ : 1.0, 1.5, 1.75, 2.0) and randomly choose one of the densities of OMT data for training.
- Choose the OMT data with $\gamma=1.5$ for validation step.



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Data Pre-processing and Augmentation (II)

γ	WT	ТС	ET
Raw	6.49%	2.42%	1.45%
1.0	13.47%	5.03%	3.04%
1.5	18.27%	6.84%	4.14%
1.75	20.93%	7.84%	4.75%
2.0	23.72%	8.90%	5.40%

Table 1: Average proportion of tumor.

- Use min-max normalization to rescale the grayscale to [0, 1].
- Use the data augmentation with following probability:

Transformation	Flip	Rotate	Add noise	Adjust brightness
Probability	0.25	0.25	0.1	0.1

Table 2: Probability of data augmentation.

Training Processing

- Epoch: 1000
- Batch size: 12
- Model: SegResNet [3] (MONAI package)
- Loss function: Dice Loss + Focal Loss
- Optimization: Adam + Weight decay + Cosine decay

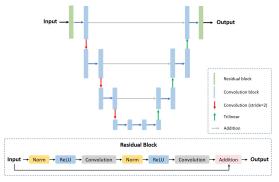
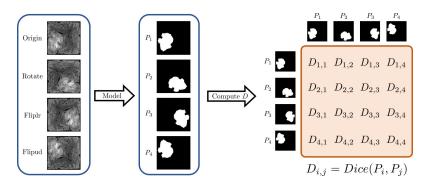


Figure 1: Architecture of SegResNet.

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Post-processing: Adaptive Ensemble



Let w be the normalized eigenvector corresponding to the spectral radius of D. Then the final prediction is

$$\widehat{P} = \sum_{i=1}^{4} w_i P_i.$$

Preprocessing	Cross-validation			Online validation		
	WT	ТС	ET	WT	ТС	ET
Random Crop	0.9340	0.9082	0.8650	0.9194	0.8542	0.8242
2P-OMT	0.9371	0.9157	0.8819	0.9214	0.8823	0.8411

Table 3: Compared to the Dice score of different preprocessing with SegResNet.

Paper [1]	Preprocessing	Model	WT	тс	ET
P. Druzhinina et al.	Random Crop	scan lite-20	0.9220	0.8680	0.8300
HY. Wu et al.	Random Crop	HarDNet-BTS	0.9260	0.8793	0.8442
J. Ma et al.	Random Crop	NnUNet	0.9259	0.8786	0.8217
H. S. Singh et al.	Random Crop	Attention UNet	0.9080	0.8600	0.8170
Proposed	2P-OMT	SegResNet	0.9214	0.8823	0.8411

Table 4: Comparison results with other participants in the BraTS2021 Challenge.

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- We developed the 2POMT framework, which significantly improved performance compared with random crop pre-processing.
- We can use density function to enlarge tumor proportion to help the model detection the tumor.
- We can generate OMT data with different densities achieving data augmentation, which can avoid over-fitting and enhance the robustness of a model.
- We proposed a novel ensemble method for post-processing, which can adaptively compute the weighted sum of predictions.

Reference

- Alessandro Crimi and Spyridon Bakas. Brainlesion: Glioma, Multiple Sclerosis, Stroke and Traumatic Brain Injuries: 7th International Workshop, BrainLes 2021 Held in Conjunction with MICCAI 2021 Virtual Event, September 27, 2021 Revised Selected Papers, Part I. 2022.
- [2] Wen-Wei Lin et al. "A novel 2-phase residual U-net algorithm combined with optimal mass transportation for 3D brain tumor detection and segmentation". In: *Sci. Rep.* 12 (2022), pp. 1–16.
- [3] Andriy Myronenko. "3D MRI brain tumor segmentation using autoencoder regularization". In: *BrainLes@MICCAI*. 2018.
- [4] Mei-Heng Yueh et al. "Projected Gradient Method Combined with Homotopy Techniques for Volume-Measure-Preserving Optimal Mass Transportation Problems". In: J. Sci. Comput. 88.3 (Sept. 2021). ISSN: 0885-7474.

Thanks for listening!