

Automatic Multi-Segmentation Method for Tumor Detection in MRI Images using Constrained kmeans Method and Region Growing-Quasi Monte Carlo Method

Abstract. Magnetic Resonance Imaging (MRI) has become an indispensable tool in the medical field, enabling the detection of critical abnormalities affecting various organs within the human body. Despite its inherent complexity, the development of automated or semi-automated detection and recognition techniques has made significant strides. In this paper, we present an innovative approach for the automatic multi and full segmentation of tumor regions within MRI scans. An enhanced region-growing method founded on the Quasi-Monte Carlo sampling and constrained k-means algorithm is presented in this paper, we define distinct classes to facilitate precise segmentation. The efficacy of our technique is evaluated through a range of metrics, demonstrating its robust performance. The proposed fully automated multi-segmentation method showcases superior results and holds potential to supplant conventional techniques for tumor detection in MRI images.

Streszczenie. Rezonans magnetyczny (MRI) stał się niezastąpionym narzędziem w medycynie, umożliwiającym wykrycie krytycznych nieprawidłowości wpływających na różne narządy w organizmie człowieka. Pomimo swojej nieodłącznej złożoności, rozwój zautomatyzowanych lub półautomatycznych technik wykrywania i rozpoznawania przyczynił znaczne postępy. W artykule przedstawiamy innowacyjne podejście do automatycznej wieloetapowej segmentacji obszarów nowotworowych w obrazach MRI. W artykule przedstawiono ulepszoną metodę powiększania regionów opartą na próbkowaniu Quasi-Monte Carlo i ograniczonym algorytmie k-średnich. Definiujemy odrębne klasy, aby ułatwić precyzyjną segmentację. Skutecznie naszą technikę oceniamy za pomocą szeregu wskaźników, co pokazuje jej solidne działanie. Proponowana w pełni zautomatyzowana metoda wielosegmentacyjna zapewnia doskonałe wyniki i może zastąpić konwencjonalne techniki wykrywania nowotworów na obrazach MRI. (Automatyczna metoda wielosegmentacyjna do wykrywania nowotworu w obrazach MRI przy użyciu metody ograniczonych kmean i metody Quasi Monte Carlo wzrostu regionu)

Keywords: Brain tumor, Multi Segmentation, Region growing constrained k-means, Quasi Monte carlo, naive Baye.

Słowa kluczowe: Guz mózgu, wielosegmentacja, rosnące w regionie średnie bayesowskie k, Quasi Monte Carlo.

Introduction

Magnetic Resonance Imaging (MRI) has revolutionized medical diagnosis and treatment by enabling non-invasive visualization of internal structures. Detecting abnormalities in organs is crucial for timely medical interventions. Automation of this process has gained prominence due to the sheer volume and complexity of MRI data. This paper introduces an automatic method for the multi-segmentation of tumor regions in MRI scans, leveraging an innovative combination of the quasi-Monte Carlo method and the Expectation Maximization algorithm. MRI segmentation can be done using three different methods, such as manual, semi-automatic and full automatic techniques [6]. For manual MRI segmentation, which is the most common technique, the segmentation is done by a doctor or an expert, and its accuracy depends on the performance and the knowledge of the doctor. Full automatic segmentation technique is an autonomous process and which need evolved algorithms for calculation and recognition. Medical image processing methods, used for full automatic segmentation, are classified into four main categories [6, 15]: Threshold based techniques, such as otsu and kapur thresholding, and adaptive thresholding [13] The second category is the region-based technique, such as region growing [9] and watershed [11] Third, the classification techniques that need a training phase, such as SVM and KNN and clustering methods, such as K-means and EM mixture [6] The last category is contour detection, such as ACM, GVF, VFC [16] and level set [1]. Many researchers have presented full automatic and hybridized MRI image segmentation model. Lu et al. [9] used an improved region growing algorithm initialized by the QMC method for liver segmentation. In their turn, W. Y. Zhanfang, and Hongbiao [19] used an improved PCNN method to perform automatic segmentation, however, their method was not applied for the segmented more complex areas. Kuwazuru et al. [8]

used hybrid method by combining ANN with the level-set method for segmentation of multiple sclerosis lesion (MS) of the brain. Their method is based on a concatenation of ANN and level set, but their method was unable to detect small areas. D. Veloz, and Allende [4] used modified EM to segment MRI images. In this paper, we employed enhancement and denoising filters to preprocess the image. Subsequently, we used the Kapur thresholding method to locate the region of interest (ROI). Then, we applied the quasi Monte Carlo method to generate a large number of seeds (Quasi Random Sampling). These seeds were grouped into k classes using an improved version of the K-means method, referred to as constrained K-means, where the spatial dependency of the samples is taken into account. The classification is established within a Naive Bayesian framework. After selecting the optimal seed for each cluster, we initialize our improved region growing approach.

Method

0.1 Preprocessing

As depicted in Figure 1, our approach is structured around three primary stages: preprocessing, localization, and segmentation and recognition. Initially, we employed the deformable model proposed by Rifai et al. [14] to remove the skull. Subsequently, we applied contrast enhancement to accentuate high-frequency regions using sigmoid filtering, as outlined in the work by Lu et al. [9]. The sigmoid filter modifies the distribution of gray levels to enhance dissimilarities between neighboring regions. The gray level of the resulting image is calculated using the equation:

$$p = (I_{\max} - I_{\min}) \cdot \frac{1}{1 + \exp\left(-\frac{p + \beta}{\alpha}\right)} + I_{\min}$$

Here, p signifies the gray level of the input image, I_{max} and I_{min} represent the maximum and minimum gray levels of the output image, respectively. α corresponds to the width of the intensity range of the input image, and β indicates the central point of this range.

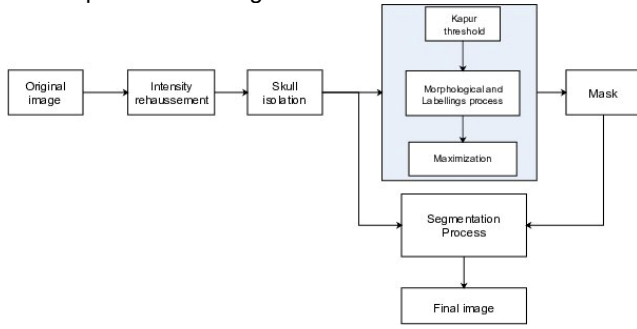


Fig. 1. Overview flowchart of image preprocessing

Subsequently, we employed the thresholding method based on the Kapur algorithm [13] to isolate the tumor region, which stands out due to its enhanced color. We computed the entropies of the object H_{ROI} and the background H_{Bg} using the following equations:

$$H_{ROI}(t) = \sum_{i=0}^t \frac{\Pr(p_i)}{\Pr(ROI)} \ln \left(\frac{\Pr(p_i)}{\Pr(ROI)} \right)$$

$$H_{Bg}(t) = \sum_{i=0}^t \frac{\Pr(p_i)}{\Pr(Bg)} \ln \left(\frac{\Pr(p_i)}{\Pr(Bg)} \right)$$

where $\Pr(ROI) = \sum_{i=0}^t P_r(p_i)$, $\Pr(Bg) = \sum_{i=0}^{+\infty} P_r(p_i)$ and

$\Pr(ROI) + \Pr(Bg) = 1$ represents the threshold value at the maximum values of $H_{ROI}(t)$ and $H_{Bg}(t)$.

Next, we applied morphological processing to decrease the number of connected regions. Afterward, we labeled regions consisting of connected pixels and identified the region with the highest pixel count as the Region of Interest (ROI).

0.2 Segmentation

In this section, we introduce a new segmentation approach comprised of three pivotal steps, which is an improved iteration of the method proposed in [23]. The initial step involves seed generation, followed by seed clustering into k -classes in the subsequent step. Ultimately, the multi-segmentation is executed after the optimal seeds are selected. The segmentation procedure is visually depicted in Figure 2. The strength of the constrained k-means method lies in its capacity to consider neighboring pixels during the classification process, in contrast to the EM algorithm. This characteristic contributes to a more homogeneous classification. The objective of the Quasi Monte Carlo method is to generate a discrepancy sequence of pixels L in our Region (ROI). To ensure the good coverage of the ROI, we generated the sequence in a rectangle (referenced as RECT) that covers the area and which has the following parameters

$$x_{min} = \min(row) \quad x_{max} = \max(row)$$

$$y_{min} = \min(col) \quad y_{max} = \max(col)$$

Where row and col are, respectively, the Cartesian coordinate vectors of the mask ROI. the Halton sequence was adopted to generate the points with:

$$x_i = [x_{min} + [(x_{max} - x_{min})] \times h_2$$

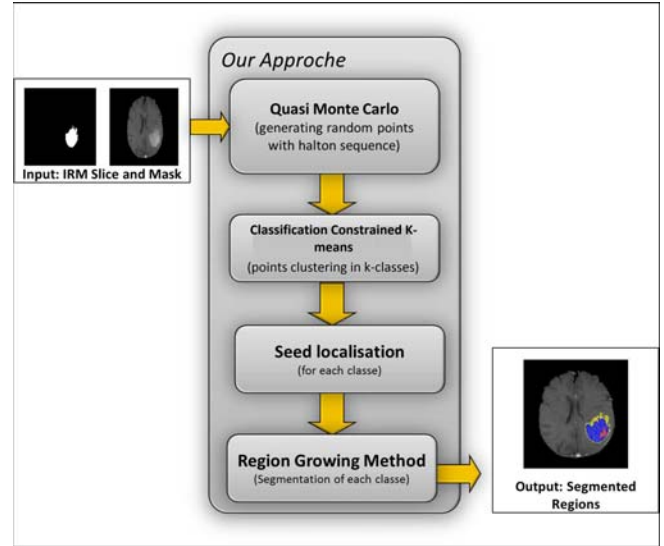


Fig. 2: segmentation flowchart

$$y_i = [y_{min} + [(y_{max} - y_{min})] \times h_3$$

with $[x_i]$ is the nearest integer to x_i , h_2 and h_3 are respectively Halton coefficient of base 2 and 3, In the end of this step we obtain a set of points S_L inside $RECT$. we used a mask M of ROI to get a subset S_i from S_L , this sets can be written as:

$S_L = \{p_1, p_2, \dots, p_L\}$ where $p_i = (x_i, y_i)$ and

$$\begin{cases} x_{min} \cdot x_i < x_{max} \\ y_{min} \cdot y_i < y_{max} \\ \forall i = 1, 2, \dots, L \end{cases}$$

with $S_i = \{p'_1, p'_2, \dots, p'_i\}$ where $p'_i \in S_i$ and $S_i \in S_L$

0.2.1 Seeds clustering

We employed a statistical method to partition the subset S_i into k classes denoted as C_i , where $i = 1, \dots, k$. This clustering process enabled the creation of pixel subsets corresponding to distinct regions within our Region of Interest (ROI). To achieve this, we utilized the constrained k-means algorithm within a naive Bayesian framework [22]. Notably, this algorithm excels in providing optimal classification by incorporating neighboring pixel information. To further enhance the effectiveness of these classes, the constrained k-means algorithm initializes the parameter vector with the state n , and subsequently, we maximize the a posteriori probability to estimate the new state class parameters $(n + 1)$.

0.2.2 K-Means Clustering

K-Means clustering is an unsupervised algorithm that is used to form different clusters of data sets so that they can be grouped together. A cluster is a collection of similar (homogeneous) data objects in one cluster and diverse (heterogeneous) data on objects in another cluster (see figure. 5). K-means is a clustering algorithm based on optimizing the criteria function. If the sample data is presented as aggregate $X = \{x_1, x_2, \dots, x_n\}$, x_i is a d -dimensional vector, and suppose the number of clusters is k , the initial K-means center is $C_i(0)$. The similarity measurement adopts Euclidean distance, as for α and β .

$$D = [\alpha - \beta] = \sqrt{(\alpha - \beta)^T (\alpha - \beta)}$$

Grouping criteria adopt the number of squared errors.

$$J = \sum_{i=1}^k \sum_{x \in C_i} [x - C_i]^2$$

The steps in the K-Means clustering as follows:

1. Initialization of parameters: specify cluster k and center, initial K-Means $C_i(0)$ are specified as random data points, where $j = 1, 2, \dots, k$.
2. Repeat revision: allocates each x_i data from the data set $X = \{x_1, x_2, \dots, x_n\}$ to class $C_p(l)$ when:

$$|x_i - C_p(l)| < |x_i - C_q(l)|$$

Where l for iterations. $p, q = 1, 2, \dots, n, p \neq q, l = 1, 2, \dots, n$.

3. Update center cluster center: new cluster center on $l+1$ calculation

$$C_j(l+1) = \frac{1}{N_j} \sum_{x_i \in C_j(l)} x_i$$

Where N_j is the amount of data in the cluster j .

4. Stop the iteration if $C_i(l+1) = C_i(l)$ or $|C_i(l+1) - C_i(l)| < \epsilon$, if not repeat to step 2.

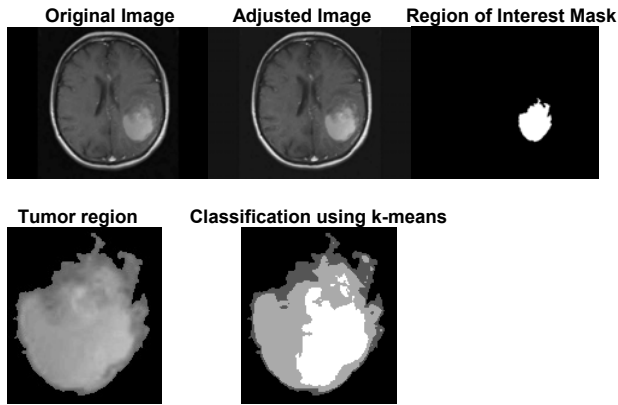


Fig. 3: Detection and classification process

0.3 Naive Bayes Classifier

To enhance the classification process, we utilize a Naive Bayes Classifier in conjunction with K-means. This combination demonstrates commendable performance when compared to other classifiers, owing to its simplicity, lower computational complexity, minimal memory demands, and strong predictive accuracy. The calculation of the Naive Bayes Classifier is delineated as follows (see figure. 4):

$$P(C_j|X) = \frac{P(X|C_j)P(C_j)}{P(X)}$$

Where, posterior probability, $P(C_j|X)$, is calculated using Class Prior Probability $P(C)$, Predictor Prior Probability $P(X)$ and Likelihood $P(X|C)$. The a posteriori estimation is achieved using iterated conditional mode [28]. The proposed method considers input data sets with attribute values as numerical and Gaussian distributions. For the Gaussian distribution the mean (μ) and standard deviation (σ) need to be calculated using the formula:

$$\mu = \frac{X_1 + X_2 + \dots + X_n}{n}$$

$$\sigma = \frac{\sum_{i=1}^n (X_i - \mu)^2}{n}$$

While the Gaussian distribution function is calculated by the

formula:

$$f(X) = \frac{1}{2\pi\sqrt{\sigma}} \exp - \frac{(X - \mu)^2}{2\sigma^2}$$

The clustering process is considered as an initiator to our segmentation method. Each class must have an initial seed P_{seed_i} . Figure. 5, the initiator seed must verify this condition

$$|I(P_{seed_i} - m_i)| \min_{1 \leq j \leq l} \{|I(P_j^i) - m_i|\}$$

with $I(P_j^i)$ is the pixel intensity of P_j^i



(a) Tumor classification using classical Kmeans
(b) Tumor classification using constrained K-means

Fig. 4: MAP estimation of image classification using iterated conditional mode

0.3.1 Multi Segmentation

After selecting the initial seeds, the segmentation process for each class will be carried out using a modified region-growing method. The primary inconveniences associated with the classical region-growing approach are related to the selection of the initial seed and the homogeneity criteria, as noted by [9].

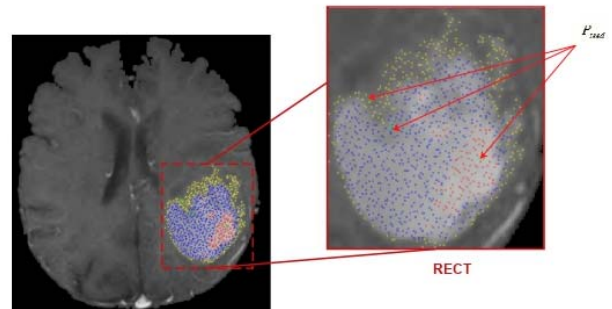


Fig. 5: distribution of points in k classes $k = 3$

The goal of the growth stage is to expand the region by incorporating neighboring pixels. This is achieved through a similarity measure that identifies connected pixels. Consequently, if the initial seed is placed in an area with significant inhomogeneity, the similarity measure may lead to substantial changes, potentially causing the growth process to halt prematurely. Thus, it is crucial to select starting points in the most homogeneous areas whenever possible.

On the other hand, a poor criterion could result in either only a partial region being covered or an excessively larger part than the region being included. Pixels are incorporated into the region based on the validity of the homogeneity criterion. To address this, we have initiated the segmentation process using an approach denoted as Pseedi for each class. To prevent overlaps between regions and maintain the integrity of segmented edges, gradient intensity information is utilized. For more comprehensive details about the segmentation process, please refer to Algorithm 1.

Algorithm 1 Algorithm: Segmentation process

Require: P_{seed_i} : initiator seed for classe i
 C_i :pixels set allowed to classe i
 δ_{int} : minimal intensity distance
 $\delta_{gradiene}$: minimal gradient distance
 m_i :initiale average of classe i
 G_{mi} :The gradient average of classe i

Ensure: R_i :Region i
 Initialize the region R_i by the pixels assigned to the class
repeat
 $V(P_{seed_i})$ =neighborhood of P_{seed_i}
 $border_{list} = border_{list} \cup V(P_{seed_i})$
 Order $border_{list}$ in ascending order compared to δ_{int} .
 P_{seed_i} = the first pixel of $border_{list}$ who check

$$(((I(p) - m_i) < \delta_{int})) \&\&$$

 (Gradient(p)- G_{mi}) < $\delta_{gradiene}$)]if P_{seed_i} exist then
 $R_i = R_i \cup P_{seed_i}$
 $border_{list} = border_{list}(P_{seed_i})$
 Update m_i
else
Break
end if
until P_{seed_i} not exist

Discussion and Experimental results

0.4 Materials and Database

We developed and tested our approach using Matlab 2018b as simulation software running on ubuntu 16.4 operating system, with cpu 2,20 GHz and processor intel core i7 with 8 Gmb of Memory (RAM). We used 2 images from Brats Miccai 2015 database [20,21], which contain low-grade and high-grade images of subjects segmented by radiologists into four sub-compositions of tumors.

0.5 Result and discussion

We used median filter to reduce the noise of the MRI images. Multi-segmenting was achieved using improved region growing methods initialized with constrained kmeans algorithm and Quasi Monte Carlo algorithms. The MRI images have been segmented into two regions, i.e., k equal to 2, in order to recognize two zones. These two classes correspond to edema (R_1) and enhanced tumor (R_2). Figure.6 (a,d) shows the MRI images of the open access database. Figure.6 (b,e) corresponds to the reference images made by the expert radiologist, and Figure6(c,f) represents the segmentation results using the method developed in this study.

In the aim to evaluate the performance of our approach. We use the confusion matrix. [17] to compute multiple Metrics in order to evaluate the performance of our approach, In order to measure the amount of overlap between the ground truth and our segmentation using three metrics (Dice, Sensitivity, Specificity) high value corresponds to a high level of overlap between the field and the reference.

$$Dice = \frac{2TP}{2TP + FP + FN}$$

$$Sensitivit y = TPR = \frac{TP}{TP + FN}$$

$$Specificit y = TNR = \frac{TN}{TN + FP}$$

where TP : true positive is the number of tumor pixels correctly detected, TN : true negative is the number of non tumor pixels correctly not detected, FP : false positive correspond to non-tumor pixels have been incorrectly detected as tumor pixels, FN : false negative is the number of non tumor pixels falsely detected. Thereafter, two metrics have been calculated, Hausdorff Distance (HD) and the mean minimum distance. Hausdorff Distance was calculated using the equation (see Table.2):

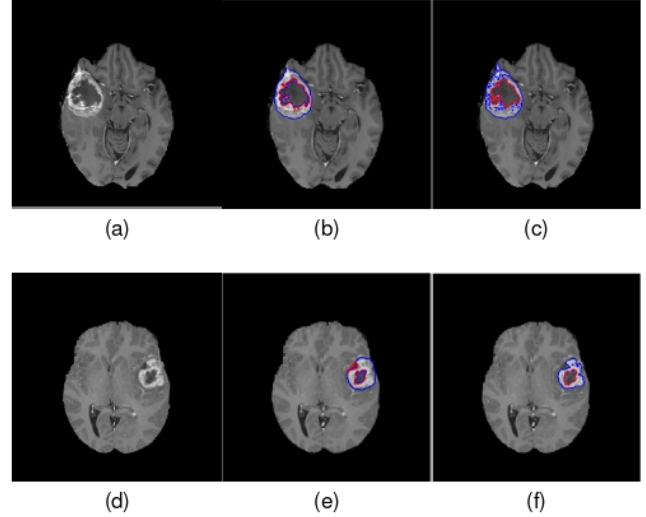


Fig. 6: (a,d): Original images, (b,e): segmentation of the expert, (c,f): segmented images using our approach.

$$HD(\text{Distance Hausdorff}) = HD(A,B) \max(h(a, b), h(b, a))$$

$$\text{with } h(a, b) = \max_{a \in A} (\min_{b \in B} \|a - b\|)$$

$$AVD = \max(d(A,B), d(B,A))$$

Table 1. illustrates the accuracy and sensitivity performance of the method developed in this research. The results showed that dice changes in the range of 0.85 to 0.91 for Edema and from 0.77 to 0.84 for Enhanced tumor and the sensitivity changes in the range of 0.75 to 0.84 for Edema and from 0.64 to 0.65 for Enhanced tumor.

Table 1. Performance analysis of our method

images	Dice		Sensitivity		Specificity	
	R_1	R_2	R_1	R_2	R_1	R_2
1	90,12	71,35	80,25	54,92	100	99
2	89,73	82,14	80,54	68,55	100	99
3	91,44	76,90	84,28	63,80	100	99
4	90,12	71,54	81,35	55,43	100	98
5	85,38	75,14	75,92	65,34	98	98

Table 2. presents the range of Average Volume Difference (AVD) for both edema and enhanced tumor, which varies between 0.010 to 0. and 0.02 to 0.25, respectively. Additionally, the Hausdorff Distance (HD) ranges from 2 to 10.44 pixels for edema and 2.23 to 10 pixels for enhanced tumor. These results serve as strong evidence affirming the excellent performance of the automatic segmentation approach developed in this study.

Table 2. Distance analysis of the fully automatic segmentation approach

		Images					
		1	2	3	4	5	6
HD	R_1	3.7454	4.0125	5.1551	1.712	1.8021	8.5421
	R_2	9.8284	4.1126	2.0084	4.7541	2.9123	9.1191
AVD	R_1	0,1207	0,1018	0,0895	0,0951	0,0152	0,0954
	R_2	0,3251	0,0124	0,0124	0,0541	0,0184	0,1542

Table 3. displays the spatial performance outcomes achieved by our approach. These results closely align with those provided by experts, validating the efficacy of the fully automated segmentation approach developed herein. Furthermore, our findings underscore the significance of

gradient information in reducing overlap between adjacent segmented regions. Notably, this technique exhibits rapid processing capabilities as it exclusively addresses areas of interest, obviating the need for tumor recognition training.

Table 3. Performance analysis of our method

Authors	Description	Dice (%)		Sensitivity(%)		Specificity(%)	
		Core	Enhanced	Core	Enhanced	Core	Enhanced
Proposed work	QMC + Constrained Kmeans + RG	0.82	80,25	0.78	0.70	0.99	1
Hachemi [23]	QMC + EM + RG	0.80	0.71	0.72	0.57	0.9	0.8
Vaidhya [24]	Multimodal image + Autoencoder	0.68	0.64	0.66	0.74	0.71	0.53
Pereira [25]	CNN	0.76	0.73	0.90	0.72	0.86	0.81
Ellwaa [27]	Random orest + Iterative training	0.72	0.73	0.73	0.75	0.99	1
Demirhan [26]	wavelets + ANN	0.77	-	0.73	-	0.95	-

Conclusion

In conclusion, this study has successfully introduced a novel fully automatic multi-segmentation technique for brain tumor detection. Our approach involved a comprehensive series of image processing steps to enhance the quality of MRI images and accurately identify regions of interest. Specifically, we applied enhancement and denoising filters to preprocess

the MRI images, followed by the application of Kapur thresholding to isolate the regions of interest. For multi-segmentation, we employed an innovative approach by initializing the region-growing method with a hybrid technique combining K-means clustering with a Naive Bayesian approach. To further refine our results, we maximized the a posteriori probability through an Iterated Conditional Mode approach, and improved the region-growing process by incorporating a Quasi Monte Carlo sampling method. The outcomes of our study have demonstrated impressive performance, suggesting that our approach has the potential to replace conventional techniques for brain tumor detection. The combination of image enhancement, advanced segmentation, and probability maximization contributes to the robustness and accuracy of our method, making it a promising advancement in the field of medical image analysis

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