

**RESEARCH ARTICLE** 

# Effective and Efficient LDA+ELM Model for Supervised Classification of Brain Tumor Types Using 2D MRI Scans

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### ABSTRACT

Application of machine learning in multiclass classification of brain tumor types has contributed to the development of computer aided diagnosis (CAD) system that can potentially enhance accuracy and speed up diagnosis of the disease. LDA+ELM model with different activation functions were investigated to achieve the optimum performances in terms of accuracy, Kappa statistic, sensitivity, precision, F-measure, training time and test time. We also proposed a user-friendly GUI in characterizing brain tumor types using MR images. First, a total of 3064 slices of CE T1-weighted brain MR images with ground truth were downloaded from a free online database. The manually segmented tumor region was augmented and then undergo several feature extraction techniques. All the feature descriptors obtained were then concatenated, followed by LDA dimensionality approach. Performance of different number of LDA features and ELM activation functions were investigated by repeated training and test. The ELM output of training data for each class was used to fit GMM and these probabilistic models used to estimate posterior probabilities of test data. LDA+ELM model with 5 LDA feature input, utilizing sigmoid function as hidden nodes activation functions achieves the best generalization performance with accuracy of 98.92% and corresponding F-scores for meningioma, glioma and pituitary tumor of 97.81%, 99.1% and 99.5% respectively. The proposed method (LDA+ELM) model performs better compared to other previous works using the same dataset and performing the same classification task.

Keywords: Multi-class classification of brain tumor, Linear Discriminant Analysis (LDA), Extreme Learning Machine (ELM), activation functions, Gaussian Mixture Model (GMM).

### 1 Introduction

Simply speaking, brain tumor is the collection of abnormal cells in central nervous system. Early detection of brain tumor is crucial for treatment planning and prognosis, thus enhancing the chance of survival of patients. Nonetheless, tumor characterization and evaluation is difficult due to the fact that neoplastic tissues often appear heterogeneous pixel-wise in biomedical images [1]. While biopsy (extraction of tissues) remains as gold standard in cancer diagnosis, it is also invasive and could not characterize brain tumor spatial and temporal heterogeneity. Analysis of medical images data can be excellent alternative as they are non-invasive, time efficient and contain rich histopathologic information. Nowadays, considerable amounts of medical image dataset are generated and collected during routine clinical practices [2], which open up the possibility of scientific knowledge discovery, while at the same time rendering manual analysis medical data implausible. of Current advancement of statistical modeling and computer vision enables semi-automated or even automated image processing in clinical practice, like image registration, tissue classification, volumetric analysis, and image segmentation. This gives rise to an emerging field of study, called radiomics, from which can be dated back to the work of [3]. Radiomics is high-throughput conversion of image data into mineable highdimensional features for the purpose of



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improved clinical diagnosis. Categorization of brain tumor types using medical images can be achieved through sequence of radiomics workflow: data acquisition, regions of interest (ROI) segmentation, feature extraction, feature reduction, followed by model construction and validation. This study can contribute to the development of computer aided diagnosis (CAD) system. One of the most important purposes of CAD is to provide objective and accurate diagnosis outcome that can complement radiologists in decision making. Additionally, it can shorten image analysis duration and solves the issue of variability of opinions among clinical experts. Ultimately, if proven accurate and consistent in real-time application, biopsy (extraction of tumor samples) can be bypassed or at least avoided if necessary in diagnosis procedure.

In view of the severity of brain tumor and lack of effective treatment, early diagnosis of the disease can be crucial for the patient recovery and treatment planning for the doctor. Advanced imaging protocols, like computed tomography (CT), positron emission tomography (PET), and MRI have provided unprecedented high resolution of medical images that make medical image analysis possible. In this paper, MRI will be the modality of choice because : 1) MRI is noninvasive, 2) produces multiple slices of images of the same tissue region with different contrast by applying different image acquisition protocols and parameters [4], 3) high contrast of soft tissues and high spatial resolution. In this paper, only contrast enhanced T1-weighted MR image will be considered as multi-spectral MR images analysis is expensive and increases odds of segmentation errors due to inconsistency and misalignment [5]. This paper aims to extract radiomic features from MRI scans to determine the correct type of tumor for MR image slices.

In recent years, there has been surge in research interest regarding supervised classification of brain tumor types using MRI scans. Early work performed by [1] investigated the use of machine learning schemes (LDA, kNN and nonlinear SVM) in binary classification of metastases and glioma as well as grades of glioma. In addition, the authors also performed multiclass classification by applying one versus all SVM model. The overall accuracy in distinguishing glioma grade II, III, IV and metastases is 63.3%. The same classification problem was explored by [6]. The highest leave-one-out cross-validation accuracy of 76.29% was achieved by Best First Search algorithm and VFI as classifier. [7] put forward a comprehensive radiomics workflow, in which classes involved are astrocytoma, glioblastoma multiforme, childhood tumormedulloblastoma, meningioma (MEN), secondary tumor-metastatic and normal regions. A total sample of 856 regions of interest and 218 features were extracted. The overall accuracy achieved is 85.23% using PCA-ANN approach. Classification problem of 6 classes, glioma grade I, II, III and IV, meningioma and metastases was performed by [8] using one against one SVM model.

Recent work of [9] utilized first order and second order statistical features from brain MR images to develop SVM model. Their experiment shows that highest accuracy of 85% was attained by SVM model trained by second-order statistical features. [10] proposed a new machine learning pipeline consisting of hybrid structure descriptor and fuzzy-based RBF SVM that categorized MR images into meningioma, metastases, glioma II and III. The overall accuracy of 96.74% was reported. Back-propagation artificial neural networks (BPANN) trained with statistical features from DWT transform and Gabor filter developed by [11] in classifying meningioma, glioma and pituitary tumor achieved overall accuracy of 91.9%.

Deep learning has now become the state-of-theart in many domains, especially in automatic image analysis, when a major breakthrough in image classification of ImageNet challenge in 2012 [12]. The reasons behind the success of deep neural networks include: 1) availability of sophisticated hardware with high computing power, 2) development of learning algorithm [13], 3) reinforcement learning, which can capture crucial semantic information pertaining class discrimination[14]. Even though deep learning model is well known for its remarkable recognition accuracy, it requires tuning of hyperparameters, sophisticated hardware and large amount of annotated data. Its performance in small data samples problem is still inconclusive.

To the best of our knowledge, most of the previous works regarding supervised classification of brain tumor types focus on development of novel machine learning pipelines and attaining high overall accuracy. In other word, only the discrete outputs of classifier (with test data as input) are being taken into account during evaluation phase of the proposed model. In addition, output scores (confidence) of classifier might not represent underlying real probability of certain class given the attributes of sample. In fact, modern neural networks are poorly calibrated, despite being accurate [15]. Accurate class probability estimates is crucial in real-world high stake decision making system, especially medical diagnosis [16], whereby there is difference in cost of misclassification and domain knowledge is required [17] because it provides likelihood or uncertainty of predictions. Since human have natural cognitive intuition for

probability [18], probabilistic outputs enhance reliability of the system. This study bridges the gap by applying Gaussian mixture model to transform the extreme machine learning (ELM) model outputs in reference to work of [19].

The goal of this paper is to address multi-class recognition of brain tumor types using 2D MRI scans by sequence of steps outlined in radiomics. Combination of linear discriminant analysis (LDA) feature reduction approach and ELM classifier, which we abbreviated as LDA+ELM is proposed. Apart from that, different activation functions, such as sigmoid, hard-limit, sine and radial basis function (RBF) are investigated. Last but not least, a user-friendly graphic interface (GUI) is put forward, with probability of respective brain tumor types (e.g. meningioma, glioma and pituitary tumor) as output.

#### 2 Research Methodology

The research framework proposed can be summarized in Fig. 1 and Fig. 2.





This workflow was performed under one to 80 LDA attributes. Additionally, for each training and test data partitioning, random stratified sampling was conducted to ensure similar class distribution for both training and test dataset. 70% of original data is allocated for training, while another 30% is partitioned for performance evaluation (test). The optimal machine learning pipeline was then determined by analyzing several performance indicators and the output of the trained ELM classifier was used to fit GMM model for each class. The resulting GMM models can be used estimate the class membership probability of test data. Lastly, a user-friendly GUI capable of performing brain tumor types classification was developed.

# 2.1 2D MRI scans acquisition

The brain MR images were downloaded from publicly available online database. A total of 3064 slices of T1-weighted contrast enhanced MR images from 233 patients was downloaded from https://figshare.com/articles/brain tumor data set/1512427. There are three kinds of brain tumor in the MR images downloaded, namely meningioma (708 slices), glioma (1426 slices), and pituitary tumor (930 slices). The brain T1weighed CE-MRI dataset was acquired from Nanfang Hospital, Guangzhou, China, and General Hospital, Tianjing Medical University, China, from year 2005 to 2010. The images have an in-plane resolution of 512×512 with pixel size  $0.49 \times 0.49 \ mm^2$ . The slice thickness is 6 mm and the slice gap is 1 mm [20, 21].

### 2.2 Tumor region augmentation

The 2D brain MR images comes with ground truth tumor region delineated manually by experienced radiologists. As pointed out in [22], tissues surrounding tumors can provide useful discriminative information about the types of tumor. Thus, augmentation of tumor region can be beneficial in extracting robust features. In this study, augmentation of tumor region was performed by morphological dilation with diskshape structuring element with radius, R of 8.

# 2.3 Feature extraction

Simply speaking, feature extraction is a dimensionality reduction method, in which instead of utilizing pixel intensity in raw images, a set of attributes, also known as feature vector is constructed to represent a certain image. This stage is crucial as it can directly impacts the generalization performance of classification models [23]. Ideal set of feature should be relevant, non-redundant, provide intrinsic dimensional representation for visualization, and increase training and inference speed of learning algorithms [24], 2003). In this paper, several feature extraction techniques were employed, including shape parameters, geometric moment invariants [25, 26], Zernike moments [27, 28], pseudo Zernike moments [29], histogram of oriented gradients (HOG) [30], linear binary pattern (LBP) [31], and bag of words (BOW) model [32]. It is worth noting that the image features will be extracted from the augmented tumor region. The number of features extracted from each method are summarized in Table 1. Therefore, the whole feature matrix dimension is  $3064 \times 1553.$ 

Table 1:	Feature	extraction	and	its	number o	of
		features.				

Feature extraction	Number of features
methods	
Binary shape parameters	11
Geometric moment	7
invariants	
Zernike moments	12
Pseudo Zernike	15
moments	
HOG	900
LBP	108
Bag of words model	500

### 2.4 LDA feature reduction

LDA is a linear dimensionality reduction method that can resolve the issue arise from high dimension data, such as high computational cost and complexity as well as undesirable modeling performance [33]. LDA is a supervised approach where class label of dataset is required. This is because, in the formulation of LDA, it tries to compute the "directions" that will maximize the separation of training samples between classes and at the same time minimizes the separation of test samples within class.

Let define some important notation,  $x_i$  be the row vector representing features of a sample,  $\overline{x_k}$ is the mean vector for k class,  $\mu$  is the overall mean vector of whole data,  $C_k$  denotes the indices of samples belong to k class and  $N_k$  is the number of class available. The steps involved for feature transformation using LDA are as follow:

- 1) Calculate the mean for each class,  $\overline{x_k}$  and overall mean,  $\mu$ .
- 2) Calculate the within class scatter matrix,  $S_w$  and between class scatter matrix,  $S_b$  using formula below:

$$S_{w} = \sum_{k=1}^{N_{k}} \sum_{i \in C_{k}} (\boldsymbol{x}_{i} - \overline{\boldsymbol{x}_{k}}) (\boldsymbol{x}_{i} \qquad (1)$$
$$- \overline{\boldsymbol{x}_{k}})^{T}$$
$$S_{b} = \sum_{k=1}^{N_{k}} n_{k} (\overline{\boldsymbol{x}_{k}} - \boldsymbol{\mu}) (\overline{\boldsymbol{x}_{k}} - \boldsymbol{\mu})^{T} \qquad (2)$$

 Maximize *w* such that between class scatter is maximized and within class scatter is minimized. Seeking optimum vector *w* is equivalent to solving generalized eigenvector system below:

$$\boldsymbol{S}_{\boldsymbol{b}}\boldsymbol{w} = \lambda \boldsymbol{S}_{\boldsymbol{w}}\boldsymbol{w} \tag{3}$$

4) After finding  $\boldsymbol{w}$  and  $\boldsymbol{\lambda}$ , rearrange the eigenvectors according to the descending order of  $\boldsymbol{\lambda}$ . The original data matrix is projected into new dimensional space,  $\boldsymbol{Z}$  by linear transformation:

$$Z = X \boldsymbol{w} \tag{4}$$

#### 2.5 ELM classifier

Recently, ELM has attracted a lot of interest among researchers due to its fast training convergence and similar prediction accuracy compared to state-of-the-art classification models like neural networks and support vector machine (SVM). The working principle of ELM is to train single layer feed forward neural networks (SLFN). Unlike conventional neural networks in which the hidden nodes parameters are iteratively tuned based on the training error, the hidden node parameters of ELM are generated randomly. It has been proven in the work of [34, 35] that without tuning hidden node parameters of SLFN with arbitrary number of hidden nodes, the system can still learn from training data. Since the hidden node parameters are randomly generated, they are independent of each other as well as of training data. The general architecture of ELM is depicted as in Fig. 3.

Consider training set containing n samples,  $(x_i, t_i)$ , where  $x_i \in \mathbb{R}^m, t_i \in \mathbb{R}^{N_k}$ , whereby  $t_1 = [1, -1, -1]^T$ ,  $t_2 = [-1, 1, -1]^T$ ,  $t_3 = [-1, -1, 1]^T$ . The output function can be expressed as below:

$$f_L(\boldsymbol{x}) = \sum_{i=1}^{L} \beta_i G(\boldsymbol{a}_i, \boldsymbol{b}_i, \boldsymbol{x})$$
 (5)

where  $\beta_i$  is the output weight for the output of hidden node activation function  $G(a_i, b_i, x)$ ,  $a_i$ and  $b_i$  are both parameters for hidden node i, Lis the number of hidden node used. Several hidden node output functions, such as sigmoid, hard limit, sine and radial basis function (rbf) activation functions in which their formula are displayed in Table 2 were experimented. It should be noted that 300 hidden nodes are chosen in this study. The above equation can also be written in matrix form:

$$H\beta = T_L$$

where,

$$H = \begin{pmatrix} G(\boldsymbol{a}_{1}, \boldsymbol{b}_{1}, \boldsymbol{x}_{1}) & \cdots & G(\boldsymbol{a}_{L}, \boldsymbol{b}_{L}, \boldsymbol{x}_{1}) \\ \vdots & \ddots & \vdots \\ G(\boldsymbol{a}_{1}, \boldsymbol{b}_{1}, \boldsymbol{x}_{n}) & \cdots & G(\boldsymbol{a}_{L}, \boldsymbol{b}_{L}, \boldsymbol{x}_{n}) \end{pmatrix}_{n \times L}$$

$$\boldsymbol{\beta}_{1}$$

$$\boldsymbol{\beta}_{2} \quad \begin{pmatrix} \beta_{1} \\ \vdots \end{pmatrix}_{L \times N_{k}}$$

$$\boldsymbol{\beta}_{L} \\ \boldsymbol{T}_{1} \\ \boldsymbol{T}_{L} = (\vdots)_{n \times N_{k}}$$

$$T_{n}$$

(6)

The objective of ELM is to minimize  $||H\beta - T||$ and  $||\beta||$ . Minimization of  $||H\beta - T||$  is equivalent to minimization of training error while minimization of  $||\beta||$  is equivalent to minimization of norm of weight of connection between hidden nodes and output layers,  $\beta$ . According to Bartlett's theory [36], for feedforward neural networks reaching smaller training error, the smaller the norm of weights is, the better generalization performance the networks tends to have. In the paper of [37], minimizing  $||\boldsymbol{\beta}||$  can be regarded as maximizing the distance of the separating margins of the two different classes in the ELM feature space. As such in order to optimize  $\boldsymbol{\beta}$ , minimal norm least square formula can be employed:

$$\boldsymbol{\beta} = \boldsymbol{H}^{\dagger} \boldsymbol{T} \tag{7}$$

where  $H^{\dagger}$  denotes generalized Moore-Penrose inverse matrix. The above equation can also be expressed as:

$$\boldsymbol{\beta} = (\boldsymbol{H}^T \boldsymbol{H})^{-1} \boldsymbol{H}^T \boldsymbol{T}$$
(8)

Nonetheless, the above formulation can become computationally implausible if inverse of  $H^T H$  do not exists and computationally unstable if the condition number of square matrix  $H^T H$  is large. To avoid these issues, diagonal elements of  $H^T H$  can be added with positive value according to ridge regression theory [38]. In our case where number of training instances are large,  $\beta$  can be solved via [39]:

$$\boldsymbol{\beta} = (\frac{l}{c} + H^T H)^{-1} H^T T$$
(9)

where c is a positive number. The author recommended the use of c in the range of  $\{2^{-24}, 2^{-23}, \dots, 2^{25}\}$ . 5-fold cross validation methods are employed to find the best c.



Figure 3: General architecture of ELM [40]

Table 2: Formula of activation functions

Activation	Formula
functions	
Sigmoid	$G(\boldsymbol{a}, \boldsymbol{b}, \boldsymbol{x}_i)$
	1
	$=\frac{1}{1+\exp\left(-(ax_i+b)\right)}$
Hard-	$G(\boldsymbol{a},\boldsymbol{b},\boldsymbol{x}_i) = \int 1, \boldsymbol{a}\boldsymbol{x}_i + \boldsymbol{b} \ge 0$
limit	$O(\mathbf{u}, b, \mathbf{x}_i) = \{0, otherwise\}$
Sine	$G(\boldsymbol{a}, \boldsymbol{b}, \boldsymbol{x}_i) = \sin\left(\boldsymbol{a}\boldsymbol{x}_i + \boldsymbol{b}\right)$
rbf	$G(\boldsymbol{a}, \boldsymbol{b}, \boldsymbol{x}_i) = \exp\left(-\boldsymbol{b} \ \boldsymbol{x}_i - \boldsymbol{a}\ ^2\right)$

#### 2.6 GMM

Simply speaking, GMM is a type of unsupervised soft clustering method that assign the likelihood of cluster membership for each data instance, with assumption that data are generated by finite number of Gaussian distributions. Being a parametric probability density function, the probability of certain data point is represented as:

$$p(\boldsymbol{x}) = \sum_{k=1}^{K} \pi_k \mathcal{N}(\boldsymbol{x}|\boldsymbol{\mu}_k, \boldsymbol{\Sigma}_k)$$
(10)

where  $\pi_k$  is mixing coefficient,  $\mathcal{N}$  represents multivariate probability density function,  $\mu_k$  is mean vector,  $\sum_k$  is covariance matrix, and K is the number of clusters. Parameters involving  $\pi_k$ ,  $\mu_k$  and  $\sum_k$  are optimized by maximizing log likelihood function as expressed below:

$$\ln (p(\mathbf{x})) = \sum_{i=1}^{n} \ln \left( \sum_{k=1}^{K} \pi_i \mathcal{N}(\mathbf{x}_n | \mu_k, \Sigma_k) \right)$$
(11)

Since there is no elegant closed form solution to equation (11), it can be solved using iterative algorithm, called expectation-maximization (EM) algorithm. To produce reliable probability estimates, separate GMM is used to fit the output vectors (from training data) of ELM that are correct for each class. Each GMM can model conditional probability of ELM output given respective class, p(t|C). By applying Bayes' theorem, we can estimate the class conditional probability given a certain ELM output from test data:

$$p(C_{i}|t) = \frac{p(t|C_{i})p(C_{i})}{p(t)} = \frac{p(t|C_{i})p(C_{i})}{\sum_{i=1}^{N_{k}} p(t|C_{i})p(C_{i})}$$
(11)

The only arbitrary parameter is the number of clusters, K. We employ Akaike Information Criterion (AIC) to select the best K. We utilize the idea in the work of [19] by fitting GMM with only the correctly identified samples during training phase and calculate the class conditional probability based on the above formula during test phase.

# 2.7 Performance evaluation

In order to evaluate the performance of LDA+ELM classification model under different number of features, 7 commonly accepted

performance measures were utilized, including test accuracy (%), Kappa statistic, sensitivity (%), precision (%), F-measure (%), training time (s) and test time (s). Mean and standard deviation of each performance metrics were computed for unbiased analysis and comparison among methods employed. This process is necessary as robust performance evaluation can provide clear picture of the strengths and weaknesses of the machine learning paradigms being analyzed. Table 3 shows some notations in confusion matrix. Table 4 describes the performance measures and its definitions.

Table 3: Notations	: in	confusion	matrix.
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		Predicted classes		
		A (meningioma)	B (glioma)	C (pituitary
				tumor)
True classes	A (meningioma)	$TP_A$	$E_{AB}$	$E_{AC}$
	B (glioma)	$E_{BA}$	$TP_B$	$E_{BC}$
	C (pituitary tumor)	E <sub>CA</sub>	E <sub>CB</sub>	$TP_{C}$

Performance measures	Definition	Formula
Accuracy	Overall efficiency and generalizability of classifier [41]. However, its use in performance comparison among classifiers is limited [42].	$Acc = \frac{TP}{n_T},$ $n_T = number of test samples$
Kappa statistic	Measure the degree of agreement between the predicted labels and the ground truth [43].	$Kappa = \frac{Acc - P(E)}{1 - P(E)},$ P(E)=expected agreement between classifier and ground truth by chance
Sensitivity	Compute the proportion of samples of class 'A' label that are correctly predicted as class 'A' in the test phase.	For class A, Sensitivity of A $= \frac{TP_A}{TP_A + E_{AB} + E_{AC}}$
Precision	Indicate the proportion of test samples that are predicted to be class 'A' that match the known true class label.	For class A, $Precision of A$ $= \frac{TP_A}{TP_A + E_{BA} + E_{CA}}$
F-measure/F-score	Harmonic mean of precision and sensitivity.	For class A, $F - score \ of \ A$ $= \frac{2TP_A}{2TP_A + E_{AB} + E_{AC} + E_{BA} + E_{CA}}$
Training time	Quantify the convergence speed of training.	-
Test time	The rate at which the output labels of test samples are generated.	-

**Table 4:** Performance measures and its respective formula.

### 3 Results

In this paper, we investigated different activation functions of ELM under different number of LDA features. Fig. 4-7 shows the performance measures (e.g. accuracy, Cohen' Kappa, training time and test time) of sigmoid, hard-limit, sine and rbf hidden node activation functions

100 95 90 average accuracy (%) 85 80 75 70 65 60 55 10 20 30 40 50 60 70 80 number of feat ires 0.9 0.8 a statistics 9.0 Kappa 0.5 0.4 0.3 0.2 10 20 30 40 50 60 70 0 80 number of features 0.05 0.045 0.04 0.035 0.035 (s) 0.03 0.025 0.02 training time (s) test time (s) 0.015 0.0 0.005 0 30 50 0 10 20 40 60 70 80 numbe of features

Figure 4: Performance measures versus number of features for sigmoid activation function.

respectively. Empirical observation of these figures shows that the performances of sigmoid and hard-limit transfer function are relatively more stable and insensitive to changes in number of features (more resistant to overfitting issue). Accuracies and Kappa statistics for both sine and rbf decline after number of features increase beyond a threshold.



Figure 5: Performance measures versus number of features for hard-limit activation function.



Figure 6: Performance measures versus number of features for sine activation function.



Figure 7: Performance measures versus number of features for rbf activation function.

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Table 6 shows the performance measures of activation function under optimal number of features. The best generalization performance is achieved by sigmoid function, with accuracy of 98.92%, using just 5 LDA attributes, Interestingly, very high standard deviation in performance measures of sine activation function indicates that its performance is very sensitive to sampling of data. Table 7 shows performance

metrics of sigmoid function before and after applying GMM on ELM outputs. There is no significant difference in performance measures before and after applying GMM on ELM outputs, suggesting that GMM can approximate the probability of classes well.

Fig. 8 shows the GUI developed and the function of each buttons clearly explained.

Activation functions		sigmoid	hardlim	sine	rbf
number of feature	s	5	23	8	9
Accuracy (%)		98.923±0.378	98.680±0.361	87.715±11.912	98.306±0.518
Kappa statistics		0.983±0.006	0.979±0.006	0.791±0.218	97.337±0.819
	А	97.536±0.971	96.186±1.447	64.232±27.192	95.088±2.684
Sensitivity (%)	В	99.509±0.360	99.509±0.360	99.953±0.113	99.392±0.595
	С	99.081±0.513	99.308±0.351	86.838±22.503	99.093±0.846
	А	98.081±0.905	98.861±0.687	99.280±1.618	98.060±1.488
Precision (%)	В	98.694±0.503	98.045±0.632	81.970±13.132	97.792±1.199
	С	99.928±0.197	99.547±0.373	99.710±0.490	99.366±0.944
F-measure (%)	А	97.805±0.740	97.498±0.763	73.947±24.859	96.514±1.085
	В	99.099±0.370	98.770±0.364	89.466±8.667	98.579±0.558
	С	99.502±0.263	99.426±0.246	90.496±20.066	99.223±0.481
training time (s)		0.029±0.003	0.031±0.003	0.031±0.002	0.067±0.004
test time (s)		0.005±0.000	0.005±0.000	0.006±0.001	0.006±0.000

**Table 6:** Performance measures of activation function under optimal number of features.

A-meningioma, B-glioma, C-pituitary tumor

**Table 7:** Performance metrics (mean  $\pm$  standard deviation) of sigmoid function before and after applying GMM<br/>on ELM outputs.

Performance measures		Using classifier scores	After using GMM (fitted with all training samples)	After using GMM (fitted with correctly identified training samples)
Accuracy (%	)	98.923±0.378	98.669±0.431	98.799±0.439
Kappa statis	tics	0.983±0.006	0.979±0.007	0.981±0.007
S	А	97.536±0.971	97.552±1.088	97.474±1.074
(%)	В	99.509±0.360	98.893±0.563	99.353±0.548
(70)	С	99.081±0.513	99.176±0.525	98.961±0.519
D · ·	А	98.081±0.905	97.010±1.452	97.640±1.256
(%)	В	98.694±0.503	99.002±0.455	98.693±0.542
(70) C	С	99.928±0.197	99.453±0.464	99.868±0.221
F-measure A (%) C	А	97.805±0.740	97.272±0.906	97.551±0.894
	В	99.099±0.370	98.946±0.392	99.021±0.424
	С	99.502±0.263	99.312±0.268	99.412±0.289
training time	(s)	0.029±0.003	0.278±0.035	0.556±1.349
test time (s)		0.005±0.000	0.007±0.001	0.007±0.002

A-meningioma, B-glioma, C-pituitary tumor

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This push button displays the 2D brain MR image and its file pathname.

- 2) This push button displays display ground truth tumor region.
  - This push button perform feature extraction mentioned in section 2.3.
  - This push button perform LDA linear dimensionality reduction. The dimension of resulting attribute vector is 5.
  - This push button feed the trained ELM model (300 hidden neurons with sigmoid activation function) with the attribute vector and output probability estimates of each brain tumor types.

承 UI Figure	- 🗆 ×
Load brain tumor image     File pathna	me C:\Users\User\Dr
2D brain MR image	
	3 Extract features Status Done
	4 LDA feature reduction Status Done
2 visualize ground truth (tumor)	5 Classify
Ground truth tumor region	Displays probability of respective tumor types in (%)
	meningioma 97.3893 %
	glioma 2.6107 %
	pituitary tumor 3.0532e-10 %

Figure 8: Details of GUI developed.

#### 4 Discussion

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Our experiment shows that sigmoid function achieves best performance compared to other non-linear activation functions, which agrees with the previous work of [44]. This probably can be attributed to meaning feature mappings that is helpful for brain tumor types discrimination.

It is also noted that sine and rbf activation function can become unstable if the number of features increases beyond a certain threshold. Below are likely justifications for the poor performances of these activation functions:

- According to [45], RBF/Gaussian activation function can only achieve good results if the data approximately follows Gaussian distribution. Additionally, the values of hidden nodes (a and b) can have significant impact of the elements of H (matrix of hidden nodes' outputs). If vector a is too far from x, or when b is too large, the zero percentage of H can be high.
- Activation functions typically used in neural networks is monotonic. As pointed out in [46], neurons learn to respond to particular regularity, in which

stronger positive correlation with the input corresponds to stronger (or equal) activation and vice versa. Sine function, being a sinusoidal (non-monotonic) function, activations of hidden nodes may oscillate between stronger and weaker activations. This condition may not be desirable if for example, two similar instances (close to each other in Euclidean space) that belong to same class may have very different projection in H. This helps to explain the large variation seen in performance measures when different set of data is used for training and test.

The GUI developed do not require any user input parameters, but the images loaded must be attached with segmented tumor region. The probability estimates of each class is computed based on Bayes' theorem stated in section 2.6, using the  $p(t|C_i)$  calculated from fitted GMM during training phase. Table 8 shows comparison with other findings in literature using same dataset and performing the same classification tasks. Effective and Efficient LDA+ELM Model for Supervised Classification of Brain Tumor Types Using 2D MRI Scans

Authors	Classifiers	Feature input	Accuracy
[47]	Deep neural	Downsampled	95.6%
	network	window from	
		imagos	
[49]	Convolutional	Original	04 20%
[40]	Convolutional	Unginal	94.270
	neurai	innages	
[40]	General	Derrorated	0 C E C 0 /
[49]	Capsule	Downsampled	80.30%
	networks	$64 \times 64$ raw	
		image patches.	
[50]	Capsule	Downsampled	87.5%
	netwoks	$64 \times 64$ raw	
	(DCNet ++)	image patches.	
[11]	ANN	Statistical	91.9%
		features from	
		DWT and	
		Gabor filter	
[51]	Kernel ELM	Representation	93.68%
		learning using	
		CNN with	
		$28 \times 28$	
		image patches	
		as input.	
[20]	SVM	BoW model	91.28%
		with region	
		partitioning	
Our	ELM	5 LDA	98.92%
work		features	
		extracted	
		augmented	
		tumor region	

**Table 8:** Previous works using same dataset and performing the same classification tasks.

### 5 Conclusions

Multi-class brain tumor categorization was performed by LDA+ELM model, alongside multiple feature descriptors. Several activation functions were experimented and it was found that sigmoid activation function achieves the highest generalization performance with accuracy of 98.92% by using only 5 LDA attributes. The performance of our proposed model is comparable or better than previous works on the same data, performing the same classification task. By fitting GMM model on training data, posterior probability estimates for test data can be obtained. Eventually, a user-friendly GUI is developed based on the optimal pipeline. Further study should be oriented towards development of statistical models (supervised learning models) that can perform online incremental learning, as medical image data normally come one by one or chunk by chunk.

# 6 Declarations

## 6.1 Study Limitations

This study just focuses on contrast enhanced T1weighted MR images, thus the results may not be extended to other imaging modalities. Another limitation of this research is that no external validation dataset to evaluate performance of proposed model on unknown data from other source (e.g. different MRI machine and its parameters as well different patients).

# 6.2 Acknowledgements

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# 6.3 Competing Interests

Not applicable.

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