**Introduction**: Cancer of the nervous system is one of the most common types of cancer in the world and mostly due to presence of a tumour in the brain. The symptoms and severity of the brain tumour depend on its location. The tumour within the brain may develop from nerves, dura (meningioma), pituitary gland (pituitary adenoma), or from the brain tissue itself (glioma).

Material and methods: In this study we proposed a feature engineering approach for classification magnetic resonance imaging (MRI) of 3 kinds of most common brain tumour, i.e. glioma, meningioma, pituitary, and no-tumour. Here 5 machine learning classifiers were used, i.e. support vector machine, K-nearest neighbour (KNN), Naive Bayes, Decision Tree, and Ensemble classifier with their paradigms. Results: The handcrafted features such as histogram of oriented gradients, local binary pattern features, and grey level co-occurrence matrix are extracted from the MRI, and the feature fusion technique is adopted to enhance the dimension of feature vector. The Fine KNN outperforms among the classifiers for recognition of 4 kinds of MRI: glioma, meningioma, pituitary, and no tumour, and achieved 91.1% accuracy and 0.95 area under the curve (AUC).

**Conclusions**: The proposed method, i.e. Fine KNN, achieved 91.1% accuracy and 0.96 AUC. Furthermore, this model has the possibility to integrate in low-end devices unlike deep learning, which required a complex system.

**Key words**: brain tumour, machine learning, feature extraction, feature fusion, classification.

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# Brain tumor magnetic resonance images classification based machine learning paradigms

## Baby Barnali Pattanaik<sup>1</sup>, Komma Anitha<sup>2</sup>, Shanti Rathore<sup>3</sup>, Preesat Biswas<sup>4</sup>, Prabira Kumar Sethy<sup>5</sup>, Santi Kumari Behera<sup>6</sup>

<sup>1</sup>Department of Electronics, Sambalpur University, Burla, India <sup>2</sup>ECE Department, PVP Siddhartha Institute of Technology, Vijayawada, India <sup>3</sup>Department of ET and T, Dr. C. V. Raman University Bilaspur Chhattisgarh, India <sup>4</sup>Department of ET and T, GEC Jagdalpur, Chhattisgarh,India <sup>5</sup>Department of Electronics, Sambalpur University, Burla, India <sup>6</sup>Department of Computer Science and Engineering, VSSUT Burla, India

## Introduction

A brain tumour is a mass of abnormally growing cells in the brain or skull. Some brain tumours are harmless, while others are dangerous. Tumours can start in the brain itself (primary) or the cancer can spread from another part of the body to the brain (metastasis). Treatment options depend on the type of tumour, its size, and its location. The World Health Organization made a classification and grading system to make it easier to converse about brain tumours, plan treatments, and predict how they will turn out. Figure 1 shows how tumours are categorised by the type of cells they are made of or where they started.

This research aims to develop automated approaches that will assist medical professionals in making diagnoses, to reduce the likelihood of incorrect diagnoses and give higher priority to complex patient diagnoses. In particular, the goal of this research is to automate the process of classifying different types of brain tumours based on images of patients' brains. When analysing brain scans, a radiologist needs to look at several image slices



Fig. 1. Anatomical view of brain with location of tumour

to properly diagnose any health problems. This process is time consuming. We want to be able to confidently identify the many types of brain cancer to increase the efficiency with which care is provided, while leaving the most difficult diagnoses to medical specialists.

It is difficult to automatically detect different forms of brain tumours. Previous studies have resulted in the development of specialised algorithms for the automated classification of brain tumours. Paul et al. suggested a deep learning system for the purpose of classifying brain images containing several types of tumours, including meningioma, glioma, and pituitary [1]. In the present study, 989 T1-weighted axial pictures were taken into consideration, and 91.43% accuracy was attained with 5-fold cross validation. A convolutional neural network (CNN) was developed by Abiwinanda et al. to automatically classify the 3 most frequent forms of brain tumours, namely gliomas, meningiomas, and pituitary tumours, without the need for any region-based pre-processing processes [2]. A total of 3064 T1-weighted contrast-enhanced magnetic resonance images (MRI) that are available to the public on Figshare [3] were used in this study. The results of the experiment showed that the accuracy of the training was 98.51%, while the accuracy of the validation was 84.19%. Kader et al. presented a differential deep CNN model (differential deep CNN) to categorise different kinds of brain tumours, including abnormal and normal MRI pictures [4]. This model is based on deep convolutional neural networks. To obtain more differential feature maps, differential operators are utilised in the extraction process.

The differential deep CNN model has a number of benefits, one of which is the ability to analyse the pixel directional pattern of images by applying contrast calculations. Another benefit is the model's high capacity to categorise a large image database. The dataset that was utilised in this study consisted of 25.000 MRI scans of the brain. This dataset included both aberrant and normal images, and it resulted in an accuracy of 99.25%. Ali et al. introduced a technique called extreme learning machine local receptive fields for classifying tumorous and non-tumorous MRIs of the brain [5]. In this location, coronal views of 9 patients were acquired for training purposes, while 7 patients were imaged for testing purposes. The information regarding the dimensions of the dataset is lacking. The author asserted that their work was accurate 97.18% of the time. A fully automatic model for the segmentation and classification of brain tumours has been presented by Díaz-Pernas et al. [6], and it makes use of a deep CNN that takes a multiscale approach. Meningiomas, gliomas, and pituitary tumours are the 3 forms of tumours that are categorised in this section. The dataset contains 3064 images with sagittal, coronal, and axial perspectives, and the accuracy rate was calculated to be 97.3%. For the purpose of 3-way (meningioma, glioma, and pituitary) classification of brain MRI, Saleh et al. analysed and compared the performance of 5 pre-trained models: Xception, ResNet50, InceptionV3, VGG16, and MobileNet [7]. The dataset contained 4480 pictures that have been separated into the following 3 categories: training, validation, and testing (unseen images). The training group contained a total of 2880 pictures, including 520 representations of each form of brain tumour. The validation group comprised a total of 800 photos, with each form of brain tumour accounting for 200 of the total number of images. The testing group had access to a total of 800 photos, with each form of brain tumour comprising 200 photographs. The F1-scores obtained as a consequence indicate that Xception had a score of 98.75%, resnet50 had a score of 98.50%, inceptionv3 had a score of 98.00%, vgg16 had a score of 97.50%, and MobileNet had a score of 97.25%. Glioma, pituitary, and meningioma are the 3 categories of brain pictures that may be classified using an approach proposed by Deepak et al. [8]. This approach is a combination of deep features and GoogLeNet. In this case, 3096 photos were employed, and with 5-fold cross validation, an accuracy rate of 98% is reached on average. The vgg19 was improved by Swati et al. using a block-wise fine-tuning method [9]. Under 5-fold cross-validation, the modified model was tested using T1-weighted pictures that are publicly available on the Figshare repository, and it attained an average accuracy of 94.82%. In both traditional machine learning [10–13] and the more recently utilised deep learning models [14–18], a large number of studies have been reported for classification and segmentation. Again, for brain tumour classification a deep learning model also achieved satisfactory results [19–22]. Furthermore, deep learning works only with large amounts of data. Training it with large and complex data models can be expensive. It also needs extensive hardware to perform complex mathematical calculations; hence, we chose machine learning rather than deep learning approaches.

The main objective of this study is to propose a medical diagnostic support system for brain tumour classification, i.e. to set up an automated system that can accurately classify the types of brain tumour from MRI using machine learning, and to show that with feature engineering, we can find comparable accuracy to deep learning.

The primary benefit of utilizing machine learning models is that they allow for a better interpretability of the classification model because they are based on feature engineering. Feature extraction is a particular kind of dimensionality modification. The chief purpose of this technique is to capture the important characteristics of the raw data and interpret this character in a less dimensionality space [23]. In this work, we used 3 methods for feature extraction, which involved local binary pattern, the histogram of oriented gradient, and grey level co-occurrence matrix. Local binary patterns (LBP) are an effective descriptor in tasks of recognition and computer vision [24]. The local binary patterns coding the grey levels of an image by comparing the central pixel with its neighbours and the result counted as a binary number converted to decimal number substitutes the central pixel value. A histogram of oriented gradients (HOG) is an introduced object descriptor, focalizing on the structure or appearance of an object in an image. The histogram of oriented gradients provides distinguishing features when lighting variation and background noise, so it is an effective descriptor [24]. The grey level co-occurrence matrix (GLCM) is one of several popular texture examination techniques that calculate the occurrence

of specific grey levels about other grey levels. The grey level co-occurrence matrix measures the frequency of various sequences of grey level values that appear in a region of interest [25]. This technique examines the association among adjacent pixels; the original pixel is identified as a reference, and the other is a neighbour pixel. Furthermore, we have taken the most prominent discriminatory features of each extraction technique. Deep learning models, on the other hand, are black box networks whose workings are extremely difficult to understand due to the complex design of the network. As a matter of fact, feature engineering is an essential part of the medical and diagnostic field for doctors because it gives them the ability to know the importance and impact of each feature on the classification and identification of cancer types, in contrast to deep learning models, which are black box networks. The main aim of this research was to find the optimal performance in brain tumour classification using multiclass data with numerous training models by means of machine learning with its paradigms. Again, comparative analysis can be done within the machine learning models with their paradigms for brain tumour classification.

The major contributions of this work are as follows:

- for the first time, the concept of feature engineering is applied to the 4-class brain tumour classification problem;
- three set of features such as GLCM, LBP, and HOG are extracted from brain MRIs, and then these features are used in various classifiers, namely support vector ma-

chine, K-nearest neighbour (KNN), naïve Bayes, tree, and ensemble classifier;

- all the classification methods are evaluated in single set feature and combined set of features;
- the combined set of features, i.e. GLCM + LBP + HOG, contributed 91.1% accuracy and 0.95 area under the curve (AUC) in fine KNN;
- the proposed method gives very good performance even with a small dataset, and it is comparable to the deep learning approach.

Structure of the article: The second section discusses the resources and approaches that were utilised. The results of the experiment are presented in Section 3, along with a commentary of them. The conclusion can be found in section 4, which comes last.

## Material and methods

This section details the dataset and the adapted methodology.

## About the dataset

The brain dataset investigated in this study is collected from the Figshare repository [3]. The dataset comprises TI-weighted MRI of no tumour and 3 different types of tumours: meningioma, glioma, and pituitary. Image resolution of 512  $\times$  512 with different views such as axial (transverse plane), coronal (frontal plane), or sagittal (lateral plane) planes was used in this dataset. The sample distribution based on the number of classes consisted of 826, 822, 827, and 395 sample instances of glioma,



Fig. 2. Samples of brain magnetic resonance imaging. 1<sup>st</sup> line: axial, 2<sup>nd</sup> line: coronal, 3<sup>rd</sup> line: ssagittal, glioma (A), meningioma (B), pituitary (C), no tumour (D)



**Fig. 3.** Overall work flow of methodology for classification brain tumour magnetic resonance imaging GLCM – grey level co-occurrence matrix, HOG – histogram of oriented gradients, KNN – K-nearest neighbour, LBP – local binary patterns, MRI – magnetic resonance imaging, SVM – support vector machine

meningioma, pituitary, and no tumours, respectively. The sample of 3 types of brain tumour is shown in Figure 2.

## Methodology

Figure 3 provides an overview of the methodology used to classify brain tumours. A total of 2870 T1-weighted MRIs were utilised throughout all phases of this study. The features of the images were extracted before the machine learning system classified the images according to their class. 20% of the dataset was utilised as test data, and 80% was used for training data (randomly chosen). The machine learning algorithm was trained using the training set's visual features. Finally, image attributes of the testing set were utilised to evaluate the model's performance. The models were run on an HP pavilion i5, Windows 10, 8 GB RAM, MATLAB 2021a platform.

Features were measurable quantities that could be useful for a predictive analysis for classification. The features contained in MRI are vital for disease diagnosis, and efficient feature extraction is crucial for improving diagnostic accuracy and cancer classification.

The extracted image characteristics were fed into machine learning algorithms. Support vector machine, KNN, tree, naive Bayes, and ensemble classifier were the machine learning techniques employed. These machine learning algorithms were trained with the training set's visual characteristics.

In this paper we extracted 13 GLCM, 36 HOG, and 59 LBP features. Here, to enhance the performance of classification models, the feature fusion technique was introduced. The combination of GLCM + LBP, GLCM + HOG, HOG + LBP and GLCM + HOG + LBP were fed into the classifiers and the performance was recorded.

## Result

This section presents the results obtained from the 5 classifiers under investigation in this paper. Indeed, the performance of the models was determined using the test data. To construct the most effective feature extraction strategies, we compared the performance of models with respect to single set and combined set features in this study.

The performance of 5 classifiers with their paradigms are recorded in Table 1 in terms of accuracy and AUC with respect to a single feature set. Here the considered feature sets are GLCM, HOG, and LBP. The observation made from Table 1 is that the highest accuracy achieved using single feature set, i.e. HOG is 85% in subspace KNN. The AUC of subspace KNN using the HOG feature is 0.97. Again, the performance contributed by subspace KNN using GLCM feature is 60.8% accuracy and 0.81 AUC and LBP 80.5% accuracy and 0.94 AUC. Furthermore, the accuracy and AUC achieved by classifiers other than subspace KNN are approximately 80% and 0.85, respectively.

#### Discussion

In the second phase the features extracted from MRI are combined, such as GLCM + HOG + LBP. The performance of different combinations of feature sets are recorded in Table 2. It was observed from Table 2 that the fine KNN performed well for the classification of 4 types of brain MRI. In fine KNN the accuracy achieved was 88.7%, 80.3%, 88.9%, and 91.1% by the combination of feature sets like GLCM + HOG, GLCM + LBP, HOG + LBP, and GLCM + HOG + LBP, respectively. Furthermore, the AUC achieved by fine KNN using GLCM + HOG, GLCM + LBP, HOG + LBP, and GLCM + HOG + LBP is 0.94, 0.85, 0.94, and 0.96, respectively. Overall, the highest performance recorded in

Classifiers			Accuracy		AUC			
		GLCM (%)	HOG (%)	LBP (%)	GLCM	HOG	LBP	
SVM	Linear	71.8	51.6	71.4	0.88	0.70	0.87	
	Quadratic	76.0	69.5	78.9	0.91	0.85	0.92	
	Cubic	79.3	76.3	76.3	0.93	0.89	0.92	
	Fine gaussian	77.7	73.2	64.1	0.90	0.94	0.87	
	Medium gaussian	72.8	71.6	76.5	0.89	0.87	0.91	
	Coarse gaussian	64.5	51.9	64.1	0.86	0.68	0.84	
KNN	Fine KNN	76.3	82.2	76.3	0.77	0.87	0.83	
	Medium KNN	70.7	71.3	71.8	0.87	0.84	0.90	
	Coarse KNN	63.1	55.9	63.1	0.84	0.75	0.86	
	Cosine KNN	72.6	70.7	67.4	0.88	0.86	0.88	
	Cubic KNN	71.1	70.6	71.1	0.87	0.83	0.90	
	Weighted KNN	77.9	77.4	73.9	0.89	0.90	0.92	
	Naive bayes	62.4	49.5	51.6	0.83	0.66	0.73	
Tree	Fine tree	67.9	53.3	67.1	0.83	0.75	0.83	
	Medium tree	64.6	47.9	63.1	0.83	0.69	0.80	
	Coarse tree	63.8	40.1	61.3	0.81	0.54	0.76	
Ensemble	Boosted trees	66.7	52.8	68.6	0.85	0.74	0.86	
	Bagged trees	76.0	69.0	73.7	0.87	0.86	0.89	
	Subspace discriminant	65.2	51.7	65.9	0.84	0.67	0.85	
	Subspace KNN	60.8	85.0	80.5	0.81	0.97	0.94	
	Rus boosted trees	65.2	53.1	67.8	0.86	0.72	0.87	

 Table 1. Brain tumour classification using magnetic resonance imaging based on a single feature set

AUC – area under the curve, GLCM – grey level co-occurrence matrix, HOG – histogram of oriented gradients, KNN – K-nearest neighbour, LBP – local binary patterns, SVM – support vector machine

Table 2. Brain tumour classification using magn	netic resonance imaging base	ed on a combination of feature sets
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Classifiers		Accuracy			AUC				
		GLCM + HOG (%)	GLCM + LBP (%)	HOG + LBP (%)	GLCM + HOG + LBP (%)	GLCM + HOG	GLCM + LBP	HOG + LBP	GLCM + HOG +LBP
SVM	Linear	72.8	77.4	73.3	79.1	0.91	0.92	0.92	0.93
	Quadratic	80.7	82.8	82.6	87.5	0.96	0.94	0.96	0.97
	Cubic	85.0	81.0	85.4	87.8	0.97	0.94	0.97	0.97
	Fine gaussian	77.7	64.1	69.3	73.0	0.96	0.93	0.96	0.96
	Medium gaussian	83.4	81.2	80.8	85.2	0.96	0.94	0.96	0.96
	Coarse gaussian	69.7	68.5	68.3	71.4	0.90	0.88	0.90	0.89
KNN	Fine KNN	88.7	80.3	88.9	91.1	0.94	0.85	0.94	0.96
	Medium KNN	76.5	80.0	79.8	81.4	0.95	0.94	0.95	0.96
	Coarse KNN	67.6	68.5	66.2	67.6	0.91	0.88	0.89	0.88
	Cosine KNN	76.0	75.8	75.4	79.4	0.94	0.92	0.93	0.94
	Cubic KNN	76.3	78.6	77.5	80.3	0.95	0.94	0.95	0.95
	Weighted KNN	82.8	82.2	83.4	86.4	0.97	0.95	0.97	0.97
	Naive Bayes	66.7	56.1	57.5	56.8	0.86	0.77	0.78	0.77
Tree	Fine tree	65.9	69.0	67.4	74.9	0.82	0.83	0.81	0.83
	Medium tree	62.2	64.6	63.4	67.9	0.84	0.82	0.82	0.82
	Coarse tree	60.1	65.7	62.2	64.6	0.79	0.81	0.77	0.79
Ensemble	Boosted trees	68.8	72.3	70.2	72.8	0.87	0.86	0.90	0.88
	Bagged trees	79.3	75.6	75.6	81.9	0.92	0.91	0.90	0.94
	Subspace Discriminant	69.5	73.7	71.1	76.0	0.89	0.90	0.90	0.90
	Subspace KNN	63.4	62.0	85.5	65.5	0.84	0.82	0.98	0.84
	Rus boosted trees	66.9	70.7	71.3	69.0	0.86	0.87	0.90	0.86



**Fig. 4.** Performance of fine K-nearest neighbour for brain tumour classification (**A**) confusion matrix (**B**) area under the curve AUC – area under the curve, GLCM – grey level co-occurrence matrix, HOG – histogram of oriented gradients, KNN – K-nearest neighbour, LBP – local binary patterns, SVM – support vector machine

the feature engineering approach (single-set feature and combined-set feature) is 91.1% accuracy and 0.96 AUC in the case of fine KNN. The confusion matrix and AUC curve are illustrated in Figure 4. Because the result revealed the accuracy to be more then 90% and AUC more than 0.95, the proposed diagnostic system, i.e. fine KNN with a combination of features GLCM, HOG, and LBP, is a very good diagnostic model.

## Conclusions

The brain tumour classification is an exploring research for machine learning people and medical practitioners because the deep learning approach is a black box method and medical practitioners are unable to analyse the exact features of brain MRI for classification. The approach proposed in this article is challenging enough to the deep learning approach. The proposed method, i.e. fine KNN, achieved 91.1% accuracy and 0.96 of AUC. Furthermore, this model has the possibility to integrate in low-end devices unlike deep learning, which requires a complex system. Again, the performance of the classification models may improve by the introduction of optimization techniques.

The authors declare no conflict of interest.

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#### Author for correspondence

Prabira Kumar Sethy, PhD Sambalpur University Burla, India e-mail: prabirsethy.05@gmail.com

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