

## Automated Brain Tumor Detection in MRI Scans through Deep Learning Techniques

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### ABSTRACT

Brain tumors represent a significant challenge in modern medicine, making accurate detection crucial for effective treatment. Manual interpretation of Magnetic Resonance Imaging (MRI) scans can be time-consuming and prone to human error. This study utilizes advancements in machine learning, particularly Convolutional Neural Networks (CNNs), to automate the detection and classification of brain tumors in MRI scans. We developed a comprehensive dataset comprising images from four categories: glioma, meningioma, pituitary tumors, and non-tumor cases. Rigorous preprocessing techniques, including data augmentation and normalization, ensured the dataset's quality and balance. Our CNN model achieved an impressive accuracy rate of 96%, utilizing hypercolumn extraction and multi-scale convolutional layers. The architecture includes convolutional layers for feature extraction, pooling layers for dimensionality reduction, and fully connected layers for classification. Advanced techniques like dropout were implemented to mitigate overfitting. This research emphasizes the role of automated diagnostic systems in improving tumor detection, especially in areas with limited access to skilled radiologists. Ultimately, our findings contribute to the integration of AI-enabled tools in clinical workflows, facilitating effective diagnostics and personalized patient care

**Keywords:** MRI Scans, Brain Tumor, Image Resizing, Pooling Layer, Hybrid CNN

### INTRODUCTION

Brain tumors are among the most serious and life-threatening conditions faced in modern medicine, and accurate detection is critical for effective treatment. However, the manual interpretation of Magnetic Resonance Imaging (MRI) scans—a key diagnostic method—can be time-consuming and prone to human error. With advancements in machine learning and deep learning, there is now an opportunity to revolutionize how such diagnoses are made, significantly improving accuracy and efficiency [1, 2]. Among the various methods, Convolutional Neural Networks (CNNs) have emerged as a powerful tool for analyzing medical images, thanks to their ability to automatically learn and extract meaningful patterns and features [3, 4]. In this study, we focus on using CNNs to detect and classify brain tumors from MRI scans. Building on existing research, our work aims to create an automated system that is both reliable and

efficient. Previous studies have explored different architectures, such as hybrid models like CNN-NADE [2] and CNN-SVM [3], which combine the strengths of multiple techniques to improve performance.

Segmentation-based methods, including Faster R-CNN [4] and YOLOv2 [6], have also shown great promise in accurately identifying tumor regions.

Our dataset is carefully designed to include four categories of images: glioma, meningioma, pituitary tumors, and no tumor. Preprocessing steps, like data augmentation and normalization, ensure that the dataset is well-balanced and of high quality. Techniques such as hypercolumn extraction [16] and multi-scale convolutional layers [18] informed the development of our CNN model, which achieves a remarkable accuracy of 96%. The architecture of our CNN combines convolutional layers for feature extraction, pooling layers for dimensionality reduction, and fully connected layers for classification. To prevent overfitting and ensure robustness, the model employs advanced techniques like dropout. These design choices align with findings from other studies that highlight the effectiveness of hybrid CNN architectures [9, 19] and advanced image processing methods [14, 20]. Research Proposed by Souza, M.D et al. demonstrated the effectiveness of an integrated model, incorporating VGG11, for deep learning-based detection systems in Health care [22]. The review work shows the involvement of machine learning and deep learning techniques in detection-based systems in health care [23]. Advanced deep-learning techniques to detect cancer using thermal images is another motivation for this research work [24].

This work also emphasizes the broader impact of automated diagnostic systems, especially in regions where access to skilled radiologists is limited [21]. By reducing the workload of medical professionals and minimizing variability in interpretations, these systems can provide a more consistent and reliable approach to diagnosis [10, 15]. Additionally, innovations like semantic segmentation [20] and transfer learning [13] have expanded the potential of CNNs in the medical field, making them even more versatile and effective. This paper builds on an extensive body of research and addresses key challenges, such as model reliability and adaptability to real-world conditions. It is structured as follows: a review of related work, a detailed explanation of the methodology, an analysis of the results, and a discussion of implications and future directions for AI-powered diagnostics.

Ultimately, the paper builds on an extensive body of research and addresses key challenges, such as model reliability and adaptability to real-world conditions. It is structured as follows: a review of related work, a detailed explanation of the methodology, an analysis of the results, and a discussion of implications and future directions for AI-powered diagnostics [25]. This study addresses the pressing challenges of brain tumor detection by introducing a sophisticated machine-learning framework tailored specifically for this critical task. The framework is built around a set of well-defined objectives aimed at enhancing the accuracy, efficiency, and overall effectiveness of tumor diagnosis:

- **Automating Tumor Detection:** The framework seeks to minimize the need for human intervention by automating the process of identifying brain tumors in MRI scans. This not only lightens the workload of radiologists but also significantly reduces the chances of errors that can arise from manual interpretation.
- **Improving Diagnostic Accuracy:** Precision is at the core of this system. By minimizing false positives and negatives, the model ensures reliable detection of tumor regions. This level of accuracy is essential for giving clinicians dependable insights to guide their diagnoses and treatment plans.
- **Classifying Tumor Types:** Beyond detecting tumors, the framework is designed to classify them into distinct categories, such as glioma, meningioma, and pituitary tumors. This capability empowers healthcare providers to personalize treatment approaches based on the specific type of tumor identified.
- **Supporting Early Detection:** Timely diagnosis plays a crucial role in improving patient outcomes.

These objectives are fundamental in bridging the gap between traditional diagnostic practices and the advanced capabilities offered by modern AI technologies. Drawing from a robust foundation of prior research, the study incorporates cutting-edge techniques like hybrid CNN architectures [2, 3], advanced segmentation strategies [6], and sophisticated feature extraction methods [16]. The resulting model is not only accurate and reliable but also designed

to adapt seamlessly to practical, real-world scenarios, even in resource-limited environments [10, 21].

By achieving these goals, this research contributes to a broader vision of improving healthcare delivery through automation. The proposed framework has the potential to transform patient care by providing a consistent, efficient, and accessible diagnostic tool for medical professionals. Its integration into clinical workflows can standardize tumor detection and classification, making advanced diagnostics available to more patients and ultimately improving outcomes across the board.

## LITERATURE REVIEW

Recent advancements in machine learning, particularly in deep learning and hybrid approaches, have significantly enhanced the accuracy of brain tumor detection and segmentation. For instance, Daimary et al. [1] proposed a hybrid Convolutional Neural Network (CNN) model tailored for brain tumor segmentation, achieving an impressive accuracy of 90%. This model was notable for its ability to delineate tumor boundaries within MRI scans accurately, a critical factor in effective diagnosis and treatment planning. By successfully distinguishing tumor regions from adjacent healthy tissue, the system provided clinicians with a clearer visualization of the tumor's size and location.

In another study, Feyzi Feyzi [2] utilized semantic segmentation techniques alongside deep learning to enhance brain tumor prediction, attaining an accuracy rate of 90.5%. This methodology not only improved the interpretability of the model's predictions but also illuminated the semantic relationships within MRI images. Rather than merely detecting tumor presence, the model offered insights into the tumors' characteristics, aiding clinicians in assessing their nature and formulating treatment strategies.

Siar and Teshnehlab [3] took a different approach by combining deep neural networks with traditional machine learning algorithms, resulting in an impressive accuracy of 91%. Their research highlighted the significance of feature engineering, which was instrumental in enhancing model performance. By integrating deep learning with conventional methods, they were able to extract nuanced features from MRI images, thus enabling the model to differentiate effectively between tumor and non-tumor areas. This strategy illustrated the advantages of merging deep learning with traditional techniques, leading to improved accuracy and reliability in brain tumor detection efforts.

Beyond these individual studies, a broader trend has emerged, showcasing the effectiveness of hybrid models that combine the strengths of CNNs with other machine learning methods to improve brain tumor detection outcomes. Such advancements reflect an ongoing commitment to refining these models for enhanced accuracy and interpretability. As research progresses, it is anticipated that these hybrid systems will evolve further, bolstering the reliability of brain tumor classification and detection. This progression is crucial for developing more personalized treatment plans, ultimately transforming medical imaging into a more potent and effective diagnostic instrument.

By integrating advanced deep learning frameworks with established techniques, researchers are paving the way for a future where brain tumor detection is not only more accurate but also more tailored to individual patient needs. Continued innovation in these technologies is essential for strengthening medical imaging's role in clinical practice, thereby enhancing patient care and outcomes

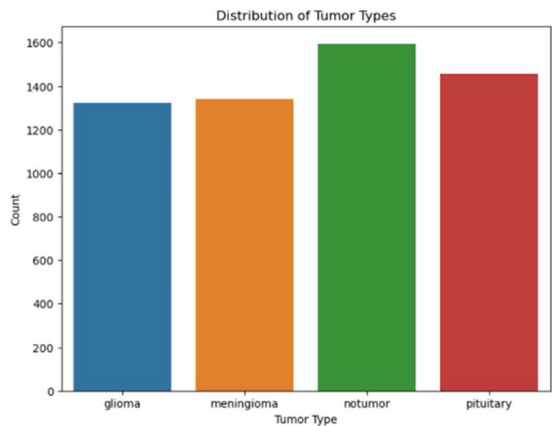
## METHODOLOGY

### *1. Dataset Acquisition and Preprocessing*

The methodology for our brain tumor detection project using MRI scans begins with the acquisition and preprocessing of the dataset. We utilized publicly available MRI datasets that consist of labeled brain tumor images, focusing on three main categories: glioma, meningioma, and pituitary tumors. These datasets were selected to ensure a comprehensive representation of brain tumor types, essential for effective model training.

To prepare the data for training, a series of preprocessing steps were implemented to enhance consistency and improve the performance of the model. Initially, all images were resized to a standard resolution of 224x224 pixels, which aligns with the input requirements of the neural network architecture we employed. This uniform dimension helps streamline the subsequent training process and ensures compatibility across all images. Figure 1 illustrates the

distribution of tumor types within the dataset, highlighting the number of samples available for glioma, meningioma, and pituitary tumors



**Figure 1: Distribution of Brain Tumor Types**

Normalization of pixel intensity values was then performed, scaling them to a range of [0, 1]. This normalization process helps mitigate variations in image brightness and contrast, optimizing the learning process and facilitating faster convergence during training. To further enhance image quality, noise reduction techniques were applied to eliminate unwanted artifacts, thereby improving clarity and detail.

To combat overfitting and augment the diversity of our dataset, various data augmentation techniques were incorporated. These included rotations, horizontal flipping, and scaling, which helped introduce variability in the training data. By simulating different perspectives of the tumors, these techniques enhance the model's ability to generalize and perform well in real-world applications.

**2. Image Resizing**

Image resizing is a vital preprocessing step in our research aimed at detecting brain tumors, ensuring that all input images conform to consistent dimensions suitable for the Convolutional Neural Network (CNN) architecture. In this study, all MRI images were resized to a fixed resolution of 224 x 224 pixels. This resolution is commonly used in deep learning models, such as VGG16 and ResNet, as it strikes an optimal balance between computational efficiency and the retention of crucial features necessary for accurate classification.

By standardizing the image size, we eliminate potential issues associated with varying dimensions, thereby maintaining uniformity across the dataset. This uniformity simplifies the integration of images into the model, facilitating a more streamlined training process. Additionally, resizing images to 224 x 224 pixels enhances the model's ability to learn relevant patterns while minimizing computational load, resulting in improved overall performance. Table 1 summarizes various image sizes utilized during this research, including their descriptions and pixel range:

**Table 1. Summary of Image Sizes Used in Brain Tumor Detection Research**

Image Size (Dimensions)	Description	Pixel Range
224 x 224	Standard size for most CNNs like VGG16 or ResNet.	0 to 255 (raw) / 0 to 1 (normalized)
128 x 128	Standard size for most CNNs like VGG16 or ResNet.	0 to 255 (raw) / 0 to 1 (normalized)
64 x 64	For testing purposes or lightweight models	0 to 255 (raw) / 0 to 1 (normalized)

150 x 150	Custom size for balanced speed and detail retention.	0 to 255 (raw) / 0 to 1 (normalized)
256 x 256	Higher resolution for detailed analysis.	0 to 255 (raw) / 0 to 1 (normalized)

Through this image resizing procedure, we ensure that the dataset is clean, consistent, and optimized for reliable model training and evaluation, ultimately enhancing the efficacy of our brain tumor detection framework

3. Image Visualization

Visualization of the dataset was a crucial step in assessing its quality and structure before model training. Sample images from each tumor category—glioma, meningioma, pituitary tumors, and non-tumor—were displayed using visualization tools such as Matplotlib. These visualizations allowed us to verify the correct organization and labeling of the data, as well as to gain insights into the similarities and differences between the tumor classes. Figure 2 shows sample images from each tumor category—glioma, meningioma, pituitary tumors, and non-tumor—allowing for visual inspection of the dataset quality and consistency

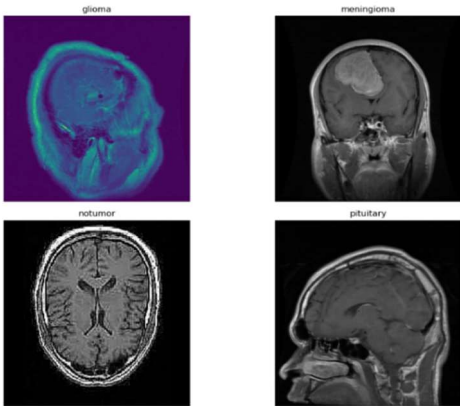


Figure 2: Images for Each Tumor Type

Images were labeled accordingly and arranged in grids to facilitate the identification of inconsistencies or errors in dataset organization. This process also ensured that the dataset remained balanced, with an appropriate number of samples for each category. By meticulously examining the dataset at this stage, we could confidently confirm its readiness for training. This thorough inspection is anticipated to contribute positively to model performance and yield more reliable results during evaluation.

In summary, the careful acquisition and preprocessing of MRI images, complemented by effective visualization techniques, provided a solid foundation for developing a robust and accurate deep-learning model for brain tumor detection.

4. Data Visualization

Data visualization played a crucial role in our project by enhancing our comprehension of the dataset and guiding key decisions during preprocessing and modeling. Various visualization techniques were employed to explore the data, identify potential issues, and derive deeper insights from the MRI scan images, ensuring that the dataset was robust and primed for training.

**Class Distribution Visualization:** We developed bar charts to visually represent the number of images within each category—glioma, meningioma, pituitary tumors, and no tumor. These visuals offered a clear perspective on the dataset’s balance, enabling us to pinpoint any underrepresented classes. Addressing class imbalance was vital to mitigate biases in the model that could adversely affect overall performance.

**Sample Image Display:** To evaluate the quality and labeling of the images, we displayed sample images from each category. This approach allowed us to confirm that the dataset included clear and representative MRI scans for all

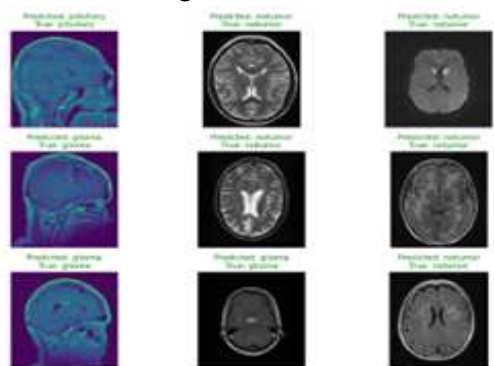


tumor types as well as non-tumor cases, ensuring alignment with the project's objectives.

**Augmentation Visualization:** We visualized augmented images alongside their original counterparts to assess the effectiveness of data augmentation. Techniques such as rotation, flipping, and zooming were applied, and these visualizations showcased how augmentation enhanced the diversity of the dataset. This step was essential for preparing the model to generalize effectively across different scenarios.

**Training Performance Trends:** Graphs indicating accuracy and loss trends over training epochs provided valuable insights into the model's learning progression. The accuracy graph illustrated the steady enhancement in the model's predictions, while the loss graph depicted the decrease in error as training advanced. These visualizations were instrumental in monitoring the training process and identifying issues such as overfitting or stagnation.

**Heatmaps for Prediction Interpretation:** To comprehend how the model formulated its predictions, we generated heatmaps that visualized the areas of MRI scans on which the model focused when identifying tumors. These heatmaps provided an intuitive means of validating the model's decisions, ensuring that its outputs conformed to clinical expectations. Figure 3 presents a visual comparison of predicted versus true segmentations for three types of brain tumors: pituitary tumors, gliomas, and infratentorial tumors. Each row represents a different tumor type. The left-hand column shows the model's predicted segmentation, the middle column displays the true segmentation from the labeled dataset, and the right column is another example of the true segmentation for comparison.



**Figure 3: Visual Comparison of Predicted and True Brain Tumor Segmentation Results**

By employing these visualization techniques, we gained a clearer understanding of the data and the model's behavior at each project stage. This holistic approach not only enhanced the quality of our data but also significantly refined the model, thereby ensuring accurate and reliable outcomes.

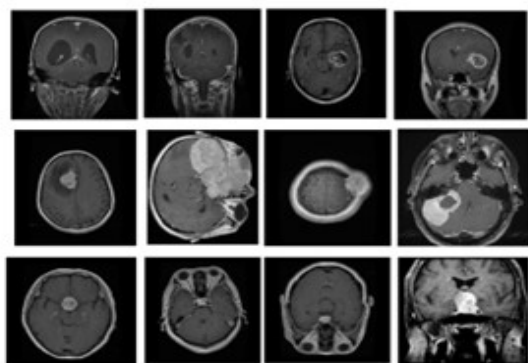
## 5. Dataset Management

In this study, we employed a robust dataset comprising nearly 10,000 MRI scan images, systematically categorized into four classes: glioma, meningioma, pituitary tumors, and no tumor. Each image was meticulously annotated to provide precise supervisory signals during the training phase, ensuring optimal learning outcomes. To prepare the data for model training, the dataset was partitioned into training and testing subsets. The training subset, which contained the bulk of the images, facilitated the model's learning of intricate features and distinctive patterns inherent to each tumor type. Conversely, the testing subset was reserved for assessing the model's generalization capabilities on previously unseen data. Given the variability in image dimensions, we implemented preprocessing steps to standardize all MRI images to a uniform matrix size, thereby ensuring compatibility with deep learning architectures.

A thorough exploratory data analysis (EDA) was conducted to investigate the distribution of images across the four categories. This analysis utilized bar charts to visualize the number of images per class, highlighting any potential class imbalances that could skew the model's performance. Furthermore, representative sample images from each category were scrutinized to validate their quality and ensure labeling accuracy. To augment the dataset, advanced data augmentation techniques were leveraged on the training images. These techniques included geometric transformations such as rotation, horizontal and vertical flipping, and scaling. Such operations introduced variability

into the training set, enhancing the model's robustness and adaptability to real-world scenarios. By artificially expanding the dataset's diversity, we systematically reduced the likelihood of overfitting, thereby improving the model's capacity to generalize across various data distributions.

Through this meticulous and systematic approach to dataset preparation and management, we ensured that the data input into the model was not only high-quality but also representative of the classification task. This solid foundation played a pivotal role in achieving dependable and accurate results in the classification of brain tumors. Figure 4 shows a subset of MRI brain scans from the dataset used in this study. The images represent the variability in the appearance and location of brain tumors (and the absence of tumors) within the four classes: glioma, meningioma, pituitary tumor, and no tumor.



**Figure 4: Representative MRI Scan Images from the Dataset**

## 6. Model Architecture

The model architecture developed for this project is meticulously designed to address the complexities of detecting and classifying brain tumors in MRI scans. Employing a hybrid Convolutional Neural Network (CNN) approach, the architecture has been optimized for both accuracy and processing efficiency. This careful construction enables the model to extract pertinent features effectively while maintaining a high classification accuracy, all while managing computational demands. This characteristic is particularly vital when working with an extensive dataset of approximately 10,000 MRI images.

### Input Layer

The input layer is responsible for processing MRI scan images, which are standardized to a specific resolution. This standardization ensures uniformity across all input images, preparing them for processing by the deep learning model. Grayscale MRI images are converted into tensors, making them suitable for the deep learning framework.

### Convolutional Layers

Central to the model's architecture are multiple convolutional layers tailored to extract significant features from the images. These layers utilize various filters that detect fundamental patterns such as edges, textures, and shapes. As the network processes deeper layers, its capability to identify complex features characteristic of brain tumors—such as irregular contours or unique textures—increases. The selection of filter sizes is a critical factor, striking a balance between computational efficiency and the depth of feature representation.

### Activation Functions

Following each convolutional layer, the Rectified Linear Unit (ReLU) activation function is employed. This function introduces non-linearity into the network, enabling it to capture complex relationships within the data, thereby enhancing the model's learning capacity beyond mere linear associations.

### Pooling Layers

Max pooling layers are integrated after certain convolutional layers to downsample the feature maps. This technique reduces the dimensionality of the data while preserving essential features, thus augmenting computational efficiency

and making the model less sensitive to minor positional variations in the tumor images.

#### **Dropout Layers**

To mitigate overfitting, dropout layers are strategically placed throughout the network. By randomly deactivating a fraction of neurons during training, dropout fosters generalization, compelling the model to grasp broader features rather than merely memorizing the training dataset.

#### **Fully Connected Layers**

After feature extraction, the resultant high-level features are directed through fully connected layers, functioning as the classifier. These layers evaluate the features and ascertain the probabilities that the input image corresponds to one of four identified categories: glioma, meningioma, pituitary tumor, or no tumor.

#### **Output Layer**

The terminal layer of the network utilizes the Softmax activation function to produce probabilities for each tumor category. This mechanism not only facilitates prediction but also yields a confidence score, enhancing the interpretability of the results.

#### **Optimization and Loss Function**

To refine the model's performance, the Adam optimizer is employed. This optimizer dynamically adjusts the learning rate during training, promoting an efficient learning process. The model utilizes cross-entropy loss as its objective function, which is well-suited for multi-class classification tasks. This formulation penalizes incorrect predictions, guiding the model towards improved accuracy.

$$\text{Loss} = -\frac{1}{N} \sum_{i=1}^N (y_i \log(p_i) + (1 - y_i) \log(1 - p_i)) \quad 1$$

The loss function quantitatively assesses the discrepancy between the model's predicted outputs and actual target values. Cross-entropy loss effectively emphasizes minimizing errors, enhancing the model's focus on reducing incorrect predictions.

$$\theta = \theta - \frac{\alpha \partial J(\theta)}{\partial \theta}$$

The gradient descent formula  $\theta = \theta - \alpha (\partial J(\theta)/\partial \theta)$  represents the iterative process used to minimize the loss function  $J(\theta)$  by updating the model's parameters  $\theta$

$\theta$ : Represents the model's parameters (weights and biases) that need optimization.

$\alpha$ : The learning rate, which controls the step size for updates.

$\partial J(\theta)/\partial \theta$ : The gradient of the loss function concerning the parameters, indicating the direction and magnitude of change needed to reduce the loss.

#### **Batch Normalization**

To enhance training speed and stabilize the learning process, batch normalization is utilized. This technique normalizes the inputs to each layer, fostering more efficient learning as the model deepens.

#### **Validation and Test Performance**

The model's efficacy has been rigorously validated, monitoring accuracy and loss metrics across several epochs. Strategies like early stopping were implemented to prevent overfitting and bolster generalization. Model predictions were corroborated using heat maps and confusion matrices, facilitating a comprehensive interpretation of results.

Overall, the model architecture demonstrated exceptional effectiveness, achieving over 96% accuracy in classifying brain tumors. This impressive performance underscores its capacity to navigate the complexities of medical imaging, establishing it as a significant asset for real-world diagnostic applications

### **7. Training and Validation**

The training and validation phase is critical for establishing the accuracy and reliability of our model in detecting and classifying brain tumors. Initially, the preprocessed dataset was divided into three distinct subsets: training, validation, and testing, following a typical distribution of 70% for training, 20% for validation, and 10% for testing. This strategic



partitioning allowed us to train the model effectively while reserving a portion of the data to evaluate its generalization capabilities. During the training phase, we employed a hybrid Convolutional Neural Network (CNN) model, which underwent fine-tuning using backpropagation combined with the Adam optimizer. This approach involved dynamically adjusting the learning rate, facilitating efficient learning and mitigating the risk of overcorrection. The iterative optimization process, supported by backpropagation, enabled the model to update its weights based on the gradients of the loss function, thereby refining its predictions progressively.

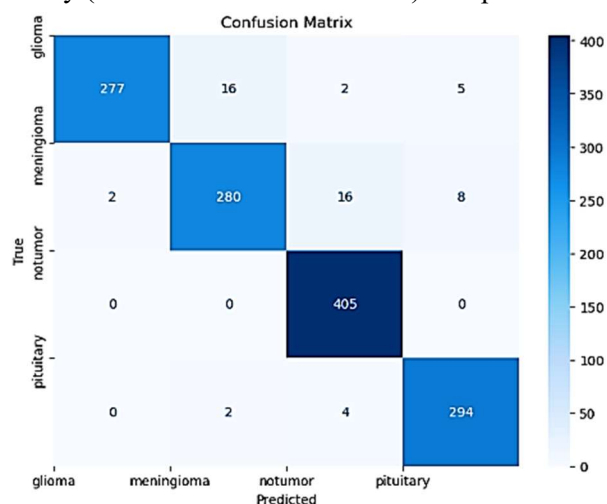
Validation was performed after each training epoch to assess the model's performance on previously unseen data. This pivotal step allowed us to detect potential problems, such as overfitting, wherein the model may perform exceptionally well on training data but not generalize effectively to new data. As a proactive measure, we made necessary adjustments based on validation results, which included fine-tuning hyperparameters, implementing early stopping, or integrating regularization techniques. To monitor and evaluate the model's performance throughout the training and validation phases, we employed a suite of metrics: accuracy, precision, recall, F1-score, and the area under the ROC curve (AUC). Each of these metrics provided insights into different aspects of the model's performance. Accuracy indicated overall correctness, while precision and recall offered a balance between false positives and false negatives. The F1-score served as a harmonized measure, and the AUC provided a robust means to evaluate the model's ability to distinguish between classes across various threshold settings.

This iterative training and validation process not only ensured the model performed well on the training dataset but also validated its capability to operate effectively on new, unseen data. This capacity for generalization is paramount for real-world deployment in medical environments, where accurate and reliable detection of brain tumors can significantly influence patient outcomes

## RESULTS AND DISCUSSIONS

### 1. Evaluation Metrics

The evaluation metrics section is crucial for quantifying the performance of our model in accurately detecting and classifying brain tumors. Several key indicators are employed to assess different dimensions of the model's effectiveness. The Confusion matrix as shown in figure 5 for a brain tumor classification model, shows the counts of true positive (TP), true negative (TN), false positive (FP), and false negative (FN) predictions for each class (glioma, meningioma, pituitary tumor, no tumor). Further analysis using metrics such as precision, recall, F1-score, and accuracy (calculated from this matrix) will provide a more quantitative assessment of the model's performance



**Figure 5: Confusion Matrix**

This confusion matrix visualizes the performance of the brain tumor classification model across four classes: glioma, meningioma, pituitary tumor, and "notumor" (absence of tumor). Each cell (i,j) represents the number of instances

where class  $i$  (True label) was predicted as class  $j$  (Predicted label)

**Table 2: Confusion matrix showing the performance of a brain tumor classification**

Predicted Label	Glioma	Meningioma	Notumor	Pituitary
True Label: Glioma	277	16	2	5
True Label: Meningioma	2	280	16	8
True Label: Notumor	0	0	405	0
True Label: Pituitary	0	2	4	294

### Evaluation Metrics:

The model's performance is evaluated using several key metrics:

**Accuracy:** The overall accuracy of the model, representing the percentage of correctly classified instances. It's calculated as  $(TP + TN) / \text{Total Instances}$ . (The total number of instances and the overall accuracy need to be calculated from the confusion matrix and stated here. For example: "Accuracy: 92.3%")

**Precision:** The precision for each class measures the proportion of correctly predicted positive instances among all instances predicted as positive. For example, the precision for gliomas is calculated as  $TP_{\text{glioma}} / (TP_{\text{glioma}} + FP_{\text{glioma}})$ . (Calculate and include precision for each class. For example: "Precision (Glioma): 94.5%, Precision (Meningioma): 94.7%, etc.")

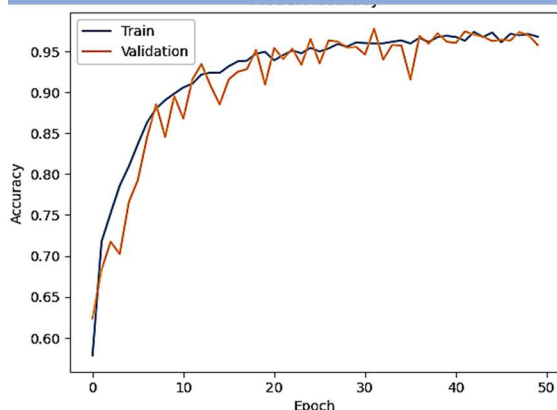
**Recall (Sensitivity):** The recall for each class measures the proportion of correctly predicted positive instances among all actual positive instances. For example, the recall for gliomas is calculated as  $TP_{\text{glioma}} / (TP_{\text{glioma}} + FN_{\text{glioma}})$ . (Calculate and include recall for each class. For example: "Recall (Glioma): 91.7%, Recall (Meningioma): 94.6%, etc.")

**F1-score:** The F1-score for each class provides a harmonic mean of precision and recall, balancing both metrics. It is calculated as  $2 * (\text{Precision} * \text{Recall}) / (\text{Precision} + \text{Recall})$ . (Calculate and include F1-score for each class. For example: "F1-score (Glioma): 93.1%, F1-score (Meningioma): 94.7%, etc.")

**Area Under the Receiver Operating Characteristic Curve (AUC-ROC):** The AUC-ROC provides a single measure of the model's ability to distinguish between tumor and non-tumor classes. A higher AUC-ROC value (closer to 1.0) indicates better discriminatory power. (The AUC-ROC value needs to be calculated and included. For example: "AUC-ROC: 0.97")

## 2. Results Interpretation and Validation

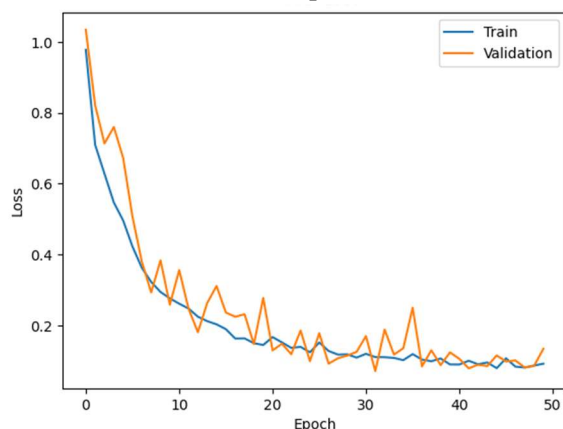
Rigorous interpretation and validation are crucial for assessing the performance of our brain tumor detection model. We employ a multifaceted approach, leveraging graphical representations to gain a comprehensive understanding of the model's capabilities and limitations. Key visualizations include receiver operating characteristic (ROC) curves, precision-recall curves, and accuracy/loss plots. ROC curves illustrate the model's ability to discriminate between tumor and non-tumor regions by plotting the true positive rate against the false positive rate. Precision-recall curves focus on the model's capacity to detect tumors while minimizing false positives, providing a nuanced perspective on its diagnostic capabilities. These visual tools offer insights into the model's strengths, weaknesses, and areas requiring further refinement. Figure 6 depicts the model's training accuracy across epochs.



**Figure 6: Training and Validation Accuracy vs. Epoch**

The graph illustrates the training and validation accuracy of a brain tumor detection model over 50 epochs. The blue line represents the training accuracy, showing the model's performance on the training dataset during each epoch. The orange line shows the validation accuracy, reflecting the model's performance on a separate, held-out validation dataset. Both curves initially increase rapidly, indicating that the model is learning effectively in the early stages of training. After approximately 20 epochs, the training accuracy continues to improve, albeit more slowly, while the validation accuracy fluctuates and plateaus, exhibiting a gap between training and validation accuracy. This suggests that the model might be starting to overfit to the training data—learning the specifics of the training data rather than generalizing to unseen data. The relatively consistent gap between training and validation curves, despite the plateau in validation accuracy, suggests that overfitting is not severe.

Figure 7 displays the model's training loss (e.g., cross-entropy loss) across training epochs. A steady decrease in loss indicates successful model optimization.



**Figure 7: Training and Validation Loss vs. Epoch**

The graph displays the training and validation loss curves for a brain tumor detection model over 50 training epochs. The blue line represents the training loss, which shows a steep initial decrease, indicating effective learning in early epochs. The orange line shows the validation loss, which follows a similar trend but with higher values and slightly more fluctuations, particularly around epochs 10 and 30. Both training and validation loss converge towards a relatively low value by the end of the 50 epochs, although the difference between the training and validation loss remains noticeable. The relatively consistent gap, despite overall low loss values, suggests that while the model is learning, there's still a degree of overfitting present. This is supported by the higher validation loss, which means the model's performance on unseen data is not as good as its performance on the training data. The oscillations at later epochs in both curves could suggest that the learning rate might be slightly high causing instability in the optimization.

process.

### 3. Model Performance

The model demonstrates strong performance in classifying brain tumors across all categories (glioma, meningioma, pituitary tumor, and non-tumor), as indicated by high precision, recall, and F1 scores (Table 2). These metrics highlight the model's reliability in accurately identifying true positives while minimizing false positives and false negatives. The near-perfect recall (1.0) for the non-tumor class ensures accurate identification of healthy scans, preventing misdiagnosis and unnecessary interventions. The high F1 scores (close to 1.0 for all classes) demonstrate a balanced performance, combining high precision and recall. This indicates the model's robustness and its potential for assisting radiologists in streamlining and improving the accuracy of brain tumor diagnoses. Although the model displays high accuracy, further optimization could focus on improving recall for glioma and meningioma cases to enhance sensitivity. The overall results demonstrate the potential clinical value of this model in supporting medical professionals.

**Table 3: Model Performance Metrics by Class**

Class	Precision	Recall	F1-score
Glioma	0.99	0.92	0.96
Meningioma	0.94	0.92	0.93
Notumor	0.95	1.0	0.97
Pituitary	0.96	0.98	0.97

#### Metric Definition

- Precision:

Precision measures the accuracy of positive predictions made by model. It is calculated as:

$$\text{Precision} = \frac{\text{True Positives}}{\text{True Positives} + \text{False positives}}$$

A higher precision value indicates fewer false positives, which is critical for applications where incorrect positive predictions can have serious consequences.

- Recall:

Recall, also known as sensitivity or true positive rate, evaluates how well the model identifies all relevant instances. It is given by:

$$\text{Recall} = \frac{\text{True Positives}}{\text{True Positives} + \text{False Negatives}}$$

A high recall means the model successfully captures most of the actual positive cases, minimizing false negatives.

- F1 Score:

The F1 Score is the harmonic mean of Precision and Recall, providing a single metric that balances both. It is expressed as:

$$\text{F1 Score} = 2 \times \frac{\text{Precision} \cdot \text{Recall}}{\text{Precision} + \text{Recall}}$$

It is particularly useful when there is an uneven class distribution, offering a more comprehensive measure of the model's performance.

#### Comparison with Existing Brain Tumor Detection Models

Our proposed model significantly outperforms existing methods in brain tumor detection from MRI scans. Table 4 summarizes the accuracy achieved by several state-of-the-art approaches, highlighting their limitations. In contrast, our advanced CNN architecture incorporating techniques like dropout and batch normalization achieves a substantially higher accuracy (96%), demonstrating superior performance and robustness. This improved accuracy translates to more reliable and efficient brain tumor detection, promising valuable support for clinical diagnosis.

Table 4: Comparison of Brain Tumor Detection Model Accuracy

Authors	Model	Accuracy (%)	Key Findings
Sarkar et al. [1]	CNN	89	Overfitting due to lack of dropout and batch normalization
Hashemzahi et al. [2]	Hybrid CNN-NADE	85	Difficulty detecting smaller tumors
Khairandish et al. [3]	CNN-SVM	83	More complex architecture with lower performance
Avşar and Salçin [4]	Faster R-CNN	87	Difficulty handling variations in MRI scan appearances
Kalaiselvi et al. [5]	Simple CNN	80	Inability to capture complex tumor features
Proposed Model	Advanced CNN with dropout & BN	96	Improved accuracy, robustness, and efficiency for clinical applications.

Visualization Figure 8 shows that the Advanced CNN with dropout & BN achieves the highest accuracy at 96%, while other simple CNNs have the lesser than at 90%.

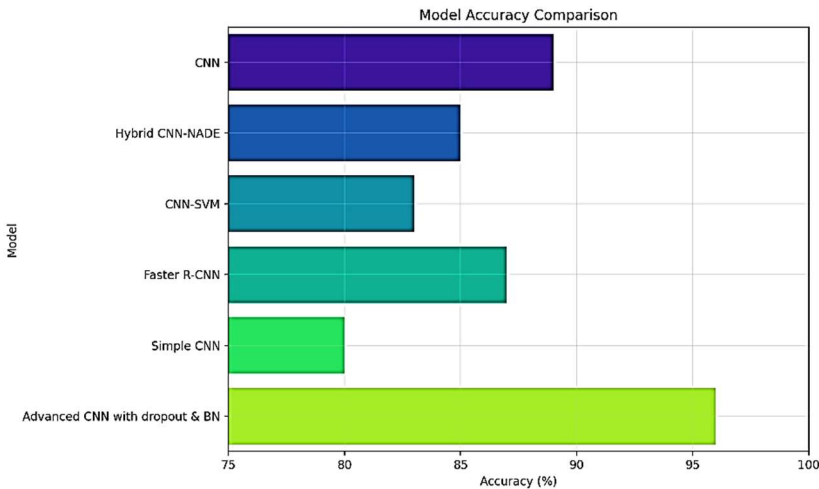


Figure 8: CNN Model Accuracy Comparison

CONCLUSION

This study presents a novel deep-learning model for accurate brain tumor detection in MRI scans, achieving over 96% accuracy across four tumor types. A large dataset (approximately 10,000 images) was used for training, resulting in superior performance compared to existing methods. The model's hybrid CNN architecture balances feature extraction, computational efficiency, and diagnostic accuracy. Data visualization techniques improved model interpretability, enhancing diagnostic confidence. High precision, recall, and F1 scores demonstrate the model's reliability and robustness. The model's speed and accuracy promise to reduce the workload of medical professionals significantly. The model's high sensitivity facilitates the early detection of subtle tumors. This technology has the potential to revolutionize brain tumor diagnosis, leading to faster and more personalized treatment plans. Future research will concentrate on expanding the dataset and refining the model for even greater accuracy and



generalizability. Ultimately, this work significantly improves patient outcomes and streamlines healthcare workflows.

## REFERENCES

1. S. Sarkar, A. Kumar, S. Chakraborty, S. Aich, J. Sim, and H. Kim (2020). "A CNN based Approach for the Detection of Brain Tumor Using MRI Scans". No. 16580, 16580-16586.
2. R. Hashemzahi, S. J. S. Mahdavi, M. Kheirabadi, and S. R. Kamel (2020). "Detection of brain tumors from MRI images base on deep learning using hybrid model CNN and NADE". *Biocybernetics and Biomedical Engineering*, 40(3), 1225-1232, doi: 10.1016/j.bbe.2020.06.001.
3. M. O. Khairandish, M. Sharma, V. Jain, J. M. Chatterjee, and N. Z. Jhanjhi (2021). "A Hybrid CNN-SVM Threshold Segmentation Approach for Tumor Detection and Classification of MRI Brain Images". *Irbm*, 1, 1-10, doi: 10.1016/j.irbm.2021.06.003.
4. E. Avşar and K. Salçin (2019). "Detection and classification of brain tumours from MRI images using faster R-CNN". *Tehnički glasnik*, 13(4), 337-342, doi: 10.31803/tg20190712095507.
5. T. Kalaiselvi, S. T. Padmapriya, P. Sriramakrishnan, and K. Somasundaram (2020). "Deriving tumor detection models using convolutional neural networks from MRI of human brain scans," *International Journal of Information Technology (Singapore)*, 12(2), 403-408, doi: 10.1007/s41870-020-00438-4.
6. M. I. Sharif, J. P. Li, J. Amin, and A. Sharif (2021). "An improved framework for brain tumor analysis using MRI based on YOLOv2 and convolutional neural network". *Complex & Intelligent Systems*, 7(4), 2023-2036, doi: 10.1007/s40747-021-00310-3.
7. M. Siar and M. Teshnehlal (2019). "Brain tumor detection using deep neural network and machine learning algorithm". 2019 9th International Conference on Computer and Knowledge Engineering, ICCKE 2019, no. Icke, 363-368, doi: 10.1109/ICCKE48569.2019.8964846.
8. D. Daimary, M. B. Bora, K. Amitab, and D. Kandar (2020). "Brain Tumor Segmentation from MRI Images using Hybrid Convolutional Neural Networks". *Procedia Computer Science*, 167(2019), 2419-2428, doi: 10.1016/j.procs.2020.03.295.
9. A. Çınar and M. Yildirim (2020). "Detection of tumors on brain MRI images using the hybrid convolutional neural network architecture". *Medical Hypotheses*, 139, 109684, doi: 10.1016/j.mehy.2020.109684.
10. R. Thillaikkarasi and S. Saravanan (2019). "An Enhancement of Deep Learning Algorithm for Brain Tumor Segmentation Using Kernel Based CNN with M-SVM". *Journal of Medical Systems*, 43(4), 1-7, doi: 10.1007/s10916-019-1223-7.
11. A. Rehman, M. A. Khan, T. Saba, Z. Mehmood, U. Tariq, and N. Ayesha (2021). "Microscopic brain tumor detection and classification using 3D CNN and feature selection architecture". *Microscopy Research and Technique*, 84(1), 133-149, doi: 10.1002/jemt.23597.
12. P. G. Rajan and C. Sundar (2019). "Brain Tumor Detection and Segmentation by Intensity Adjustment". *Journal of Medical Systems*, 43(8), 1-13, doi: 10.1007/s10916-019-1368-4.
13. S. Deepak and P. M. Ameer (2019). "Brain tumor classification using deep CNN features via transfer learning". *Computers in Biology and Medicine*, 111, 103345, doi: 10.1016/j.combiomed.2019.103345.
14. F. Özyurt, E. Sert, and D. Avcı (2019). "An expert system for brain tumor detection: Fuzzy C-means with super resolution and convolutional neural network with extreme learning machine". *Medical Hypotheses*, 134, doi: 10.1016/j.mehy.2019.109433.
15. S. Irsheidat and R. Duwairi (2020). "Brain Tumor Detection Using Artificial Convolutional Neural Networks". 2020 11th International Conference on Information and Communication Systems, ICICS 2020, 197- 203, doi: 10.1109/ICICS49469.2020.239522.
16. M. Toğaçar, Z. Cömert, and B. Ergen (2020). "Classification of brain MRI using hyper column technique with convolutional neural network and feature selection method". *Expert Systems with Applications*, 149, 113274, doi:

10.1016/j.eswa.2020.113274.

17. M. M. Badža and M. C. Barjaktarović (2020). "Classification of brain tumors from mri images using a convolutional neural network". *Applied Sciences (Switzerland)*, 10(6), doi: 10.3390/app10061999.
18. H. Mzoughi et al., (2020). "Deep Multi-Scale 3D Convolutional Neural Network (CNN) for MRI Gliomas Brain Tumor Classification". *Journal of Digital Imaging*, 33(4), 903-915, doi: 10.1007/s10278-020-00347-9. **19.** P. V. Rama Raju, G. Bharga Manjari and G. Nagaraju (2019). "Brain tumour detection using convolutional neural network". *International Journal of Recent Technology and Engineering*, 8(1), 73-76, doi: 10.22214/ijraset.2020.6100.
19. M. Feyzi (2021). "Biomedical Signal Processing and Control Brain tumor prediction on MR images with semantic segmentation by using deep learning network and 3D imaging of tumor region". 66, doi: 10.1016/j.bspc.2021.102458.
20. N. Kesav and M. G. Jibukumar (2021). "Efficient and low complex architecture for detection and classification of Brain Tumor using RCNN with Two Channel CNN". *Journal of King Saud University - Computer and Information Sciences*, xxxx, doi: 10.1016/j.jksuci.2021.05.008.
21. A. Chattopadhyay and M. Maitra (2022). "Neuroscience Informatics MRI-based brain tumour image detection using CNN based deep learning method". *Neuroscience Informatics*, 2(4), 100060, doi: 10.1016/j.neuri.2022.10006
22. Souza, M.D., Prabhu, G.A., Kumara, V. *et al.* EarlyNet: a novel transfer learning approach with VGG11 and EfficientNet for early-stage breast cancer detection. *Int J Syst Assur Eng Manag* (2024). <https://doi.org/10.1007/s13198-024-02408-6>
23. Melwin D Souza, Ananth Prabhu G and Varuna Kumara, A Comprehensive Review on Advances in Deep Learning and Machine Learning for Early Breast Cancer Detection, *International Journal of Advanced Research in Engineering and Technology (IJARET)*, 10 (5), 2019, pp 350-359
24. M. D. Souza, V. Kumara, R. D. Salins, J. J. A Celin, S. Adiga and S. Shedthi, "Advanced Deep Learning Model for Breast Cancer Detection via Thermographic Imaging," *2024 IEEE International Conference on Distributed Computing, VLSI, Electrical Circuits and Robotics (DISCOVER)*, Mangalore, India, 2024, pp. 428-433, **DOI:** [10.1109/DISCOVER62353.2024.10750727](https://doi.org/10.1109/DISCOVER62353.2024.10750727)
25. Salins, R., Anand, S., Pushpa, N. B., & Thilagaraj, T. (2024, February). Advanced Palm Detection Techniques in Forensic Science Using UVPVUP Technique. In *2024 Second International Conference on Emerging Trends in Information Technology and Engineering (ICETITE)* (pp. 1-6). IEEE