Deep Learning and Multi-Objective Evolutionary Fuzzy Classifiers: A Comparative Analysis for Brain Tumor Classification in MRI Images

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- Keywords: Brain Tumor Classification, Explainable Artificial Intelligence, Deep Learning Learning, Fuzzy Rule-Based Classifiers, Multi-Objective Fuzzy Systems.
- Abstract: This paper presents a comparative analysis of Deep Learning models and Fuzzy Rule-Based Classifiers (FBRCs) for Brain Tumor Classification from MRI images. The study considers a publicly available dataset with three types of brain tumors and evaluates the models based on their accuracy and complexity. The study involves VGG16, a convolutional network known for its high accuracy, and FBRCs generated via a multi-objective evolutionary learning scheme based on the PAES-RCS algorithm. Results show that VGG16 achieves the highest classification performance but suffers from overfitting and lacks interpretability, making it less suitable for clinical applications. In contrast, FBRCs, offer a good balance between accuracy and explainability. Thanks to their straightforward structure, FRBCs provide reliable predictions with comprehensible linguistic rules, essential for medical decision-making.

1 INTRODUCTION

The use of Machine Learning (ML) and Artificial Intelligence (AI) for Magnetic Resonance Imaging (MRI) scan analysis is revolutionizing the tools supporting physicians for brain cancer detection, diagnosis, and prognosis. This disease affects approximately 24,000 people annually in the U.S.¹ and 22,000 people in Europe². With around 18,000 and 17,000 deaths per year in the U.S. and in Europe, respectively, these advanced diagnostic tools are crucial for improving accuracy and timeliness in patient treatment (Khalighi et al., 2024).

The first generation of methods for automatic medical image analysis was based on classical ML models. Currently, Deep Learning (DL) models have become the state-of-the-art approach because of their ability to automatically learn complex features from raw image data (Zhou et al., 2023). Convolutional neural networks (CNNs), in particular, have shown remarkable performance in accurately classifying brain tumor images, often surpassing traditional ML methods (Al-Zoghby et al., 2023). However, despite their high accuracy, DL models are often criticized for being "black boxes" with limited transparency, in a field

where the decision-making process is as important as the accuracy of the prediction (Hulsen, 2023). As a result, proposals in the specialized literature are presented to make medical decision-making questionable, understandable, and explainable to the different stakeholders. As discussed in (Wang et al., 2024), the transparency and explainability requirements are fundamental due to the critical and high-risk nature of AI-based medical imaging applications.

To address the explainability requirements, posthoc techniques have been developed to provide insights into the predictions made by DL models (Van der Velden et al., 2022). Methods such as saliency maps, Grad-CAM, and SHAP are commonly used to highlight the regions of an image that most influence the model's decision. However, post-hoc explanations are often approximations that may not fully capture the model's reasoning process, and they can be computationally intensive, adding complexity to the analysis pipeline.

Alternatively, tools such as radiomics (Saidak et al., 2024), that extracts quantitative features from medical images, can be used in combination with interpretable by-design classifiers, such as decision trees (Du et al., 2023). However, radiomics-based models still require careful design and feature selection, which can be laborious and time-consuming.

108

Miglionico, G., Ducange, P., Marcelloni, F. and Pedrycz, W.

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¹https://seer.cancer.gov/statfacts/html/brain.html ²https://ecis.jrc.ec.europa.eu

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Fuzzy rule-based classifiers (FRBCs) are capable of meeting the demands of explainability and transparency in critical health applications (Cao et al., 2024), since they utilize a set of human-readable rules to make decisions. Compared to DL models with post-hoc explainability methods, FRBCs are simpler, faster to generate, and involve fewer parameters, especially if combined with radiomics features (Zhang et al., 2022).

One advanced approach in the realm of FR-BCs is represented by Multi-Objective Evolutionary Fuzzy Classifiers (MOEFCs) (Antonelli et al., 2016). MOEFCs deal with FRBCs designed using a Multiobjective evolutionary learning (MOEL) scheme to generate models characterized by good trade-offs between accuracy and interoperability.

This paper presents an experimental analysis comparing a DL model with FRBCs, generated by using PAES-RCS, in brain tumor classification (BTC) from MRI scans. As regards the PAES-RCS method, we discuss its advantages in generating models that are almost as accurate as those generated using DL. Moreover, we argue on the fact that the FBRCs generated by PAES-RCS not only offer competitive classification performance but also provide superior interpretability and transparency than DL models, making them a valuable tool in the clinical decision-making process.

The rest of the paper is organized as follows: in Section 2 we review the most recent state-of-the-art on ML and DL models used for brain tumor image classification. In Section 3 we describe the architecture of the DL model adopted in the experimental analysis and we introduce MOEFCs. Section 4 presents a comprehensive overview of the dataset involved in the experiments and shows details on how radiomic features are extracted from MRI brain images included in the dataset that we selected for our experimental analysis. Section 5 argues on the experiments and presents a detailed description of the achieved results, focusing on accuracy metrics and model complexity. Finally, Section 6 offers some concluding remarks.

2 RELATED WORK

Over the last years, methods based on DL have been the most adopted ones for dealing with brain tumor classifications from MRI images, (Kaifi, 2023). Most of them consider Convolutional Neural Networks (CNNs), which are well known for their ability to perform automatic feature extraction (Al-Zoghby et al., 2023). When dealing with DL, authors mainly consider two general approaches, with or without a preliminary segmentation step (Muhammad et al., 2020). Recent works (Ghamry et al., 2023), (Unde and Rathore, 2024), directly adopt DL models, such as VGG16, AlexNet, ResNet50, and R-CNN. On the contrary, the authors of the works (Akter et al., 2024) and (Khan et al., 2023), first adopt a segmentation stage based on U-Net model and Fuzzy C-Means, respectively. Then, a DL model, such as VGG and Efficient-Net, is applied considering the segmented portion of the image as an input.

To ensure some degree of explainability, the authors of (Maqsood et al., 2022) and (Chmiel et al., 2023) adopted a post-hoc procedure to derive an explicator for the classification decision. Specifically, the authors considered Grad-CAM technique to identify regions of the MRI image that contribute significantly to the final prediction of different types of CNNs, such as VGG16, ResNet50, and Efficient-NetB7.

In addition to DL approaches, traditional ML models, such as decision trees and support vector machines, have also been employed for BTC (Muhammad et al., 2020). To use these models, it is necessary to manually extract features from the images. In (Decuyper et al., 2018), a CNN is used for feature extraction, which is then combined with a Random Forest classifier. In contrast, in (Cho et al., 2018), qualitative radiomic features are extracted and then used as inputs of ML classifiers such as logistic regression, support vector machine, and random forest.

Few works discuss the use of fuzzy classifiers for BTC. Specifically, in (Kalam et al., 2023) authors propose the use of the well-known Adaptive Neuro Fuzzy Inference System (ANFIS) adapted for classification tasks. Both papers leverage the ANFIS model for classifying segmented portions of the MRI. To this aim, radiomic features are extracted from the segmented images. It is worth noting that although AN-FIS belongs to the category of interpretable models by design, its level of interpretability is much lower than that of MOEFC. Indeed, frequently the integrity of the fuzzy partitions is compromised, especially due to the high overlapping of the fuzzy sets. Moreover, ANFIS considers all rules to make inferences, instead of just one as in the case of FRBCs generated with PAES-RCS. Thus, ANFIS local interpretability, namely the capability of explaining the decision taken for a specific input, may be compromised, especially if the rule base contains several rules.

3 PRELIMINARIES

3.1 Workflow of the Traditional BTC

The methodology commonly adopted in CAD systems for brain tumor classification using MRI, detailed in (Muhammad et al., 2020), involves the steps discussed in the following.

The first two steps regards the acquisition and data collection of brain images using different MRI scanning sequences. The preprocessing stage improves image quality with noise reduction and intensity correction techniques.

Segmentation is a critical step to identify regions of interest (ROIs) in the image, namely suspicious areas of brain tumor. ROIs can be detected manually by experienced radiologist or automatically by using specific algorithms or AI models.

Once the ROIs have been segmented, various types of features may be extracted. Usually quantitative values, such as radiomics features, describing morphological and geometrical aspects may be extracted from the image. When dealing with DL models feature extraction is automatically carried out by the convolutional layers. However, these features, are not easy to interpret.

Feature selection and dimensionality reduction stages involve techniques for enhancing model performance by reducing overfitting, improving computational efficiency, and highlighting the most relevant features, thereby potentially increasing the accuracy and interpretability of the model.

The classification stage may involve both supervised or unsupervised models, for categorizing the segmented image into malignant or benign lesions or to distinguish the tumor among different types or gravity levels.

Finally, the CAD system may provide a possible brain tumor diagnosis and tumor grade classification, presenting the results visually to facilitate clinical interpretation.

3.2 VGG Models for Image Classification

The DL model that we adopted in our experimental analysis is the VGG16 network. It has been designed and developed by the Visual Geometry Group (VGG) of the University of Oxford (Simonyan and Zisserman, 2014). It has been recently experimented for addressing the BTC task in (Muhammad et al., 2020). In this study, the VGG16, appropriately modified for dealing with the specific dataset, achieved the best results among other DL models in BTC tasks in segmented MRIs. It is worth noticing that in our experimental analysis, we adopted the same dataset used in (Muhammad et al., 2020) and discussed in Section 4. We built our specific BTC model from an VGG16 network pre-trained on the ImageNet dataset and publicly available in the TorchVision library³ and properly fine tuned on the selected dataset.

3.3 Multi-Objective Evolutionary Fuzzy Classifiers

Over the past decades, Multi-Objective Evolutionary Algorithms (MOEAs) have been extensively employed to design the architecture of FRBCs. The combination of MOEAs and FRBCs led to the so-called MOEFCs (Antonelli et al., 2016).

We recall that an FRBC comprises a rule base (RB), a database (DB) and an inference engine for the classification. The RB is composed by linguistic *if*-*then* rules: the antecedent part of each rule includes fuzzy conditions. These conditions, contained in the DB, are defined for each input variables by properly partitioning them with fuzzy sets. In this work, the output of the FRBC is generated by using the *max*-*imum matching* inferencing method: the rule which is fired the most by an input pattern provides the estimated class. Details on fuzzy rules and inference methods can be found in (Antonelli et al., 2016).

In our experimental analysis, we adopted the Pareto Archived Evolution Strategy (PAES) for Rule and Condition Selection (RCS) method as MOEL scheme for concurrently learning the RB and the DB of a set of FRBC. PAES-RCS generates a set of FRBC characterized by different trade-off between accuracy and complexity. PAES-RCS has been successfully experimented in (Antonelli et al., 2016) for classification tasks with tabular datasets. The adopted MOEL scheme starts from a set of candidate rules generated by using the multi-way fuzzy decision tree for classification tasks described in (Segatori et al., 2018). Once generated the initial set of candidate rules, the evolutionary process will select only the most relevant ones, along with their most relevant conditions. Simultaneously, the algorithm tunes the strong fuzzy partitions that define the DB by applying a lateral displacement of the core. The optimization is guided by two conflicting objective functions, namely the Total Rule Length (TRL) and the accuracy computed in terms of classification rate. At the end of the evolutionary process, an approximation of the Pareto Front is achieved. Details on chromosome coding, crossover and mutation operators, the scheme

³https://pytorch.org/vision/stable/index.html

of PAES-RCS, and the parameters to set for running the algorithm can be found in (Antonelli et al., 2016).

4 DATASET DESCRIPTION & FEATURE EXTRACTIONS

In our experimental analysis, we considered the "Brain Tumor Public Data Set" introduced in (Cheng et al., 2015) that includes T1-weighted and Contrast-Enhanced MRI images of 233 patients, retrieved from two different hospitals in China between 2005 and 2010. It consists of 3064 imaging sections or slices. Each image has a size of 512 x 512, a thickness from 6 to 1 mm, and a space between sections of 6 mm, and can be associated with one of the following labels: meningioma (708 images), glioma (1426 images), and pituitary tumor (939 images). The tumor area is segmented by three experienced radiologists. An example of an MRI image and tumor mask is shown in Figure 1.

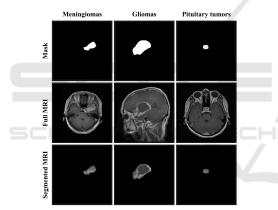


Figure 1: Images of brain tumors from the public dataset. The first row presents the segmentation masks, the second row shows the complete images, and the last row illustrates the segmented tumors.

To provide images as inputs of FRBCs, we need to transform them into numerical vectors. To this aim, we use radiomics algorithms that describes image characteristics such as pixel intensities, relationships, shapes, and textures (Bera et al., 2022).

In this work, we adopted the pipeline for extracting radiomic features proposed in (Carré et al., 2020). We used PyRadiomics⁴, a flexible open-source Python library, to extract a number of features from MRI images. Adhering to the guidelines of the Imaging Biomarker Standardization Initiative (Zwanenburg et al., 2020), PyRadiomics ensures standardization and reproducibility of radiomic features extracted from medical images. Specifically, Z-Score normalization, combined with absolute discretization, was used for the extraction of radiomic features. The extracted features include: first-order features, such as the mean, standard deviation, skewness, and kurtosis of pixel values; second-order features, such as the gray-level co-occurrence matrix, which measures the frequency of pixel pairs with specific gray values; and higher-order features, such as the gray-level run-length matrix and the gray-level size zone matrix, which assess the length of pixel sequences and the size of homogeneous gray-level zones, respectively.

A total of 110 features were extracted from each ROI of the MRI images. A decision tree-based procedure was adopted for feature selection using the same cross-validation scheme discussed in Section 5. The 15 most relevant features were selected. The selected features are described in Table 1.

5 EXPERIMENTAL ANALYSIS

5.1 Experimental Setup

In our analysis, we adopted a five-fold crossvalidation procedure. During the creation of the folds, attention was focused on two crucial aspects: each fold contains images from distinct groups of patients and images of the same patient are not included in different folds.

To handle the insufficient number of data for network model identification, we used the transfer Learning technique, discussed in (Muhammad et al., 2020): the final convolutional set of layers, the fully connected layers, and the softmax layer were finetuned, while the other layers were kept frozen. The fine-tuning was conducted by showing the network with the images of segmented regions of tumor tissue. No augmentation techniques were employed. VGG16 network fine-tuning sessions lasted 50 epochs, employing a validation mechanism to prevent overfitting and selecting the optimal set of weights. The batch size was set to 32, the maximum learning rate factor to 0.001, and the optimizer used was Adam.

When running PAES-RCS, we considered the tabular dataset of images described in the radiomic features space outlined in Section 4. We adopted a publicly available PAES-RCS implementation⁵. The values of the parameters used for running PAES-RCS are the same than the ones in (Antonelli et al., 2016). Some parameters, namely C_R and C_T , underwent a tuning procedure to achieve an optimal balance be-

⁴https://pyradiomics.readthedocs.io/en/latest

⁵https://github.com/GionatanG/skmoefs

No.	Feature Name	Description
X_1	Skewness	Measures the asymmetry of the distribution of gray levels in the image.
X_2	Maximum	Maximum pixel value in the original image.
X3	Contrast	Quantifies the contrast based on the gray level co-occurrence matrix.
X_4	Mean	Mean pixel value in the original image.
X5	Minimum	Minimum pixel value in the image.
X ₆	Small Dependence High Gray Level Emphasis	Emphasizes small dependencies with high gray levels.
X7	10th Percentile	The 10th percentile of gray levels in the image.
X ₈	Range	Range between the maximum and minimum pixel values in the image.
X9	Root Mean Squared	The square root of the mean of squared gray level values.
X10	Gray Level Non-Uniformity	Measures the non-uniformity of gray levels based on the gray level de-
		pendence matrix.
X ₁₁	Large Dependence Low Gray Level Emphasis	Emphasizes large dependencies with low gray levels.
X ₁₂	Median	Median pixel value in the image.
X ₁₃	Kurtosis	Measures the "peakedness" of the distribution of gray levels.
X ₁₄	Long Run High Gray Level Emphasis	Emphasizes long runs with high gray levels in the gray level run length
		matrix.
<i>X</i> ₁₅	Energy	Sum of squared gray level values, representing the energy of the image.

Table 1: Selected Feature Descriptions.

tween exploring and exploiting the solution space, taking into account the dataset's distinctive features for brain cancer classification.

5.2 **Results and Discussions**

As regards PAES-RCS, for each fold we run ten trials (each with a different seed of the random number generator). For each fold and each trial of the crossvalidation we generated an approximation of the optimal Pareto front. We report the average results, considering 50 trials in total, in terms of classification performance and model complexity, of three representative solutions. As discussed in (Antonelli et al., 2016), we sorted the FRBCs in each Pareto front approximation in ascending order of accuracy. Then, we extracted the First (the most accurate and the less explainable), the Median, and the Last solution (the less accurate and the most explainable).

5.2.1 Classification Performance Analysis

Table 2 presents the mean and standard deviation of the accuracy achieved by each model, along with precision, recall, and F1-score metrics for each class.

It is easy to notice that VGG16 achieves the highest average values of accuracy both on the training and test set. However, it suffers the most from overfitting.

Glioma is the tumour that is best recognised by all models. Indeed, the VGG16 model achieves an F1-score of 84% on the test set. In comparison, the First, Median, and Last FRBCs attain, respectively, 86%, 83%, and 72%. For meningiomas, VGG16 achieves an F1-score of 65%, while the First, Median, and Last FRBCs accomplish 65%, 62%, and 46%, respectively. For pituitary tumors, VGG16 attains an F1-

score of 86%, whereas the First, Median, and Last FRBCs achieve, respectively, 72%, 68%, and 50%.

In a nutshell, for Meningioma and Glioma tumors, the First and Median FRBCs perform similarly to VGG16. As regards Pituitary tumor, VGG16 outperforms all three FRBCs. However, in this class, VGG16 achieves an F1-score of 99% on the training set, which drops to 86% on the test set suggesting that the model suffers from overfitting, likely due to the underrepresentation of the Pituitary tumor class in the training data.

5.2.2 Complexity Analysis

Table 3 presents the mean and standard deviation of some complexity metrics for each model. Specifically, for all models, we show the total number of parameters (NP), the model weight in terms of memory occupancy in kBs, and the number of input variables F. As regards the FRBCs, we also show the TRL and the total number of rules in the RB (M). We recall that PAES-RCS performs also feature selection during the optimization process, thus the total number of input variables considered in the FRBCs in the Pareto front approximation may be lower than 15, i.e. lower than the number of features that we extracted and selected using the procedure described in Section 4.

The total number of parameters NP for representing an FRBC is the sum of the parameters of its DB, equal to the total number of real numbers adopted for representing all the fuzzy sets of each input variable (in our case 3), and of its RB, equal to the total number of conditions in the antecedents and the total number of class labels of each rule. Thus, the value of *NP* can be calculated as follows:

$$NP = F \times \sum_{f=1}^{F} T_f \times 3 + TRL + M \tag{1}$$

Model		Accuracy	Precision	Meningioma Recall	F1-score	Precision	Glioma Recall	F1-score	P Precision	ituitary Tumo Recall	r F1-score
FRBC-First	Train Test	$\begin{array}{c} 0.83 \pm 0.02 \\ 0.78 \pm 0.04 \end{array}$	$\left \begin{array}{c} 0.76 \pm 0.05 \\ 0.65 \pm 0.09 \end{array} \right $	$\begin{array}{c} 0.73 \pm 0.05 \\ 0.64 \pm 0.12 \end{array}$	$\begin{array}{c} 0.75 \pm 0.03 \\ 0.65 \pm 0.09 \end{array}$	$\begin{array}{c} 0.87 \pm 0.03 \\ 0.82 \pm 0.04 \end{array}$	$\begin{array}{c} 0.92 \pm 0.03 \\ 0.90 \pm 0.04 \end{array}$	$\begin{array}{c} 0.89 \pm 0.02 \\ 0.86 \pm 0.03 \end{array}$	$\begin{array}{c} 0.80 \pm 0.03 \\ 0.75 \pm 0.08 \end{array}$	$\begin{array}{c} 0.76 \pm 0.06 \\ 0.69 \pm 0.09 \end{array}$	$\begin{array}{c} 0.78 \pm 0.03 \\ 0.72 \pm 0.07 \end{array}$
FRBC-Median	Train Test	$\begin{array}{c} 0.79 \pm 0.03 \\ 0.75 \pm 0.05 \end{array}$	$\left \begin{array}{c} 0.74 \pm 0.06 \\ 0.62 \pm 0.11 \end{array} \right $	$\begin{array}{c} 0.68 \pm 0.07 \\ 0.59 \pm 0.13 \end{array}$	$\begin{array}{c} 0.70 \pm 0.04 \\ 0.62 \pm 0.10 \end{array}$	$\begin{array}{c} 0.84 \pm 0.04 \\ 0.81 \pm 0.04 \end{array}$	$\begin{array}{c} 0.90 \pm 0.03 \\ 0.87 \pm 0.07 \end{array}$	$\begin{array}{c} 0.87 \pm 0.03 \\ 0.83 \pm 0.04 \end{array}$	$\begin{array}{c} 0.77 \pm 0.05 \\ 0.72 \pm 0.11 \end{array}$	$\begin{array}{c} 0.72 \pm 0.09 \\ 0.66 \pm 0.10 \end{array}$	$\begin{array}{c} 0.74 \pm 0.05 \\ 0.68 \pm 0.08 \end{array}$
FRBC-Last	Train Test	$\begin{array}{c} 0.65 \pm 0.09 \\ 0.63 \pm 0.10 \end{array}$	$ \begin{vmatrix} 0.69 \pm 0.14 \\ 0.51 \pm 0.22 \end{vmatrix} $	$\begin{array}{c} 0.49 \pm 0.22 \\ 0.44 \pm 0.23 \end{array}$	$\begin{array}{c} 0.52 \pm 0.19 \\ 0.46 \pm 0.19 \end{array}$	$\begin{array}{c} 0.71 \pm 0.11 \\ 0.69 \pm 0.10 \end{array}$	$\begin{array}{c} 0.83 \pm 0.05 \\ 0.82 \pm 0.22 \end{array}$	$\begin{array}{c} 0.76 \pm 0.06 \\ 0.72 \pm 0.14 \end{array}$	$\begin{array}{c} 0.68 \pm 0.15 \\ 0.68 \pm 0.19 \end{array}$	$\begin{array}{c} 0.51 \pm 0.22 \\ 0.47 \pm 0.24 \end{array}$	$\begin{array}{c} 0.54 \pm 0.16 \\ 0.50 \pm 0.18 \end{array}$
VGG16	Train Test	$\begin{array}{c} 0.99 \pm 0.00 \\ 0.80 \pm 0.00 \end{array}$	$ \begin{vmatrix} 0.99 \pm 0.01 \\ 0.69 \pm 0.09 \end{vmatrix} $	$\begin{array}{c} 1.00 \pm 0.00 \\ 0.63 \pm 0.06 \end{array}$	$\begin{array}{c} 0.99 \pm 0.01 \\ 0.65 \pm 0.06 \end{array}$	$\begin{array}{c} 1.00 \pm 0.00 \\ 0.83 \pm 0.05 \end{array}$	$\begin{array}{c} 1.00 \pm 0.00 \\ 0.85 \pm 0.04 \end{array}$	$\begin{array}{c} 1.00 \pm 0.00 \\ 0.84 \pm 0.02 \end{array}$	$\begin{array}{c} 0.99 \pm 0.01 \\ 0.84 \pm 0.04 \end{array}$	$\begin{array}{c} 1.00 \pm 0.00 \\ 0.87 \pm 0.07 \end{array}$	$\begin{array}{c} 0.99 \pm 0.01 \\ 0.86 \pm 0.03 \end{array}$

Table 2: Average performance results achieved by PAES-RCS and VGG16.

Table 3: Average complexity results achieved by PAES-RCS and VGG16.

Model	NP	Weight (kB)	TRL	М	F
FRBC-First	309.42 ± 58.48	13.59 ± 0.07	125.06 ± 46.34	21.16 ± 7.43	10.88 ± 0.85
FRBC-Median	235.32 ± 70.26	11.36 ± 0.03	72.54 ± 49.67	12.48 ± 8.09	10.02 ± 1.24
FRBC-Last	138.82 ± 24.26	4.05 ± 0.03	17.02 ± 9.69	3.30 ± 1.61	7.90 ± 1.18
VGG16	$123M\pm0.0$	540471.00	-	-	16

where *F* is the number of the input variables of the FRBC, T_f is the number of fuzzy sets adopted for each input variable X_f , *TRL* is the total number of parameters considered in the RB and *M* is the number of rules in the RB.

As shown in Table 3, FRBC models have significantly fewer parameters than the VGG16 model. In particular, the average number of parameters of FRBC-First, FRBC-Median, and FRBC-Last is 300.42, 235.32, and 183.82, respectively. In contrast, VGG16 has 123 million parameters, so it has a higher complexity that can have an impact both on the computational resources required for training and inference and on the suffering from overfitting. We verified that the fine-tuning process of VGG16 takes around 26 minutes on the hardware previously discussed. In comparison, PAES-RCS takes only around 2 minutes and the radiomic feature extraction process takes around 6 minutes. Table 3 also shows that the complexity of the model in terms of NP is closely related to the weight of the model. Indeed, FRBC models are lightweight, while VGG16 has a substantially larger memory footprint making FRBC models more suitable for deployment on devices with limited memory resources.

5.2.3 Accuracy-Complexity Tradeoff Analysis

Figure 2 presents a projection in the accuracy and complexity (expressed in terms of NP) plane of the mean values associated with the FRBCs generated by PAES-RCS and the VGG16 network. Each model is depicted by a point, illustrating the trade-off between accuracy and complexity.

Using the notion of non-dominance, adopted in multi-objective optimization, none of the models dominates the others. This means that all models represent a different trade-off between the accuracy and

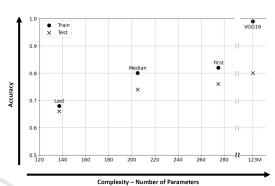


Figure 2: Performance-complexity trade-offs of the models obtained on the "Brain Tumor Public Data Set" dataset.

the complexity. It is worth to notice that, while differences in the accuracy dimension are on the same order of magnitude, the overall complexity of the FR-BCs is four orders of magnitude smaller than the one of VGG16. Moreover, on the one hand, FRBC-First demonstrates competitive classification performance compared to the VGG16, achieving an average overall accuracy of 78%, only 2 percentage points lower than the VGG16. On the other hand, the FRBC-Median represents an excellent compromise between accuracy and explainability. The TRL is reduced from 125 to 72, the number of rules from 21 to 12, and it incurs only a 4% decrease in accuracy compared to the FRBC-First. Finally, The FRBC-Last represents the most interpretable solution, reducing the TRL from 125 to 17 and the number of rules from 21 to 3, but its precision is 11 percentage points lower than the FIRST solution, resulting in an overall accuracy of 63%. This solution is particularly suitable when having a highly interpretable model is a mandatory requirement, even if it means sacrificing some accuracy.

5.2.4 Some Discussions the Explainability of Fuzzy Rules

In Fig. 3 we show some examples of fuzzy rules extracted from the RB of an FRBC-First, picked from one of the Pareto front approximations generated by PAES-RCS. The linguistic rules are formulated in terms of the optimized strong fuzzy partitions of each input variable. Each variable has been described using 5 fuzzy set labeled as follows: VL (very low), L (low), M (medium), H (high), and VH (very high). We verified that, at the end of the optimization process, the fuzzy partitions still maintain a good level of integrity in terms of ordering, coverage, and distinguishability.

It is worth to notice that the RB of the chosen FRBC-first is very compact and the rules include a reduced number of conditions. Indeed, the average rule length of the entire RB is equal to 6.0. This aspect also supports the high explainability level of the generated FRBCs. Indeed, the lower the number of conditions in each rule, the higher the local explainability of the decision associated with a specific rule. In conclusion, the extracted rules are highly interpretable and easy-to-understand by humans. In the example, R_3 , if activated, explains that an image has been classified as Meningioma because the value of the intensity of most of the pixel(information related to X_4 and X_{12}) is high (prevalence of white level) and the ROI is averagely jagged (information related to X_6 and X_{10}). In contrast, the deep neural network structure of VGG16 makes it difficult to explain individual predictions or understand the contribution of specific input variables without the a posterior application of sophisticated tools and techniques, such as layer-wise relevance propagation or saliency maps (Mandloi et al., 2024). It is worth highlighting that, as regards FRBCS generated by PAES-RCS, no extra computation, such as in the post-hoc explainability procedure adopted for DL models, is necessary. Indeed, FRBCs are interpretable by design and all the elements necessary for the explanations are already available in the DB and in the RB.

 R_1 : IF X_1 is M AND X_2 is H AND X_7 is M AND

 X_8 is L AND X_{12} is M THEN Class is *Glioma*

 R_2 : IF X_4 is H AND X_6 is L AND X_{12} is H

THEN Class is *Pituitarytumor*

 R_3 : IF X_4 is H AND X_6 is M AND X_{10} is M AND

X₁₂ is H THEN Class is Meningioma

Figure 3: Some examples of fuzzy rules extracted from an FRBC-first.

6 CONCLUSION

In this paper, we presented a comparative analysis between two different types of artificial intelligence models for approaching the BTC task from MRI images. Specifically, we considered Deep Learning models and FBRCs. We carried out an experimental campaign considering a publicly available dataset composed by MRI images including 3 different types of brain tumors. The comparison was performed along the accuracy and the complexity of the models. We considered the VGG16 convolutional network and FBRCs based on the PAES-RCS algorithm. VGG16 directly takes the MRI image in input, whereas FBRCs take a representation of the image expressed in terms of quantitative features extracted using the radiomics methodology.

Results have shown that even though VGG16 achieves the highest classification performance, it suffers from overfitting, its architecture is very complex, characterized by 123 millions of parameters, and the lack of transparency and interpretability limits its clinical applicability. In contrast, PAES-RCS has generated a set of FRBCs characterized by different trade-offs between accuracy and complexity. The most complex FRBCs, composed by hundreds of parameters and able to provide explanations in terms of simple linguistic rules, are characterized by a low loss of classification performance in comparison with VGG16.

Despite the promising results, there exist several directions for future research. In particular, it is necessary to improve the feature selection process to enhance interpretability and make the explanations more intuitive. In addition, exploring advanced data augmentation and re-balancing techniques could reduce the overfitting issues and improve the classification performance of the different classification models, particularly for the recognition of underrepresented classes.

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REFERENCES

- Akter, A., Nosheen, N., Ahmed, S., Hossain, M., Yousuf, M. A., Almoyad, M. A. A., Hasan, K. F., and Moni, M. A. (2024). Robust clinical applicable CNN and u-net based algorithm for MRI classification and segmentation for brain tumor. *Expert Systems with Applications*, 238:122347.
- Al-Zoghby, A. M., Al-Awadly, E. M. K., Moawad, A., Yehia, N., and Ebada, A. I. (2023). Dual deep

Deep Learning and Multi-Objective Evolutionary Fuzzy Classifiers: A Comparative Analysis for Brain Tumor Classification in MRI Images

CNN for tumor brain classification. *Diagnostics*, 13(12):2050.

- Antonelli, M., Ducange, P., Lazzerini, B., and Marcelloni, F. (2016). Multi-objective evolutionary design of granular rule-based classifiers. *Granular Computing*, 1:37–58.
- Bera, K., Braman, N., Gupta, A., Velcheti, V., and Madabhushi, A. (2022). Predicting cancer outcomes with radiomics and artificial intelligence in radiology. *Nature reviews Clinical oncology*, 19(2):132–146.
- Cao, J., Zhou, T., Zhi, S., Lam, S., Ren, G., Zhang, Y., Wang, Y., Dong, Y., and Cai, J. (2024). Fuzzy inference system with interpretable fuzzy rules: Advancing explainable artificial intelligence for disease diagnosis—a comprehensive review. *Information Sciences*, 662:120212.
- Carré, A., Klausner, G., Edjlali, M., Lerousseau, M., Briend-Diop, J., Sun, R., Ammari, S., Reuzé, S., Alvarez Andres, E., Estienne, T., et al. (2020). Standardization of brain MR images across machines and protocols: bridging the gap for MRI-based radiomics. *Scientific reports*, 10(1):12340.
- Cheng, J., Huang, W., Cao, S., Yang, R., Yang, W., Yun, Z., Wang, Z., and Feng, Q. (2015). Enhanced performance of brain tumor classification via tumor region augmentation and partition. *PloS one*, 10(10):e0140381.
- Chmiel, W., Kwiecień, J., and Motyka, K. (2023). Saliency map and deep learning in binary classification of brain tumours. *Sensors*, 23(9):4543.
- Cho, H.-h., Lee, S.-h., Kim, J., and Park, H. (2018). Classification of the glioma grading using radiomics analysis. *PeerJ*, 6:e5982.
- Decuyper, M., Bonte, S., and Van Holen, R. (2018). Binary glioma grading: radiomics versus pre-trained cnn features. In Medical Image Computing and Computer Assisted Intervention–MICCAI 2018: 21st International Conference, Granada, Spain, September 16-20, 2018, Proceedings, Part III 11, pages 498–505. Springer.
- Du, P., Wu, X., Liu, X., Chen, J., Chen, L., Cao, A., and Geng, D. (2023). The application of decision tree model based on clinicopathological risk factors and pre-operative mri radiomics for predicting short-term recurrence of glioblastoma after total resection: a retrospective cohort study. *American Journal of Cancer Research*, 13(8):3449.
- Ghamry, F. M., Emara, H. M., Hagag, A., El-Shafai, W., El-Banby, G. M., Dessouky, M. I., El-Fishawy, A. S., El-Hag, N. A., and El-Samie, F. E. A. (2023). Efficient algorithms for compression and classification of brain tumor images. *Journal of Optics*, 52(2):818–830.
- Hulsen, T. (2023). Explainable artificial intelligence (XAI): Concepts and challenges in healthcare. *AI*, 4(3):652–666.
- Kaifi, R. (2023). A review of recent advances in brain tumor diagnosis based on AI-based classification. *Diagnostics*, 13(18):3007.
- Kalam, R., Thomas, C., and Rahiman, M. A. (2023). Brain tumor detection in MRI images using adaptive-anfis classifier with segmentation of tumor and edema. *Soft Computing*, 27(5):2279–2297.

- Khalighi, S., Reddy, K., Midya, A., Pandav, K. B., Madabhushi, A., and Abedalthagafi, M. (2024). Artificial intelligence in neuro-oncology: advances and challenges in brain tumor diagnosis, prognosis, and precision treatment. *NPJ Precision Oncology*, 8(1):80.
- Khan, M. A., Khan, A., Alhaisoni, M., Alqahtani, A., Alsubai, S., Alharbi, M., Malik, N. A., and Damaševičius, R. (2023). Multimodal brain tumor detection and classification using deep saliency map and improved dragonfly optimization algorithm. *International Journal of Imaging Systems and Technol*ogy, 33(2):572–587.
- Mandloi, S., Zuber, M., and Gupta, R. K. (2024). An explainable brain tumor detection and classification model using deep learning and layer-wise relevance propagation. *Multimedia Tools and Applications*, 83(11):33753–33783.
- Maqsood, S., Damaševičius, R., and Maskeliūnas, R. (2022). Multi-modal brain tumor detection using deep neural network and multiclass SVM. *Medicina*, 58(8):1090.
- Muhammad, K., Khan, S., Del Ser, J., and De Albuquerque, V. H. C. (2020). Deep learning for multigrade brain tumor classification in smart healthcare systems: A prospective survey. *IEEE Transactions on Neural Networks and Learning Systems*, 32(2):507–522.
- Saidak, Z., Laville, A., Soudet, S., Sevestre, M.-A., Constans, J.-M., and Galmiche, A. (2024). An MRI radiomics approach to predict the hypercoagulable status of gliomas. *Cancers*, 16(7):1289.
- Segatori, A., Marcelloni, F., and Pedrycz, W. (2018). On distributed fuzzy decision trees for big data. *IEEE Transactions on Fuzzy Systems*, 26(1):174–192.
- Simonyan, K. and Zisserman, A. (2014). Very deep convolutional networks for large-scale image recognition. *arXiv preprint arXiv:1409.1556*.
- Unde, M. and Rathore, A. S. (2024). Brain mri image analysis for alzheimer's disease diagnosis using mask rcnn. *International Journal of Intelligent Systems and Applications in Engineering*, 12(13s):137–149.
- Van der Velden, B. H., Kuijf, H. J., Gilhuijs, K. G., and Viergever, M. A. (2022). Explainable artificial intelligence (xai) in deep learning-based medical image analysis. *Medical Image Analysis*, 79:102470.
- Wang, A. Q., Karaman, B. K., Kim, H., Rosenthal, J., Saluja, R., Young, S. I., and Sabuncu, M. R. (2024). A framework for interpretability in machine learning for medical imaging. *IEEE Access*.
- Zhang, Y., Yang, D., Lam, S., Li, B., Teng, X., Zhang, J., Zhou, T., Ma, Z., Ying, T.-C., and Cai, J. (2022). Radiomics-based detection of covid-19 from chest Xray using interpretable soft label-driven TSK fuzzy classifier. *Diagnostics*, 12(11):2613.
- Zhou, S. K., Greenspan, H., and Shen, D. (2023). Deep learning for medical image analysis. Academic Press.
- Zwanenburg, A., Vallières, M., Abdalah, M. A., Aerts, H. J., Andrearczyk, V., Apte, A., Ashrafinia, S., Bakas, S., Beukinga, R. J., Boellaard, R., et al. (2020). The image biomarker standardization initiative: standardized quantitative radiomics for high-throughput image-based phenotyping. *Radiology*, 295(2):328– 338.