Comparative Analysis of Machine Learning Methods for Breast Cancer Classification in Genetic Sequences

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1. Introduction

Cancer stands second as the cause of death worldwide. 10 million people die of cancer, the most threatening disease, every year. Some of the causes of cancer include internal factors such as genetic mutations, hormone changes, less immunity, and external factors namely eating practices and environmental changes as well as population rate. For the prediction of any disease, next-generation sequencing plays a vital role for few decades.

Machine learning and artificial intelligence have a promising future in every technological development, especially in the healthcare industry. Early detection of cancer and due strategies for preventing the disease can save many lives. For the purpose of breast cancer prognosis, the latest machine learning methods ease the prediction, prevention, and cure. Next generation sequencing using machine learning methods resumes by extraction of genetic sequences, both benign and malignant from any resource, such as the National Centre for Biotechnology Information (NCBI) or Wisconsin. Features are extracted from these DNA sequences for classification purposes. Analysis of features is done with the box method to find the outliers, histogram for data distribution, and scatter matrix for revealing the feature relationship. The distinction between benign and malignant sequences is done. Training and testing datasets are derived in the ratio 80:20. Classification is done by various traditional as well as boosting classifiers. Classification accuracy is calculated for various machine learning models, and the performance is evaluated using the F1 score. An optimal method is selected based on the accuracy of classification, and hence, the distinction between benign and malignant becomes much easier.

1.1. Related Work. A plethora of research has been carried out on cancer prognosis using various machine learning methods. It is very challenging to diagnose cancer at an early stage and thus do the needed treatment since it is a dangerous disease. Combining artificial intelligence and NGS has research scope in the diagnosis and cure of BC. Many
researchers have implemented several ML methods for making prediction easier. [1] compared several machine learning algorithms in detecting disease as well as finding metastasis. The methods were evaluated for performance with specificity, accuracy total, and ratio of likelihood. In order to differentiate between malignant and benign tumors, genetic programming techniques were applied by using [2], and the best features as well as parameters of the classifiers were selected. Decision tree and gradient boosting were applied together for the distinction between negative breast cancer and positive breast cancer, and predictive performance was conducted [3]. Gradient boosting has achieved better accuracy than the decision tree technique. Transparent breast cancer management is developed for identifying major risk components in the occurrence of BC with the decision tree as well as the neural network [4].

This random forest model is also utilized in cancer prediction with measures such as the F metric and the curve of ROC [5]. An ensemble method for breast cancer detection which was an efficient technique was conducted with two machine learning algorithms, the random forest algorithm and the gradient boosting algorithm [6]. While classifying with 12 features, the random forest algorithm achieved a classification accuracy of 74.73% and XGBoost achieved 73.63%. Nine supervised machine learning techniques including boosting algorithms were applied for breast cancer prediction by extracting 10 features from the genetic sequences of Homo sapiens, BRCA1, and BRCA2 [7]. The decision tree algorithm outperformed other models with 94.03% accuracy.

A genetic algorithm was combined with an online gradient boosting algorithm for the detection of breast cancer which was an efficient method because of its incremental way [8]. A hierarchical clustering-based random forest algorithm was used for calculating the similarity between all decision trees [9]. In order to build the hierarchical clustering random forest, the representative trees were chosen from divided clusters. Classifiers are made by a protocol using the AdaBoost algorithm, and frequently occurring breast tumor patterns were considered for disease prognosis [10]. A breast cancer classification model that combined random forest and AdaBoost algorithms to differentiate between malignant and benign data was developed [11].

1.2. System Description. Breast cancer prognosis is conducted with the help of four classifiers namely the decision tree technique, random forest as well as boosting algorithms such as AdaBoost and gradient boosting. The overall cancer prediction consists of three data retrieval, classifying data, and optimal classifier selection. Data/genetic sequences are extracted from the NCBI database in the form of FASTA files. The next step in disease prediction is classification, which consists of feature extracting, construction of machine learning models, performance evaluation as well as comparative analysis of classifiers. The final step is the best classifier selection process that is based on the accuracy of classification. The architecture diagram is depicted in Figure 1.

1.3. Data Extraction. Various normal human genetic sequences as well as cancerous sequences such as BRCA1 and BRCA2 datasets were derived as data instances in the form of FASTA files from NCBI. Though the sequences vary in their length, the average of the nucleobases was considered, and hence, the dataset reliability is conserved. A genetic sequence comprises various occurrences of nucleobases such as adenine, guanine, cytosine as well as thymine. The sequences derived vary in their length from 648 to 12386. Random sequences were selected for classification because the human genome comprises of millions of nucleobases. The resilience and stability of the DNA sequences make the work more promising than RNA sequences. DNA information is better protected and can be easily repaired compared to RNA sequences. The sequences stored in a variable are fed as input to the immediate classification phase.

1.4. Data Classification. Data classification makes use of the class or labels for forecasting an unlabelled dataset. The classification in the breast cancer prediction work consists of the extraction of features, construction of classifiers for the purpose of classification, and selection of classifiers that are optimized.

1.5. Features Extraction. The classification of benign as well as malignant breast cancer is performed with various features extracted related to breast cancer. The features derived for the purpose include the occurrence of G-quadruplex, count of ORF, GC content, class value, and mutation rate. The features were selected based on their relevance to cancer acquisition. The class value is used as the classification target that comprises values 0, 1, and 2. The occurrence of G-quadruplex and ORF contributed more to the prediction of breast cancer because it increases the probability of malignancy. The features strength was calculated using the histogram, scatter matrix as well as box plot graph. The box plot graph represents the data outliers. Outliers were identified for data using the box plot graph. Table 1 shows all 5 features along with their corresponding classes.

The extraction of features is conducted by the following algorithms.

1. G-Quadruplex Occurrence

$$\text{avg}_{G4} = \frac{C}{\text{length}(S_j)}$$

(i) Let the count of 'GGGG' be C.

(ii) Calculate the average count of G4.

C - Total count of 'GGGG' in the sequence.

Avg$_{G4}$ - Average count of 'GGGG'.

length($S_j$) - $j^{th}$ sequence length.

(2) Open Reading Frame (ORF) Measure
(i) $S_{\text{Length}} \leftarrow$ length of $\text{SeqDNA}$
(ii) initial_codon = ATG; final_codon = TAG, TAA, TGA
(iii) for $i$ varies from 1 to $S_{\text{Length}}$ Till $\text{EoS}_S \neq \text{True}$
   (a) Convert $S_i$ to string
   (b) $\text{cdn}_S \leftarrow$ Divide the sequence into 3 continuous nucleobases
   (c) initial_val$_{S_i} \leftarrow$ start codon points from $\text{cdn}_S$
   (d) final_val$_{S_i} \leftarrow$ stop codon points from $\text{cdn}_S$
   (e) $m \leftarrow \text{len}(\text{initial_val}_{S_i}); n \leftarrow \text{len}(\text{final_val}_{S_i})$
   (f) $j \leftarrow 1; k \leftarrow 1; \text{ORF}_{S_i} \leftarrow 0$
   (g) If ($j < m$) and ($k < n$)
      If (initial_val$_{S_{ij}} < \text{final_val}_{S_{ik}}$)
         $\text{ORF}_{S_i} \leftarrow \text{ORF}_{S_i} + 1$
      Else if (initial_val$_{S_{ij}} < \text{final_val}_{S_{ik}} - 1$)
         Move $j$ until (initial_val$_{S_{ij}} < \text{final_val}_{S_{ik}}$)
      (i) ORF$_{S_i} \leftarrow \text{ORF}_{S_i} + 1$
      (ii) Measure $p1$:
         $p1 = \frac{\text{Match}(\text{Seq}_i)}{\text{len(Seq}_i)} \times 100$. (3)
   (i) Calculate Mutation Rate

(3) GC- Content

$$\text{Avg}_{\text{GC}} = \frac{\text{Count}_G + \text{Count}_C}{2 \times \text{len}(S_i)}$$ (2)

Avg of GC occurrence is calculated as above
Count$_G$ - Total count of Guanine.
Count$_C$ - Total count of Cytosine.
Len($S_i$) - $i^{th}$ sequence length.

(4) Class Value

If Normal Homo sapiens Then $\text{Targ}_{S_i} = 0$
else if BRCA1 then $\text{Targ}_{S_i} = 1$
else $\text{Targ}_{S_i} = 2$
where Seq$_i$ - $i^{th}$ sequence.

(5) Mutation Rate

$$p2 = \frac{\text{Ins}(\text{Seq}_i) + \text{Del}(\text{Seq}_i)}{\text{Al.len(Seq}_i)} \times 100.$$ (4)

(i) Homo Sapien reference DNA, $R$ of nucleobase range 52861230 is extracted from NCBI.
(ii) Employ paired alignment technique “GlobalAlignment (1)” to find the align sequence length of $S_i$, no: of matches, no: of mismatches, no: of insertion and number of deletion w.r.t to the reference genome.
(iii) For Seq$_i$
(iv) Measure $p1$:
   (i) Measure $p2$:
      $$p2 = \frac{\text{Ins}(\text{Seq}_i) + \text{Del}(\text{Seq}_i)}{\text{Al.len(Seq}_i)} \times 100.$$ (4)

(i) Calculate Mutation Rate
1.6. Construction of the Machine Learning Model. Classification of breast cancer is performed by construction after the selection of features. Four classifiers such as the decision tree technique, random forest, AdaBoost algorithm as well as gradient boosting algorithm were used for differentiation between benign as well as malignant sequences, and their comparative classification performance was evaluated. For every class of sequences, 4 different sets of instances are derived ranging from 50 to 200 in groups of 50 genetic instances. Features such as G-quadruplex, count of ORF, GC content, and mutation rate are applied to all the four classifiers. These models derive the model class named from the class label. Training and testing genetic sequences are divided with an 80:20 ratio. Testing is carried out in the absence of the target value. 

1.7. Selection of the Optimal Classifier Model. The selection of an optimal model is done based on the performance metrics. Statistical methods such as classification metrics and error matrices are used for this purpose. With the help of the confusion matrix, parameters for performance measurement are calculated. The performance of classification is evaluated by calculating the F1 score, precision, recall, and support values. The accuracy of breast cancer classification can be enhanced by including more features such as copy number variations.

Among the four classifiers, the best model is chosen for efficient sequence classification. For this purpose, statistical measures such as classification measurement and error representation matrix are generated. With the help of the confusion matrix, performance measurement parameters are calculated. Based on the performance parameters, an optimal classification model is generated.

2. Results and Discussion

Three types of benign and malignant instances were extracted under categories, class 0, 1, and 2, respectively. In each class, the size of sequences ranges from 50 to 200 in groups of 50. The length of the genetic sequences greatly influences the execution time. The extraction time of all three categories of NGS sequences is given in Table 2.

Five machine learning models such as the decision tree technique, random forest, the AdaBoost algorithm as well as the gradient boosting model were made with training and testing data sequences. Training and testing datasets are following the ratio of 80:20 for the breast cancer classification process. For all 3 classes of genetic sequences, the performance of classification is represented by Table 3.

The number of classes used for cancer classification is represented by a 3×3 confusion matrix. Three classes, C1, C2, and C3 constitute the 1st, 2nd, and 3rd row/column, respectively. Testing data detected correctly in the corresponding class is denoted as the diagonal values in the matrix and is characterized as C_i. The row summation in the confusion matrix represents the sum of testing instances in every class. The 1st, 2nd, and 3rd rows’ total denote the entire instances for the test in the classes C1, C2 as well as C3, respectively.

The accuracy rate of breast cancer classification is measured as a percentage of classes correctly found and the total data tested. The accuracy of classification for all classifiers is shown in Table 4.

For the dataset sizes of 50, 150, and 200, the classification accuracy report depicts that the gradient boosting classifier has achieved a maximum accuracy of 67.50, 95.82, 90.72, and 95.39, respectively. The comparative classification accuracy of the traditional models such as the random forest learning algorithm and decision tree technique as well as boosting algorithms such as AdaBoost and gradient boosting is shown in Figure 2.

The classification performance is measured with parameters of performance measurement. Table 5 represent the performance parameters of gradient boosting.

The above table shows that the F1 score of the gradient boosting model is .95, the same as the accuracy value of the corresponding model calculated using the confusion matrix. Hence, the gradient boosting model has performed better than all the other three models. The inference clearly shows...
that the boosting model could perform better than traditional classifiers.

3. Conclusion

Since the real causes of breast cancer are still unclear and vary from person to person, the prediction and diagnosis of breast cancer are complex. In our research, various genetic sequences, namely, benign human sequences and BRCA1 as well as BRCA2 as three classes are extracted from the NCBI data repository, and classification between benign and malignant data was performed. From all three classes, the datasets were categorized as groups of 50 DNA sequences ranging from 50 to 200, totalling 2640 sequences. Four classifiers namely the decision tree technique, random forest, and the AdaBoost model as well as the gradient boosting model were constructed with five features relevant to cancer and compared based on classification accuracy. Gradient boosting outperformed all three models and was selected as the optimal model with a classification accuracy of 95% for the distinction of datasets. For the prediction of COVID-19, the work could be extended where extraction of RNA sequence features could be used for classification purposes.

Data Availability

All the required data used to support the findings of the study are available within the article.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

References

[1] L. Tapak, N. Shirmohammadi-Khorram, P. Amini, B. Alafchi, O. Hamidi, and J. Poorolajal, “Prediction of survival and metastasis in breast cancer patients using machine learning classifiers,” *Clinical Epidemiology and Global Health*, vol. 7, no. 3, pp. 293–299, 2019.

[2] H. Dhahri, E. Al Maghayreh, A. Mahmood, W. Elkilani, and M. Faisal Nagi, “Automated breast cancer diagnosis based on machine learning algorithms,” *Journal of healthcare engineering*, vol. 2019, pp. 1–11, Article ID 4253641, 2019.

[3] T. A. Assegie, R. L. Tulasi, and N. K. Kumar, “Breast cancer prediction model with decision tree and adaptive boosting,” *IAES International Journal of Artificial Intelligence*, vol. 10, no. 1, p. 184, 2021.

[4] A.K. Verma, M. Chakraborty, and S. K. Biswas, “Breast cancer management system using decision tree and neural network,” *SN Computer Science*, vol. 2, no. 3, pp. 1–15, 2021.

[5] Y. Li and Z. Chen, “Performance evaluation of machine learning methods for breast cancer prediction,” *Applied and Computational Mathematics*, vol. 7, no. 4, pp. 212–216, 2018.

[6] S. Kabiraj, M. Raihan, N. Alvi et al., “Breast cancer risk prediction using XGBoost and random forest algorithm,” in *Proceedings of the 2020 11th International Conference on Computing, Communication and Networking Technologies (ICCCNT)*, pp. 1–4, IEEE, Kharagpur, India, July 2020.

[7] B. Kuriian and V. L. Jyothi, “Breast cancer prediction using an optimal machine learning technique for next generation sequences,” *Concurrent Engineering*, vol. 29, no. 1, pp. 49–57, 2021.

[8] H. Lu, H. Wang, and S. W. Yoon, “A dynamic gradient boosting machine using genetic optimizer for practical breast cancer prognosis,” *Expert Systems with Applications*, vol. 116, pp. 340–350, 2019.

[9] Z. Huang and D. Chen, “A breast cancer diagnosis method based on VIM feature selection and hierarchical clustering
random forest algorithm,” *IEEE Access*, vol. 10, pp. 3284–3293, 2022.

[10] Q. Huang, Y. Chen, L. Liu, D. Tao, and X Li, “On combining biclustering mining and AdaBoost for breast tumor classification,” *IEEE Transactions on Knowledge and Data Engineering*, vol. 32, no. 4, pp. 728–738, 2020.

[11] D. Yifan, L. Jialin, and F. Boxi, “Forecast model of breast cancer diagnosis based on RF-AdaBoost,” in *Proceedings of the 2021 International Conference on Communications, Information System And Computer Engineering (CISCE)*, pp. 716–719, IEEE, Beijing, China, May 2021.