



Artificial Intelligence Enabled Decision Support System on E-Healthcare Environment

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Abstract: In today's digital era, e-healthcare systems exploit digital technologies and telecommunication devices such as mobile devices, computers and the internet to provide high-quality healthcare services. E-healthcare decision support systems have been developed to optimize the healthcare services and enhance a patient's health. These systems enable rapid access to the specialized healthcare services via reliable information, retrieved from the cases or the patient histories. This phenomenon reduces the time taken by the patients to physically visit the healthcare institutions. In the current research work, a new Shuffled Frog Leap Optimizer with Deep Learning-based Decision Support System (SFLODL-DSS) is designed for the diagnosis of the Cardiovascular Diseases (CVD). The aim of the proposed model is to identify and classify the cardiovascular diseases. The proposed SFLODL-DSS technique primarily incorporates the SFLO-based Feature Selection (SFLO-FS) approach for feature subset election. For the purpose of classification, the Autoencoder with Gated Recurrent Unit (AEGRU) model is exploited. Finally, the Bacterial Foraging Optimization (BFO) algorithm is employed to fine-tune the hyperparameters involved in the AEGRU method. To demonstrate the enhanced performance of the proposed SFLODL-DSS technique, a series of simulations was conducted. The simulation outcomes established the superiority of the proposed SFLODL-DSS technique as it achieved the highest accuracy of 98.36%. Thus, the proposed SFLODL-DSS technique can be exploited as a proficient tool in the future for the detection and classification of CVD.

Keywords: E-healthcare; decision support system; cardiovascular disease; feature selection; deep learning

1 Introduction

The prevalence of non-communicable chronic diseases has drastically increased across the globe since the lifestyles of people have changed dramatically in the past few years due to technological advancements [1]. According to the World Health Organization (WHO), chronic obstructive lung illness, Ischemic Heart



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Disease (IHD), lower respiratory infection, and stroke are the top most diseases that contribute to a high mortality rate over the past few decades in both developed as well as the developing countries [2]. In recent times, Artificial Intelligence (AI) techniques have been widely applied in the healthcare industry, especially in the following domains such as prevention, diagnosis, medical payment systems, treatment and so on [3]. Mobile internet networks, big data and other innovative information technologies are also leveraged in healthcare-based AI applications. Thus, the AI-related Medical (M)-healthcare and electronic (E)-healthcare solutions have gained much attention to obtain information, develop processes and offer a clear output for different types of audiences such as the caregiver, physician and patients. AI-related M-healthcare and E-healthcare solutions are also helpful in achieving the objectives and the needs of medical treatments in an efficient manner [4]. Being a data-driven technology, the AI technique brought a paradigm shift from information domain to application domain in intelligent treatment and diagnosis, intelligent payment, etc. [5]. In the domain of Information and Communication Technology (ICT), several authors have investigated the applications of AI in healthcare industry.

Cardiovascular Disease (CVD) is one of the critical health problems with a high mortality rate in both developing as well as developed countries [6]. Some typical symptoms of CVD include body weakness, swollen feet and shortness of breath. In literature, various researchers attempted to develop an effective method that can diagnose cardiac disease at its early stages. This is because the existing diagnostic methods for heart diseases are either inaccurate or not fast enough to predict the onset of the disease or its progression [7]. Both prognosis as well as the treatment of cardiac diseases are highly challenging to accomplish, especially in the absence of modern technology and highly-experienced medical specialists. Some common reasons behind the occurrence of CVD include the intake of high-calorie diet, sugars, saturated fats and physical inactivity. Further, these causes are linked with atherosclerosis development and other metabolic disturbances, too, like hypertension, metabolic syndrome and Diabetes Mellitus (DM). CVD patients mostly suffer from the above-mentioned conditions over the course of their life [8]. Machine Learning (ML) prediction methods require proper data for both trainings as well as testing purposes. The performance of ML methods can be improved only if a balanced data set is used for training and testing purposes. Moreover, the prediction abilities of a model can be enhanced further with the help of the appropriate features relevant to the data [9,10]. Hence, Feature Selection (FS) and the data balancing processes are crucial phases that enhance the performance of a model.

In the current study, a new Shuffled Frog Leap Optimizer with a Deep Learning-based Decision Support System (SFLODL-DSS) is designed to diagnose Cardiovascular Disease (CVD). The proposed SFLODL-DSS technique primarily incorporates SFLO-based Feature Selection (SFLO-FS) approach for feature subset election. For the purpose of classification, the Autoencoder with Gated Recurrent Unit (AEGRU) model is exploited. Finally, the Bacterial Foraging Optimization (BFO) algorithm is employed to finetune the hyperparameters involved in the AEGRU technique. To demonstrate the enhanced performance of the proposed SFLODL-DSS technique, a series of simulations was conducted.

2 Literature Review

In the study conducted earlier [11], five selection strategies were implemented for multi-label active learning. These strategies were utilized to mitigate the labelling costs via an iterative selection of the most appropriate data and by querying the labels. The hyperparameters of the feature selection techniques, with ranks for each label, were maximized with the help of the grid search. This was accomplished to execute the prediction models for every scenario in the heart disease dataset. In literature [12], an intelligent healthcare structure was devised to predict the occurrence of heart diseases based on the Swarm-Artificial Neural Network (Swarm-ANN) approach. The presented Swarm-ANN approach arbitrarily produced a set

of pre-defined Neural Network (NN) numbers to evaluate and train the structures related to the solution's consistency. At last, the weight of the neuron got altered by sharing the global optimum weight with the rest of the neurons. Then, the accuracy of the heart disease prediction was estimated.

In literature [13], the researchers compared the performance of the conventional mechanisms against the presented system that predicts cardiac disease with the help of traditional ML classifier. The mechanism modelled in this study was helpful in fine-tuning the hyperparameters for the five classification methods considered in this study, with the help of grid search method. The authors in literature [14] introduced an ML-related prediction method for the prediction of multiple-and binary-classification heart disease datasets concurrently. Initially, the researchers devised a Fuzzy-GBDT approach by integrating Gradient Boosting Decision Tree (GBDT) and Fuzzy Logic (FL) techniques to reduce the complexities in the data and increase the generalization of the binary classifier's prediction. Then, the Fuzzy-GBDT was compiled with bagging to avoid the issue of overfitting.

A web-related decision support structure was presented in the study conducted earlier [15]. This structure was able to generate a pre-guidance report based on the decisions taken from the Bayesian network analysis results on disease dataset. Further, the report was produced in adherence to the mined disease paradigms over medical and non-medical factors of the patients. The mined disease paradigms were retrieved from the patient's past medical reports. The outcomes of the report inferred the likelihood of getting affected by a disease for the provided health metrics. In the study conducted earlier [16], a comparison was made among various computational intelligence methods in terms of heart disease identification. In this study, two computational intelligence approaches such as the k-Nearest Neighbour (KNN) and Decision Tree (DT), were compared and contrasted. Further, the Autoencoder (AE) feature extraction method was used in this study to reduce the number of attributes required to describe the heart disease dataset.

3 The Proposed Model

An intelligent SFLODL-DSS technique is designed in this study to diagnose the CVD. The proposed SFLODL-DSS technique primarily incorporates the SFLO approach for feature subset election. For the purpose of classification, the AEGRU model is exploited. Finally, the BFO algorithm is employed to fine-tune the hyperparameters involved in the AEGRU technique. The overall processes involved in the proposed model are shown in Fig. 1.



Figure 1: Working process of the SFLODL-DSS method

3.1 Data Pre-Processing

In this study, the Min-Max scaler is exploited to scale the dataset in the range of [0,1]. This range guarantees fast convergence for gradient learning techniques and is formulated using the following equation.

$$f_{scaled} = \frac{f - f_{\min}}{f_{\max} - f_{\min}} \tag{1}$$

Here, f_{\min} and f_{\max} represent the lower and higher limits of the features, correspondingly.

3.2 Processes Involved in Feature Selection Technique

In this stage, the SFLODL-DSS technique primarily incorporates the SFLO approach for feature subset election [17]. SFLO, a memetic metaheuristic technique, is generally applied to find a global solution by executing an informed heuristic search via a heuristic function. The SFLO technique is a populationbased algorithm inspired from frogs of similar characteristics. In this technique, all the frogs are regarded as the solutions. The overall population of the frogs is categorized under several subclasses called 'memeplex'. Further, different subcategories are appreciated as disparate frog memes. The whole set of the memeplexes is held accountable in case of a constrained exploration. In every memeplex, the frogs tend to affect the activities of the other frogs, whereas the memeplexes are forced to combine. This phenomenon results in the generation of novel memeplexes due to the shuffling technique. The shuffling procedure promotes unbiased traditional progress in the direction of a certain interest. The end condition is met when the shuffling procedure and the local search alternate are chosen.

- (1) This process contains a population 'p' that corresponds to a possible number of solutions and is constrained by a collection of virtual frogs (n).
- (2) The population is divided into subsections known as 'memeplexes' (m). Here, a memeplex is regarded as a group of the corresponding frog cultures that try to accomplish certain objectives.
- (3) Frog *i* is demonstrated as $X_i = (X_{i1}, X_{i2}, \ldots, X_{is})$ in which the value *S* corresponds to the number of variables.
- (4) Within every memeplex, the frog culture searches for a space in dissimilar directions and independently exchanges ideas. The frog with the worst fitness value is represented by X_b , whereas the frog with the best fitness value is denoted by X_w .
- (5) The frog with the global optimum fitness is recognized as X_g .
- (6) The frog with the worst fitness value is determined as follows.

$$D_i = rand \left(X_b - X_w \right) \tag{2}$$

$$X_{neww} = X_{oldW} + D_i \left(-D_{\max} \le D_i \le D_{\max} \right) \tag{3}$$

Now, the rand function produces an arbitrary value in the range of [0, 1], while D_i denotes the leaping step size of the frog *i* and D_{max} shows the maximal value allowed if the location of the frog is altered. Once the fitness value X_w becomes highly efficient than the present value of X_w , it is accepted. If the fitness is not adapted, then the computation is iterated by replacing X_b with X_g . In case there is no prospect for improvement, X_w is arbitrarily generated. Both local searches as well as shuffling processes continue until the convergence conditions produce suitable outcomes [68, 70, 71].

In FS problem, all the solutions are constrained to binary numbers such as 0 and 1. In the SFLO application algorithm, the SFLO-FS approach is developed by originating a binary form that was determined earlier. In the SFLO-FS technique, a solution is described through 1D vector, whereas the length of the vector depends on the count of the features in the new dataset. All the cells in a vector have

separate values, i.e., either 1 or 0. The value '1' signifies that the resultant features are carefully chosen; otherwise, the value is defined by 0.

$$Z_{mn} = \begin{cases} 1 & \text{if } X_{mn} > 0.5 \\ 0 & \text{otherwise} \end{cases}$$
(4)

Here, Z_{mn} represents a dissimilar form of the solution vector X, whereas X_{mn} specifies the continuous position of the searching agent *m* at dimension *n*.

The FS process is modelled as a multi-objective optimization problem, in which two conflicting objectives are fulfilled, such as high classification performance and low FS quantity. Here, the classification performance is applied as a Fitness Function (FF) value to assess the efficacy of the entire searching agent. To create a balance between classification performance and the number of FS for every solution, the FF value is applied in both SFLO algorithm as well as the whole approach to determine the searching agents.

$$Fitness = \rho Err(D) + \varphi \frac{|F|}{|T|}$$
(5)

In Eq. (5), Err (D) signifies the classification error rate, ρ and φ are constants that are used to control the reduction feature and classification accuracy respectively, |F| characterizes the size of the known feature set and |T| symbolizes the overall quantity of the features.

3.3 Classification Using Optimal AEGRU Model

In current study, the AEGRU model is exploited for the purpose of classification. The GRU model is a specific case of Long Short Term Memory (LSTM) model and is established to reduce the extended training time taken by LSTM [18]. In comparison with LSTM, the GRU model is too simple since it encompasses only two gates as update gate and the reset gate that control the flow of the data inside the unit. The modification function amongst the GRU neurons is given herewith.

$$r(n) = \sigma(w_r x(n) + u_r h(n-1) + b_r) \tag{6}$$

$$z(n) = \sigma(w_z x(n) + u_z h(n-1) + b_z) \tag{7}$$

$$\hat{h}(n) = \sigma(w_h x(n) + u_h(r(n)^* h(n-1)) + b_h)$$
(8)

$$h(n) = (1 - z(n))^* h(n-1) + z(n)^* \hat{h}(n)$$
(9)

Now r(n) indicates the reset gate, z(n) represents the update gate and w and u imply the variable matrices of the GRU model. Further, h(n), h(n), and b correspondingly denote the candidate output, output and bias [18]. The activation function is denoted by σ .

The structure of the GRU model is shown in Fig. 2. The suggested AEGRU technique is similar to AE with LSTM (AELSTM) method. Here, the AE technique is leveraged to shrink the medical data by describing the architecture of the information and attaching the encoder information for GRU network. The encoders of the medical data and its past data are given to GRU model for the purpose of training and to make the neuron fit into the model. This phenomenon results in the prediction of the desired output. Particularly, the AEGRU methodology is trained using a collection of past encoder medical datasets. In this final stage, the BFO algorithm is employed for optimal fine-tuning of the hyperparameters involved in the AEGRU approach. The conventional BFO method is initialized as two major phases that are briefed herewith.



Figure 2: GRU model

- 1) Initiation of the solution space: The spatial dimension D, range and the mapping function f(x) are employed.
- 2) Initiation of the bacteria: The bacterial count is designated as S. The position of the *i-th* bacterium in optimization space is specified as $P_i(j, k, l)$, which corresponds to the finest parameter of the solution, i.e., $P_i(j, k, l) = [m_1, m_2, \dots, m_D]$.

Therefore, the fitness of the *i*-th bacterium in optimization space is specified as $J_i(j, k, l)$ and is determined by the function of bacterium's location.

$$J_i(j, k, l) = f(P_i(j, k, l)) = f_{ij,k,l}(m_1, m_2, \dots, m_D).$$
(10)

Now, if the function has a low value, it denotes high fitness. *i* represents the *i*-th bacterium, whereas j, k, and l are related to the centralized method of the BFO technique in terms of reproduction, dispersal, elimination, and chemotaxis, respectively.

Chemotaxis

The Chemotaxis process comprises a massive number of flipping and swimming motions [19]. In *j*-th chemotaxis technique, the motion of the *i*-th bacterium is represented by Eq. (11).

$$P_i(j+1, k, l) = P_i(j, k, l) + \frac{\Delta(i)}{\sqrt{\Delta^T(i)\Delta(i)}}C(i)n,$$
(11)

The swimming step length of the *i*-th bacterium is classified based on the amount of swimming n and a single swimming step size, C(i). Here, (i) denotes the vector direction of the *i*-th bacterium in p dimensional optimization space. Each component of (i) is a mathematical value in the range of [-1, 1] in which initiation is set as an arbitrary number. When the *i*-th bacterium distinguishes the highest fitness position from a favourable environment in *j*-th chemotaxis, it moves in the same direction according to the time. Instead, (i) selects a novel arbitrary direction.

Swarming

The bacteria are classified into attractive and repulsive bacteria, for which the numerical relationship is established as follows.

$$J_{cc}(P_i) = \sum_{i=1}^{s} \left[-d_{att} \exp\left(-\omega_{att} \sum_{m=1}^{p} \left(P_{i,m} - \overline{P_m}\right)^2\right) \right] + \sum_{i=1}^{s} \left[h_{rep} \exp\left(-\omega_{rep} \sum_{m=1}^{p} \left(P_{i,m} - \overline{P_m}\right)\right) \right],$$
(12)

In Eq. (3), d_{att} signifies the depth at which the attracted material is produced by the *i-th* bacterium and ω_{att} characterizes the width of the attracted material. Since two bacteria cannot co-exist in parallel positions, the repulsion is adopted by both h_{rep} and ω_{rep} . Subsequently, the swarming process and the fitness of the *i-th* bacterium are shown below.

$$J_i(j+1, k, l) = J_i(j, k, l) + J_{cc}(P_i(j, k, l))$$
(13)

Reproduction

The bacteria tend to replicate once it accomplishes an optimal environment; otherwise, it dies. Consequently, after swarming and chemotaxis techniques, the fitness of the whole bacteria is calculated and sorted as defined below.

$$J_{i,health} = \sum_{j=1}^{N_c} J_i(j, k, l).$$
(14)

Here, one half of the bacteria reaches the optimal state $S_r = \left(\frac{S}{2}\right)$ and are selected to survive, whereas the rest of the bacteria are left to die. The bacteria that survive this environment reproduce via two colonies that are located in the corresponding regions and retain the total number of bacteria i.e., S set.

Elimination and Dispersal

After reproduction, each bacterium is disseminated through the possibility of P_{ed} , though the total amount of bacteria remains unchanged. As soon as a bacterium gets detached, it is distributed arbitrarily toward the original location.

$$r = random [0, 1];$$

$$P_{i}(j, k, l) = \begin{cases} P_{i}(j, k, l) & r > P_{ed}, \\ m'_{1}, m'_{2}, \dots m'_{p} & r < P_{ed}. \end{cases}$$
(15)

Now, the removal process is followed when $r_i < P_{ed}$. The novel position of the *i*-th bacterium P_i is replaced using an original position, $P'_i = (m'_1, m'_2, \ldots, m'_D)$. As a result, the *m* optimal parameter is upgraded towards an *m'* arbitrary parameter since it can be solved during optimization phase.

4 Experimental Validation

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[n 1]

In this section, the performance of the proposed SFLODL-DSS method was experimentally validated using a dataset sourced from UCI repository. The dataset holds 303 samples, and the proposed model selected seven features, as shown in Table 1.

Table	1:	Dataset	detai	ls
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Class	No. of samples (attributes = 303)
Absence	164
Presence	139
Total no. of samples	303

Fig. 3 shows the confusion matrices generated by the proposed SFLODL-DSS method using distinct Training (TR) and Testing (TS) datasets. With 80% of TR data, the proposed SFLODL-DSS method classified 123 and 105 samples under absence and presence classes, respectively. Moreover, with 20% of

TS data, the SFLODL-DSS method segregated 32 and 28 samples under absence and presence, respectively. In parallel, with 70% of TR data, the proposed SFLODL-DSS method categorized 116 and 82 samples under absence and presence classes correspondingly. Similarly, with 30% of TS data, the SFLODL-DSS method recognized 48 and 41 samples as absence and presence classes, correspondingly.



Figure 3: Confusion matrix of SFLODL-DSS method

Table 2 and Fig. 4 provide the overall classification results accomplished by the proposed SFLODL-DSS method on the applied dataset. With 80% of TR data, the proposed SFLODL-DSS method achieved average $accu_y$, $prec_n$, $reca_l$, F_{score} and $G_{measure}$ values such as 94.21%, 94.10%, 94.32%, 94.18% and 94.20% respectively. At the same time, with 20% of TS data, the proposed SFLODL-DSS method gained average $accu_y$, $prec_n$, $reca_l$, F_{score} and $G_{measure}$ values such as 98.36%, 98.48%, 98.28%, 98.35% and 98.37% correspondingly. In addition, with 70% of TR data, the SFLODL-DSS algorithm attained average $accu_y$, $prec_n$, $reca_l$, F_{score} and $G_{measure}$ values such as 93.40%, 94.62%, 92.71%, 93.22% and 93.44% correspondingly. At last, with 30% of TS data, the proposed SFLODL-DSS approach acquired average $accu_y$, $prec_n$, $reca_l$, F_{score} and $G_{measure}$ values such as 97.80%, 98%, 97.67%, 97.79% and 97.81% correspondingly.

Labels	Accuracy	Precision	Recall	F-Score	G-measure
Training phase	(80%)				
Absence	94.21	96.09	93.18	94.62	94.63
Presence	94.21	92.11	95.45	93.75	93.76
Average	94.21	94.10	94.32	94.18	94.20
Testing phase (20%)					
Absence	98.36	96.97	100.00	98.46	98.47
Presence	98.36	100.00	96.55	98.25	98.26
Average	98.36	98.48	98.28	98.35	98.37
Training phase (70%)					
Absence	93.40	89.23	100.00	94.31	94.46
Presence	93.40	100.00	85.42	92.13	92.42
Average	93.40	94.62	92.71	93.22	93.44
Testing phase (30%)					
Absence	97.80	96.00	100.00	97.96	97.98
Presence	97.80	100.00	95.35	97.62	97.65
Average	97.80	98.00	97.67	97.79	97.81

Table 2: Overall classification outcomes of SFLODL-DSS method



Figure 4: Average classification results of the SFLODL-DSS method

Both Training Accuracy (TA) and Validation Accuracy (VA) values, obtained by the proposed SFLODL-DSS method on test dataset, are demonstrated in Fig. 5. The experimental outcomes imply that the proposed SFLODL-DSS approach attained the maximal TA and VA values whereas the VA values were higher than the TA values.



Figure 5: TA and VA analyses results of the SFLODL-DSS method

Both Training Loss (TL) and Validation Loss (VL) values, gained by the proposed SFLODL-DSS method on test dataset, are shown in Fig. 6. The experimental outcomes implicitly explain that the proposed SFLODL-DSS algorithm exhibited the minimal TL and VL values whereas the VL values were lower than the TL values.



Figure 6: TL and VL analyses results of the SFLODL-DSS method

A clear precision-recall study was conducted upon the proposed SFLODL-DSS method using the test dataset and the results are shown in Fig. 7. The figure signifies that the proposed SFLODL-DSS method produced enhanced precision-recall values under all the classes. Fig. 8 shows the detailed ROC analysis results achieved by the presented SFLODL-DSS algorithm on test dataset. The results infer that the proposed SFLODL-DSS method exhibited its supreme capability in terms of categorizing the test dataset under distinct classes.



Figure 7: Precision-recall analyses results of the SFLODL-DSS method



Figure 8: ROC analysis results of the SFLODL-DSS method

Table 3 and Fig. 9 show the comprehensive comparison analysis outcomes accomplished by the proposed SFLODL-DSS model and other recent models such as Random Forest (RF), Bidirectional Long Short Term Memory (BiLSTM), Inception v3, Extreme Gradient Boosting (XGBoost), Deep Belief Network (DBN), Support Vector Classification (SVC) and Iterative Dichotomiser 3 (ID3). The experimentation outcomes showcase the overall improvements achieved by the proposed SFLODL-DSS model. With respect to $accu_v$, the proposed SFLODL-DSS model offered a maximum $accu_v$ of 98.36%, whereas RF, BiLSTM, Inception v3, XGBoost, DBN, SVC and ID2 models reported low $accu_v$ values such as 95.52%, 96.38%, 95.78%, 94.19%, 94.28%, 94.00% and 96.53% respectively. Also, with respect to $prec_n$, the proposed SFLODL-DSS model rendered a maximum $prec_n$ of 98.48%, whereas other models such as RF, BiLSTM, Inception v3, XGBoost, DBN, SVC and ID2 reported the least $prec_n$ values such as 96.11%, 96.04%, 94.57%, 94.87%, 96.26%, 96.57% and 95.39% correspondingly. In contrast to these, with respect to reca₁, the proposed SFLODL-DSS model accomplished a maximum recal of 98.28%, whereas RF, BiLSTM, Inception v3, XGBoost, DBN, SVC and ID2 models reported the least recal values such as 97.52%, 96.15%, 94.70%, 97.32%, 95.59%, 97.86% and 97.81% correspondingly. Moreover, with respect to F_{score} , the SFLODL-DSS model attained a maximum F_{score} of 98.35%, whereas RF, BiLSTM, Inception v3, XGBoost, DBN, SVC and ID2 models achieved the least F_{score} values such as 97.04%, 95.43%, 94.88%, 94.43%, 96.67%, 95.10% and 96.46% correspondingly.

Methods	Accuracy	Precision	Recall	F-Score
SFLODL-DSS	98.36	98.48	98.28	98.35
RF Algorithm	95.52	96.11	97.52	97.04
Bi-LSTM Model	96.38	96.04	96.15	95.43
Inception-V3 algorithm	95.78	94.57	94.70	94.88
XGBoost	94.19	94.87	97.32	94.43
DBN model	94.28	96.26	95.59	96.67
SVC model	94.00	96.57	97.86	95.10
ID3 algorithm	96.53	95.39	94.81	96.46

Table 3: Classification outcomes of the proposed SFLODL-DSS method and other existing models



Figure 9: Comparative classification results of the proposed SFLODL-DSS method

Finally, the SFLODL-DSS model was compared with existing models in terms of Execution Time (EXET) and the results are shown in Table 4 and Fig. 10. The experimental values imply that the rest of the models such as RF, XGBoost and DBN demanded high EXET such as 2.230, 2.040 and 2.100 s respectively.

Methods	Execution time (sec)
SFLODL-DSS	0.012
RF algorithm	2.230
Bi-LSTM model	0.560
Inception-V3 algorithm	1.660
XGBoost	2.040
DBN model	2.100
SVC model	1.990
ID3 algorithm	1.770

Table 4: EXET of the proposed SFLODL-DSS method and other existing models



Figure 10: Comparative EXET results of SFLODL-DSS method

However, the Inception v3 and SVC models required slightly lesser EXET values such as 1.660 and 1.990 s respectively. Though the BiLSTM model demanded a reasonable EXET of 0.560 s, the proposed SFLODL-DSS model achieved a superior outcome with the least EXET of 0.012 s. Based on the analytical results and discussions made above, it is evident that the proposed SFLODL-DSS model is an excellent performer than the existing models.

5 Conclusion

In this article, an intelligent SFLODL-DSS has been designed for the diagnosis of Cardiovascular Diseases. The proposed SFLODL-DSS technique primarily incorporates the SFLO approach for feature subset election. For the purpose of classification, the AEGRU model is exploited. At the final stage, the BFO algorithm is employed for optimal fine-tuning of the hyperparameters related to the AEGRU approach. To demonstrate the enhanced performance of the proposed SFLODL-DSS technique, a series of simulations was conducted and the results established the superiority of the proposed SFLODL-DSS technique over other techniques. Thus, the SFLODL-DSS technique can be exploited as a proficient tool in the future for detection and the classification of CVD. Further, the performance of the SFLODL-DSS technique can be improved in the future using outlier detection models.

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