

based on the analogy between the simulation of the annealing of solids and the problem of solving large combinatorial optimization problems. In this work the hybridized GA-SA perform optimization to find both the most significant feature sub set and the kernel parameters of SVM.

3 Datasets Used

The proposed method includes four common diseases, namely diabetes, breast cancer, hepatitis and cardiac arrhythmia. All the data sets are available in the UCI machine learning repository.

The Pima Indian Diabetes (PID) dataset ,there are 268 instances in class '1' and 500 instances in class '0', where '1' means a positive test for diabetes and '0' is a negative test for diabetes. The total number of attributes is 8.

In the Wisconsin Breast Cancer Dataset (WBCD), there are 569 instances in which 357 cases are benign and 212 cases are malignant. Total number of attributes is 32.

In the hepatitis disease dataset, the presence or absence of hepatitis disease is predicted by using the results of various medical tests carried out on a patient. Hepatitis dataset contains 155 instances belonging to two different classes, 'die' with 32 instances and 'live' with 123 instances. The total number of attributes is 19.

In the arrhythmia dataset, there are 452 ECG records described by 279 attributes, out of which, 206 attributes are linear and the remaining 73 attributes are nominal. The first four attributes are age, sex, height and weight. The rest of the attributes are extracted from patients' ECG signals.

4 Proposed Scheme

The overall process of the proposed system is depicted in fig.1 which contains Data preprocessing, feature selection, classification and optimization.

4.1 Data Preprocessing

Data preprocessing is a data mining strategy that transforms raw data into a clear format. Real-world data is often incomplete, inconsistent, and/or lacking in certain behaviors or trends, and is likely

to contain numerous errors. By using mean algorithm, the incomplete data is filled.

4.2 Feature Selection

In feature selection, a subset of the original variables is discovered with mean to reduce and dispose the noise dimension. The principle behind feature selection is to pick a subset of input variables by taking out features with little or no predictive information. It transforms high-dimensional data into lower dimensions. When the input to an algorithm is too large to be processed and is suspected to be extremely redundant (much data, but not much information), then the input data is transformed into a reduced representation set of features.

Dimensionality reduction can be achieved by either taking out data that is nearly related to other data in the set, or joining data to make a small set of features [12]. In machine learning, the issue of supervised classification is concerned with utilizing labeled samples to impel a model that classifies objects into a limited set of known classes. The samples are described by a vector of numeric or nominal features. Some of these features may be unimportant or redundant. Avoiding unimportant or redundant features is essential because that they may have a negative impact on the accuracy of the classifier.

Selection of the most significant subset of features is an optimization problem. In this system, the feature selection is done using hybrid GA-SA local search mechanism which is explained in section 4.4.

4.3 Classification

Classification is a data mining function that assigns items in a collection to target classes. The objective of classification is to accurately predict the target class for each case in the data. Classification models are tested by comparing the predicted values to known target values in a set of test data. The historical data for a classification project is typically divided into two sets: one for building the model (training) and the other for testing the model. SVM is a class of machine learning algorithms that can perform pattern

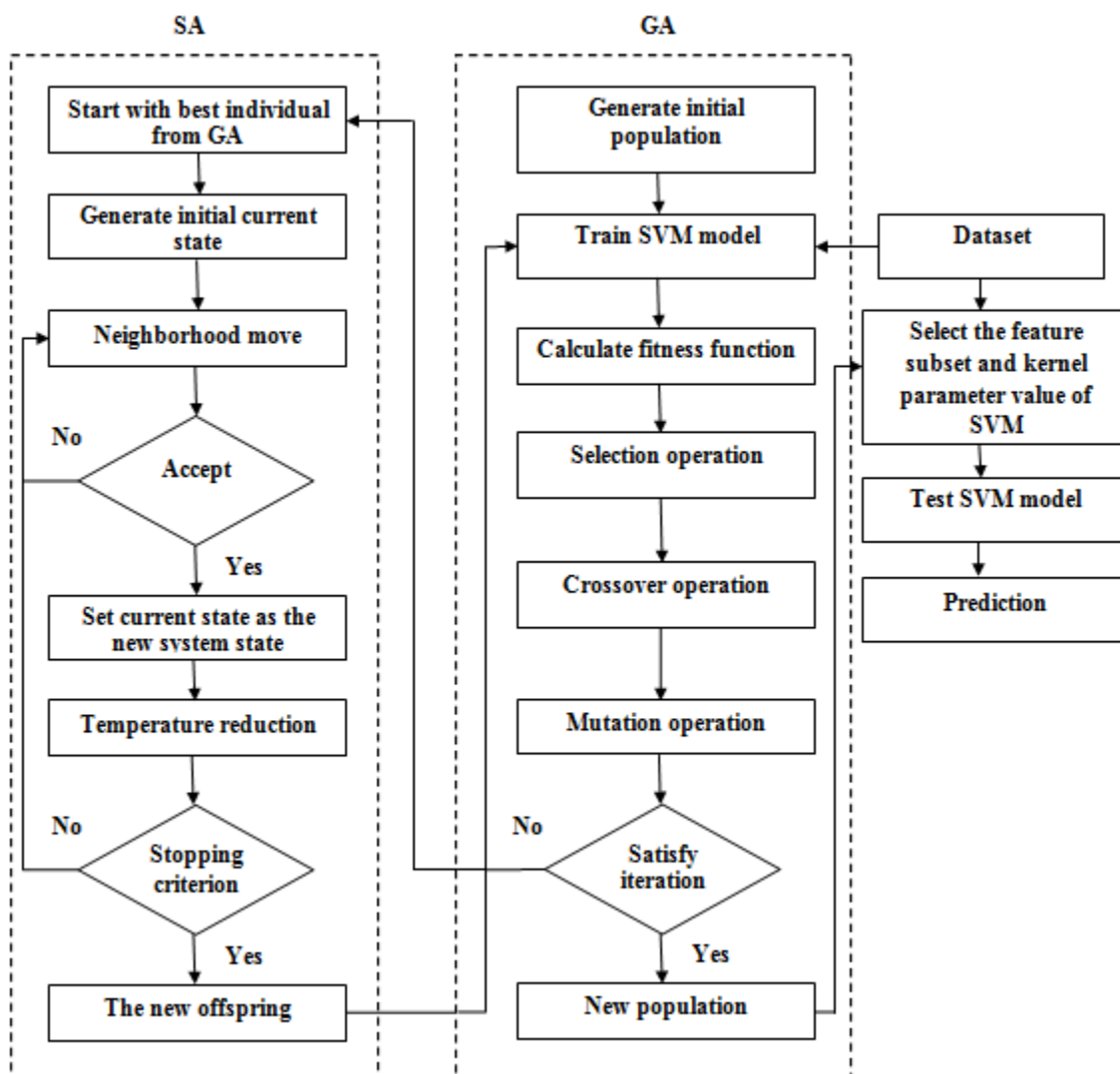


Fig.1 A flow diagram of the proposed GASA-SVM model

recognition and regression based on the theory of statistical learning and the principle of structural risk minimization [15].

Vladimir Vapnik invented SVM for searching a hyperplane that separates a set of positive examples from a set of negative examples with maximum margin [16]. The margin is defined by the distance of the hyperplane to the nearest of the positive and negative examples [17].

Given the training sample of instance-label pairs (x_i, y_i) , $i = 1, \dots, l$, $x_i \in R^n, y_i \in \{1, -1\}$, SVMs require the solution of the following (primal) problem [18]:

$$\min_{w,b,\epsilon} \frac{1}{2} W^T W + C \sum_{i=1}^l \epsilon_i$$

$$\text{Subject to } y_i(W^T Z_i + b) \geq 1 - \epsilon_i, \quad (1)$$

$$\epsilon_i \geq 0, i = 1, \dots, l,$$

where the training vector x_i is mapped onto a high dimension space by mapping function ϕ as $z_i = \phi(x_i)$. $C > 0$ is the penalty parameter of the error term.

Usually, equation (1) is solved by solving the following dual problem:

$$\min_{\alpha} F(\alpha) \frac{1}{2} \alpha^T Q \alpha - e^T \alpha$$

$$\text{Subject to } 0 \leq \alpha_i \leq C, i = 1, \dots, l \quad (2)$$

$$Y^T \alpha = 0,$$

where 'e' is the vector of all 1's and 'Q' is an l by l positive semi definite matrix.

The (i, j)th element of Q is given by

$$Q_{i,j} \equiv y_i y_j K(x_i x_j),$$

The kernel function is given by equation 3

$$K(x_i, x_j) \equiv \phi^T(x_i) \phi(x_j) \quad (3)$$

$\{\alpha_j\}_{j=1}^l$ is Lagrange multipliers, and

$W = \sum_{i=1}^l \alpha_i y_i \phi(x_i)$ is the weight vector.

The classification decision function is,

$$\text{sgn}(W^T \phi(x) + b) = \text{sgn}(\sum_{i=1}^l \alpha_i y_i K(x_i, x) + b) \quad (4)$$

The kernel function $K(x_i, x_j)$ has manifold forms. In this paper, the Gaussian Kernel Function is used. Gaussian kernel function is expressed as follows:

$$K(x, x_i) = \exp(-\gamma \|x - x_i\|^2), \quad (5)$$

(or)

$$K(x, x_i) = \exp\left(-\frac{1}{\sigma^2} \|x - x_i\|^2\right) \quad (6)$$

Both the equations (5) and (6), which are in the same context, can transform parameter γ and parameter σ^2 . The Gaussian kernel parameter c is determined by $\gamma = \frac{1}{\sigma^2}$. The parameters of SVMs with Gaussian RBF kernel refer to the pair: the error penalty parameter 'C' and the Gaussian kernel parameter γ , usually depicted as (C, γ).

4.4 Optimization

The performance of a SVM classifier depends mainly on the values of the kernel function parameter, Gamma (γ) and penalty function parameter (C). Finding the best values of these two parameters, in order to achieve maximum classification accuracy of the SVM classifier, is an optimization problem. A hybrid GA-SA algorithm is used to solve this problem and to find the optimal values of C and Gamma.

4.4.1 Genetic Algorithm (GA)

GA is a model of machine learning which derives its behavior from a metaphor of some of the mechanisms of evolution in nature. The individuals represent candidate solutions to the optimization problem being solved. In GAs, the individuals are typically represented by n-bit binary vectors [13]. The resulting search space corresponds to an n-dimensional boolean space. It is assumed that the quality of each candidate solution can be evaluated using a fitness function. GAs use some form of fitness-dependent probabilistic selection of individuals from the current population to produce individuals for the next generation. The fitness function is used to search the best result. The selected individuals are submitted to the action of genetic operators to obtain new individuals that constitute the next generation.

Mutation and crossover are two of the most commonly used operators that are used with GAs to represent individuals as binary strings. Mutation works on a single string and generally changes a bit

at random while crossover operates on two parent strings to deliver two off springs. Other genetic representations require the use of appropriate genetic operators. The process of fitness-dependent selection and application of genetic operators to generate successive generations of individuals is repeated many times until a satisfactory solution is found.

- **Initial population:** Like in any step by step optimization drawback, the data of great starting parameter advantages the convergence speed of the algorithm. But such kind of data is rarely accessible. This leads to the coverage of large parts of the solution space due to the generation of the random initial population. Hence, to make the exploration of the solution space easier, a heterogeneous initial population is suitable.
- **Selection Step:** According to the fitness value, the individuals are positioned and most elevated rank is given to the best one. At that point, one solution is kept in the next generation.
- **Crossover Step:** The objective of this step is to assemble interesting features of a few solutions in new individuals by making combination of already held solutions.
- **Mutation Step:** This step brings the vital risk for effectively explore the solution space. Any purpose of this space can be reached, it is guaranteed. In addition, if a local optimum is obtained, then a too quick convergence to this neighborhood optimum will be kept away by mutation. Initially, it is set to a maximum value, then decreases to permit convergence and it expands again to avoid local optima.

Algorithm Steps:

Step 1: Randomly generate an initial source population of 'n' chromosomes.

Step 2: Calculate the fitness function $f(x)$ of all chromosomes in the source population using equation (7).

$$\min f(x) = 100 * (x(1)^2 - x(2))^2 + (1 - x(1))^2 \quad (7)$$

Step 3: Create an empty successor population and then repeat the following steps until 'n' chromosomes have been created.

Step 4: Using fitness, select two chromosomes x_1 and x_2 from the source population.

Step 4.1: Apply crossover to x_1 and x_2 to obtain a child chromosome 'n'.

Step 4.2: By applying mutation to 'n', it deviant new offspring.

Step 4.3: Place new offspring in a new population.

Step 4.4: Replace the source population with the successor population.

Step 5: If the termination criterion is satisfied, stop the algorithm; otherwise go to Step 2.

4.4.2 Simulated Annealing (SA)

Annealing is a procedure in metallurgy, in which metals are gradually cooled to make them achieve low energy where they are very strong. In general, SA is a methodology where the temperature is reduced gradually, beginning from a random search at high temperature eventually getting pure greedy descent as it approaches zero temperature.

In optimization, the randomness should be appropriate to jump out of local minima and find regions that have a low heuristic worth. It is a probabilistic meta-heuristic for the global optimization issue of placing a good estimate to the global optimum of a given capacity in a large search space. It is regularly utilized when the search space is discrete [14].

Acceptance Function

The calculation chooses which answers to accept so we can better understand how it's able to avoid these local optimums.

Initially, the neighbor solution is checked to see whether it improves the current solution. If so, the current solution is accepted. Else, the neighbour solution is not better and hence the following couple of factors are considered.

- How greatly worse the neighbour solution is?
- How high the current temperature of our system is?

At high temperatures, the system is more likely to accept solutions that are worse.

Algorithm Steps:

Terminology:

X = Design Vector

fc = System Energy (i.e. Objective Function Value)

t = Temperature

Δ = Difference in system energy between two configuration vectors

Simulated Annealing Algorithm:

Step 1: Choose a random X_i , select the initial temperature t_1 and specify the cooling schedule.

Step 2: Evaluate $f_c(X_i)$ using a simulation model.

Step 3: Perturb X_i to get a neighboring Design Vector (X_{i+1}).

Step 4: Evaluate $f_c(X_{i+1})$ using a simulation model.

Step 5: If $f_c(X_{i+1}) < f_c(X_i)$, X_{i+1} is the new current solution.

Step 6: If $f_c(X_{i+1}) > f_c(X_i)$, then accept X_{i+1} as the new current solution with a probability using equation (8)

$$\exp(-\Delta/t) \text{ where } \Delta = f_c(X_{i+1}) - f_c(X_i) \quad (8)$$

Step 7: Reduce the system temperature according to the cooling schedule.

Step 8: Terminate the algorithm.

Neighborhood Search (Hybrid GA –SA)

In the hybrid SA algorithm, the best obtained solution in each GA generation is transferred to SA in order to improve the quality of solution through neighborhood search to produce a solution close to the current solution in the search space, by randomly choosing one gene in a chromosome, removing it from its original position and inserting it at another random position in the same chromosome. According to this criterion, even when the value of the next solution is worse, the solution can be accepted based on current temperature to avoid algorithm to stick in a local optimum. In the cooling phase the new temperature is determined by the decrement function t.

5 Experimentation and Results

5.1 Performance Metrics

To evaluate the prediction performance of SVM classifier, the classification accuracy, sensitivity and specificity are defined and computed.

Confusion Matrix: Confusion matrix shows classifications and predicted. A confusion matrix for a classification problem with two classes is of size 2×2 , as given in Table 1.

Table 1 Confusion Matrix

Predicted	Actual	
	Positive	Negative
Positive	TP (True Positive)	FP (False Positive)
Negative	FN (False Negative)	TN (True Negative)

- TP represents an instance, which is actually positive and predicted by the model as positive.
- FN represents an instance, which is actually positive but predicted by the model as negative.
- TN represents an instance, which is actually negative and predicted by the model as negative.
- FP represents an instance, which is actually negative but predicted by the model as positive.

Accuracy, Sensitivity and Specificity:

$$\text{Accuracy} = \frac{TP+TN}{TP+FP+FN+TN} \times 100\% \quad (9)$$

Sensitivity is the true positive rate and specificity is the true negative rate. They are defined as follows:-

$$\text{Sensitivity} = \frac{TP}{TP+FN} \times 100\% \quad (10)$$

$$\text{Specificity} = \frac{TN}{FP+TN} \times 100\% \quad (11)$$

5.2 Simulation Results

The system was implemented using MATLAB 7. In this paper, four diseases are diagnosed by using MATLAB software. The UCI machine learning repository datasets namely, Pima Indian Diabetes (PID) dataset, Wisconsin Diagnostic Breast Cancer (WDBC) dataset, Hepatitis disease dataset and Cardiac arrhythmia disease dataset are used to validate the proposed system with 80 % tuples for training and the remaining 20 % for testing.

In GA, an average of 30 generations is taken. The best fitness value found is 0.1481 and the mean fitness values for the four dataset are also calculated as 0.17 for PID, 0.19 for breast cancer, 0.18 for hepatitis and 0.18 for cardiac arrhythmia. The creation of generation and convergence of GA for all the four dataset- PID, breast cancer, hepatitis and cardiac arrhythmia is given in fig 2, fig 4, fig 6 and fig 8 respectively.

For SA, the initial temperature is set as 6.455 and the final temperature as 0.333. The temperature of SA is gradually reduced from initial value to the final in 50 cycles. The stopping criterion for the algorithm is given in equation (8). The GA receives the best chromosome with the help of SA. The working of SA using non-convex function for the four medical dataset- PID, breast cancer, hepatitis and cardiac arrhythmia is shown in fig 3, fig 5, fig 7 and fig 9 respectively.

The values of C and gamma achieved is C=400 and gamma = 0.0524.

In the PID dataset, the total number of attributes is 8, out of which 5 attributes are selected. The selected attributes list is shown in the Table 2.

Table 2 Selected features of Diabetes Dataset

Selected Attributes
Number of times pregnant
Plasma glucose concentration
Body mass index
Diabetes pedigree function
Age

In the WDBC dataset, the total number of attributes is 32, from which 12 attributes are selected. The selected attributes list is shown in the Table 3.

Table 3 Selected features of Breast Cancer Dataset

Selected Attributes
Texture
Perimeter
Area
Smoothness
Compactness
Concavity
Concave Points
Symmetry
Radius SE
Texture SE
Perimeter SE
Smoothness SE

In the hepatitis disease dataset, the total number of attributes is 19, out of which 10 attributes are selected. The selected attributes list is shown in the Table 4.

Table 4 Selected features of Hepatitis Dataset

Selected Attributes
Sex
Steroid
Antivirals
Fatigue
Anorexia
Liver Big
Spleen Palpable
Spiders
Ascites
Varices

In the arrhythmia dataset, there are 279 attributes. The first four attributes are namely age, sex, height, and weight, and the rest are attributes extracted from patients' ECG signals. The selected attributes list is shown in the Table 5.

Table 5 Selected features of Cardiac arrhythmia Dataset

Selected Attributes
QRS duration: Average of QRS duration
T interval: Average duration of T wave, linear
P interval: Average duration of P wave, linear

Heart rate: Number of heart beats, linear
Of channel DI: Q wave
Of channel DI: R' wave, small peak just after R
Number of intrinsic deflections, linear
Existence of diphasic derivation of T wave, nominal
Of channel AVR: Q wave
Of channel V1: R wave
Of channel V3: Q wave
Of channel V4: Existence of ragged P wave, nominal
Of channel V6: R wave
Of channel DI: Q wave, linear
Of channel AVF: Q wave

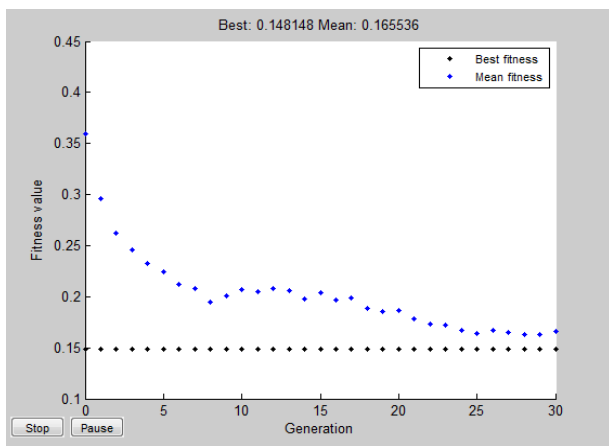


Fig.2 Generation creation using Genetic Algorithm for Diabetes dataset

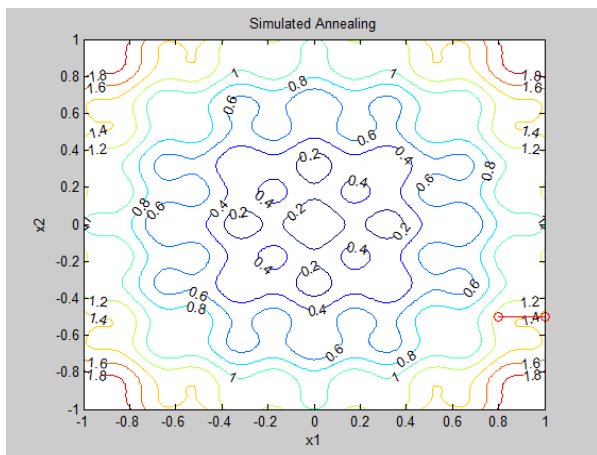


Fig.3 Neighborhood Search for Diabetes Dataset

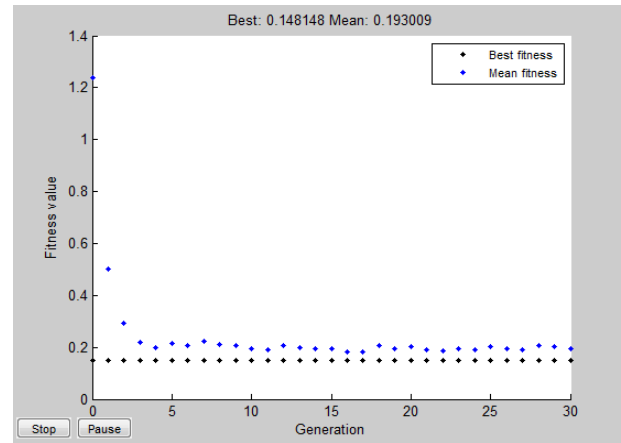


Fig.4 Generation creation using Genetic Algorithm for Breast Cancer Dataset

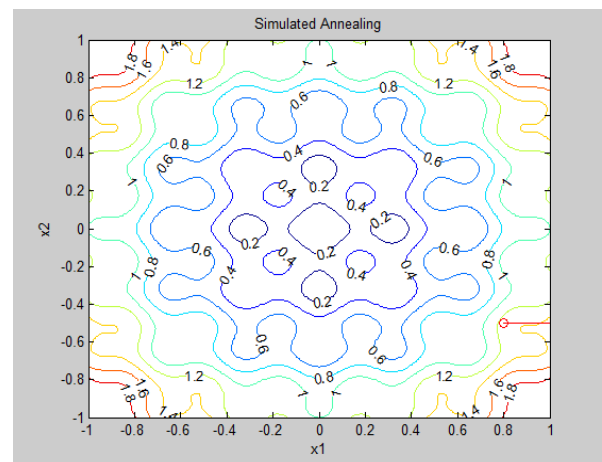


Fig.5 Neighborhood Search for Breast Cancer Dataset

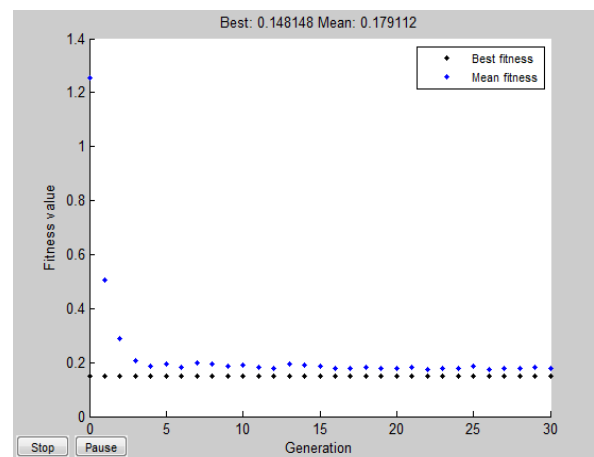


Fig.6 Generation creation using Genetic Algorithm for Hepatitis Dataset

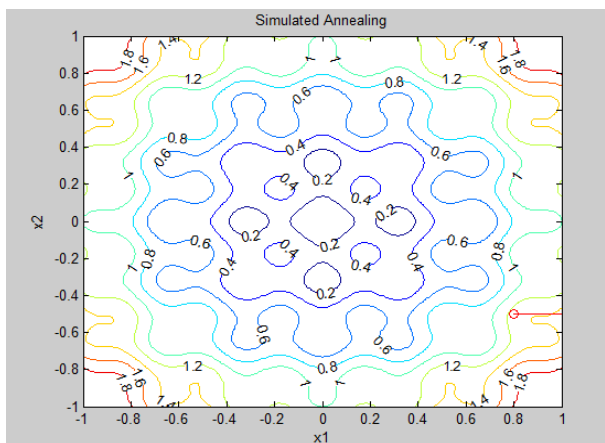


Fig.7 Neighborhood Search for Hepatitis Dataset

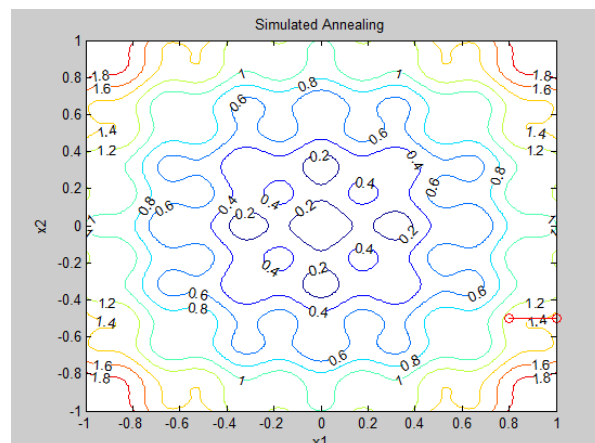


Fig.9 Neighborhood Search for Cardiac Arrhythmia Dataset

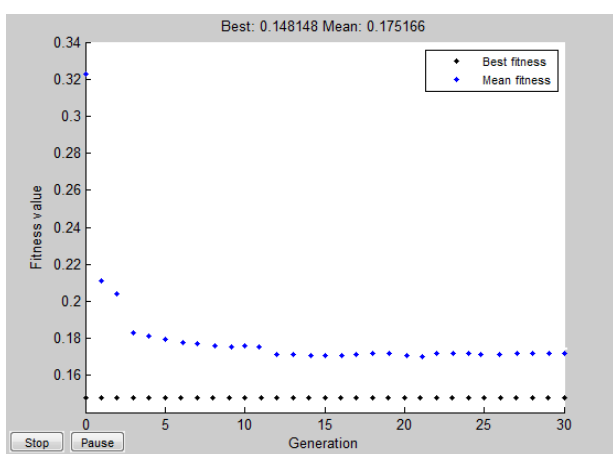


Fig.8 Generation creation using Genetic Algorithm for Cardiac Arrhythmia Dataset

6 Results and Discussion

Using 80-20 train-test combination, the proposed system is constructed and tested. The hybrid GA-SA optimization module is used to perform feature selection. The SVM classifier kernel parameters are also optimized using the GA-SA algorithm. The optimized kernel parameters ‘C’ and gamma ensure the best performance of the classifier. The performance of the system is measured using the metrics like classification accuracy, sensitivity and specificity. The details of the performance metrics obtained for all the four medical dataset is listed in table 6. The comparison of the performance of the proposed system, in terms of the classification accuracy, with the existing systems are given in table 7, table 8, table 9 and table 10.

Table 6 Accuracy, Specificity and Sensitivity of the Proposed System

Dataset	GASA-SVM		
	Accuracy (%)	Specificity (%)	Sensitivity (%)
Diabetes	91.2	89.3	97.5
Breast Cancer	93.60	88.32	97.21
Hepatitis	87	82	93
Cardiac Arrhythmia	89.3	83.41	94.72

Table 7 Comparison of Accuracy between the Proposed Method and the Other Methods for PID

Diabetes	SVM NSGA-II [19]	ANN [20]	SVM [21]	GA-SVM [22]	LDA-ANFIS [23]	Proposed GASA-SVM
Accuracy%	86.13	73.4	94	82.98	84.61	91.2

Table 8 Comparison of Accuracy between the Proposed Method and the Other Methods for Breast Cancer

Breast Cancer	SVM [29]	GA-SVM [29]	RST [30]	Grid Algorithm [22]	CART [32]	Proposed GASA-SVM
Accuracy%	75.87	76.57	85	90.78	93.5	93.60

Table 9 Comparison of Accuracy between the Proposed Method and the Other Methods for Hepatitis

Hepatitis	GA-SVM [27]	SVM-SA [28]	SVM [28]	KNN [29]	C4.5 [31]	Proposed GASA-SVM
Accuracy%	86.12	88.08	74	75.0	83.60	87.0

Table 10 Comparison of Accuracy between the Proposed Method and the Other Methods for Cardiac Arrhythmia

Cardiac Arrhythmia	ANN [24]	CFS-LM [25]	RF [11]	MNN Model [26]	GFFNN [26]	MLP [26]	Proposed GASA-SVM
Accuracy%	86.67	87.71	90	82.22	82.35	86.67	89.3

7 Conclusion

An expert system based on GASA-SVM is proposed for the diagnosis of various diseases. The hybrid GA-SA is used for the selection of the most significant feature subset of the dataset and also this module optimizes the kernel parameters of the SVM. Achieving the optimal values of C and Gamma leads to the improved classification accuracy of SVM. The SVM uses the Gaussian

RBF. The performance of the system is evaluated using various performance metrics like accuracy, sensitivity and specificity. The classification accuracy of the proposed system is compared with that of existing systems. In the future the system can be used to perform diagnosis on real life clinical data.

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