A Cooperative Coevolutionary Approach to Discretization-Based Feature Selection for High-Dimensional Data

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Abstract: Recent discretization-based feature selection methods show great advantages by introducing the entropy-based cut-points for features to integrate discretization and feature selection into one stage for high-dimensional data. However, current methods usually consider the individual features independently, ignoring the interaction between features with cut-points and those without cut-points, which results in information loss. In this paper, we propose a cooperative coevolutionary algorithm based on the genetic algorithm (GA) and particle swarm optimization (PSO), which searches for the feature subsets with and without entropy-based cut-points simultaneously. For the features with cut-points, a ranking mechanism is used to control the probability of mutation and crossover in GA. In addition, a binary-coded PSO is applied to update the indices of the selected features without cut-points. Experimental results on 10 real datasets verify the effectiveness of our algorithm in classification accuracy compared with several state-of-the-art competitors.

Keywords: feature selection; genetic algorithms; particle swarm optimization; cooperative coevolutionary; entropy-based cut-points

1. Introduction

Feature selection (FS) is an important task in machine learning, aiming to find an optimal subset of features to improve the performances of classification [1] or clustering [2,3]. By removing those redundant and irrelevant features, the model complexity is reduced and the overfitting in the training process can be avoided. Current FS algorithms can be generally categorized into wrapper and filter methods [4]. The filter approaches select the feature subset by scoring the features, and set a threshold to select those features that meet the conditions to form the feature subset. The wrapper methods treat FS as a search problem, generating different feature subsets and then evaluating the feature subsets until a certain feature subset reaches the expected standard. In general, the wrapper methods can achieve better and more robust results than the filter methods, but suffering from heavy computational burden.

For wrapper feature selection, it is essentially a combinatorial optimization problem, where the search space for data with \( N \) features is \( 2^N \). For high-dimensional data, the search space rises exponentially [5]. In the past few decades, it has been empirically demonstrated that evolutionary algorithms achieved their success in many different applications, for example soft sensors [6] and compressed sensing [7]. To overcome the shortage of traditional search methods [8,9], evolutionary algorithms were introduced into FS, among which, particle swarm optimization (PSO) [10–12] and genetic algorithms (GA) [13,14] are most widely adopted, due to their fast convergence [15] and...
powerful search capabilities [16], respectively. The core task of FS performed by evolutionary algorithms (EAs) is to identify the indices of selected features, so discretization-based encoding in evolutionary algorithm is desirable. The most commonly-used strategy is to apply a binary coding that indicates whether a feature is selected or not (one or zero), such as binary PSO (BPSO) [17]. However, for high-dimensional data, the search process is easily trapped into local optima. Another approach is to encode the indices of selected features directly. However, this often suffers from different encoding lengths and complicated updating rules during the search procedure, which is not very appropriate for the high-dimensional FS problem.

Recently, discretization-based feature selection algorithms have received much attention due to their good performance in high-dimensional data classification [18–21]. These algorithms map the search features into that of cut-points in PSO, which are generated by the univariate discretization algorithm MDL [22], combining feature discretization and feature selection into one stage. However, current discretization-based FS methods treat each feature independently without taking into account feature interaction, where the features without cut-points are not involved in the search process, leading to information loss and limiting the classification accuracy.

In recent years, the idea of coevolution has been successfully applied in different applications involving a large number of design variables, such as classification [23], artificial neural networks [24], function optimization [25], and image processing [26]. Cooperative coevolutionary algorithms (CCAs) decompose the original problem into subproblems with less decision variables. During the optimization process, CCAs are usually composed of two or more populations, which evolve simultaneously by applying different objectives or search methods and allow interaction, trying to obtain a global solution after combining the respective final solutions together. More related works along CCA can be found in [27].

Inspired by this methodology, in this paper, we propose a cooperative coevolutionary discretization-based FS algorithm (CC-DFS), which searches the subset of features with cut-points and without cut-points simultaneously. In our method, at first, the discretization technique is used to obtain the features with and without cut-points. Then, GA is applied to search the features with cut-points where a reset operation is used to jump out of local optima, and an individual scoring mechanism is introduced to control the probability of crossover and mutation. For features without cut-points, a binary-coded PSO is applied to update the indices of the selected features without cut-points. The final selected feature subset is composed of the results obtained by evolving both populations. Experimental studies on 10 real-world datasets demonstrate the superiority of our proposed approach in classification accuracy compared with some state-of-the-art discretization-based FS methods.

2. Background

2.1. Particle Swarm Optimization

PSO [28] is a population-based stochastic algorithm that searches the decision space by simulating the flight of birds in nature. In PSO, the population maintains a set of particles, each of which represents a feasible solution in the decision space. A fitness function of these particles is used to guide the search. The particle with the best fitness value in the population is selected as $g_{best}$, and the best individual fitness value achieved by each particle will be recorded as $p_{best}$. Each particle has a position vector and a velocity parameter, which are randomly initialized. The velocity parameter is updated according to $p_{best}$ and $g_{best}$ in each iteration as follows:

\[
v_{id}^{t+1} = w \times v_{id}^t + c_1 \times \text{rand} \times (p_{id}^t - x_{id}^t) + c_2 \times \text{rand} \times (g_{id}^t - x_{id}^t) \tag{1}\n\]

\[
x_{id}^{t+1} = x_{id}^t + v_{id}^{t+1} \tag{2}\n\]

where $x_{id}^t$ and $v_{id}^t$ represent the position and velocity of particle $i$ in dimension $d$ at the $t^{th}$ iteration, $w$ represents the inertia weight, which indicates the effect of the current velocity on the updated
velocity, $c_1$ and $c_2$ are acceleration coefficients used to control the effects of $p_{best}$ and $g_{best}$, and $\text{rand}$ is a function used to generate random numbers between zero and one. The velocity is usually limited by a preset threshold $V_{\text{max}}$, which limits the velocity to $[-V_{\text{max}}, V_{\text{max}}]$.

2.2. Genetic Algorithm

The genetic algorithm (GA) [29] is a heuristic search algorithm inspired by natural selection and biological inheritance, which provides a powerful search capability to find near optimal solutions in complex and large search spaces [16]. GA maintains a set of candidate solutions in a population, which are used to explore the decision space. Individuals (or chromosomes) in a population represent a set of feasible solutions to the problem, and next-generation individuals are produced through operators such as crossover and mutation. In addition, a fitness function is used to evaluate the quality of the individual, and those individuals with better fitness function values are selected to participate in the next iteration process. The bit-flip mutation [30] and discrete crossover [31] operations are shown in Figure 1 and Figure 2, respectively. In bit-flip mutation, a parent with binary encoding randomly selects a gene to flip it from zero to one, and vice versa. In discrete crossover, two parents are randomly selected to generate one offspring, and the offspring randomly selects genes from both parents.

2.3. Minimum Description Length Principle

The minimum description length (MDL) [22] is a supervised multivariate discrete algorithm, which finds the cut-points that satisfy the minimum description length principle (MDLP) to discretize the data. For each feature $A$, the features are sorted according to the feature value, and the algorithm selects the candidate cut-points, the feature value of which lies between instances of different classes. The formula for calculating the information gain of the cut-point $T$ is as follows:

$$\text{Gain}(T, A; S) = E(S) - \frac{|S_1|}{|S|} E(S_1) - \frac{|S_2|}{|S|} E(S_2)$$

where $E(S)$ denotes the entropy of dataset $S$ and $|S_1|$ and $|S_2|$ represent the number of samples of each part after the dataset $S$ is divided into two sample subsets by the cut-point $T$.

The algorithm divides the dataset recursively until the cut-point cannot pass the MDLP criterion. Feature values that satisfy the MDLP criteria are used as our cut-points to discrete datasets. MDLP criteria are calculated based on Equation (4).
\[ \text{Gain}(T, A; S) > \frac{\log_2(|S| - 1)}{|S|} + \frac{\delta(T, A; S)}{|S|} \]  

(4)

\[ \delta(T, A; S) \text{ is calculated by (5).} \]

\[ \delta(T, A; S) = \log_2(3^{k_s} - 2) - [k_sE(S) - k_sE(S_1) - k_sE(S_2)] \]  

(5)

where \( K_s \) denotes the number of classes present in \( S \) and \( S_1 \) and \( S_2 \) represent the two sample subsets after the sample \( S \) is divided.

### 2.4. Discretization-Based FS Algorithms

In the past few years, discretization-based algorithms have demonstrated great potential to deal with high-dimensional data, such as EPSO [19] and PPSO [18].

In EPSO, individuals in the population are encoded as real numbers that are within the range of feature values. As shown in Figure 3, the particle’s position of \( F_1 \) is \(-21\), falling within its feature value range, which represents selecting feature \( F_1 \) and using this value to discretize the corresponding feature.

During the update process, if the value of the particle in a dimension exceeds the upper or lower limit of the feature, the value is set to the corresponding upper or lower limit. When the value of the particle in a dimension is equal to the upper or lower limit, the feature is discarded, because when the feature is discretized with the upper limit or the lower limit of the feature value, the discrete values of the feature are equal, which means that it cannot contribute to classification.

EPSO directly uses features to evolve the cut-points, which results in a huge search space. PPSO introduced the MDL algorithm to select the features with cut-points and encode the position of cut-points in the search space of PSO. Individuals in the population are encoded by the cut-point table, as shown in Figure 4. The MDL algorithm was initially used to calculate the cut-point table of features as shown on the left side, where \( #P \) represents the number of cut-points for each feature, and \( C_1 \), \( C_2 \), and \( C_3 \) represent the index of the cut-point for the features. If the value exceeds the range of the cut-point index during the particle update process, it is set to zero, which means that the feature is not selected. For instance, the value of \( F_3 \) is two, which indicates that the cut-point \( C_2 \) for \( F_3 \) is selected to discretize the feature; the value for \( F_1 \) is zero, which means that the feature is discarded.

Figure 3. The particle representation of EPSO.

Figure 4. The particle representation of PPSO.
3. Proposed Method

Although discretization-based FS algorithms have achieved good results in high-dimensional data, the absence of features without cut-points causes information loss, which affects the classification accuracy negatively. The discretization process treats each feature independently, and some groups or pairs of weak features may have a greater impact on the classifying performances than one individual strong feature; therefore, features that do not have cut-points or participate in the discrete process are likely to contribute to the classification task. In this section, we introduce the details of our proposed cooperative coevolutionary discretization-based FS (CC-DFS) method, which considers searching the feature subset from the features with and without cut-points simultaneously. The pseudocode of CC-DFS is presented in Algorithm 1, and Figure 5 shows a flowchart of our algorithm.

![Figure 5. Overview of our proposed method.](image)

**Algorithm 1:** The pseudocode of the proposed CC-DFS.

**Input:** Training set  
**Output:** Selected feature subset and the cut-points table

begin

Cut-points table ← calculated by MDL;  
Initialize the population G and P of GA and PSO, respectively;  
while Stopping criteria are not met do

if Meet the reset condition then

if rand < 0.5 then

Set individuals that perform operations and cross operations on G to zero vectors of equal length.  
for Each individual i do

\[ G_i \leftarrow \text{Offspring from parent through mutation (15) and crossover.} \]

\[ P_i \leftarrow \text{Particle updating by using Equations (1) and (2)} \]

else

Set gbest of P to zero vectors of equal length.  
for Each individual i do

\[ G_i \leftarrow \text{Offspring from parent through mutation (15) and crossover.} \]

\[ P_i \leftarrow \text{Particle updating by using Equations (1) and (2)} \]

else

for each individual i do

\[ G_i \leftarrow \text{Offspring from parent through mutation (15) and crossover.} \]

\[ P_i \leftarrow \text{Particle updating by using Equations (1) and (2)} \]

Update pbest of G and P.  
Select N individuals from G and form a feature subset PG with P.  
Update gbest of PG.  
Output the selected feature subset and the cut-points table.

end
3.1. Representation of an Individual

CC-DFS was designed to search both the features with and without cut-points, simultaneously. For simplicity, we denote the features with cut-points by discrete features and the features without cut-points by continuous features. To achieve this, the whole decision space \( A \) is divided into two parts, where decision space \( D \) contains discrete features, while decision space \( C \) contains continuous features. The population is encoded using two different encoding methods where the discrete features use the encoding way of PPSO and the continuous features use binary encoding. As shown in Figure 6, for feature \( F_5 \), one means to select the cut-point with Index 1 in the cut-point table to discretize the data and for \( F_6 \); it means selecting the cut-point with Index 2 in the cut-point table. The features with a value of zero mean not selecting a cut-point to discretize the feature and for continuous features, and zero means discarding the feature, while one means selecting the feature. At last, these two parts are combined into one vector to represent a feature subset.

![Figure 6. The particle’s representation of our algorithm.](image)

3.2. Fitness Function

The fitness function plays a vital role in the population update process. It is used to evaluate and guide the population update. In the FS problem, we want to use fewer features to maintain a better or competitive classification accuracy than all the features. Considering the above issues, we use the distance [32] and balanced _error to guide individual updates. These two objective functions use weight aggregation [33–35] as the fitness function as shown in (6), where the smaller the fitness value, the better the individual’s performance.

\[
\text{fitness} = (\beta \times \text{balanced}\_\text{error} + (1 - \beta) \times \text{distance})
\]  
(6)

where \( \beta \) is a weight coefficient to combine the balanced _error and distance. balanced _error and distance are calculated by (7) and (8), respectively.

\[
\text{balanced}\_\text{error} = \frac{1}{n} \sum_{i=1}^{n} \frac{FP_i}{|S_i|}
\]  
(7)

\[
distance = \frac{1}{1 + \exp^{-5(D_W - D_B)}}
\]  
(8)

\[
D_B = \frac{1}{|N|} \sum_{i=1}^{N} \min_{(j\neq i,\text{class}(V_i)\neq\text{class}(V_j))} \text{Dis}(V_i, V_j)
\]  
(9)

\[
D_W = \frac{1}{|N|} \sum_{i=1}^{N} \max_{(j\neq i,\text{class}(V_i)=\text{class}(V_j))} \text{Dis}(V_i, V_j)
\]  
(10)

where \( n \) denotes the number of the classes, \( FP_i \) represents the number of misclassified samples in class \( i \), \( |S_i| \) means the number of samples in each class, \( |N| \) denotes the number of samples, \( D_B \) represents
the distance between each sample in the data and its nearest sample of different classes, and $D_W$ represents the distance between each sample in the data and its farthest sample of the same class.

In CC-DFS, since two different encoding methods for FS are performed for two different decision spaces, the calculation of $Dis(V_i, V_j)$ is also different from each other. As $Dis(V_i, V_j)$ represents the distance between two feature subset vectors $V_i$ and $V_j$, for discrete features, the Hamming distance is applied for discrete features, while the Euclidean distance is calculated for continuous features. In addition, the historically optimal positions of discrete and continuous feature subset individuals are recorded as $pbest$. The population optimal record of the feature subset of discrete and continuous feature combinations is $gbest$, where its discrete feature part is called $dbest$ as the $pbest$ of discrete individuals and the continuous part $cbest$ as the $pbest$ of continuous feature individuals.

3.3. Updating Strategy

For discrete features and continuous features, due to the easy implementation and fast convergence, GA and PSO are employed to search their feature subset, respectively.

3.3.1. Search in Discrete Features

Different from the traditional GA algorithm, we propose a novel mutation and crossover operation. In our method, each individual in the population is scored using a ranking function, where individual scores are used to control their probability of mutation and crossover. Inspired by the ranking function in [36], considering that the population’s variability becomes lower with the increase of iterations, an adaptive scoring mechanism is introduced, which aims to reduce the mutation and crossover probability of each dimension in the later stage shown as follows:

$$Pc_i = L_{min} + L_{max} \times \frac{\exp{\left(\frac{10(r-1)}{S-1}\right)} - 1}{\exp{10} - 1}$$

$$L_{max} = L_{max} - \frac{t}{It} \times L_{max}$$

where $L_{min}$ and $L_{max}$ control the upper and lower limits of the score, $S$ represents the number of populations, $r$ is the individual’s ranking, $t$ is the number of current iterations, and $It$ is the maximum number of iterations. In this paper, the lower the fitness value is, the higher the individual ranks. Each dimension of individuals in the population takes its ranking as a probability to decide whether to perform mutation and crossover operations. It is guaranteed that individuals with high rankings have a lower probability of performing mutation and crossover in each dimension, and individuals with low rankings perform a higher probability of mutation and crossover.

The particle encoding range of discrete features is an integer between $[0, \# C]$; therefore, for the mutation operation, traditional bit-flip mutation is not suitable. For each dimension of the individual, we randomly select two other individuals in the population, and the one with the better fitness value performs the mutation operation with the current individual. The mutation operation is similar to the update operation in [37]. The formula is as follows:

$$ch_{id}^{t+1} = \begin{cases} N(\mu, \sigma) & \text{rand} < 0.5 \\ pa_{id}^{t} & \text{otherwise} \end{cases}$$

where $ch_{id}^{t+1}$ represents the value of the individual $i$ in dimension $d$ after $t$ iterations, $pa_{id}^{t}$ represents the value of parent $i$ in dimension $d$ at the $t^{th}$ iteration, and the mean of the two values performing the mutation operation as $\mu$; the absolute value of the gap as $\sigma$, rand is to generate a random number between $[0,1]$, which represents a 50% probability that each dimension of the individual chooses to use the Gaussian function, and there is a 50% probability that the value of the offspring is directly set to the value of the parent at the current position. In the cross operation, we use the ranking function
of each individual of the population as the cross probability of the individual. When each gene of
the individual performs the cross operation, two other individuals are randomly selected, and then,
the highest ranked individual and the current individual perform a crossover operation. In this way,
for higher ranking individuals, the crossover probability is lower; while for lower ranking individuals,
the crossover probability is higher.

3.3.2. Search in Continuous Features

For continuous feature search, we use binary coding and then use BPSO to search for the continuous
feature subset. The difference between BPSO and traditional PSO is that the position update of the
particles no longer depends on the position of the previous moment, but the sigmoid function is
used to map the velocity to the probability to determine the value of the particle’s current dimension.
The formula is as follows:

\[ S_{id}^t = \frac{1}{1 + e^{-v_{id}^t}} \]  \hspace{1cm} (14)

\[ x_{id}^t = \begin{cases} 
1 & \text{rand} < S_{id}^t \\
0 & \text{otherwise}
\end{cases} \]  \hspace{1cm} (15)

where \( x_{id}^t \) and \( v_{id}^t \) represent the velocity of particle \( i \) in dimension \( d \) at the \( t \)th iteration. \( S_{id}^t \) represents
the probability of the \( i \)th feature being selected, and \( \text{rand} \) is used to generate random numbers between
zero and one.

3.3.3. Combination of Continuous and Discrete Features

For discrete features, 2N individuals are generated by crossover and mutation of N individuals,
and N individuals of continuous features are still N individuals after updating. Therefore, we need
to select N individuals in the GA population to form a new feature subset with continuous features.
For these discrete features, we use a distance function to measure the quality of these individuals.
The first N individuals with better distance measures are selected, and these individuals and continuous
feature individuals form a new feature subset.

3.3.4. Reset Operation

When the optimal fitness value of the population has not been improved after three consecutive
updates, we consider that the population has fallen into a local optimum. The reset operation is
introduced, and in the next update, GA and PSO will randomly choose one to perform the reset
operations. If GA is selected, for each individual in the population, the individual who performs
mutation and crossover operations with it will be a zero vector of equal length. If PSO is selected,
\( \text{gbest} \) will be set to zero in the next update. Moreover, if the particle’s optimal fitness value has not
been improved after \( 11 \) updates, the iterative process will stop to reduce the program’s running time.

4. Experimental Results and Analysis

This section introduces the details of the experiment and parameters in the algorithm. Besides,
the baseline methods for comparison with our method are introduced.

4.1. Datasets

Ten real genetic data were used to test the performances of different algorithms, which can be
downloaded from https://github.com/primekangkang/Genedata. Table 1 shows the number of
features, samples, classes, and the proportion of the smallest class and largest class. It can be observed
that most of the data were small samples and high-dimensional, and there were class imbalances.
Data were normalized before input.
Table 1. Datasets.

| Dataset      | # of Features | # of S\(^2\) | # of C\(^3\) | # of Small | # of Big |
|--------------|---------------|--------------|--------------|------------|----------|
| SRBCT        | 2308          | 83           | 4            | 13         | 35       |
| DLBCL        | 5469          | 77           | 2            | 25         | 75       |
| 9Tumor       | 5726          | 60           | 9            | 3          | 15       |
| Leukemia 1   | 5327          | 72           | 3            | 13         | 53       |
| Leukemia 2   | 11,225        | 72           | 3            | 28         | 39       |
| Brain Tumor1 | 5920          | 90           | 5            | 4          | 67       |
| Brain Tumor2 | 10,367        | 50           | 4            | 14         | 30       |
| Prostate     | 10,509        | 102          | 2            | 49         | 51       |
| Lung Cancer  | 12,600        | 203          | 5            | 3          | 68       |
| 11Tumor      | 12,533        | 174          | 11           | 4          | 16       |

\(^1\) # means; \(^2\) S means the number of samples; \(^3\) C means the number classes.

4.2. Parameter Settings and Comparison Method

The parameters of our algorithm are shown in Table 2.

Table 2. Parameter setting.

| Parameter          | Setting                                                                 |
|--------------------|-------------------------------------------------------------------------|
| Population         | No. of features/20 (Limited to 300 and no less than 100)                |
| Maximum iteration  | 100                                                                     |
| c1 and c2          | 1.49445                                                                |
| \(\beta\)          | 0.5                                                                    |
| Lmin               | 0.25                                                                   |
| Lmax               | 0.5                                                                    |
| Stopping criterion | The fitness value of \(g_{best}\) not improved for 11 iterations        |

To test the performance of the algorithm, we used the K-NN algorithm with \(K = 1\) as the classifier. The feature subsets selected by the algorithm were input into the classifier to obtain the average classification accuracy on the test set. In the experiment, we conducted a two layer 10-fold cross-validation. At first, the whole dataset was divided into ten portions where one portion was used for testing. Then, the other nine were divided into ten portions again, nine for training and one for validation to calculate the training accuracy. The average testing accuracy was recorded after running the algorithms 30 times.

The classification result of all feature input KNN was used for comparison with our algorithm. EPSO, deep multilayer perceptron (DMLP), PPSO, and PSO-FS [18] were used to compare with our algorithm, and the parameters of the algorithms used for comparison were the parameters recommended in [18]. For DMLP, the number of neurons in the input layer was set to the number of features, and the number of neurons in the output layer was fixed at 128 as the number of selected features. In addition, the cooperative coevolutionary using bare-bone PSO instead of GA, called CCB-DFS, was also used for comparison, and the parameter setting was the same as CC-DFS.

4.3. Results and Analysis

In this section, we compare and analyze the results of our algorithm with respect to other algorithms. The results are shown in Table 3, where Full means we used all feature input KNN for classification and S represents the result of the statistical Wilcoxon significance test with a 5% significance level, and the result of the statistical Wilcoxon significance test was a comparison of our method with other methods, where “+” indicates that our algorithm results were better than the compared algorithms, “−” indicates that our algorithm results were worse than the compared algorithms, and “=” indicates that the results of the two algorithms were not significantly different.
Table 3. Experimental results. DMLP, deep multilayer perceptron; FS, feature selection; CCB-DFS, coevolutionary discretization-based using bare-bone PSO FS. (Bold indicates the best value).

| Dataset   | Method       | # of Features | Best (%) | Avg (std) | S     |
|-----------|--------------|---------------|----------|-----------|-------|
|           | Full         | 2308.0        | 87.08    | +         |       |
|           | DMLP         | 128           | 100.00   | 97.72 (1.67) | +     |
|           | PSO-FS       | 150.0         | 97.50    | 91.31 (2.71) | +     |
| SRBCT     | EPSO         | 137.3         | 100.00   | 96.89 (1.64) | +     |
|           | PSO          | 108.5         | 100.00   | 95.78 (1.96) | +     |
|           | CCB-DFS      | 168.1         | 100.00   | 99.26 (0.89) | =     |
|           | CC-DFS       | 220.6         | 100.00   | 98.90 (1.05) | =     |
| DLBCL     | Full         | 5469.0        | 83.67    | +         |       |
|           | DMLP         | 128           | 97.92    | 93.26 (3.35) | −     |
|           | PSO-FS       | 101.8         | 96.67    | 80.03 (6.13) | +     |
|           | EPSO         | 42.8          | 94.17    | 85.18 (5.46) | +     |
|           | PSO          | 44.0          | 94.17    | 86.22 (3.58) | +     |
|           | CCB-DFS      | 150.0         | 100.00   | 99.26 (0.89) | =     |
|           | CC-DFS       | 180.6         | 100.00   | 98.90 (1.05) | =     |
| 9Tumor    | Full         | 5726.0        | 83.00    | +         |       |
|           | DMLP         | 128           | 97.92    | 91.64 (3.99) | +     |
|           | PSO-FS       | 150.0         | 92.22    | 81.60 (4.72) | +     |
|           | EPSO         | 138.5         | 65.00    | 58.22 (3.12) | −     |
|           | PSO          | 118.1         | 65.00    | 59.28 (2.08) | −     |
|           | CCB-DFS      | 314.2         | 58.20    | 52.66 (3.64) | =     |
|           | CC-DFS       | 278.0         | 61.48    | 53.76 (3.39) | =     |
| Leukemia1 | Full         | 11,225.0      | 84.44    | +         |       |
|           | DMLP         | 128           | 97.92    | 91.64 (3.99) | +     |
|           | PSO-FS       | 150.0         | 92.22    | 81.60 (4.72) | +     |
|           | EPSO         | 139.9         | 95.56    | 93.37 (1.83) | =     |
|           | PSO          | 80.4          | 95.42    | 94.37 (1.36) | =     |
|           | CCB-DFS      | 128.6         | 96.67    | 94.14 (1.35) | =     |
|           | CC-DFS       | 166.4         | 97.50    | 94.02 (1.45) | =     |
| Leukemia2 | Full         | 5920.0        | 72.08    | +         |       |
|           | DMLP         | 128           | 97.92    | 91.64 (3.99) | +     |
|           | PSO-FS       | 150.0         | 92.22    | 81.60 (4.72) | +     |
|           | EPSO         | 135.9         | 95.56    | 93.37 (1.83) | =     |
|           | PSO          | 86.7          | 100.00   | 96.74 (1.64) | =     |
|           | CCB-DFS      | 220.6         | 95.17    | 90.20 (2.00) | =     |
|           | CC-DFS       | 317.1         | 95.57    | 92.09 (2.09) | =     |
| Brain Tumor1 | Full      | 10,367.0      | 62.50    | +         |       |
|            | DMLP         | 128           | 81.81    | 73.93 (3.25) | =     |
|            | PSO-FS       | 317.3         | 78.75    | 71.00 (3.06) | =     |
|            | EPSO         | 150.0         | 93.89    | 86.11 (3.97) | +     |
|            | PSO          | 86.7          | 100.00   | 95.17 (2.00) | =     |
|            | CCB-DFS      | 130.8         | 73.48    | 75.90 (2.49) | =     |
|            | CC-DFS       | 187.4         | 83.08    | 76.57 (3.47) | =     |
| Brain Tumor2 | Full      | 10,509.0      | 85.33    | +         |       |
|            | DMLP         | 128           | 83.40    | 74.25 (3.21) | =     |
|            | PSO-FS       | 777.4         | 90.33    | 85.20 (2.35) | +     |
|            | EPSO         | 54.9          | 90.33    | 83.74 (3.55) | +     |
|            | PSO          | 65.6          | 95.17    | 91.82 (1.77) | −     |
|            | CCB-DFS      | 129.8         | 92.50    | 89.06 (2.08) | =     |
|            | CC-DFS       | 180.6         | 92.17    | 88.52 (1.63) | =     |
| Prostate   | Full         | 12,533.0      | 71.42    | +         |       |
|            | DMLP         | 128           | 79.36    | 73.69 (3.19) | +     |
|            | PSO-FS       | 1638.8        | 86.07    | 82.62 (1.70) | +     |
|            | EPSO         | 149.9         | 83.68    | 79.29 (2.11) | +     |
|            | PSO          | 167.0         | 83.20    | 76.83 (2.91) | +     |
|            | CCB-DFS      | 1422.4        | 89.09    | 85.27 (1.92) | =     |
|            | CC-DFS       | 1380.2        | 87.77    | 84.84 (2.47) | =     |
| 11Tumor    | Full         | 12,600.0      | 78.05    | +         |       |
|            | DMLP         | 128           | 81.21    | 73.76 (3.73) | +     |
|            | PSO-FS       | 686.2         | 85.73    | 81.72 (2.08) | =     |
|            | EPSO         | 150.8         | 85.58    | 80.60 (2.42) | =     |
|            | PSO          | 203.0         | 84.11    | 79.38 (3.26) | +     |
|            | CCB-DFS      | 433.9         | 86.92    | 82.88 (2.13) | =     |
|            | CC-DFS       | 155.6         | 89.71    | 81.40 (3.10) | =     |
4.3.1. CC-DFS VS Full

As can be seen from Table 3, our algorithm selected fewer features, and the classification performance was higher than using all the features on all datasets. For SRBCT, CC-DFS selected 9.5% of all features, with the average classification accuracy of 98.9%, which was an 11.1% improvement over Full, and the best classification accuracy reached 100%. For Leukemia1, three-point-one percent of the features were selected, and the average classification accuracy was 94.02%. Compared with Full, the average classification accuracy was improved by 21.94%, and the best classification accuracy was 97.50%. For 11Tumor, the average classification accuracy increased by 13.42%, and the best classification accuracy increased by 16.35%.

In general, CC-DFS obtained ten “+” compared to Full, which meant that the classification performance was better than Full on all datasets.

4.3.2. CC-DFS vs. DMLP

Compared with DMLP, on eight datasets, our algorithm’s performance was better than DMLP in average classification accuracy. For Prostate, Compared with DMLP, the number of features increased by 52.6, but the average classification accuracy and the best classification accuracy were improved by 14.27% and 8.77%, respectively. For Leukemia1, although the best classification accuracy of DMLP was improved by 0.42% compared to CC-DFS, the average classification accuracy was reduced by 2.38%. For Brain2, although the average classification accuracy of DMLP was reduced by 1.71% compared to CC-DFS, the best classification accuracy was improved by 1.94%. Compared with DMLP, CC-DFS obtained eight “+”, one “=”, and one “−”.

4.3.3. CC-DFS vs. PSO-FS

Compared with PSO-FS, for seven datasets, our algorithm selected fewer features, and the classification performance was higher. For the remaining three datasets, although the feature size obtained by our algorithm was slightly higher, there was a significant improvement in classification performance. For DLBCL, CC-DFS selected 1.4% of all features, with the average classification accuracy of 90.37%, which was an 10.34% improvement over PSO-FS. Although the algorithm achieved the same best classification accuracy, our algorithm reduced the number of features by 24.2. For Leukemia1, three-point-one percent of the features were selected, and the average classification accuracy was 94.02%. Compared with PSO-FS, the average classification accuracy was improved by 12.42%, and the best classification accuracy was improved by 5.28%. For Lung Cancer, the average classification accuracy was 81.40%, which was slightly lower than PSO-FS. but the best classification accuracy increased by 3.98%. Compared with PSO-FS, CC-DFS obtained nine “+” and one “=”, without any “−”.

4.3.4. CC-DFS vs. EPSO

Compared with EPSO, in terms of classification performance, CC-DFS obtained seven “+”, one “−”, and two “=”. The lowest average classification accuracy improvement was 0.65%, appearing on Leukemia1, and the highest improvement was 5.64%, appearing on Leukemia2. The maximum classification accuracy improvement was 5.56%, appearing on Leukemia2. For Brain Tumor1, in terms of average classification accuracy, three-point-seven-eight percent improvement was obtained, and the best classification accuracy increased by 3.91%. Except Leukemia2, on the other datasets, the selected feature subset length was higher than EPSO. In summary, compared with EPSO, at the cost of a small number of features, a good improvement in classification performance was achieved.

4.3.5. CC-DFS vs. PPSO

For eight datasets, CC-DFS achieved better or similar best classification accuracy compared to PPSO. For six datasets, the average classification accuracy of CC-DFS was higher than PPSO. For Lung
Cancer, an average of 155.6 features was selected, which was 47.4 fewer than PPSO, while the average classification accuracy was improved by 2.02%, and the best classification accuracy was increased by 5.6%.

4.3.6. CC-DFS vs. CCB-DFS

Compared with CCB-DFS, similar classification performance was achieved on nine datasets. On six datasets, CC-DFS selected fewer features than CCB-DFS. For 9Tumor, in terms of average classification accuracy, one-point-one-two percent improvement was obtained, and the best classification accuracy increased by 3.28%, but the number of selected features dropped by 36.2.

In summary, for 50 comparisons, CC-DFS obtained 31 “+”, 5 “−”, and 14 “=”, which meant that among 45 comparisons, CC-DFS obtained better or similar classification performance, and only five times, the classification performance was lower than the compared algorithm.

4.4. Effectiveness of Reset Operation

To verify the performance of the reset operation, in this section, the algorithm with (denoted by W) and without the reset operation (denoted by W/R) were compared on ten datasets. Figure 7 shows the change in the algorithm’s fitness value during the iteration during the update process. Table 4 shows the running time of the algorithm. As shown in Table 4, the running time of the algorithm using the reset operation was much smaller than the running time of the algorithm without the reset algorithm on the 10 datasets. This result was because the reset operation stopped the iteration in advance when the fitness value of gbest was not updated for 11 consecutive times. It can also be seen in Figure 7 that the gbest fitness value of the algorithm using the reset operation was better than the value without the reset algorithm when the iteration was stopped, and because of the early termination mechanism, the number of iterations was also much smaller than W/R. In addition, with fewer iterations, a better fitness value was obtained.

Table 4. Running time (s). W, with the reset operation; W/R, without the reset operation.

| Dataset    | Time (W) | Time (W/R) |
|------------|----------|------------|
| SRBCT      | 157.1    | 352.2      |
| DLBCL      | 397.1    | 782.3      |
| 9Tumor     | 260.4    | 637.9      |
| Leukemia1  | 475.1    | 778.9      |
| Leukemia2  | 1465.4   | 2283.8     |
| Brain Tumor1| 909.8   | 1445.1     |
| Brain Tumor2| 1591.6  | 2041.4     |
| Prostate   | 884.4    | 1334.8     |
| 11Tumor    | 2062.2   | 4885.1     |
| Lung Cancer| 4756.9   | 12,075.0   |

4.5. Computational Time

In this section, we measure the time of feature selection algorithms in Figure 8, since all these approaches were based on evolutionary algorithms and an offline cut-point table.

From Figure 8, we can see that PSO-FS had a shorter running time on most datasets. This was mainly because PSO-FS only discretized the data once before searching while the remaining algorithms needed to discretize the data during the search process. Although the running time of CC-DFS was much higher than that of EPSSO and PPSO, its classification accuracy was superior to these two algorithms. As can be seen from Figure 8, the greater the feature size, the greater the gap between the running time of CC-DFS and the running time of EPSSO and PPSO. We considered continuous features and selected more features, which in turn led to a much higher running time required by the fitness evaluation process. The reason CC-DFS spent more time than CCB-DFS was that the mutation and crossover operations of the GA consumed more runtime than PSO, but CC-DFS improved the
best classification accuracy compared to CCB-DFS. Despite the long running time of our algorithm, considering continuous features significantly improved the classification accuracy.

![Figure 7. With reset operation and without reset operation.](image)

Figure 7. With reset operation and without reset operation.

![Figure 8. Comparison of the running time.](image)

Figure 8. Comparison of the running time.

5. Conclusions

In this paper, a new cooperative coevolutionary discretization-based FS method, CC-DFS, was proposed, where both discrete and continuous features were searched to reduce the information loss by only considering the discrete features. During the update process, GA and PSO were applied to search for discrete and continuous feature subsets, respectively. Average classification error and distance measures were used as individual evaluation indicators. The ranking mechanism was introduced to control the mutation and crossover probability of individuals in GA, each dimension of which allowed the mutation and crossover operation with different individuals. Through the distance measure, we selected $N$ individuals from discrete features to form a feature subset with continuous features. In addition, a reset operation was used to jump out of the local optimum. Experimental results showed that our method was able to improve the classification accuracy on the benchmark datasets. Compared with using all features, the feature subset selected by CC-DFS could achieve higher classification accuracy. Compared with some state-of-the-art discretization-based methods, it demonstrated the advantages of considering both discrete and continuous features in FS.
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References
1. Blum, A.L.; Langley, P. Selection of relevant features and examples in machine learning. Artif. Intell. 1997, 97, 245–271. [CrossRef]
2. Dash, M.; Choi, K.; Scheuermann, P.; Liu, H. Feature Selection for Clustering—A Filter Solution. In Proceedings of the 2002 IEEE International Conference on Data Mining, Maebashi City, Japan, 9–12 December 2002.
3. Alelyani, S.; Tang, J.; Liu, H. Feature Selection for Clustering: A Review. In Data Clustering: Algorithms and Applications; CRC Press: New York, NY, USA, 2013.
4. Guyon, I.; Elisseeff, A. An introduction to variable and feature selection. J. Mach. Learn. Res. 2003, 3, 1157–1182.
5. Dash, M.; Liu, H. Feature Selection for Classification. Intell. Data Anal. 1997, 1, 131–156. [CrossRef]
6. Souza, F.; Araujo, R.; Mendes, J. Review of soft sensor methods for regression applications. Chemom. Intell. Lab. Syst. 2016, 152, 69–79. [CrossRef]
7. Zhou, Y.; Kwong, S.; Guo, H.; Zhang, X.; Zhang, Q. A Two-Phase Evolutionary Approach for Compressive Sensing Reconstruction. IEEE Trans. Cybern. 2017, 47, 2651–2663. [CrossRef]
8. Guan, S.U.; Liu, J.; Qi, Y. An incremental approach to contribution-based feature selection. J. Intell. Syst. 2004, 13, 15–42. [CrossRef]
9. Reunanen, J. Overfitting in making comparisons between variable selection methods. J. Mach. Learn. Res. 2003, 3, 1371–1382.
10. Inbarani, H.H.; Azar, A.T.; Jothi, G. Supervised hybrid feature selection based on PSO and rough sets for medical diagnosis. Comput. Methods Programs Biomed. 2014, 113, 175–185. [CrossRef]
11. Wang, X.; Yang, J.; Teng, X.; Xia, W.; Jensen, R. Feature selection based on rough sets and particle swarm optimization. Pattern Recognit. Lett. 2007, 28, 459–471. [CrossRef]
12. Samraj, A.; Inbarani, H.H.; Banu, N. Unsupervised hybrid PSO-Quick reduct approach for feature reduction. In Proceedings of the 2012 International Conference on Recent Trends in Information Technology, Tamil Nadu, India, 19–21 April 2012.
13. Dong, H.; Li, T.; Ding, R.; Sun, J. A novel hybrid genetic algorithm with granular information for feature selection and optimization. Appl. Soft Comput. 2018, 65, 33–46. [CrossRef]
14. Nakariyakul, S.; Casasent, D. An improvement on floating search algorithms for feature subset selection. Pattern Recognit. 2009, 42, 1932–1940. [CrossRef]
15. Engelbrecht, A.P. Computational Intelligence: An Introduction, 2nd ed.; John Wiley & Sons: Chichester, UK, 2007.
16. Mendes, J.; Araujo, R.; Matias, T.; Seco, R.; Belchior, C. Automatic extraction of the fuzzy control system by a hierarchical genetic algorithm. Eng. Appl. Artif. Intell. 2014, 29, 70–78. [CrossRef]
17. Kennedy, J.; Eberhart, R.C. A discrete binary version of the particle swarm algorithm. In Proceedings of the 1997 IEEE International Conference on Systems, Man, and Cybernetics, Computational Cybernetics and Simulation, Orlando, FL, USA, 12–15 October 1997.
18. Tran, B.; Xue, B.; Zhang, M. A New Representation in PSO for Discretization-Based Feature Selection. IEEE Trans. Cybern. 2018, 48, 1733–1746. doi:10.1109/TCYB.2017.2714145. [CrossRef] [PubMed]
19. Tran, B.; Xue, B.; Zhang, M. Bare-Bone Particle Swarm Optimisation for Simultaneously Discretising and Selecting Features for High-Dimensional Classification. In Proceedings of the European Conference on the Applications of Evolutionary Computation, Porto, Portugal, 30 March–1 April 2016; pp. 701–718.
20. Huang, X.; Chi, Y.; Zhou, Y. Feature Selection of High Dimensional Data by Adaptive Potential Particle Swarm Optimization. In Proceedings of the 2019 IEEE Congress on Evolutionary Computation (CEC), Wellington, New Zealand, 10–13 June 2019; pp. 1052–1059.

21. Lin, J.; Zhou, Y.; Kang, J. An Improved Discretization-Based Feature Selection via Particle Swarm Optimization. In Knowledge Science, Engineering and Management; Douligeris, C., Karagiannis, D., Apostolou, D., Eds.; Springer International Publishing: Cham, Switzerland, 2019; pp. 298–310.

22. Fayyad, U.; Irani, K.B. Multi-Interval Discretization of Continuous-Valued Attributes for Classification Learning. Mach. Learn. 1993, 2, 1022–1027. Available online: http://hdl.handle.net/2014/35171 (accessed on 27 May 2020).

23. Li, M.; Wang, Z. A hybrid coevolutionary algorithm for designing fuzzy classifiers. Inf. Sci. 2009, 179, 1970–1983. [CrossRef]

24. Garcia-Pedrajas, N.; Hervas-Martinez, C.; Ortiz-Boyer, D. Cooperative coevolution of artificial neural network ensembles for pattern classification. IEEE Trans. Evol. Comput. 2005, 9, 271–302. [CrossRef]

25. Yang, Z.; Tang, K.; Yao, X. Large scale evolutionary optimization using cooperative coevolution. Inf. Sci. 2008, 178, 2985–2999. [CrossRef]

26. Gong, M.; Li, H.; Luo, E.; Liu, J.; Liu, J. A Multiobjective Cooperative Coevolutionary Algorithm for Hyperspectral Sparse Unmixing. IEEE Trans. Evol. Comput. 2017, 21, 234–248. [CrossRef]

27. Ma, X.; Li, X.; Zhang, Q.; Tang, K.; Liang, Z.; Xie, W.; Zhu, Z. A Survey on Cooperative Co-Evolutionary Algorithms. IEEE Trans. Evol. Comput. 2019, 23, 421–441. [CrossRef]

28. Kennedy, J.; Eberhart, R. Particle swarm optimization. In Proceedings of the ICNN’95—International Conference on Neural Networks, Perth, Australia, 27 November–1 December 1995.

29. Holland, J.H. Bibliography. In Adaptation in Natural and Artificial Systems: An Introductory Analysis with Applications to Biology, Control, and Artificial Intelligence; MIT Press: Cambridge, MA, USA, 1992; pp. 203–205.

30. Chicano, F.; Sutton, A.M.; Whitley, L.D.; Alba, E. Fitness probability distribution of bit-flip mutation. Evol. Comput. 2015, 23, 217–248. [CrossRef]

31. Umbarkar, A.J.; Sheth, P.D. Crossover Operators in Genetic Algorithms:A Review. ICTACT J. Soft Comput. 2015, 6, 1083–1092.

32. Al-Sahaf, H.; Zhang, M.; Johnston, M.; Verma, B. Image descriptor: A genetic programming approach to multiclass texture classification. In Proceedings of the 2015 IEEE Congress on Evolutionary Computation (CEC), Sendai, Japan, 25–28 May 2015; pp. 2460–2467.

33. Tran, B.Q.; Zhang, M.; Xue, B. A PSO based hybrid feature selection algorithm for high-dimensional classification. In Proceedings of the 2016 IEEE Congress on Evolutionary Computation (CEC), Vancouver, BC, Canada, 24–29 July 2016; pp. 3801–3808.

34. Martino, A.; Giuliani, A.; Rizzi, A. (Hyper)Graph Embedding and Classification via Simplicial Complexes. Algorithms 2019, 12, 223. [CrossRef]

35. Martino, A.; Giuliani, A.; Todde, V.; Bizzarri, M.; Rizzi, A. Metabolic networks classification and knowledge discovery by information granulation. Comput. Biol. Chem. 2020, 84, 107187. [CrossRef] [PubMed]

36. Liang, J.J.; Qin, A.K.; Suganthan, P.N.; Baskar, S. Comprehensive learning particle swarm optimizer for global optimization of multimodal functions. IEEE Trans. Evol. Comput. 2006, 10, 281–295. [CrossRef]

37. Kennedy, J. Bare bones particle swarms. In Proceedings of the 2003 IEEE Swarm Intelligence Symposium, Indianapolis, IN, USA, 26–26 April 2003.