

Chapter 7

Unsupervised Feature Selection

Feature Selection (FS) is a remedy that can reduce the dimensionality of data without degrading the computational efficiency of the problem. Conventional feature subsets are selected based on some evaluation metric that is estimated based on its dependency or relation to decision class value in the data. However, not all the real world problems have label or class attached to them thereby creating the need for FS in unsupervised domain. FS in unsupervised domain, in layman terms, selects those feature subsets that produce best groupings or clusters among individuals [42]. Some of the unsupervised approaches that utilized clustering as a criteria to evaluate feature subsets discussed previously were based on expectation maximisation [41], sequential FS [30], entropy based techniques [7], FS based on genetic algorithm [84], multi cluster feature selection [19]. Similarity based unsupervised FS was employed by Mitra et. al. [109] to avoid redundancy. Unsupervised FS was proposed using ant colony optimization technique in [139]. Adaptive and embedding learning based unsupervised FS adaptively learned the embeddings that preserved manifold structure was successfully used by Wu [161] while Liu et. al. [97] employed neighbourhood embedding for FS. Lim et. al. [93] employed pairwise dependence among features

for FS. Principal component and Linear Discriminant Analysis change the semantics of data while performing dimensionality reduction. However, all these approaches ignore the fuzziness existing in the dataset.

Fuzzy rough set theory offers a methodological solution to reduce the dataset dimensionality by selecting the most informative features. It efficiently reduces the dataset size as only set operations are involved, computational overload is also reduced. The entire process is accomplished without requiring any human intervention or any prior information. Most importantly, the underlying semantics of the dataset is preserved. Only some researchers have worked on FS using fuzzy rough set theory in unsupervised learning case, where class labels are unknown or missing. Parthala et. al. [116] employed fuzzy rough set theory technique for FS on datasets without class information. Wang et. al. [159] combined the idea of fuzzy and sparse learning for unsupervised FS. While embeddings of fuzzy membership was used by Zhang et. al. [172]. The performance of fuzzy rough set theory model is dependent on the quality of feature subsets selected for evaluation while maintaining the computation time to a low value. However, none of the researchers have discussed this issue. Swarm intelligence models the social behaviour of animals by having a population of artificial agents that perform simple task while co-operatively solving hard optimization problem. This chapter proposes the novel method of selecting relevant, non redundant high quality and information rich subset of features employing a metaheuristic earthworm optimization thereby taking into consideration the fuzziness existing in the real world datasets.

7.1 Feature Selection based on Fuzzy Rough Set in Unsupervised Domain

In case of supervised approach, a decision class is associated with each instance. Based on the value of degree of dependency, the quality of feature subset can be evaluated. The reduct comprises of the minimal feature subset that maintains the quality of original dataset. Any search technique like greedy hill climbing algorithm can be applied to construct feature subset.

Extending the idea to unsupervised domain, decision class D can be replaced by non-intersecting subset of features \bar{B} i.e $A \cap B = \phi$ to compute dependency of A as:

$$\gamma_A(B) = \frac{\sum_{x_i \in U} Pos_A(B)(x_i)}{|U|} \quad (7.1)$$

The positive region and thereby the lower and upper approximations [116] are defined as:

$$Pos_A(B)(x_i) = sup_{z \in U} R \downarrow_A R_{B_z}(x_i) \quad (7.2)$$

where R_{B_z} is the fuzzy similarity relation for sample z (say).

$$R \downarrow_A R_{B_z}(x_i) = inf_{x_j} I(R_A(x_i, x_j), R_B(x_j, z)) \quad (7.3)$$

$$R \uparrow_A R_{B_z}(x_i) = sup_{x_j} T(R_A(x_i, x_j), R_B(x_j, z)) \quad (7.4)$$

The lower approximation describes the certainty with which an instance belongs to a set. The upper approximation gives the possibility of an object belonging to a set. Usually, lower approximation is used to evaluate the quality of feature subsets, while the information contained in upper approximation is not considered. For two subsets having same lower approximation, the one containing lower upper approximation is more accurate reflection of original content. Therefore, the information contained in upper approximation should be utilized, boundary region (which is given by difference between upper and lower approximation) is such a measure that undertakes both lower and upper approximation into account. The value of boundary region

decreases as search progresses towards optimal subset position until the lowest value is reached. The total certainty degree is employed instead of dependency degree to evaluate feature subset quality as the following:

$$\gamma_A(B) = 1 - \frac{\sum_{z \in U} \sum_{x_i \in U} BND_A(B)(x_i)}{|U|^2} \quad (7.5)$$

This measure can then be used to guide the feature selection task.

7.1.1 Feature subset quality evaluation

The quality of feature subset must be determined to guide the search towards optimal feature subset selection. The quality of subset of features is examined using dependency and boundary region measure in this work.

7.1.1.1 Dependency measure

Let A be the subset of attributes. if the non-selected feature subset $B = C - A$ depends on the selected subset A , then B can be effectively eliminated as the information content in B would be redundant. Dependency degree can be utilized to compute dependence of B on A using the following formula:

$$\gamma_A(B) = \frac{\sum_{x_i \in U} Pos_A(B)(x_i)}{|U|} \quad (7.6)$$

The positive region is defined in the same way as equation (7.2) considering that each object belongs to its own class:

$$Pos_A(B)(x_i) = R_A \downarrow R_B(x_i) \quad (7.7)$$

The modified lower and upper approximation are defined as:

$$R_A \downarrow R_B(x_i) = \inf_{x_j} I(R_A(x_i, x_j), R_B(x_i, x_j)) \quad (7.8)$$

$$R_A \uparrow R_B(x_i) = \sup_{x_j} T(R_A(x_i, x_j), R_B(x_i, x_j)) \quad (7.9)$$

On the similar lines, boundary region measure can also be defined as:

$$BND_A(B)(x_i) = R_A \uparrow R_B(x_i) - R_B \downarrow R_B(x_i) \quad (7.10)$$

The corresponding measure to evaluate feature quality is given as follows:

$$\gamma_A(B) = 1 - \frac{\sum_{x_i \in U} BND_A(B)(x_i)}{|U|} \quad (7.11)$$

We illustrate the concept via a toy example shown in Table 7.1. Here, standard t-norm and implicator are applied for the computation. The similarity measure given below equation is employed for this example.

$$R_a(x_i, x_j) = \max\left(\min\left(\frac{(a(x_j) - (a(x_i) - std_a))}{std_a}, \frac{((a(x_i) + std_a) - a(x_i))}{std_a}, 0\right)\right)$$

where std_a is the standard deviation of feature a . Suppose features $A = a_1, a_5, a_6$ are selected, then the similarity values between each pair of instances are computed and shown in Table 7.2. On similar lines, the similarity values of non-selected features $B = a_2, a_3$ are noted down in Table 7.3. Based on these values, the lower approximation is evaluated as:

$$\begin{aligned} R_A \downarrow R_B(x_1) &= \inf(I(1, 1), I(0, 0.801175), I(0, 0.801175), \\ &I(0, 0.92047), I(0.242991, 0), I(0, 0.045641), \\ &I(0, 0), I(0, 0)) = 0.7570 \\ R_A \downarrow R_B(x_2) &= \inf(I(0, 0.801175), I(1, 1), I(0, 0.602351), \\ &I(0, 0.880705), I(0, 0), I(0, 0), I(0, 0), I(0, 0)) = 1 \end{aligned}$$

Similarly, the method is iterated for all the instances and the corresponding positive region is calculated as:

$$\begin{aligned} Pos_A(B)(x_3) &= R_A \downarrow R_B(x_3) = 0.9147, \\ Pos_A(B)(x_4) &= \overline{R_C} \downarrow R_B(x_4) = 1, \\ Pos_A(B)(x_5) &= R_A \downarrow R_B(x_5) = 0.7570, \\ Pos_A(B)(x_6) &= \overline{R_C} \downarrow R_B(x_6) = 0.9147, \\ Pos_A(B)(x_7) &= R_A \downarrow R_B(x_7) = 1, \\ Pos_A(B)(x_8) &= \overline{R_C} \downarrow R_B(x_8) = 1 \end{aligned}$$

TABLE 7.1: Example dataset

Features \ Instances	a_1 a_2			a_3	a_4	a_5
	x_1	0.08	0.08	0.1	0.24	0.9
x_2	0.06	0.06	0.05	0.25	0.33	
x_3	0.1	0.1	0.15	0.65	0.3	
x_4	0.08	0.08	0.08	0.98	0.24	
x_5	0.09	0.15	0.4	0.1	0.66	
x_6	0.15	0.02	0.34	0.4	0.01	
x_7	0.24	0.75	0.32	0.18	0.86	
x_8	0.276	0.225	0.81	0.27	0.33	

TABLE 7.2: Fuzzy similarity values for selected features

R_A	x_1	x_2	x_3	x_4	x_5	x_6	x_7	x_8
x_1	1	0	0	0	0.242991	0	0	0
x_2	0	1	0	0	0	0	0	0
x_3	0	0	1	0	0	0.08528	0	0
x_4	0	0	0	1	0	0	0	0
x_5	0.242991	0	0	0	1	0	0	0
x_6	0	0	0.08528	0	0	1	0	0
x_7	0	0	0	0	0	0	1	0
x_8	0	0	0	0	0	0	0	1

TABLE 7.3: Fuzzy similarity values for non-selected features

R_B	x_1	x_2	x_3	x_4	x_5	x_6	x_7	x_8
x_1	1	0.801175	0.801175	0.92047	0	0.045641	0	0
x_2	0.801175	1	0.602351	0.880705	0	0	0	0
x_3	0.801175	0.602351	1	0.721645	0.005876	0.244466	0	0
x_4	0.92047	0.880705	0.721645	1	0	0	0	0
x_5	0	0	0.005876	0	1	0.452148	0	0
x_6	0.045641	0	0.244466	0	0.452148	1	0	0
x_7	0	0	0	0	0	0	1	0
x_8	0	0	0	0	0	0	0	1

Hence, the value of dependency degree is given as:

$$\gamma_A(B) = \frac{\sum_{x_i \in U} Pos_A(B)(x_i)}{|U|} = \frac{7.3434}{8} = 0.9179 \quad (7.12)$$

7.1.2 Search strategy

The previous section sets the platform for evaluating the quality of feature subset. There is a need for a search strategy to select optimal feature subset that would lead towards good performance. This is advantageous as whole dataset need not be evaluated for achieving good performance.

Binary earthworm optimization proposed in this study, is inspired by reproductive behaviour of earthworms. Each earthworm produces two kinds of offsprings using variants of reproduction namely reproduction 1 and reproduction 2. The child earthworm is of same size as that of parent. Consider the population of N earthworms. Each individual earthworm is represented by m dimensional binary vector representing presence or absence of attributes with m being the total number of attributes in the dataset. Let the i^{th} earthworm in generation t be denoted by $e^{i,t}$. For moving on to next generation, the earthworms reproduce using reproduction 1 or 2 to produce offsprings. The two reproductions are described as follows:

7.1.2.1 Reproduction 1

Earthworms are hermaphrodites allowing a single parent to generate child earthworm. The offspring is generated in the following way as:

$$e_1^{i,t+1} = 1 - \alpha_o e^{i,t} \quad (7.13)$$

where $e^{i,t}$ denotes the i^{th} earthworm in the present generation t . Likewise, $e_1^{i,t+1}$ denotes the newly generated i^{th} earthworm in generation $t+1$. To allow the value at any position of earthworm to oscillate in 0 and 1, the parameter α_o is chosen such that $\alpha_o \in [0, 1]$ indicating the similarity between child and parent. If $\alpha_o = 0$, the child earthworm has all the values as 1 while when $\alpha_o = 1$ the newly generated earthworm has all the features that are not possessed by parent earthworm. Therefore, the value

of α_o is set to 1 for further experimentation. This leads to global search promoting exploration.

7.1.2.2 Reproduction 2

Reproduction 2 is an improvised version of crossover operator. In this paper, multi-point crossover is employed wherein two earthworms are used to generate a offspring. The two parents $e^{p_1,t}$ and $e^{p_2,t}$ are randomly selected from the population. The i^{th} child earthworm $e_2^{i,t+1}$ is generated using the following equation:

$$e_2^{i,t+1} = [e^{p_1,t}(1 : r_1), e^{p_2,t}(r_1 + 1 : r_2), e^{p_1,t}(r_2 + 1, m)] \quad (7.14)$$

where r_1 and r_2 are two randomly generated integers lying in range $(1, m)$.

Using these two kinds of reproduction, the i^{th} earthworm is generated as:

$$e^i = \begin{cases} 1, \beta_o e_1^{i,t+1} + (1 - \beta_o) e_2^{i,t+1} \geq 0.5 \\ 0, \beta_o e_1^{i,t+1} + (1 - \beta_o) e_2^{i,t+1} < 0.5 \end{cases} \quad (7.15)$$

where the parameter β_o is used to adjust the contributions of the two reproductions.

The value of β_o at $t + 1$ generation is updated using:

$$\beta_o^{t+1} = \sigma \beta_o^t \quad (7.16)$$

where σ is like a cooling factor and has a constant value. For $t = 0$, initial value of β_o is set to 1. The value of β_o is decreased as the number of generation increases implying that the contribution from reproduction 1 decreases with increment of generations and the impact of reproduction 2 becomes more and more dominant. Thus, local search is more significant when the optimization technique is about to reach the end of generations maintaining the balance between exploration and exploitation. Further, in order to head towards optimal solution the worst fitness earthworm at current generation is replaced with best fitness earthworm of previous generation. The fitness of individual earthworm is formulated using the following

equation:

$$Fit_{e^i} = \alpha \times \gamma_A(B) + \beta \times \frac{|m| - |A|}{|m|} \quad (7.17)$$

where $|m|$ and $|A|$ denote the number of total and selected features respectively. α and β are the constants governing the importance of classification performance and subset length respectively, such that $\alpha = 1 - \beta, \alpha \in [0, 1]$. The entire approach is described in Algorithm 7.1.2.2 and illustrated via Figure 7.1.

Algorithm 7.1.2.2 Earthworm search strategy

Input: *generation*: number of generation; N : number of earthworms; α_o : similarity parameter of reproduction 1; β_o : adjustment parameter used for varying the impact of two reproductions; $\sigma = 0.9$: cooling factor used for computing β ; $e^i = m$ bit vector generated randomly; $i = 1, 2, \dots, N$

Evaluate the fitness of each earthworm

$t = 0$;

while $t < \textit{generation}$ **do**

for \forall earthworm $e^{i,t}$ **do**

 # Implement Reproduction 1

 Generate $e_1^{i,t+1}$ using Reproduction 1 using Equation (7.13)

 # Implement Reproduction 2

 Randomly select two parent earthworms $e^{p_1,t}$ and $e^{p_2,t}$

 Generate offspring $e_2^{i,t+1}$ using Equation (7.14)

 Update the i^{th} earthworm at $t + 1$ generation using Equation (7.15)

end for

 Replace the worst earthworm at $t + 1$ generation with the best fitness earthworm at t generation

$t \leftarrow t + 1$;

end while

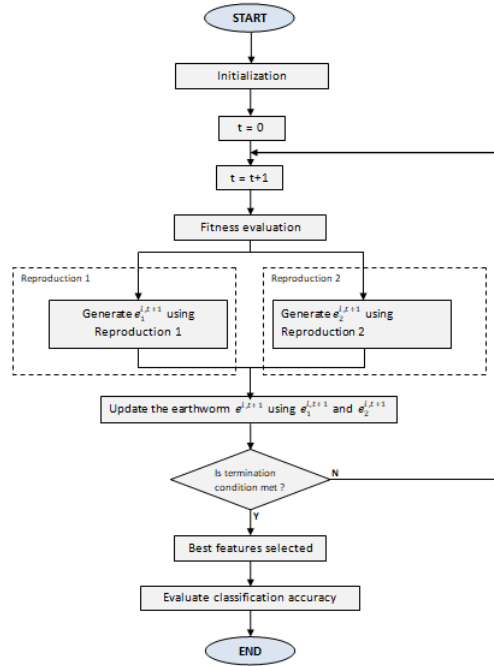


FIGURE 7.1: The flowchart of entire methodology for obtaining a reduced representation of the dataset

return Best fitness earthworm

7.2 Experimentation

The effectiveness of the proposed approach can be illustrated by conducting proper experimentation for the same. The various parameter settings employed for experimentation are as follows: The parameters α and β used in fitness calculation are set to 0.9 and 0.1 respectively. The similarity factor $\alpha_o = 1$, the initial proportional factor β_o is set to 1 and cooling factor σ as 0.9. The value of generation and the number of earthworms are set to 50. These default values as given in [156] are used for further experimentation.

7.2.1 Results

Sixteen different benchmark datasets used for experimental evaluation are taken from UCI repository [3] (as shown in Table 7.4). Performance of proposed approach and its variants is illustrated by splitting the experimentation into four sections namely:

1. The proposed approach (UFRESO) and its variant based on boundary measure (UFRBESO).
2. Comparative analysis of the proposed work with existing dependency based approach.
3. Comparison with previous non dependency based approaches.
4. Applying feature selection taking the class labels into consideration i.e. supervised approach based on fuzzy rough sets and performing the comparative study of the same. It might be unreasonable to compare with supervised approaches as class labels brings in additional information. However, that could help in verifying the fact that the valuable information are retained in case of unsupervised approach also.

The comparison is done in aspects of reduct size, classification accuracy and classification error. Ten fold cross validation is used for performance computation. So, the results are averages of 10 folds. The significance are established using statistical tests namely two tailed student's t test by calculating difference between average accuracy of two models statistically. The level of significance is set to 0.05 for the experiment. The probability p-val associated with t test is also depicted in the tables. The small p-val values (less than 0.05) illustrates the significant differences amongst the algorithms. Various symbols like +, -, o are used to show corresponding statistical win,

TABLE 7.4: Benchmark dataset

Dataset	Instance	Feature	Class	Classification accuracy			
				KNN		SVM	
				Acc	Std	Acc	Std
Auto-univ-au1_1000	1000	20	2	71.70	4.21	74.10	3.57
German	1000	24	2	70.80	6.35	76.60	3.53
Cardiotocography-3class	2126	36	3	98.86	0.71	92.07	10.37
Diabetes	768	8	2	72.50	5.70	64.86	6.35
Leaf	340	14	30	66.47	5.03	48.23	9.32
Abalone-11class	3842	9	11	23.20	2.10	27.11	1.98
Heart-cleveland	303	14	5	55.12	6.74	57.00	6.13
Hepatitis	155	19	2	81.33	7.56	84.00	7.16
Ionosphere	351	34	2	84.28	4.71	88.00	5.00
Fertility-diagnosis	100	9	2	88.00	1.35	88.00	12.29
Flags-religion	194	28	8	48.42	10.46	44.21	9.98
Lung-cancer	32	57	3	43.33	22.49	46.67	32.20
Lymphography	148	18	4	85.00	10.35	80.71	12.16
Trains	10	26	2	50.00	52.70	70.00	48.30
Wine	178	13	3	95.29	6.67	95.29	5.40
Semeion	1593	257	10	90.94	2.26	93.63	1.91

loss and tie respectively of the respective approach at 5% level of significance. Two classifiers namely KNN (with 3 nearest neighbours) [91] and SVM [81] are used for learning.

7.2.1.1 Using variants of proposed approach

The various performance measures i.e. reduct size, accuracy along with standard deviation and error are shown in Tables 7.5, 7.6 and 7.7 for both the dependency and boundary based measures. It could be clearly seen that there is difference in number of features selected by UFRESO and UFRBESO with latter selecting fewer features than former for most of the datasets. The accuracy measures values are nearly same as that obtained from original unreduced dataset except for leaf and lymphography for which low value is obtained which can be justified by unsupervised

TABLE 7.5: Number of features selected by employing variants of proposed approach

Dataset	UFRESO	UFRBESO
Auto-univ-au1_1000	15.1	6.3
German	17.3	8.5
Cardiotocography-3class	22.1	11.2
Diabetes	6.6	1.5
Leaf	10.9	3.2
Abalone-11class	7.1	1.7
Heart-cleveland	10.5	3.1
Hepatitis	14.3	5.3
Ionosphere	16.0	17.4
Fertility-diagnosis	8.2	2.1
Flags-religion	19.6	8.6
Lung-cancer	19.1	33.6
Lymphography	13.0	4.9
Trains	8.9	18.8
Wine	10.9	2.6
Semeion	103.9	149.9

nature of dataset. However, hepatitis, fertility-diagnosis and trains yielded higher performance than benchmark ones. On comparing UFRESO and UFRBESO, the classification accuracy of UFRESO is more or comparable with UFRBESO for all the datasets except lung-cancer. The bit lower performance may be attributed to the use of boundary (or possible instances in calculation of approximations). A similar trend could be observed for classification error also.

7.2.1.2 Comparison with state of art dependency based approach

A comparative analysis of the proposed approach with existing state of art algorithm is undertaken in this section. Unsupervised fuzzy rough set based dimensionality reduction (UFRFS) [102] is used for comparison (as shown in Tables 7.8, 7.12 and 7.9). There is increase in classification accuracy for UFRESO for all datasets except diabetes and lung-cancer for which the decrease in insignificant. Less number of

TABLE 7.6: Classification accuracy by employing variants of proposed approach

Dataset		UFRESO		UFRBESO	
		Acc	Std	Acc	Std
Auto-univ-au1_1000	KNN	70.73	2.21	67.86	1.18
	SVM	71.65	2.50	72.81	0.98
German	KNN	73.27	2.61	64.21	4.18
	SVM	77.44	1.48	71.81	1.08
Cardiotocography-3class	KNN	96.21	1.38	93.78	1.53
	SVM	83.18	3.29	76.87	7.35
Diabetes	KNN	70.14	2.58	67.48	0.84
	SVM	63.32	3.43	66.64	0.99
Leaf	KNN	67.74	4.90	32.49	3.61
	SVM	39.14	5.64	11.25	5.94
Abalone-11class	KNN	22.78	0.46	21.81	0.54
	SVM	28.74	1.02	23.46	1.31
Heart-cleveland	KNN	56.20	2.55	53.70	5.17
	SVM	60.28	2.98	55.58	1.81
Hepatitis	KNN	88.21	3.05	78.39	4.8
	SVM	91.26	2.38	76.95	4.11
Ionosphere	KNN	87.86	1.24	87.57	2.51
	SVM	85.23	0.99	86.63	2.91
Fertility-diagnosis	KNN	88.29	1.61	88.78	4.18
	SVM	93.36	2.54	91.14	3.67
Flags-religion	KNN	51.26	2.71	38.03	4.09
	SVM	39.90	1.89	38.02	4.53
Lung-cancer	KNN	37.33	3.69	57.29	5.42
	SVM	44.51	9.28	49.71	6.56
Lymphography	KNN	78.98	4.01	61.97	2.83
	SVM	83.89	1.46	68.13	3.14
Trains	KNN	72.32	22.12	74.13	19.11
	SVM	90.18	10.73	90.18	10.73
Wine	KNN	94.54	2.34	76.48	2.54
	SVM	93.74	2.66	82.17	5.14
Semeion	KNN	89.20	0.06	88.72	0.29
	SVM	89.81	0.05	92.30	0.79

TABLE 7.7: Classification error results for variants of proposed approach

Dataset		Classification error	
		UFRESO	UFRBESO
Auto-univ-au1_1000	KNN	0.29	0.33
	SVM	0.26	0.27
German	KNN	0.29	0.33
	SVM	0.24	0.28
Cardiotocography-3class	KNN	0.03	0.07
	SVM	0.17	0.36
Diabetes	KNN	0.28	0.32
	SVM	0.34	0.34
Leaf	KNN	4.07	7.96
	SVM	6.47	11.25
Abalone-11class	KNN	1.59	1.83
	SVM	1.39	1.54
Heart-cleveland	KNN	0.71	0.86
	SVM	0.63	0.75
Hepatitis	KNN	0.14	0.18
	SVM	0.10	0.20
Ionosphere	KNN	0.11	0.12
	SVM	0.14	0.13
Fertility-diagnosis	KNN	0.10	0.15
	SVM	0.07	0.12
Flags-religion	KNN	1.47	1.55
	SVM	1.79	1.70
Lung-cancer	KNN	0.70	0.50
	SVM	0.73	0.53
Lymphography	KNN	0.21	0.40
	SVM	0.17	0.32
Trains	KNN	0.50	0.30
	SVM	0.20	0.20
Wine	KNN	0.05	0.25
	SVM	0.05	0.24
Semeion	KNN	0.50	0.50
	SVM	0.41	0.27

TABLE 7.8: Number of features selected on comparison with other state of art feature selection algorithm

Dataset	UFRFS	UFRESO
Auto-univ-au1_1000	19.4	15.1
German	17.3	17.3
Cardiotocography-3class	20.6	22.1
Diabetes	8.0	6.6
Leaf	7.2	10.9
Abalone-11class	8.0	7.1
Heart-cleveland	11.0	10.5
Hepatitis	15.1	14.3
Ionosphere	19.4	16.0
Fertility-diagnosis	8.9	8.2
Flags-religion	1.0	19.6
Lung-cancer	32.1	19.1
Lymphography	15.2	13.0
Trains	10.1	8.9
Wine	6.0	10.9
Semeion	50.9	103.9

features are selected by UFRESO for most of the datasets along with high performance. A lower classification error is observed by UFRESO by most of the datasets. Statistical results illustrate that UFRESO has achieved better or comparable performance and it loses only for one dataset namely diabetes for KNN classifier while for SVM, UFRESO loses for three and wins for seven. The statistical results lay down the superiority of UFRESO.

7.2.1.3 Comparison with state of art non dependency based approach

Four state of art algorithms namely Spectral feature selection for supervised and unsupervised learning (USFS) [175], Unsupervised feature selection for multi cluster data (UFSMC) [19], Laplacian score for feature selection (LSFS) [59] and Unsupervised Feature Selection using Feature Similarity (FSFS) [109] has been employed for comparison and the results are noted down in Tables 7.13, 7.14 and 7.15. Cai et. al.

TABLE 7.9: Classification error comparison with other state of art feature selection algorithm

Dataset		Classification error	
		UFRFS	UFRESO
Auto-univ-au1_1000	KNN	0.35	0.29
	SVM	0.26	0.26
German	KNN	0.34	0.29
	SVM	0.23	0.24
Cardiotocography-3class	KNN	0.03	0.03
	SVM	0.29	0.17
Diabetes	KNN	0.30	0.28
	SVM	0.33	0.34
Leaf	KNN	0.04	4.07
	SVM	0.06	6.47
Abalone-11class	KNN	0.14	1.59
	SVM	0.16	1.39
Heart-cleveland	KNN	0.18	0.71
	SVM	0.27	0.63
Hepatitis	KNN	0.26	0.14
	SVM	0.15	0.10
Ionosphere	KNN	0.11	0.11
	SVM	0.13	0.14
Fertility-diagnosis	KNN	0.17	0.10
	SVM	0.12	0.07
Flags-religion	KNN	0.20	1.47
	SVM	0.20	1.79
Lung-cancer	KNN	0.38	0.70
	SVM	0.34	0.73
Lymphography	KNN	0.16	0.21
	SVM	0.27	0.17
Trains	KNN	0.51	0.50
	SVM	0.20	0.20
Wine	KNN	0.12	0.05
	SVM	0.26	0.05
Semeion	KNN	0.08	0.50
	SVM	0.16	0.41

[19] showing UFSMC to be effective in preserving cluster structure. All these works are parameter driven while the proposed approach is data driven. The number of features selected by UFRESO are significantly less than obtained by UFEMC, USFS, LSFS and FSFS. The significant increase in accuracy for all the datasets except a minute decrease for auto-univ-au1_1000, cardiocography-3class and diabetes can also be observed from paired t-test results. But regardless of a bit increase for three datasets, the need for supplying parameter values beforehand makes the USFS, UFSMC, LSFS and FSFS approaches user dependent. Further, the number of wins and losses clearly indicate that UFRESO has won the match. Exactly same pattern is observed for classification error also.

7.2.1.4 Comparison with supervised approach

In this section, fuzzy rough set based feature selection [72] using ant colony (FRASO) and particle swarm (FRPSO) search heuristic is used for comparison. Although the comparison of supervised and unsupervised approach is unreasonable, however that would be effective in laying the supremacy of the proposed work. Tables 7.10, 7.16 and 7.11 depicts the recorded results. The number of features selected by supervised approaches is comparatively less than that obtained by UFRESO. The presence of class labels adds some information that is utilized by supervised approaches, which can not be taken as an advantage in case of unsupervised methods. Considering classification accuracy, there is similar trend in accuracy for both supervised and unsupervised approaches for most of datasets, again because of discrimination information supplied by decision class. However, datasets namely heart-cleveland, fertility-diagnosis, flags-religion, trains and semeion have produced high performance with UFRESO that may have resulted because of super set of features selected by

TABLE 7.10: Number of features selected on comparison with other state of art supervised feature selection algorithms

Dataset	FRPSO	FRASO	UFRESO
Auto-univ-au1_1000	16.3	1.0	15.1
German	13	16.5	17.3
Cardiotocography-3class	10.8	19.4	22.1
Diabetes	8.0	8.0	6.6
Leaf	11.7	13.1	10.9
Abalone-11class	8.0	8.0	7.1
Heart-cleveland	8.1	7.7	10.5
Hepatitis	7.1	6.9	14.3
Ionosphere	7.0	7.8	16.0
Fertility-diagnosis	6.9	6.9	8.2
Flags-religion	1.0	2.0	19.6
Lung-cancer	5.4	5.5	19.1
Lymphography	7.3	7.3	13.0
Trains	1.0	2.0	8.9
Wine	4.8	4.8	10.9
Semeion	22.5	18.4	103.9

UFRESO. Statistical results demonstrates the supremacy of UFRESO. Classification error further reflects the better performing behaviour of UFRESO.

7.3 Summary

Two fuzzy rough set based approaches employing dependency degree and boundary measure are proposed in this study for feature selection in unsupervised domain. The two phases of feature selection namely feature subset selection and subset quality evaluation has been dealt in this work. The candidate feature subsets are selected using proposed earthworm search strategy. The use of search strategies have further enhanced the performance. The feature subset quality is evaluated considering fuzziness arising in real world applications.

TABLE 7.11: Classification error comparison with other state of art supervised feature selection algorithms

Dataset		Classification error		
		FRPSO	FRASO	UFRESO
Auto-univ-au1_1000	KNN	0.36	0.38	0.29
	SVM	0.26	0.26	0.26
German	KNN	0.32	0.33	0.29
	SVM	0.33	0.29	0.24
Cardiotocography-3class	KNN	0.04	0.03	0.03
	SVM	0.23	0.23	0.17
Diabetes	KNN	0.30	0.30	0.28
	SVM	0.35	0.35	0.34
Leaf	KNN	0.03	0.03	4.07
	SVM	0.06	0.06	6.47
Abalone-11class	KNN	0.14	0.14	1.59
	SVM	0.16	0.16	1.39
Heart-cleveland	KNN	0.19	0.20	0.71
	SVM	0.27	0.27	0.63
Hepatitis	KNN	0.22	0.22	0.14
	SVM	0.15	0.15	0.10
Ionosphere	KNN	0.12	0.15	0.11
	SVM	0.17	0.18	0.14
Fertility-diagnosis	KNN	0.18	0.20	0.10
	SVM	0.12	0.12	0.07
Flags-religion	KNN	0.20	0.20	1.47
	SVM	0.20	0.20	1.79
Lung-cancer	KNN	0.40	0.41	0.70
	SVM	0.36	0.33	0.73
Lymphography	KNN	0.13	0.13	0.21
	SVM	0.27	0.27	0.17
Trains	KNN	0.60	0.54	0.50
	SVM	0.20	0.10	0.20
Wine	KNN	0.05	0.06	0.05
	SVM	0.24	0.25	0.05
Semeion	KNN	0.08	0.10	0.50
	SVM	0.16	0.16	0.41

TABLE 7.12: Classification accuracy comparison with other state of art feature selection algorithm

Dataset		UFRFS			UFRESO	
		Acc	Std	p-val	Acc	Std
Auto-univ-au1_1000	KNN	72.6	3.53	0.17 o	70.73	2.21
	SVM	74.1	0.53	0.01 +	71.65	2.5
German	KNN	69.3	3.95	0.02 -	73.27	2.61
	SVM	76.9	4.09	0.70 o	77.44	1.48
Cardiotocography-3class	KNN	96.66	1.53	0.50 o	96.21	1.38
	SVM	77.85	0.14	0.00 -	83.18	3.29
Diabetes	KNN	73.69	3.03	0.01 +	70.14	2.58
	SVM	67.31	2.97	0.01 +	63.32	3.43
Leaf	KNN	51.47	7.63	0.00 -	67.74	4.9
	SVM	26.76	6.71	0.00 -	39.14	5.64
Abalone-11class	KNN	23.2	2.1	0.54 o	22.78	0.46
	SVM	26.92	2.18	0.03 -	28.74	1.02
Heart-cleveland	KNN	56.46	6.51	0.91 o	56.2	2.55
	SVM	58.14	5.32	0.28 o	60.28	2.98
Hepatitis	KNN	78.71	8.93	0.01 -	88.21	3.05
	SVM	85.21	9.32	0.06 o	91.26	2.38
Ionosphere	KNN	89.46	3.31	0.17 o	87.86	1.24
	SVM	86.9	5.07	0.32 o	85.23	0.99
Fertility-diagnosis	KNN	89	7.38	0.77 o	88.29	1.61
	SVM	88	4.22	0.00 -	93.36	2.54
Flags-religion	KNN	30.95	0.82	0.00 -	51.26	2.71
	SVM	30.95	0.82	0.00 -	39.9	1.89
Lung-cancer	KNN	48.33	22.84	0.15 o	37.33	3.69
	SVM	57.5	15.91	0.04 +	44.51	9.28
Lymphography	KNN	72.81	9.6	0.08 o	78.98	4.01
	SVM	83.05	10.26	0.80 o	83.89	1.46
Trains	KNN	40	51.64	0.09 o	72.32	22.12
	SVM	80	42.16	0.47 o	90.18	10.73
Wine	KNN	84.77	10.37	0.01 -	94.54	2.34
	SVM	87.03	9.6	0.05 -	93.74	2.66
Semeion	KNN	64.15	3.85	0.00 -	89.2	0.06
	SVM	63.58	4.31	0.00 -	89.81	0.05
Loss/Win/Tie	KNN	6/1/9				
	SVM	7/3/6				

TABLE 7.13: Number of features selected on comparison with other state of art non dependency based feature selection algorithms

Dataset	UFSMC	USFS	LSFS	FSFS	UFRESO
Auto-univ-au1_1000	17.7	20.0	16.8	18.0	15.1
German	21.3	23.0	17.9	19.5	17.3
Cardiotocography-3class	30.0	19.4	35.0	31.0	22.1
Diabetes	8.0	8.0	8.0	7.0	6.6
Leaf	14.0	14.0	13.7	13.0	10.9
Abalone-11class	8.0	8.0	8.0	7.0	7.1
Heart-cleveland	12.9	12.9	11.2	12.0	10.5
Hepatitis	19.0	17.7	19.0	16.3	14.3
Ionosphere	28.4	28.1	32.0	30.0	16.0
Fertility-diagnosis	8.9	9.0	9.0	8.0	8.2
Flags-religion	28.0	27.6	28.0	25.0	19.6
Lung-cancer	54.3	41.9	36.3	46.9	19.1
Lymphography	17.9	13.8	15.9	15.9	13.0
Trains	13.3	21.7	10.3	20.6	8.9
Wine	11.3	11.8	13.0	12.0	10.9
Semeion	171.9	166.9	197.7	195.4	103.9

TABLE 7.14: Classification accuracy comparison with other state of art non dependency based feature selection algorithms

Dataset		UFSMC			USFS			LSFS			FSFS			UFRESO	
		Acc	Std	p-val	Acc	Std	p-val	Acc	Std	p-val	Acc	Std	p-val	Acc	Std
Auto-univ-au1_1000	KNN	70	4.47	0.65 o	70.89	4.29	0.52 o	70.3	4.89	0.80 o	69.14	1	0.05 -	70.73	2.21
	SVM	73.5	2.41	0.11 o	74.3	2.53	0.03 +	74	2.68	0.06 o	71.64	1.52	0.99 o	71.65	2.5
German	KNN	71.5	3.93	0.25 o	67.1	2.73	0.00 -	67.9	3.36	0.00 -	67.67	1.14	0.00 -	73.27	2.61
	SVM	70.8	4.77	0.00 -	68.7	7.45	0.00 -	66	13.94	0.02 -	74.73	0.8	0.00 -	77.44	1.48
Cardiotocography-3class	KNN	83.06	10.71	0.00 -	80.57	9.68	0.00 -	79.3	11.38	0.00 -	98.39	0.34	0.00 +	96.21	1.38
	SVM	89.73	12.42	0.12 o	73.83	20.24	0.17 o	70.67	19.73	0.06 o	82.71	8.79	0.88 o	83.18	3.29
Diabetes	KNN	69	4.64	0.51 o	69	4.64	0.51 o	69	4.64	0.51 o	68.14	3.79	0.18 o	70.14	2.58
	SVM	67.19	3.29	0.02 +	67.19	3.29	0.02 +	67.19	3.29	0.02 +	62.55	2.23	0.56 o	63.32	3.43
Leaf	KNN	5.88	6.16	0.00 -	5.88	6.16	0.00 -	4.7	4.4	0.00 -	58.91	1.74	0.00 -	67.74	4.9
	SVM	2.64	3.07	0.00 -	2.64	3.07	0.00 -	2.64	3.07	0.00 -	37.68	2.11	0.45 o	39.14	5.64
Abalone-11class	KNN	23.89	6.2	0.58 o	23.89	6.2	0.58 o	23.89	6.2	0.58 o	21.79	0.49	0.00 -	22.78	0.46
	SVM	27.38	7.32	0.57 o	27.38	7.32	0.57 o	27.38	7.32	0.57 o	28.01	0.35	0.05 -	28.74	1.02
Heart-cleveland	KNN	47.82	8.02	0.01 -	49.16	9.74	0.04 -	48.5	8.4	0.01 -	56.85	2.03	0.54 o	56.2	2.55
	SVM	47.69	21.83	0.09 o	53.4	7.02	0.01 -	37.06	17.41	0.00 -	54.47	1.22	0.00 -	60.28	2.98
Hepatitis	KNN	73.33	11.63	0.00 -	73.33	11.63	0.00 -	73.33	11.63	0.00 -	85.92	4.02	0.17 o	88.21	3.05
	SVM	78.7	19.4	0.06 o	77	16.07	0.01 -	79.7	11.55	0.01 -	88.69	6.16	0.23 o	91.26	2.38
Ionosphere	KNN	84.92	9.99	0.37 o	84.36	10.76	0.32 o	81.5	11.4	0.10 o	81.43	4.65	0.00 -	87.86	1.24
	SVM	85.23	9.22	1.00 o	82.36	9.1	0.33 o	81.51	8.56	0.19 o	83.35	4.6	0.22 o	85.23	0.99
Fertility-diagnosis	KNN	86	11.13	0.53 o	86	11.13	0.53 o	86	11.13	0.53 o	81.13	2.65	0.00 -	88.29	1.61
	SVM	88	11.66	0.17 o	88	11.66	0.17 o	88	11.66	0.17 o	81.47	2.49	0.00 -	93.36	2.54
Flags-religion	KNN	38.1	8.13	0.00 -	38.63	8.85	0.00 -	38.1	8.13	0.00 -	37.07	6.22	0.00 -	51.26	2.71
	SVM	28.34	8.3	0.00 -	36.05	14.87	0.43 o	33.1	9.83	0.05 -	37.2	10.19	0.42 o	39.9	1.89
Lung-cancer	KNN	35.83	22.98	0.84 o	39.16	32.5	0.86 o	33.33	27.13	0.65 o	26.95	11.85	0.02 -	37.33	3.69
	SVM	35	21.66	0.22 o	40	21.66	0.55 o	39.16	32.92	0.63 o	24.88	11.16	0.00 -	44.51	9.28
Lymphography	KNN	79.66	6.2	0.77 o	77.71	5.93	0.58 o	78.33	8.96	0.84 o	73.83	3.18	0.01 -	78.98	4.01
	SVM	81.04	13.63	0.52 o	79	10.64	0.17 o	81	13.07	0.50 o	77.93	0.98	0.00 -	83.89	1.46
Trains	KNN	60	48.98	0.48 o	60	48.98	0.48 o	70	45.82	0.89 o	68.75	14.5	0.67 o	72.32	22.12
	SVM	50	50	0.02 -	50	50	0.02 -	80	40	0.45 o	79.71	13.66	0.07 o	90.18	10.73
Wine	KNN	87.05	14.88	0.13 o	85.35	13.79	0.05 -	71.96	13.84	0.00 -	91.03	2.41	0.00 -	94.54	2.34
	SVM	91.01	7.54	0.29 o	87.51	9.99	0.07 o	86.99	12	0.00 -	89.87	2.08	0.00 -	93.74	2.66
Semeion	KNN	89.76	1.92	0.37 o	88.25	4.18	0.48 o	89.13	3.45	0.95 o	87.44	0.65	0.00 -	89.2	0.06
	SVM	87.38	2.27	0.00 -	87.25	1.66	0.00 -	85.56	3.19	0.00 -	92.37	0.87	0.00 +	89.81	0.05
Loss/Win/Tie	KNN	5/0/11			7/0/9			7/0/9			11/1/4				
	SVM	5/1/10			6/2/8			7/1/8			7/1/8				

TABLE 7.15: Classification error comparison with other state of art non dependency based feature selection algorithms

Dataset		Classification error				
		UFSMC	USFS	LSFS	FSFS	UFRESO
Auto-univ-au1_1000	KNN	0.30	0.29	0.29	0.30	0.29
	SVM	0.26	0.25	0.26	0.27	0.26
German	KNN	0.28	0.32	0.32	0.31	0.29
	SVM	0.29	0.31	0.33	0.25	0.24
Cardiotocography-3class	KNN	0.20	0.26	0.27	0.01	0.03
	SVM	0.11	0.40	0.36	0.10	0.17
Diabetes	KNN	0.30	0.30	0.30	0.29	0.28
	SVM	0.32	0.32	0.32	0.36	0.34
Leaf	KNN	11.19	11.19	11.87	4.52	4.07
	SVM	11.72	11.72	12.21	7.05	6.47
Abalone-11class	KNN	1.56	1.56	1.56	1.62	1.59
	SVM	1.37	1.37	1.37	1.35	1.39
Heart-cleveland	KNN	0.91	0.90	0.97	0.72	0.71
	SVM	0.86	0.78	1.05	0.66	0.63
Hepatitis	KNN	0.26	0.26	0.26	0.15	0.14
	SVM	0.21	0.22	0.20	0.16	0.10
Ionosphere	KNN	0.15	0.15	0.16	0.16	0.11
	SVM	0.14	0.17	0.18	0.14	0.14
Fertility-diagnosis	KNN	0.14	0.14	0.14	0.20	0.10
	SVM	0.12	0.12	0.12	0.19	0.07
Flags-religion	KNN	1.70	1.72	1.70	1.70	1.47
	SVM	1.88	1.88	1.80	1.63	1.79
Lung-cancer	KNN	0.67	0.64	0.70	0.70	0.70
	SVM	0.68	0.63	0.64	0.66	0.73
Lymphography	KNN	0.20	0.24	0.22	0.25	0.21
	SVM	0.20	0.23	0.20	0.25	0.17
Trains	KNN	0.40	0.40	0.30	0.30	0.50
	SVM	0.50	0.50	0.20	0.20	0.20
Wine	KNN	0.15	0.17	0.35	0.07	0.05
	SVM	0.08	0.12	0.13	0.07	0.05
Semeion	KNN	0.43	0.52	0.46	0.50	0.50
	SVM	0.51	0.51	0.59	0.35	0.41

TABLE 7.16: Classification accuracy comparison with other state of art supervised feature selection algorithms

Dataset		FRPSO			FRASO			UFRESO	
		Acc	Std	p-val	Acc	Std	p-val	Acc	Std
Auto-univ-au1_1000	KNN	71.1	2.73	0.74 o	74.1	0.32	0.00 -	70.73	2.21
	SVM	74.1	0.32	0.01 +	74.1	0.32	0.01 +	71.65	2.5
German	KNN	72.2	5.22	0.57 o	71.9	4.07	0.38 o	73.27	2.61
	SVM	66.7	12.03	0.01 -	71.2	10.48	0.08 o	77.44	1.48
Cardiotocography-3class	KNN	95.58	1.35	0.32 o	96.47	1.76	0.72 o	96.21	1.38
	SVM	95.81	0.81	0.00 +	96.61	1.58	0.00 +	83.18	3.29
Diabetes	KNN	73.69	3.03	0.01 +	73.69	3.03	0.01 +	70.14	2.58
	SVM	65.11	0.36	0.12 o	65.11	0.36	0.12 o	63.32	3.43
Leaf	KNN	55.88	8.43	0.00 -	57.65	9.53	0.01 -	67.74	4.9
	SVM	47.94	6.8	0.01 +	47.35	8.02	0.02 +	39.14	5.64
Abalone-11class	KNN	23.63	1.71	0.15 o	23.63	1.71	0.15 o	22.78	0.46
	SVM	27.2	1.4	0.01 -	27.2	1.4	0.01 -	28.74	1.02
Heart-cleveland	KNN	54.15	5.9	0.33 o	55.19	8.62	0.73 o	56.2	2.55
	SVM	59.8	6.08	0.83 o	58.14	5.55	0.30 o	60.28	2.98
Hepatitis	KNN	81.29	8.26	0.02 -	83.04	10	0.14 o	88.21	3.05
	SVM	85.13	6.26	0.01 -	84.96	10.49	0.08 o	91.26	2.38
Ionosphere	KNN	88.89	4.14	0.46 o	86.33	4.39	0.30 o	87.86	1.24
	SVM	82.62	4.76	0.11 o	82.35	5.29	0.11 o	85.23	0.99
Fertility-diagnosis	KNN	88	4.22	0.84 o	86	8.43	0.41 o	88.29	1.61
	SVM	88	4.23	0.00 -	88	4.22	0.00 -	93.36	2.54
Flags-religion	KNN	30.95	0.82	0.00 -	30.87	9.28	0.00 -	51.26	2.71
	SVM	30.95	0.82	0.00 -	31.37	9.16	0.01 -	39.9	1.89
Lung-cancer	KNN	43.33	32.82	0.57 +	34.17	23.06	0.67 o	37.33	3.69
	SVM	54.17	28.12	0.32 +	66.67	23.9	0.01 +	44.51	9.28
Lymphography	KNN	78.38	8.75	0.85 -	80.33	12.11	0.74 o	78.98	4.01
	SVM	84.43	10.97	0.88 +	83.1	5.72	0.68 o	83.89	1.46
Trains	KNN	0	0	0.00 -	40	51.64	0.09 o	72.32	22.12
	SVM	80	42.16	0.47 o	90	31.62	0.99 o	90.18	10.73
Wine	KNN	94.41	5.24	0.94 o	92.71	5.33	0.33 o	94.54	2.34
	SVM	92.78	9.82	0.77 o	88.69	8.49	0.09 o	93.74	2.66
Semeion	KNN	70.44	2.97	0.00 -	62.27	4.09	0.00 -	89.2	0.06
	SVM	71.25	2.77	0.00 -	61.57	4.03	0.00 -	89.81	0.05
Loss/Win/Tie	KNN	6/2/8			4/1/11				
	SVM	6/5/5			4/4/8				
