Innovative Approach to Cancer Diagnosis through Hybrid Wrapper-Filter Feature Selection

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ABSTRACT

Feature selection ,as a preprocessing step to machine learning, is effective in reducing

dimensionality, removing irrelevant data and increasing learning accuracy. The development of microarray dataset technology has supplied a large volume of data to many fields. In particular, it has been applied to prediction and diagnosis of cancer, so that it helps us to exactly predict and diagnose cancer. To precisely classify cancer we have to select genes related to cancer. The challenging task in cancer diagnosis is how to identify salient expression genes from thousands of genes in microarray data because extracted genes from microarray dataset have many unwanted datas not related to cancer. In this project we attempt explore a novel hybrid wrapper and filter feature selection algorithm for classification problem using a memetic framework i.e. ,a combination of genetic algorithm (GA) and local search (LS) has been proposed.. The LS is performed using correlation based filter methods are discritize, ranking and redundancy elimination with symmetrical uncertainty (SU) measure .using this hybrid method we can able find can correlated gene, From the larger amount of gene datas .using that smaller dataset doctors can able to find the affected gene and provide better treatment. The efficiency and the effectiveness of the method are demonstrated through extensive comparisons with other methods using real-world datasets of high dimensionality

Keywords: Feature Selection, Memetic Algorithm, Filters ,wrappers, genetic algorithm, symmetrical uncertainty.

I. INTRODUCTION:

Cancer classification, which can help to improve health care of patients and the quality of life of individuals, is essential for cancer diagnosis and drug discovery. An accurate prediction of cancer has greatvalue in providing better treatment and response to therapy varying from different aspects . However, traditional diagnostic methods are mainly based on the morphological and clinical appearance of cancer. They have limited contributions because cancers usually result from many environmental factors, and even the same tumor may have different symptoms under different conditions. Thus, it is necessary to inject systemic approaches into the problemofcancerdiagnosisandprediction.Fromthebio-medicalperspective,eachkindofdiseaseisassociated with certain genes in tissues and the mutation of genes may give rise to the occurrence of certain diseases. Fortunately, the advent of DNA microarray technique, which allows simultaneously measure the expression levels of thousands of genes in a single experiment, makes the accurate prediction of cancer possible and easier. Since it is capable of comparing the gene expression levels in tissues under different conditions, the microarray technique may bring many advantages to cancer prediction and make the diagnosis result more objective, accurate and reliable. During past years, this method has drawn a great deal of attention from both biological and engineering fields

2. Discretization

Thediscretizationistheprocessofdividingthecontinuousdataintoadiscreteone,with help of information theory using **MDL algorithm** based on minimum description length principle.

MDL Algorithm for Discretization

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MDLM(D)

(1) $MDL = \infty$

(2) For all feature subsets L

  1.1 Compute $Length_L = \sum_{i=1}^{i=q} \frac{p_i}{2} \log \frac{|D_L(i)|}{|D_L|} + h_L$

    where $h_L = \frac{1}{2}(N - M)(N + M + 3) \log P + \sum_{i=1}^{i=q} M(M + 3) \log P_i$,

    N – total number of features,

    M – number of features in the candidate subset,

    P – total number of instances in D,

    P_i – number of instances with class label i,

    q – total number of class labels,

    D_L – covariance matrix formed from all the useful feature vectors,

    $D_L(i)$ – covariance matrix formed from the useful feature vectors,

  of class i,

    $|.|$ – denotes determinant.

  1.2 If $Length_L < MDL$ then

    $T = L$, $MDL = Length_L$

(3) Return T

3. RANKING

Applying ranking filter we can obtain a number of top ranked genes. The ranking is done with help of SU which is based on **Information theory.** information in the theory is known as **entropy**,

3.1 Entropy

The **entropy**, H, of a discrete random variable X is a measure of the amount of *uncertainty* associated with the value of X.

(1)

$$H(X) = -\sum_i p(x) \log_2 p(x)$$

3.2 Crossentropy

The **cross entropy** between two probability distributions measures the average number of bits needed to identify an event from a set of possibilities, if a coding scheme is used based on a given probability distribution q, rather than the "true" distribution p.

$$H(p,q) = -\sum_x p(x) \log q(x).$$

3.3 Mutualinformation(transformation)

Mutualinformationmeasurestheamountofinformationthatcanbeobtainedaboutonerandom variable by observing another. The mutual information of X relative to Y is given by:

$$I(X,Y) = \sum_{x,y} p(x,y) \log_2 \frac{p(x,y)}{p(x)p(y)} \quad (3)$$

Whenonevariablebecomescompletelyredundantwiththeknowledgeoftheother.Anothersymmetrical measure is the *symmetric uncertainty* [19], given by (4)
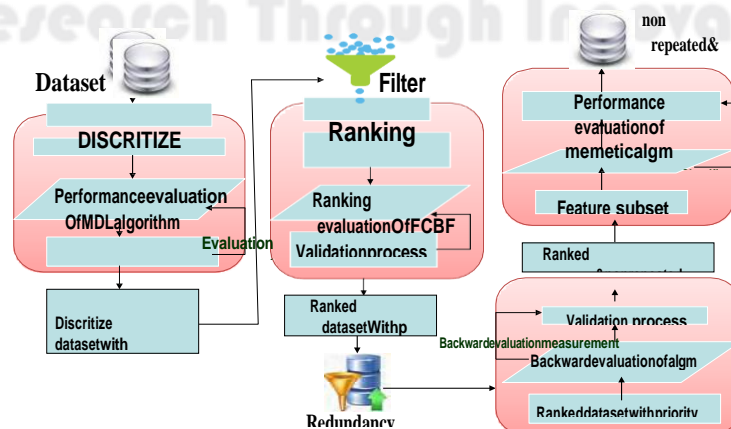
I. SYSTEMDESIGN

Fig1:systnemdesig

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4. RedundancyElimination

Applying redundancy filter to eliminate repetitive gene in the data set. FCBF is an efficient and fast algorithm which uses interdependence of features together with the dependence to the class. FCBF achieves this goal by giving every feature a temporary predominance in the elimination process and making them start eliminating features from the features which are least correlated with the class

FCBFAlgorithm

Input:$S(F1,F2,…,FN,C)$//atrainingdataset
δ //apredefinedthreshold
Output:S_{best} //anoptimalsubset
begin
DiscretizeusingMDLmethod
for i=1to Ndobegin
calculateSUi,cforFi;
if($SUi,c≥\delta$)
append$F$itoS'_{list};
end;
orderS'_{list}indescendingSUi,cvalue;
$Fp=getFirstElement(S'_{list})$;
dobegin
$Fq=getNextElement(S'_{list},Fp)$;
if($Fq<>$NULL)
do begin
$F'q= Fq$;
if($SUp,q≥SUq,c$)
removeFqfromS'_{list};
$Fq=getNextElement(S'_{list,}F'q)$;
else$Fq=getNextElement(S'_{list,}Fq)$;
enduntil($Fq==$NULL);
$Fp=getNextElement(S'_{list,}Fp)$;
enduntil($Fp==$ NULL)
$S_{best}=S'_{list}$;
end;

5. Classification:

the proposed Correlation based Memetic feature selection Algorithm (MA-C) for classification problems which is depicted in the GA population is randomly initialized with each chromosome encoding a candidate feature subset. Subsequently, a local search (LS) is performed. The LS is performed on all gene or portion of the gene, to reach a local optimal solution or to improve the feature subset.
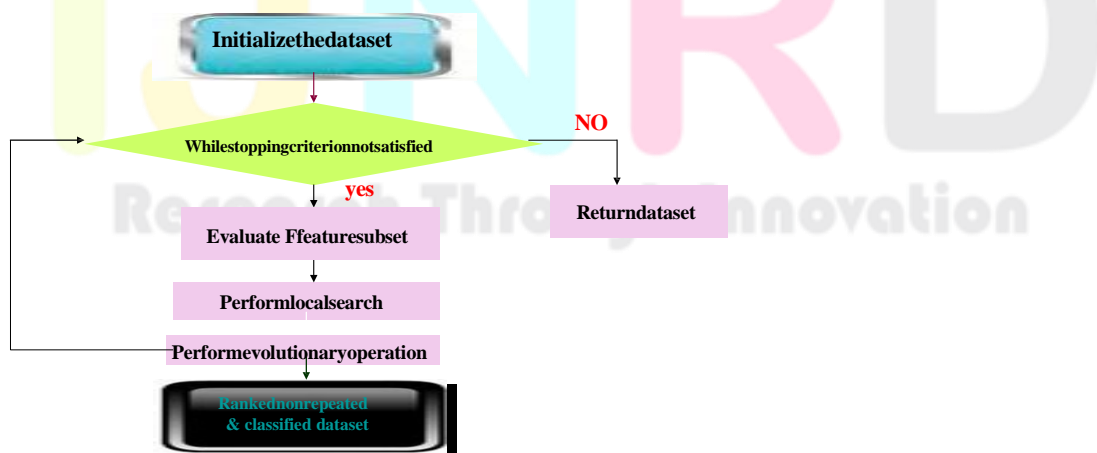

Fig2:memeticsearch

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Dataset	No of genes	No of Samples	No of classes	Max no of genes
CNS	7129	60	2	100
Breast	24481	97	4	100
lung	7129	96	2	100
MLL	10289	102	2	100
LEUKEMIA	7219	72	2	75

Table1:microarraydatasetsusedintheexperiment

Dataset	No of genes	After GA	AfterMA- C
CNS	7120	24	22
Breast	2309	24	23
lung	12601	27	25
MLL	12583	19	19
LEUKIMIA	7130	24	21

Table2:no of instances after global search and memetic search

CONCLUSION

The goalis to improve classification performance and to accelerate the search to identify important feature subsets. In particular, the filter method fine-tunes the population of GA solutions by adding or deleting features based on SU measure. Hence, our focus here is on filter methods that are able to assess the goodness of the individual features. Empirical study of MA-C on several commonly used datasets from the UCI repository , indicates that it outperforms recent existingmethods in theliterature in terms of classification accuracy, selected feature size and efficiency. Further, we also investigate the balance between local and genetic search to maximize the search quality and efficiencyfind the correct gene which is affected. So Implementing this future Enhancement may take a considerable amount of time. Implementing Causes a Very effective system and an error free data pattern.

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