COMPARISON OF K-MEANS AND PAM ALGORITHMS USING CANCER DATASETS

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Keywords: Data mining, Clustering, k-means, PAM.

Abstract: Data mining is a search for relationship and patterns that exist in large database. Clustering is an important datamining technique. Because of the complexity and the high dimensionality of gene expression data, classification of a disease samples remains a challenge. Hierarchical clustering and partitioning clustering is used to identify patterns of gene expression useful for classification of samples. In this paper, we make a comparative study of two partitioning methods namely k-means and PAM to classify the cancer dataset.

1 INTRODUCTION

According to Usama Fayyad et al. (Fayyad, 1996), “Data mining is a step in the KDD (Knowledge Discovery in Databases) process that consists of applying data analysis and discovery algorithms that produce a particular enumeration of patterns (or models) over the data”.

According to Guha et al. (Guha, 1998), “Clustering problem is about partitioning a given data set into groups (clusters) such that the data points in a cluster are more similar to each other than points in different clusters”. A mathematical definition of clustering is the following: let \( X = \{ x_1, x_2, x_3, \ldots, x_{m-1}, x_m \} \subset \mathbb{R}^n \) set of data items representing a set of \( m \) points \( x_i \) in \( \mathbb{R}^n \) where \( x_i = \{ x_{i1}, x_{i2}, x_{i3}, \ldots, x_{in} \} \). The goal is to partition \( X \) into k-groups \( \{ C_i : 1 \leq i \leq k \} \) such that data belong to the same group are more “alike” than data in different groups. Each of the k-groups is called a cluster. The result of the algorithm is an injective mapping of data items \( x_i \) to groups \( C_k \).

Partitional clustering algorithms divide the whole data set into a set of disjoint clusters directly. These algorithms attempt to determine an integer number of clusters that optimise a certain objective function through an iterative procedure.

To classify the various types of cancer into its different subcategories, different data mining techniques have been used over gene expression data. Gene expression data, obtained using gene expression monitoring by DNA microarrays, provides an important source of information that can help in understanding many biological processes. A common aim, then, is to use the gene expression profiles to identify groups of genes or samples in which the members behave in similar ways. One might want to partition the data set to find naturally occurring groups of genes with similar expression patterns. Golub et al (Golub, 1999), Alizadeh et al (Alizadeh, 2000), Bittner et al (Bittner, 2000) and Nielsen et al (Nielsen, 2002) have considered the classification of cancer types using gene expression datasets. There are many instances of reportedly successful applications of both hierarchical clustering and partitioning clustering in gene expression analyses. Yeung et al (Yeung, 2001) compared k-means clustering, CAST (Cluster Affinity Search Technique), single-, average- and complete-link hierarchical clustering, and totally random clustering for both simulated and real gene expression data. And they favoured k-means and CAST. Gibbons and Roth (Gibbons, 2001) compared k-means, SOM (Self-Organizing Map), and hierarchical clustering of real temporal and replicate microarray gene expression data, and favoured k-means and SOM.

In this paper, we make a comparative study of two clustering algorithms namely k-means and PAM to classify the cancer datasets and is based on accuracy and ability to handle high dimensional data.
2 K-MEANS ALGORITHM

The k-means algorithm is a partitioning clustering algorithm. The k-means algorithm is very simple and most popular clustering algorithm. The k-means algorithm is a squared error-based clustering algorithm.

The k-means is given by MacQueen (MacQueen, 1967) and aim of this clustering algorithm is to divide the dataset into disjoint clusters by optimizing an objective function that is given below

\[
\text{Optimize } E = \sum_{i=1}^{k} \sum_{x \in C_i} d(x, m_i)
\]

Here \(m_i\) is the center of cluster \(C_i\), while \(d(x, m_i)\) is the euclidean distance between a point \(x\) and cluster center \(m_i\). In k-means algorithm, the objective function \(E\) attempts to minimize the distance of each point from the cluster center to which the point belongs.

Consider the data set with ‘n’ objects, i.e.,

\[ S = \{x_i : 1 \leq i \leq n\} \]

1) Initialize k-partitions randomly or based on some prior knowledge.

i.e., \(\{C_1, C_2, C_3, \ldots, C_k\}\).

2) Calculate the cluster prototype matrix \(M\) (distance matrix of distances between \(k\)-clusters and data objects).

\[ M = \{m_1, m_2, m_3, \ldots, m_k\} \]

where \(m_i\) is a column matrix \(1 \times n\).

3) Assign each object in the data set to the nearest cluster - Cm i.e. \(x_j \in C_m\) if \(d(x_j, C_m) \leq d(x_j, C_i) \forall 1 \leq j \leq k, j \neq m\) where \(j=1,2,3,\ldots,n\).

4) Calculate the average of cluster elements of each cluster and change the k-cluster centers by their averages.

5) Again calculate the cluster prototype matrix \(M\).

6) Repeat steps 3, 4 and 5 until there is no change for each cluster.

3 PAM ALGORITHM

The purpose for the partitioning of a data set into \(k\) separate clusters is to find groups whose members show a high degree of similarity among themselves but dissimilarity with the members of other groups. The objective of PAM (Partitioning Around Medoids) (Kaufman, 1990) is to determine a representative object (medoid) for each cluster, that is, to find the most centrally located objects within the clusters. Initially a set of \(k\)-items is taken to be the set of medoids. Then, at each step, all objects from the input dataset that are not currently medoids are examined one by one if they should be medoids. That is the algorithm determines whether there is an object that should replace one of the existing medoids. Swapping of medoids with other non-selected objects is based on the value of total cost of impact \(T_{ab}\). The PAM represents a cluster by a medoid so PAM is also known as k-medoids algorithm.

The PAM algorithm consists of two parts. The first build phase follows the following algorithm:

Phase-1:

Consider an object \(i\) as a candidate. Consider another object \(j\) that has not been selected as a prior candidate. Obtain its dissimilarity \(d_i\) with the most similar previously selected candidates. Obtain its dissimilarity with the new candidate \(i\). Call this \(d(j; i)\):

1) If the difference is positive, then object \(j\) contributes to the possible selection of \(i\). Calculate \(C_{ji} = \max (d_i - d(j; i); 0)\) where \(d_i - \) Euclidian distance between \(j^{th}\) object and most similar previously selected candidate and \(d(j; i) - \) Euclidian distance between \(j^{th}\) and \(i^{th}\) object.

2) Sum \(C_{ji}\) over all possible \(j\).

3) Choose the object \(i\) that maximizes the sum of \(C_{ji}\) over all possible \(j\).

4) Repeat the process until \(k\) objects have been found.

Phase-2:

The second step attempts to improve the set of representative objects. This does so by considering all pairs of objects \((i; h)\) in which \(i\) has been chosen but \(h\) has not been chosen as a representative. Next it is determined if the clustering results improve if object \(i\) and \(h\) are exchanged. To determine the effect of a possible swap between \(i\) and \(h\) we use the following algorithm:

Consider an object \(j\) that has not been previously selected. We calculate its swap contribution \(C_{jih}\):

1) If \(j\) is further from \(i\) and \(h\) than from one of the other representatives, set \(C_{jih}\) to zero.

2) If \(j\) is not further from \(i\) than any other representatives (\(d(j; i) = d_j\)), consider one of two situations:

a) \(j\) is closer to \(h\) than the second closest representative & \(d(j; h) < E_j\) where \(E_j\) is the Euclidian distance of between \(j^{th}\) object and the second most similarly representative. Then \(C_{jih} = d(j; h) - d(j; i)\).

Note: \(C_{jih}\) can be either negative or positive depending on the positions of \(j, i\) and \(h\). Here only if
j is closer to i than to h is there a positive influence at implies that a swap between object i and h are a disadvantage in regards to j.

b) j is at least as distant from h than the second closest representative \( d(j; h) \geq E_j \). Let \( C_{jih} = E_j - d_j \). The measure is always positive, because it not wise to swap i with h further away from j than second closest representative.

3) If j is further away from i than from at least one of the other representatives, but closer to h than to any other representative, \( C_{jih} = d(i; h) - d_j \) will be the contribution of j to the swap.

4) Sum the contributions over all j. \( T_{ih} = \sum C_{jih} \). This indicates the total result of the swap.

5) Select the ordered pair \((i; h)\) which minimizes \( T_{ih} \).

6) If the minimum \( T_{ih} \) is negative, the swap is carried out and the algorithm returns to the first step in the swap algorithm. If the minimum is positive or 0, the objective value cannot be reduced by swapping and the algorithm ends.

4 CANCER DATASETS USED FOR COMPARISON OF K-MEANS AND PAM

We used three different datasets to make a comparison study between k-means and PAM algorithms. Brief description is given below:

The Leukemia data set is a collection of gene expression measurements from 72 leukemia (composed of 62 bone marrow and 10 peripheral blood) samples reported by Golub.

It contains an initial training set composed of 47 samples of acute lymphoblastic leukemia (ALL) and 25 samples of acute myeloblastic leukemia (AML).

The Colon dataset is a collection of gene expression measurements from 62 Colon biopsy samples from colon-cancer patients reported by Alon. Among them, 40 tumor biopsies are from tumors (labelled as "negative") and 22 normal (labelled as "positive") biopsies are from healthy parts of the colons of the same patients. Two thousand out of around 6500 genes were selected.

The Lymphoma dataset is a collection of gene expression measurements from 96 normal and diffused malignant lymphocyte samples reported by Alizadeh. It contains 42 samples of diffused large B-cell lymphoma (DLBCL) and 54 samples of other types. The Lymphoma data set contains 4026 genes.

5 RESULTS

5.1 Comparison of k-means and PAM for Gene-leukemia Dataset

Here we apply k-means and PAM algorithms on leukemia data set to classify it into two equivalent classes. We use two variations of leukemia data set one with 50-genes and another with 3859-genes.

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We observe that k-means algorithm converges fast in comparison to PAM algorithm. In this case, accuracy for k-means is also better than the accuracy of PAM algorithm.

Graphical representation of two cluster centers of 50-gene-leukemia data set using k-means and PAM algorithm is shown below:

Figure 1: Graphical representation of Leukemia dataset.

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When we apply these algorithms on 3859-gene-leukemia dataset results are different as compared to results with 50-gene-leukemia dataset. In this case PAM algorithm’s accuracy is better than k-means algorithm’s accuracy. This shows that PAM perform better when we increase number of attributes.

Table 2: Results for 3859-gene-leukemia dataset.

<table>
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<th>Clustering Algorithm</th>
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<tr>
<td>k-means</td>
<td>61</td>
<td>84.72</td>
</tr>
<tr>
<td>PAM</td>
<td>68</td>
<td>94.44</td>
</tr>
</tbody>
</table>

5.2 Comparison of K-means and PAM for 2000-Gene-colon Dataset

Analysis of 2000-gene-colon data set is also done with the help of these two partitioning algorithms i.e. k-means and PAM algorithm. In this case PAM algorithm performs better then k-means method. But accuracy difference between these algorithms over colon data set is significantly low. Average accuracy remains low.

Table 3: Results for 2000-gene-colon dataset.

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<tr>
<td>k-means</td>
<td>33</td>
<td>53.22</td>
</tr>
<tr>
<td>PAM</td>
<td>34</td>
<td>54.84</td>
</tr>
</tbody>
</table>

5.3 Comparison of K-means and PAM for 4026-Gene-lymphoma Dataset

Using these algorithms, we divide the whole dataset into two different clusters which are used to differentiate between normal and diffused samples. Here PAM algorithm correctly classifies 77 records out of 96 whereas k-means algorithm correctly classifies 71 records.

Table 4: Results for 4026-gene-dlbcl dataset.

<table>
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<th>Correctly Classified</th>
<th>Average Accuracy</th>
</tr>
</thead>
<tbody>
<tr>
<td>k-means</td>
<td>71</td>
<td>73.96</td>
</tr>
<tr>
<td>PAM</td>
<td>77</td>
<td>80.21</td>
</tr>
</tbody>
</table>

So it is clear that PAM performs better when we increase the number of genes.

6 SUMMARY

Algorithm’s comparision shows that accuracy of PAM is better from accuracy of k-means as number of objects in the dataset increases. In case of k-means intial selection of cluster centres plays a very important role. So there is a possibility to improve both algorithms by using some good intial selection technique. Here in this paper PAM performs better in the classification of cancer types using cancer datasets than k-means.

REFERENCES


