



Published in final edited form as:

Knowl Inf Syst. 2014 December ; 41(3): 667–696. doi:10.1007/s10115-013-0684-0.

Efficient Mining of Discriminative Co-clusters from Gene Expression Data

Omar Odibat and Chandan K. Reddy

Department of Computer Science, Wayne State University, Detroit, MI, 48202

Abstract

Discriminative models are used to analyze the differences between two classes and to identify class-specific patterns. Most of the existing discriminative models depend on using the entire feature space to compute the discriminative patterns for each class. Co-clustering has been proposed to capture the patterns that are correlated in a subset of features, but it cannot handle discriminative patterns in labeled datasets. In certain biological applications such as gene expression analysis, it is critical to consider the discriminative patterns that are correlated only in a subset of the feature space. The objective of this paper is two-fold: first, it presents an algorithm to efficiently find arbitrarily positioned co-clusters from complex data. Second, it extends this co-clustering algorithm to discover discriminative co-clusters by incorporating the class information into the co-cluster search process. In addition, we also characterize the discriminative co-clusters and propose three novel measures that can be used to evaluate the performance of any discriminative subspace pattern mining algorithm. We evaluated the proposed algorithms on several synthetic and real gene expression datasets, and our experimental results showed that the proposed algorithms outperformed several existing algorithms available in the literature.

Keywords

Co-clustering; biclustering; discriminative pattern mining; gene expression data; negative correlation

1 Introduction

Discriminative models are used to extract patterns that are highly correlated in one class compared to another class. Mining such discriminative patterns can provide valuable knowledge toward understanding the differences between two classes and identifying class-specific patterns. For example, discriminative mining of gene expression data can lead to the identification of cancer-associated genes by comparing the expression patterns of the genes between healthy and cancerous tissues [13]. However, these genes can be correlated only in a subset of the cancerous samples due to the heterogeneity in the sample space [27]. Since the existing discriminative models are based on using all the features to find the discriminative patterns, it is crucial to develop a model that can identify discriminative patterns that are correlated in a subset of the feature space.

Co-clustering has been proposed to identify subsets of objects that are inter-related under subsets of features (*co-clusters*) [10], [16], [3], [27], [1]. However, co-clustering is an unsupervised procedure that does not consider the class labels to find the discriminative patterns in labeled datasets. In order to capture the subspace discriminative patterns (or *discriminative co-clusters*), discriminative co-clustering is being proposed in this paper by incorporating the class labels into the co-clustering process.

1.1 Co-clustering

Given a data matrix with two entities such as (*genes, samples*) in gene expression data [26], a subset of rows may be inter-related under a subset of columns forming blocks of substructures (or *co-clusters*) [14]. Applying traditional clustering techniques, such as k-means and hierarchical clustering, will not capture such co-clusters [10], [16], [22], [4], [32]. However, co-clustering (or biclustering)¹ has been proposed to simultaneously cluster both dimensions of a data matrix by utilizing the relationship between the two entities [10], [16], [26], [34], [27]. Co-clusters have several characteristics that should be considered in the search process. Here, we describe the important characteristics of the co-clusters in the gene expression domain. However, many of these characteristics are applicable to several other domains. **1. Arbitrarily positioned co-clusters.** Due to the heterogeneity of the samples, a subset of genes can be correlated in any subset of the samples. Hence, the co-clusters can be arbitrarily positioned in the matrix [27]. **2. Overlapping.** A gene can be involved in several biological pathways. Hence, that gene can belong to more than one co-cluster [27], [15]. **3. Positive and negative correlations.** In a positive (negative) correlation, genes show similar (opposite) patterns [39]. **4. Noisy data.** The expression data contains a huge amount of noise [23]. Hence, the co-clustering algorithms should be robust against noise.

Recently, the (κ, ℓ) co-clustering model has been proposed to simultaneously find $\kappa\ell$ co-clusters [4], [15]. This model was shown to perform well in various applications [4], [15]. However, the main limitation of this model is that it assumes a grid structure comprised of $\kappa \times \ell$ co-clusters as shown in Fig. 1(a). The assumption here is that the rows in each row cluster should be correlated under each of the ℓ column clusters. Such an assumption may not hold when a subset of rows is correlated in a limited subset of columns (or vice versa). To overcome this limitation, we propose a novel co-clustering algorithm that is able to identify arbitrarily positioned co-clusters as shown in Fig. 1(b). This algorithm is extended to efficiently find the discriminative co-clusters in the data.

1.2 Discriminative Co-clustering

Discriminative models aim to extract patterns that are differentially correlated between two classes [19]. Figure 2 shows the correlations between three objects in two classes. These objects are highly correlated in a subset of the features in class A, but they are not correlated in class B. Such discriminative patterns cannot be discovered using standard discriminative models that use all the features. In order to capture these patterns, discriminative co-clustering is also being proposed in this paper.

¹To be consistent, we will be using the term 'co-clustering' throughout the paper. The Bioinformatics research community preferably calls it as 'biclustering'.

In addition to the above mentioned characteristics of the co-clusters, the discriminative co-clusters must possess the following characteristics: **1. High discriminative coherence.** Coherence (or correlation) is a measure of similarity between a set of objects [27]. The discriminative co-clustering algorithms should identify the set of co-clusters with the maximum difference in the coherence between the two classes. The co-clusters that have the same correlation in both of the classes should be ignored. **2. Low inter-class overlapping.** The discriminative co-clusters discovered in one class should have a minimal number of rows that are common with the co-clusters discovered in the other class. **3. High discriminative power.** Incorporating the class labels can improve the performance of classification algorithms [21]. Discriminative co-clusters must be able to make more accurate predictions.

Example. Fig. 3 shows an example of discriminative and non-discriminative co-clusters. The width of each co-cluster (X) indicates the number of features in it, and its shade represents its correlation score, which is also displayed as a percentage inside each co-cluster. The correlation score can be measured by various functions such as the mean-squared residue (MSR) [10]. In this example, the higher the percentage (or the darker the shade), the stronger the correlation. The co-cluster properties (shade and width) are the main criteria used to distinguish between discriminative and non-discriminative co-clusters. A co-cluster is considered as a discriminative co-cluster if it is correlated only in one class (such as $X1$ and $X5.b$), if it is highly correlated in one class and less correlated in the other class (such as $X4$) or if it is correlated in relatively higher percentage of features (such as $X3$ and $X6$). The co-clusters $X2$ and $X5.a$ are not considered as discriminative co-clusters because they are similarly correlated in both classes.

Can any co-clustering algorithm be used to identify the discriminative co-clusters?—A naive solution to this problem is to co-cluster each class separately and then identify the co-clusters that appear in only one class. However, there are many limitations in following such a procedure: (i) Standard co-clustering algorithms focus on identifying the most correlated co-clusters. Therefore, discriminative co-clusters that have low correlation score (such as $X1$ and $X6$) will not be discovered. (ii) Since the standard co-clustering algorithms do not detect *all* the co-clusters, it is possible that co-cluster $X2$ is discovered only in one class and considered as a discriminative co-cluster. (iii) Most co-clustering algorithms prefer large co-clusters. Therefore, the complete co-cluster $X5$ may be considered as a discriminative co-cluster because part a is not discovered in class B due to its size limitation. In this paper, we develop a novel algorithm that directly optimizes an objective function to efficiently identify the discriminative co-clusters, and we propose two metrics to score the discriminative co-clusters based on their correlation scores and the number of features in them.

1.3 Our Contributions

The primary contributions of this paper are as follows:

1. A novel co-clustering algorithm, **Ranking-based Arbitrarily Positioned Overlapping Co-Clustering (RAPOCC)**, to efficiently extract significant co-clusters.

- Propose a novel ranking-based objective function to find arbitrarily positioned co-clusters.
 - Extract large and overlapping co-clusters containing both positively and negatively correlated rows.
2. A novel discriminative co-clustering algorithm, **Discriminative RAPOCC (Di-RAPOCC)**, to efficiently extract the discriminative co-clusters from labeled datasets.
 - Find the discriminative co-clusters from labeled datasets efficiently by incorporating the class information into the co-clustering process.
 - Propose three new evaluation metrics to quantify the results of the discriminative co-clustering algorithms on both synthetic and real gene expression datasets. Two metrics are used to measure the discriminative coherence property of the discriminative co-clusters, and the third one measures the inter-class overlap property.
 3. In addition to summarizing some of the widely used co-clustering algorithms, we categorize the state-of-the-art approaches for discriminative co-clustering and characterize each category. We also empirically compare the performance of these categories with the proposed algorithm.

The rest of this paper is organized as follows: Section 2 presents an overview of the related work. Section 3 introduces the coherence measure and formulates the problems of co-clustering and discriminative co-clustering. Section 4 describes the *RAPOCC* algorithm. Section 5 presents the *Di-RAPOCC* algorithm. Section 6 presents the results of the proposed algorithms on synthetic and real datasets along with the comparisons with other algorithms available in the literature. Finally, we conclude our discussion in Section 7.

2 Related Work

In this section, we describe some of the widely-used co-clustering algorithms and categorize the state-of-the-art approaches for discriminative co-clustering.

2.1 Co-clustering Algorithms

Cheng and Church (CC) [10] proposed the first co-clustering algorithm that produces one co-cluster at a time. The obtained co-cluster is replaced with random numbers, which typically reduces the quality of the co-clusters. The Order-Preserving Submatrices (OPSM) algorithm [7] finds one co-cluster at a time in which the expression levels of all genes induce the same linear ordering of the experiments. This algorithm does not capture the negatively correlated genes. The Iterative Signature Algorithm (ISA) [22] defines a co-cluster as a co-regulated set of genes under a set of experimental conditions. It starts from a set of randomly selected rows that are iteratively refined until they are mutually consistent. The Robust Overlapping Co-clustering (ROCC) algorithm [15] finds $\kappa \times \ell$ co-clusters using the Bregman co-clustering algorithm [6]. This algorithm does not handle the negative correlations. Our proposed co-clustering algorithm overcomes all of the above limitations by

(i) capturing arbitrarily positioned co-clusters, (ii) handling overlapping and positive and negative correlations and (iii) being robust against noise.

2.2 Discriminative Co-clustering Algorithms

In general, the co-clustering algorithms work in an unsupervised manner. However, some algorithms incorporate a priori knowledge in the co-clustering process. For example, in constrained co-clustering, some information can be incorporated such as the must-link and cannot-link constraints [30], [36], [35]. In discriminative co-clustering, the class labels are incorporated to find class-specific co-clusters. As illustrated in Fig. 4, the existing discriminative co-clustering approaches can be categorized as two-step or one-step approaches.

2.2.1 Two-step approaches—There are two sub-categories of these approaches: (i) first co-clustering, and then discriminative analysis. In [29], differentially expressed gene modules are identified by applying co-clustering each class separately, then the identified co-clusters are ranked based on their discrimination between the two classes. (ii) first discriminative analysis, and then co-clustering. The *DeBi* algorithm [33] uses two steps to identify differentially expressed co-clusters. The first step is to find the up or the down regulated genes using fold change analysis. In the second step, the *MAFIA* algorithm [8] is used to find the co-clusters from the up-regulation and the down-regulation data. There are two limitations for the two-step approaches: (i) the co-clustering is done for each class separately, and (ii) the discriminative analysis step is independent of the co-clustering step. Therefore, the one-step approaches have been proposed to overcome these limitations.

2.2.2 One-step approaches—The subspace differential co-expression (SDC) algorithm [18] uses the Apriori search algorithm to identify the discriminative patterns. The Apriori approach depends on using thresholds to define the discriminative patterns [19]. For example, a given pattern is considered as a discriminative pattern if the difference between the correlations of this pattern in the two classes is above a fixed threshold. Otherwise, this pattern will be split into smaller patterns to be tested again using the same threshold. Therefore, the *SDC* method suffers from the following limitations: (i) It generates very small patterns [18]. (ii) The number of the discovered patterns dramatically grows with the size of the datasets, and it significantly varies with the threshold value [19]. (iii) It has computational efficiency problems and does not scale well to large-scale datasets. In addition, the *SDC* method does not identify the subset of columns in which a given pattern shows the maximum correlation. In our previous work [28], we proposed a discriminative co-clustering algorithm to analyze the differences in the biological activities of several genes between two classes. Although this algorithm generated large co-clusters compared to the *SDC* method, this algorithm does not scale to large datasets because it maintains, for each pair of rows (genes), the set of columns under which the two rows are differentially correlated. Recently, locally discriminative co-clustering was proposed in [40] to explore the inter-sample and inter-feature relationships, but it does not find discriminative co-clusters as defined in our paper. To overcome all of the above limitations of the existing approaches, we propose a novel discriminative co-clustering algorithm that directly optimizes an objective function to efficiently identify the discriminative co-clusters from a given labeled

dataset. It should be noted that we primarily focus on co-clusters in our work rather than other concepts such as emerging patterns or contrast set patterns [17]. While we acknowledge the fact that these approaches are generic and probably can be modified for our problem, we emphasize that there is no existing work in the area of discriminative co-clustering and hence this is an exciting direction of future research.

3 Preliminaries

In this section, we introduce the coherence measure that can be used to measure the quality of the co-clusters, and we formulate the problems of co-clustering and discriminative co-clustering. The notations used in this paper are described in Table 1.

3.1 Measuring the Coherence of Co-clusters

Coherence is a measure of how similar a set of gene expression profiles are. Cheng and Church proposed the mean-squared residue (MSR) score as a measure of coherence [10]. Since the overall shapes of gene expression profiles are of greater interest than the individual magnitudes of each feature [23], we normalize the expression values of each gene to be between 0 and 1. As a result, the value of the objective function will also be bounded between 0 and 1.

Definition 1. (Coherence measure H)—The coherence of a co-cluster X of $|I|$ rows and $|J|$ columns is measured as

$$H(X) = 1 - \frac{1}{|I||J|} \sum_{i \in I, j \in J} (X_{ij} - X_{i\cdot} - X_{\cdot j} + X_{\cdot\cdot})^2$$

where X_{ij} is the value in row i and column j in co-cluster X , $X_{i\cdot} = \frac{\sum_{j \in J} X_{ij}}{|J|}$ is the row mean, $X_{\cdot j} = \frac{\sum_{i \in I} X_{ij}}{|I|}$ is the column mean and $X_{\cdot\cdot} = \frac{\sum_{i,j} X_{ij}}{|I||J|}$ is the overall mean of X .

Using Definition 1, a perfect co-cluster will have a score = 1. Given two rows (x and y) and J columns, the coherence measure can be re-written as follows:

$$\begin{aligned} h(x, y, J) &= 1 - \frac{1}{2|J|} \sum_{j \in J} \left(x_j - \bar{x} - \frac{x_j + y_j}{2} + \frac{\bar{x} + \bar{y}}{2} \right)^2 \\ &\quad - \frac{1}{2|J|} \sum_{j \in J} \left(y_j - \bar{y} - \frac{x_j + y_j}{2} + \frac{\bar{x} + \bar{y}}{2} \right)^2 \\ &= 1 - \frac{1}{|J|} \sum_{j \in J} \left(\frac{(x_j - \bar{x}) - (y_j - \bar{y})}{2} \right)^2 \end{aligned} \tag{1}$$

where \bar{x} and \bar{y} represent the mean of the values for the rows x and y , respectively. An optimal co-cluster has a value of $H(X) = 1$, which results from the case where $(x_j - \bar{x}) = (y_j - \bar{y}), \forall j \in J$. This type of correlation is positive ($h_+(x, y, J)$). In the negative correlation, the rows have opposite patterns (i.e. the two negatively correlated rows will get a perfect score

when $(x_j - \bar{x}) = -(y_j - \bar{y}) \forall j \in J$. Figure 5 shows an example these correlations. In a positive correlation, genes show similar patterns while in a negative correlation, genes show opposite patterns. The positive and negative correlations are defined in Definition 2.

Definition 2. (Positive and negative correlations)—Given two rows (x and y) and J columns, the positive correlation between them is defined as

$$h_+(x, y, J) = 1 - \frac{1}{|J|} \sum_{j \in J} \left(\frac{(x_j - \bar{x}) - (y_j - \bar{y})}{2} \right)^2$$

and the negative correlation is defined as

$$h_-(x, y, J) = 1 - \frac{1}{|J|} \sum_{j \in J} \left(\frac{(x_j - \bar{x}) + (y_j - \bar{y})}{2} \right)^2$$

Definition 3. (Pairs-based Coherence HP)—Given a co-cluster X of $|I|$ rows and $|J|$ columns, the coherence of this co-cluster is measured based on all the pairs in X :

$$HP(X) = \frac{2}{|I|(|I| - 1)} \sum_{x, y \in X} (h_o(x, y, J))$$

where $o \in \{-, +\}$.

The type of correlations between any two rows, referred to as o in Definition 3, is maintained for each pair of rows in each co-cluster in the proposed algorithms.

3.2 Problem Formulations

Here, we formally define the problems of co-clustering and discriminative co-clustering.

Definition 4. (Co-clustering)—Let $D \in \mathbb{R}^{M \times N}$ denote a data matrix; the goal of co-clustering is to find a row mapping (ρ) that maps the rows to the κ row clusters and a column mapping (γ) that maps the columns to the ℓ column clusters

$$\rho: \{1, 2, \dots, M\} \rightarrow \{1, 2, \dots, \kappa\}$$

$$\gamma: \{1, 2, \dots, N\} \rightarrow \{1, 2, \dots, \ell\}$$

such that the coherence of the top- K co-clusters is maximized.

$$\arg \max_{X_1, X_2, \dots, X_K} \sum_{i=1}^K HP(X_i)$$

The problem of finding the co-clusters is an NP-hard problem [10]. In this paper, we propose a novel co-clustering algorithm to efficiently find arbitrarily positioned co-clusters from a given data matrix.

Definition 5. (Discriminative Co-clustering)—If $HP^A(X_i)$ measures the coherence of the co-cluster X_i in class A, the goal is to find the set of co-clusters that has maximal discriminative coherence

$$\arg \max_{X_1, X_2, \dots, X_{K^A}} \sum_{i=1}^{K^A} (HP^A(X_i) - \psi^B(X_i))$$

$$\arg \max_{X_1, X_2, \dots, X_{K^B}} \sum_{i=1}^{K^B} (HP^B(X_i) - \psi^A(X_i))$$

where $\psi^A(X_i)$ ($\psi^B(X_i)$) is the maximum coherence of any subset of the objects in X_i in class A (B). The challenge here is to find $\psi(X_i)$, which is similar to the NP-hard problem of finding the maximum subspace in X_i [10]. In the proposed discriminative co-clustering algorithm, we propose two approximations for computing $\psi(X_i)$ that can be used to efficiently discover discriminative co-clusters by incorporating the class labels into the co-clusters discovery process.

4 The Proposed RAPOCC Algorithm

In this Section, we describe the *RAPOCC* algorithm. This algorithm is proposed to efficiently extract the most coherent and large co-clusters that are arbitrarily positioned in the data matrix. These co-clusters can overlap and have positively and negatively correlated rows.

4.1 Ranking-based Objective Function

In the proposed iterative algorithm, the score of each of the $\kappa \ell$ co-clusters is computed at each iteration, and the overall value of the objective function is computed based on the coherence score of the top- K scores where K is the number of optimized co-clusters ($1 \leq K \leq \kappa \ell$).

$$\arg \max_{X_1, X_2, \dots, X_K} \sum_{i=1}^K HP(X_i)$$

The set of the top- K co-clusters can be any subset of the $\kappa * \ell$ co-clusters. The objective function will be computed for each possible change in the row/column mapping to maintain non-decreasing values for the objective function. The advantage of using this objective function is that it allows the discovery of arbitrarily positioned co-clusters as shown in Fig. 1(b).

4.2 The RAPOCC Algorithm

The main steps of the *RAPOCC* algorithm are shown in Fig. 6. The algorithm starts with a two-dimensional matrix (*objects* \times *features*) as an input. In the first step, Fig 6(b), a divisive approach is used for initialization. Basically, it starts with all the rows and columns in one co-cluster; then the algorithm splits the co-cluster with the largest error. This iterative procedure continues until κ row clusters and ℓ column clusters are obtained. The core co-clustering step, Fig. 6(c), finds the optimal row and column clusterings (ρ, γ). In the third step, Fig. 6(d), similar co-clusters are merged using a hierarchical agglomerative approach. In this step, more rows and columns are added to each co-cluster individually. Finally, a pruning step is used to prune the co-clusters with low coherence scores. These steps are described in Algorithm 1. In this algorithm, $H(u, v)$ and $HP(u, v)$ indicate the coherence of the co-cluster formed by the row cluster u and column cluster v . The inputs to this algorithm include the data matrix $D \in \mathbb{R}^{M \times N}$, the number of row clusters κ and the number of column clusters ℓ . These are common parameters in the co-clustering methods [15], and they can be set based on the size of the data matrix. K determines the number of the optimized co-clusters and can be set to any value between 1 and $\kappa \times \ell$. We set the parameters $\kappa=5$ and $\ell=3$. The parameter K can be set to large value (we set it to 20 in our implementation) because the *RAPOCC* algorithm will only report the most coherent co-clusters, and the remaining ones will be pruned in the last step. In addition, we had a threshold that indicates that the minimum number of rows in a co-cluster to be 20 and the minimum number of columns in

Algorithm 1

RAPOCC(D, κ, ℓ, K)

1: **Input:** Data matrix (D)
 No. of row clusters (κ)
 No. of column clusters (ℓ)
 No. of optimized co-clusters (K)

2: **Output:** A set of K co-clusters ($\{X\}$)

3: **Procedure:**

4: **Step 1** : initialization

5: $i \leftarrow 1, j \leftarrow 1$

6: $\rho(g) \leftarrow i, \forall [g]_1^m$

7: $\gamma(c) \leftarrow j, \forall [c]_1^n$

8: **while** $i < \kappa$ **or** $j < \ell$ **do**

9: **if** $i < \kappa$ **then**

```

10:    $i \leftarrow i + 1$ 
11:    $\alpha \leftarrow \arg \min_{\alpha} \sum_{j=1}^{\ell} H'(u, v) : \rho(u) = \alpha, \gamma(v) = l$ 
12:   Partition  $\alpha$  using bisecting clustering algorithm
13: end if
14: if  $j < \ell$  then
15:    $j \leftarrow j + 1$ 
16:    $\beta \leftarrow \arg \min_{\beta} \sum_{i=1}^{\kappa} H'(u, v) : \rho(u) = i, \gamma(v) = \beta$ 
17:   Partition  $\beta$  using bisecting clustering algorithm
18: end if
19: end while
20: Step 2 : core co_clustering
21: repeat
22:   /* Row clustering */
23:   for  $a = 1 : M$  do
24:      $\rho(a) = \arg \max_{u \in \{-\kappa, \dots, -1, 0, 1, \dots, \kappa\}} HP(\rho(a) = u, \gamma)$ 
25:   end for
26:   /* Column clustering */
27:   for  $b = 1 : N$  do
28:      $\gamma(b) = \arg \max_{b \in \{0, 1, \dots, \ell\}} HP(\rho, \gamma(b) = v)$ 
29:   end for
30: until convergence
31: Step 3 : Merging similar co_clusters and refinement
32: Step 4 : Pruning

```

a co-cluster to be 5. This is chosen because in biological problems, the number of conditions are usually far less compared to the number of genes.

Step 1: Initialization—Inspired by the bisecting K-means clustering technique [37], we use a deterministic algorithm for the initialization. Each row is mapped to one of the κ clusters, and each column is mapped to one of the ℓ clusters, resulting in a checkerboard structure $\kappa \times \ell$ as shown in Fig. 6(b). The initialization algorithm is a divisive algorithm that starts with the complete data assigned to one cluster as described in Algorithm 1 (lines 5-7); then, the following steps are repeated until the desired number of row clusters is obtained. (1) Find the row cluster with the lowest coherence score (a_{min}). (2) Find the two rows in a_{min} with the lowest correlation (r_1, r_2). (3) Create two new row clusters a_1 and a_2 . Add r_1 to a_1 and r_2 to a_2 . (4) Add each of the remaining rows in a_{min} to a_1 (a_2) if it is more correlated to r_1 (r_2). The column clusters are initialized in the same manner. The algorithm alternates between clustering the rows and the columns as described in Algorithm 1 (lines 8-19).

Step 2: Core Co-clustering (ρ, γ)—This step finds the optimal row and column clusterings (ρ, γ) as shown in Fig. 6(c). To update ρ , each row (r_i) is considered for one of the following three actions as described in Algorithm 1 (lines 20-30):

- Exclude r_i from any row cluster by setting ρ to 0.
- Find the best row cluster to include r_i as a **positively correlated** row $\{1, 2, \dots, \kappa\}$.
- Find the best row cluster to include r_i as a **negatively correlated** row $\{-\kappa, \dots, -2, -1\}$.

The objective function is computed for each possible action, and the action to be carried out is the one corresponding to the maximum value of the three objective function values. Within each co-cluster, there is a sign vector that determines the type of correlation (positive or negative) of each row. Therefore, a row can be positively correlated in some of the co-clusters and negatively correlated in other co-clusters. The column mapping (γ) is calculated in a similar manner, but there is no consideration for negatively correlated columns. Following this strategy, the value of the objective function is monotonically increasing, and the convergence is guaranteed as shown in Theorem 1. After convergence, the result will be a non-overlapping set of co-clusters.

Theorem 1. The Algorithm RAPOCC (Algorithm 4) converges to a solution that is a local optimum.

Proof: From Definition 3, the coherence measure HP is bounded between 0 and 1. Hence, the objective function given in Definition 4 is bounded. Algorithm 4 iteratively performs a set of update operations for the row clustering and the column clustering. In each iteration, it monotonically increases the objective function. Since this objective function is bounded for the top- K co-clusters, the algorithm is guaranteed to converge to a locally optimal solution.

Step 3: Merging the Co-clusters—The top- K co-clusters with the maximum coherence are retained from the previous step. In this step, similar co-clusters are merged as shown in Fig. 6(d) using an agglomerative clustering approach. Before merging, the co-clusters with the lowest scores are pruned. If there is no pre-determined threshold for pruning the co-clusters, the top η co-clusters will be retained, and the remaining co-clusters will be pruned. The similarity between any two co-clusters is defined using the coherence function of the union of the rows and columns of the co-clusters, and the merging is performed following an agglomerative clustering approach. The two most similar co-clusters are merged in each iteration. The goal of this step is two-fold: (i) it allows the discovery of **large** co-clusters, and (ii) it allows for **overlapping** co-clusters. In this step, the algorithm also adds more rows and columns to each co-cluster individually to obtain **larger** co-clusters and also allows for **overlapping** co-clusters. Hence, the same row/column can be added to several co-clusters.

Step 4: Pruning—In this step, we prune the co-clusters with the lowest coherence scores. To determine which co-clusters to prune, (i) sort the co-clusters based on their coherence (measured by HP), (ii) compute the difference between the consecutive scores and (iii) report the set of co-clusters just before the largest difference, and prune the remaining co-clusters. The time complexity of the RAPOCC algorithm is $O(\kappa \ell(\max(MN^2, NM^2)))$.

We now extend the RAPOCC algorithm to discover discriminative co-clusters from labeled datasets that has two classes. First, rather than looking at $\kappa \times \ell$ co-clusters, we will search for $K^A \times 1$ and $K^B \times 1$ co-clusters. Second, the initialization step and the objective functions

are changed to consider the two-classes problem. The core co-clustering, merging, refinement and pruning steps will be used with some modifications. The positive and negative correlations will be handled similarly.

5 The Proposed Di-RAPOCC Algorithm

Discriminative co-clustering aims to extract patterns that are highly correlated in a subset of the features in one class but not correlated in the other class. As illustrated in Fig. 3., the rows of a discriminative co-cluster in one class should not form a co-cluster in the other class. This implies that there are two tasks that should be performed simultaneously: (i) search for a co-cluster in one class, and (ii) find the coherence of the rows of the co-cluster in the other class ($\psi^A(X)$ or $\psi^B(X)$ in Definition 5). **The challenge is to compute $\psi^B(X)$ ($\psi^A(X)$) while searching for the co-cluster in class A (B).**

Consider X^A as a co-cluster in class A that has $|I|$ rows and $|J^A|$ columns, and consider $D^B(I, .)$ as the sub-matrix composed of the I rows and all the columns in class B . X^A will be considered as a discriminative co-cluster if there are no co-clusters in $D^B(I, .)$. An optimal solution for this would be to apply a co-clustering algorithm to find the maximal co-cluster in class $D^B(I, .)$. However, this is an NP-hard problem [10].

An alternative solution to this problem is to consider the correlations of each pair of rows in $D^B(I, .)$. Given two rows (x and y) in $D^B(I, .)$, the aim is to find the subset of columns where the coherence between the two rows is maximized. To find an exact solution, one should enumerate all possible subsets of the $|N^B|$ columns. However, this solution is computationally infeasible since it requires enumerating all the $2^{|N^B|}$ subsets, where N^B is the number of columns in class B . To avoid such an exhaustive enumeration, we propose two efficient solutions: (i) a greedy-columns-selection solution and (ii) a clustering-based solution. Table 2 demonstrates a running example to illustrate how these solutions work.

5.1 Greedy-Columns-Selection

The intuition behind this measure is to iteratively compute the coherence between x and y based on the best J^i sets of columns for $1 \leq J^i \leq N^B$ and then report a weighted average of these N^B computations. In the first iteration, all the N^B columns are used. In the second iteration, one of the columns (j) is removed, and the remaining $N^B - 1$ columns are used to compute the coherence between the two rows. These are the set of $N^B - 1$ columns that achieves the maximum coherence between the two rows. This will be repeated to compute the coherence of the two rows using the best $N^B - 2, N^B - 3, \dots, 1$ columns. The final value of this measure is a weighted average of $\{h(x, y, J^1), \dots, h(x, y, J^{N^B})\}$:

$$\frac{\sum_{i=1}^{N^B} h_+(x, y, J^i) |J^i|/N^B}{\sum_{i=1}^{N^B} |J^i|/N^B}$$

$$J^{(i+1)} = \{J^i\} - \underset{j}{\operatorname{arg\,max}} \ h(x, y, \{J^i\} - \{j\})$$

$|J^i|/N^B$ is the weight assigned to each set of columns such that larger sets of columns are assigned more weight than smaller sets of columns. This measure can be used to capture the negative correlations by applying $h_-(x, y, J)$ instead of $h_+(x, y, J)$. Since no prior knowledge about the correlations between the rows is used, h_G will be computed twice, and the final value for this measure $h_G(x, y)$ is computed as the maximum of

$$\left(\frac{\sum_{i=1}^{N^B} h_+(x, y, J^i) |J^i|/N^B}{\sum_{i=1}^{N^B} |J^i|/N^B}, \frac{\sum_{i=1}^{N^B} h_-(x, y, J^i) |J^i|/N^B}{\sum_{i=1}^{N^B} |J^i|/N^B} \right)$$

Finally, $\psi_G^B(X)$ is computed as:

$$\psi_G^B(X) = \frac{|2|}{|I|(|I| - 1)} \sum_{x, y \in X} h_G(x, y)$$

As an example, Table 3 shows the results of applying h_G on the x and y rows in Table 2. From this table, it should be noted that the two rows form a perfect co-cluster in the columns $\{c_1, c_4, c_6, c_7, c_9\}$. Fig. 7(a) shows a plot for all the three rows in all the columns, and Fig. 7(b) shows a plot for all the three rows in the identified subset of the columns. Based on the greedy-columns-selection method, the first proposed discriminative coherence measure is defined as follows:

$$\Delta_G^A(X) = \psi_G^A(X) - \psi_G^B(X). \quad \Delta_G^B(X) = \psi_G^B(X) - \psi_G^A(X)$$

The range of Δ_G^A and Δ_G^B is $(-1, 1)$.

5.2 Clustering-based Discretization

The goal of the discretization step is to create a new representation of the data using a standard one-dimensional clustering algorithm to cluster each row separately. We rank the clusters in each row, and each value in a row will be represented by the rank of the cluster it belongs to. After clustering, we estimate the coherence between any two rows using the new representation.

The intuition of using clustering is to guarantee that similar data points within each row will be represented by the same value. The basic idea is as follows: (i) Cluster the values of each row to c clusters. (ii) Rank the clusters based on the mean of the values of each cluster such that cluster 1 contains the lowest values in x , and cluster c contains the highest values in x . (iii) Map each value of x to the rank of the cluster the value belongs to.

$$\zeta: \{1, 2, \dots, N^B\} \rightarrow \{1, 2, \dots, c\}$$

As shown in Section 3, the positive correlation between two rows is defined as $(x_j - x)^- = (y_j - \bar{y})$ and the negative correlation between them is defined as $(x_j - x)^- = -(y_j - \bar{y})$. Using the new representation, the positive correlation can be represented as

$$\zeta(x_j) - \zeta(y_j) = s^+$$

where s^+ is the positive shift parameter. Since $\zeta(x_j)$ and $\zeta(y_j)$ can take any value between 1 and c , the shift parameter (s^+) can take any value from the following set: $\{-(c - 1), \dots, -1, 0, 1, \dots, c - 1\}$. Similarly, the negative correlation can be represented as

$$\zeta(x_j) + \zeta(y_j) = s^-$$

where s^- is the negative shift parameter that can take any value from the following set: $\{2, 3, \dots, 2c\}$. Now, we can efficiently estimate the correlation between any two rows by finding the values of s^+ and s^- which will have a finite number of possible values. To estimate the positive correlation between x and y , we will subtract $\zeta(x_j)$ from $\zeta(y_j)$, and the most frequent value that appears in many columns will be considered as the value for s^+ . Similarly, to estimate the negative correlation between x and y , we will add $\zeta(x_j)$ to $\zeta(y_j)$, and the most frequent value that appears in many columns will be considered as the value for s^- . To determine if the two rows are positively or negatively correlated, we compare the number of columns in which the two rows are considered positively correlated to the number of columns in which the two rows are considered negatively correlated.

$$J_{C+} = \{j \mid \zeta(x_j) - \zeta(y_j) = s^+\}$$

$$J_{C-} = \{j \mid \zeta(x_j) + \zeta(y_j) = s^-\}$$

If $|J_{C+}| > |J_{C-}|$, x and y are considered positively correlated, and their coherence is computed

as $h_c(x, y) = h_+(x, y, J_{C+}) \frac{|J_{C+}|}{|NB|}$, else, x and y are considered negatively correlated, and

their coherence is computed as $h_c(x, y) = h_-(x, y, J_{C-}) \frac{|J_{C-}|}{|NB|}$. Finally, ψ_c^B in class B can be computed as

$$\psi_c^B(X) = \frac{|2|}{|I|(|I| - 1)} \sum_{x, y \in X} h_c(x, y)$$

To illustrate how this measure works, Table 4 shows the results of clustering each row in Table 2 (Here we used k-means, $k=3$. However, any other clustering algorithm can be used). The values in this table are the rankings of the clusters. For example, 1 indicates the cluster that has the lowest values in the corresponding row, and 3 indicates the cluster that has the maximum value. As an example, consider the first two rows. Subtracting $\zeta(x)$ from $\zeta(y)$ yields the following:

$$(0, 2, -1, 0, -2, 0, 0, -1, 0, -1)$$

This means that the maximum positive correlation between x and y is in 5 columns $\{c_1, c_4, c_6, c_7, c_9\}$ with $s^+ = 0$, while adding $\zeta(x)$ to $\zeta(y)$ yields

$$(6, 4, 5, 4, 4, 2, 2, 5, 6, 5)$$

This means that the maximum negative correlation between x and y is in 3 columns: $\{2, 4, 5\}$ with $s^- = 4$ or $\{c_3, c_8, c_{10}\}$ with $s^- = 5$. Hence, the coherence between x and y is computed as follows:

$$h_C(x, y) = h_+(x, y, \{c_1, c_4, c_6, c_7, c_9\}) \frac{5}{10} = 0.5$$

As another example, the last two rows (y and z) are negatively correlated in the same set of columns:

$$h_C(y, z) = h_-(y, z, \{c_1, c_4, c_6, c_7, c_9\}) \frac{5}{10} = 0.5$$

The results here are similar to those obtained using h_G in terms of the set of columns in which the two rows have the maximum coherence, which is $\{c_1, c_4, c_6, c_7, c_9\}$. Based on the clustering-based discretization method, the second proposed discriminative coherence measures is defined as follows:

$$\Delta_C^A(X) = \psi_C^A(X) = \psi_C^B(X). \quad \Delta_C^B(X) = \psi_C^B(X) - \psi_C^A(X)$$

Similar to Δ_G^A and Δ_G^B , the range of Δ_C^A and Δ_C^B is $(-1, 1)$. Our preliminary results showed that ψ_C and ψ_G produced very similar results on some of the simulated datasets. Since the computation of ψ_C is much faster than the computation of ψ_G , ψ_C is implemented in the proposed discriminative co-clustering algorithm. However, both measures will be used for evaluation purposes to quantify the resulting discriminative co-clusters using the proposed and the existing algorithms.

The Di-RAPOCC algorithm, described in Algorithm 2, optimizes for the following objective function in order to extract the discriminative co-clusters.

Definition 6. (Discriminative Objective Function) To obtain the top- K^A discriminative co-clusters from class A, the objective function can be written as:

$arg \max_{x_1, x_2, \dots, x_{K^A}} \sum_{i=1}^{K^A} \Phi^A(X)$ where $\Phi^A(X) = (HP^A(X_i) - \psi_C^B(X_i))$. To obtain the top- K^B discriminative co-clusters from class B, the objective function can be written as:

$arg \max_{x_1, x_2, \dots, x_{K^B}} \sum_{i=1}^{K^B} \Phi^B(X)$ where $\Phi^B(X) = (HP^B(X_i) - \psi_C^A(X_i))$.

5.3 The Di-RAPOCC Algorithm

We will now explain the different steps of the proposed Di-RAPOCC algorithm.

Step 1: Initialize the K^A and K^B co-clusters—First, we compute h_C^A and h_C^B for all pairs of rows. This step is preceded by clustering the values of each class. The clustering is only used to identify the set of columns in which two rows have the maximum correlation, and the original values will be used in all the steps. Hence, there is no loss of information in this step. Then, we define $\delta_C^A(x, y)$ as follows:

$$\delta_C^A(x, y) = h_C^A(x, y) - h_C^B(x, y)$$

Similarly, we can also define $\delta_C^B(x, y)$. These will be used to identify K^A groups of rows, S^A , to be used as the seeds for the co-clusters (lines 7-12). For our experiments, we chose to have the values of both K^A and K^B equal to 5. If α is the minimum number of rows in any co-cluster (which is set to 3 in our experiments), the candidate set for each row R_x is computed as follows:

$$R_x^A = arg \max_{r_1, r_2, \dots, r_\alpha} \sum_{i=1}^{\alpha} \delta_C^A(x, r_i) \quad (2)$$

From all of the M candidate sets (since there are M rows in the data matrix, each row will be a candidate to be considered as a seed for a co-cluster), the top- K^A sets are used as the initial co-clusters for each class.

$$S^A = arg \max_{S_1^A, S_2^A, \dots, S_{K^A}^A} \sum_{i=1}^{K^A} \left(\sum_{x, y \in R_i^A} \delta_C^A(x, y) \right) \quad (3)$$

Similarly, R^B and S^B will be computed for class B. Regarding the columns, all of them will be included in each co-cluster in the initialization.

Step 2: Updating the row/column clusterings—This is an iterative step in which we consider each row/column to be added/deleted from each co-cluster (lines 13-27). For each row, there are three possible assignments $\{-1, 0, 1\}$: 1 (-1) indicates adding the row to the co-cluster as positively (negatively) correlated, and 0 indicates removing the row from the corresponding co-cluster. The assignments of the columns do not consider a negative

correlation since the definition of negative correlation only comes from the rows. The same row (or column) is allowed to be included in more than one co-cluster in this step. Similar to the *RAPOCC* algorithm, the convergence of the *Di-RAPOCC* algorithm is guaranteed since the maintained objective function is bounded and optimized to be monotonically increasing.

Algorithm 2

Di-RAPOCC(D, K, a)

1: **Input:** Data matrix (D)
 No. of co-clusters (K^A and K^B)
 Minimum No. of rows in any co-cluster (a)

2: **Output:** Two sets of discriminative co-clusters ($\{X^A\}, \{X^B\}$)

3: **Procedure:**

4: **Step 1: Compute δ_C for all the rows**

5: $\forall x, y \in \{I\} \delta_C^A \leftarrow h_C^A(x, y) - h_C^B(x, y)$

6: $\forall x, y \in \{I\} \delta_C^B \leftarrow h_C^B(x, y) - h_C^A(x, y)$

7: **Initialize each of the K co-clusters for each class**

8: Compute S^A and S^B as defined in Equation (3)
 /* Initialize rows and columns of each co-cluster */

9: **for** $k = 1 : K$ **do**

10: $\forall_{m \in S_k^A} X_k^A.r(m) = 1, \forall_{n \in N^A} X_k^A.c(n) = 1$

11: $\forall_{m \in S_k^B} X_k^B.r(m) = 1, \forall_{n \in N^B} X_k^B.c(n) = 1$

12: **end for**

13: **Step 2: Update the row and the column clusterings**

14: **repeat**

15: **for** $k = 1 : K$ **do**

16: **for** $i = 1 : M$ **do**

17: $X_k^A.r(i) = \arg \max_{u \in \{-1, 0, 1\}} \Phi(X_k^A.r(i) = u)$

18: $X_k^B.r(i) = \arg \max_{u \in \{-1, 0, 1\}} \Phi(X_k^B.r(i) = u)$

19: **end for**

20: **for** $j = 1 : N^A$ **do**

21: $X_k^A.c(j) = \arg \max_{v \in \{0, 1\}} \Phi(X_k^A.c(j) = v)$

22: **end for**

23: **for** $j = 1 : N^B$ **do**

24: $X_k^B.c(j) = \arg \max_{v \in \{0, 1\}} \Phi(X_k^B.c(j) = v)$

25: **end for**

26: **end for**

27: **until convergence**

28: Step 3: Merging similar co-clusters.

29: Step 4: Pruning.

Step 3: Merging the Co-clusters—Similar to the *RAPOCC* algorithm, the goal of this step is to merge similar co-clusters using an agglomerative clustering approach. The two most similar co-clusters, within the same class, are merged in each iteration. This step allows the discovery of **large** discriminative co-clusters, and it allows **intra-class overlapping** co-clusters.

Step 4: Pruning—In this step, we prune the co-clusters with the lowest discriminative scores. To determine which co-clusters to prune, (i) sort the co-clusters based on $\Phi^A(X)$, in class *A* and $\Phi^B(X)$, in class *B*, (ii) compute the difference between the consecutive scores and (iii) report the set of co-clusters just before the largest difference, and prune the remaining co-clusters.

6 The Experimental Results

To demonstrate the effectiveness of the proposed algorithms, several experiments were conducted using both synthetic and real-world gene expression datasets.

6.1 Experimental Setup

6.1.1 Datasets—For the synthetic datasets, a set of co-clusters were implanted in randomly generated datasets using the shifting and scaling patterns [39]. Given two rows, x and y , their relationship can be represented as:

$$x_j = y_j * s_{scale} + s_{shift}$$

where s_{shift} and s_{scale} are the shifting and scaling parameters. The sign of s_{shift} determines the correlation type: if $s_{shift} > 0$, then x and y are positively correlated, and if $s_{shift} < 0$, then x and y are negatively correlated [39]. In addition, two types of synthetic datasets were used, one without noise and the other with Gaussian noise. For the real-world datasets, we used eight expression datasets in the co-clustering experiments as described in Table 5. In the discriminative co-clustering experiments, we only used the first four datasets in Table 5 since each of these datasets has two distinct classes of biological samples.

6.1.2 Comparisons with existing methods—In the co-clustering experiments, we compared the results of the *RAPOCC* algorithm against the *CC* [10], the *OPSM* [7], the *ISA* [22] and the *ROCC* [15] algorithms. We used BiCAT software (<http://www.tik.ethz.ch/sop/bicat/>) to run *CC*, *ISA* and *OPSM* algorithms using the default parameters. The code for the *ROCC* was obtained from the authors of [15]. In the discriminative co-clustering experiments, we compared the results of the *Di-RAPOCC* algorithm against the *SDC* algorithm [18] and the *OPSM* algorithm [7]. The *OPSM* algorithm is not a discriminative co-clustering algorithm. Therefore, we used the following procedure: (i) Apply *OPSM* on each class separately, (ii) compute the inter-class overlap, (iii) remove the co-clusters that have

inter-class overlap 50%, and (iv) report the remaining co-clusters. We refer to this modified algorithm as Discriminative OPSM (*Di-OPSM*). The SDC algorithm takes as input three parameters (*SDC*, *r*, *minpattsiz*) [18], which were set to the default values: (0.2, 0.2, 3) unless otherwise stated.

6.1.3 Evaluation Measures—Several measures were used such as the **number of co-clusters**, the **average size** and the **average coherence** of the co-clusters computed using Definition 3. We also used the recovery and relevance measures as defined in [31].

Recovery determines how well each of the implanted co-clusters is discovered, and **relevance** is the extent to which the generated co-clusters represent the implanted co-clusters. Given a set of implanted co-clusters denoted by Y_{imp} and a set of co-clusters obtained by an algorithm denoted by X_{res} , the recovery and the relevance can be defined as follows:

$$Recovery = \frac{1}{|Y_{imp}|} \sum_{(Y \in Y_{imp})} \arg \max_{(X \in X_{res})} \frac{|X \cap Y|}{|X \cup Y|}$$

$$Relevance = \frac{1}{|X_{res}|} \sum_{(X \in X_{res})} \arg \max_{(Y \in Y_{imp})} \frac{|X \cap Y|}{|X \cup Y|}$$

In addition, we used the following proposed metrics to evaluate the results of the discriminative co-clustering:

- **Greedy-based discriminative coherence (Δ_G)**

$$\Delta_G = \frac{1}{(K^A + K^B)} \left(\sum_{k=1}^{K^A} \Delta_G^A + \sum_{k=1}^{K^B} \Delta_G^B \right)$$

- **Clustering-based discriminative coherence (Δ_C)**

$$\Delta_C = \frac{1}{(K^A + K^B)} \left(\sum_{k=1}^{K^A} \Delta_C^A + \sum_{k=1}^{K^B} \Delta_C^B \right)$$

- **Inter-class overlap.** If X^A (X^B) is the set of discriminative co-clusters in class A (B), the inter-class overlap is defined as the average of:

$$\left(\sum_{k=1}^{K^A} \arg \max_{X_k^B} \frac{|X_k^A \cap X_k^B|}{|X_k^A \cup X_k^B|} + \sum_{k=1}^{K^B} \arg \max_{X_k^A} \frac{|X_k^B \cap X_k^A|}{|X_k^B \cup X_k^A|} \right)$$

where the union and intersection operations are computed using the rows in the co-clusters.

6.1.4 Biological Evaluation—The biological significance was estimated by calculating the p-values using the DAVID tool (<http://david.abcc.ncifcrf.gov/>) to test if a given co-cluster is enriched with genes from a particular category to a greater extent than would be expected by chance [24]. The range of the p-values is from 0 to 1. Lower p-values indicate biological significance [11].

6.2 Co-clustering Results

In this subsection, we present the results for the co-clustering experiments.

6.2.1 Results on Synthetic Data—Two types of datasets were used, one without noise and one with 10% noise. The size of each synthetic dataset is 200×150 . Two co-clusters were implanted in each dataset, and the size of each co-cluster is 50×50 . As shown in Fig. 8, the *RAPOCC* algorithm outperformed the other algorithms because it optimizes for high-quality co-clusters. As a result, fewer random data points are added to the co-clusters obtained by our algorithm.

6.2.2 Results on Real Gene Expression Data—Fig. 9 shows two co-clusters obtained by the proposed algorithm. The first co-cluster contains positively correlated genes, while the second co-cluster contains both types of correlations. The results of the five co-clustering methods on the eight datasets are shown in Table 6 and summarized in the following observations:

- **Coherence of the co-clusters.** The *RAPOCC* algorithm outperformed all the other algorithms on all the datasets. The *OPSM* and the *ROCC* algorithms performed better than the *CC* and the *ISA* algorithms. These results confirmed one of our initial claims that the proposed *RAPOCC* algorithm was designed to identify high-quality co-clusters.
- **Size of the co-clusters.** Except for the *Leukemia* dataset, the *RAPOCC* produced either the largest or the second largest co-clusters in all the datasets. The *OPSM* and the *RAPOCC* algorithms produced the largest co-clusters in four datasets and three datasets, respectively.
- **Number of the co-clusters.** The *ROCC* algorithm produced the largest number of co-clusters in all of the datasets. However, we observed that, in most of the cases, the co-clusters generated by this algorithm were either duplicates, subsets of each other or highly overlapping. On the other hand, the *ISA* algorithm did not produce any co-cluster for three datasets: *Leukemia*, *Cho yeast* and *Causton yeast*.
- **Biological significance of the co-clusters.** Fig. 10 shows the average of the percentages of the biologically significant co-clusters using the DAVID tool from all the eight gene expression datasets. As shown in this figure, our proposed algorithm outperformed all other algorithms.

Overall, the proposed algorithm produced the higher quality, more biologically significant and relatively larger co-clusters compared to the other algorithm. Furthermore, the *RAPOCC* algorithm is more robust to noise.

6.3 Discriminative Co-clustering Results

In this subsection, we present the results for discriminative co-clustering experiments. Due to space limitations, in some of the tables we used *OPM* and *RPC* to refer to *Di-OPSM* and *Di-RAPOCC* algorithms, respectively.

6.3.1 Results on Synthetic Data—Using the shifting-and-scaling model [39], four co-clusters were generated of the size 10×10 . Half of those co-clusters were designed to be discriminative, while the remaining co-clusters were common in both classes. The structure of the synthetic datasets is similar to the structure shown in Fig. 3. In the first experiment, we implanted the synthetic co-clusters in random matrices of different sizes given by $s \times 20$, where $s = (50, 100, 300, 500)$. Fig. 11 shows the relevance and recovery results of *SDC*, *Di-OPSM* and *Di-RAPOCC* co-clustering algorithms when applied to the synthetic datasets. The noise level, η , in this set of experiments is 0. The proposed algorithm outperformed other algorithms indicating that the proposed algorithm is capable of identifying the discriminative co-clusters. Since *Di-OPSM* was not directly designed to extract discriminative co-clusters, the identified co-clusters include both discriminative and non-discriminative co-clusters. The poor performance of the *SDC* algorithm can be explained by two main reasons. (i) *SDC* generates too many patterns as shown in Table 7. As the size of the dataset increases, the number of the generated patterns generated by the *SDC* algorithm *increases dramatically*. (ii) The *SDC* algorithm generates *very small patterns* (average of 3 rows per pattern). On the other hand, the *Di-RAPOCC* algorithm prunes any non-discriminative co-cluster.

In the second experiment, different levels of noise were used, which are 0, 5%, 10%, 15% and 20%, respectively, to the synthetic dataset of size 100×20 . Fig. 12 shows the recovery and the relevance of the three algorithms. As the noise level increases in the dataset, the relevance and the recovery values are degraded. However, our algorithm is still the algorithm most robust to noise due to the use of a clustering approach to estimate the coherence of any co-cluster. Table 8 shows the average results of the discriminative measurements γ_G and γ_C for all the different synthetic datasets. Unsurprisingly, our algorithm achieved the best results in all the datasets because it primarily focuses on identifying the most discriminative co-clusters in the search process. Fig. 13 shows the inter-class overlap on synthetic datasets. The *Di-RAPOCC* algorithm achieved the best results because it avoids common patterns in both of the classes.

6.3.2 Results on Real Gene Expression Data—The *SDC* algorithm was applied on the *Medulloblastoma* and the *Scleroderma* datasets with the parameters values set to (0.3, 0.3, 3) to avoid *out of memory* problems. For the *Leukemia* datasets, out of memory errors occurred for different combinations of the parameters; therefore, there are no results for this dataset. As shown in Table 9, the *Di-RAPOCC* algorithm achieved the best results in terms of the discriminative coherence measures (γ_G and γ_C). The results were also analyzed in terms of the number of co-clusters, the inter-class overlap and the average coherence as shown in Table 10. The coherence measure cannot be applied to the results of the *SDC* algorithm because it does not report the columns in which a set of rows is correlated. Here, we make some remarks regarding the performance of the three algorithms.

- The **SDC** algorithm tends to produce a large number of small patterns. Since the *SDC* algorithm uses the Apriori approach, it has some computational efficiency problems, and the number of the discovered patterns grows dramatically with larger datasets.
- The **Di-OPSM** algorithm tends to produce co-clusters that are too large. Therefore, it does not give good results in terms of the coherence, inter-class overlap and discriminative measures. Since it is not a discriminative co-clustering algorithm, we have to run it on each class independently.
- The **Di-RAPOCC** algorithm keeps the top discriminative co-clusters and prunes the other co-clusters, and it works well on noisy and large datasets.

Fig. 14 shows the biological evaluation of the results. The *SDC* algorithm was excluded from this analysis because it produced too many patterns. The *Di-RAPOCC* algorithm outperformed the *Di-OPSM* algorithm in three datasets, while *OPSM* was better in the *Leukemia* dataset. However, for this dataset, *Di-RAPOCC* outperformed *Di-OPSM* in terms of the inter-class overlap, the coherence and the discriminative coherence measures. In a different analysis, we found several significant biological pathways that were enriched in the co-clusters produced by the proposed algorithm. For example, the *MAPK signaling pathway* which has a p-value = $4.77E - 12$ was reported as an up-regulated pathway in the metastatic tumors that is very relevant to the study of metastatic disease [25]. The summary of comparisons between the three algorithms is shown in Table 11.

6.3.3 Discussion—We will now describe how the proposed model can overcome the limitations of the existing methods and can obtain the discriminative co-clusters that have all the desired characteristics mentioned earlier. (i) The proposed model incorporates the class label within each iteration step while optimizing the objective function. This will ensure to yield patterns with the maximum discriminative coherence and discriminative power. In addition, since ψ is computed for each feasible pair in both the classes, the model will generate patterns with minimum inter-class overlap. (ii) The proposed model will generate larger patterns compared to the *SDC* and other algorithms. This will be achieved by the update operations in which each row/column will be considered to be added to each pattern in each class. (iii) The pruning step will keep only the relevant patterns and remove the irrelevant ones. This will overcome the limitation of the *SDC* algorithm which generates too many unnecessary patterns. (iv) The efficient use of the clustering-based measure ψ_C for approximating the optimization criteria makes the proposed model an order of magnitude faster than the previous algorithms. (v) The proposed algorithm allows the patterns to share some columns (or conditions). This intra-class overlap is an important property of subspace patterns as described earlier.

7 Conclusion

In this paper, we presented a novel algorithm for discovering arbitrarily positioned co-clusters, and we extended this algorithm to discover discriminative co-clusters by integrating the class label in the co-clustering discovery process. Both of the proposed algorithms are robust against noise, allow overlapping and capture positive and negative correlations in the

same co-cluster. Comprehensive experiments on synthetic and real-world datasets were carried out to illustrate the effectiveness of the proposed algorithms. The results showed that both of the proposed algorithms outperformed existing algorithms and can identify co-clusters that are biologically significant. As future work, we are interested in analyzing the discriminative power of the proposed approach and extending it to solve prediction problems. Also, we plan to extend the work to other forms of subspace clustering algorithms such as correlation clustering [5].

Acknowledgements

This work was supported in part by the National Cancer Institute of the National Institutes of Health under Award Number R21CA175974 and the U.S. National Science Foundation grants IIS-1231742 and IIS-1242304. The content is solely the responsibility of the authors and does not necessarily represent the official views of the NIH and NSF.

Biographies



Omar Odibat received his Ph.D. from the Department of Computer Science at the Wayne State University. Earlier he received his M.S. from University of Jordan in 2005 and B.S. in Computer Science from Yarmouk University in 2003. His research interests are in the areas of data mining and bioinformatics. He is a student member of ACM and SIAM.



Chandan K. Reddy is an Associate Professor in the Department of Computer Science at Wayne State University. He received his Ph.D. from Cornell University and M.S. from Michigan State University. His primary research interests are in the areas of data mining and machine learning with applications to healthcare, bioinformatics, and social network analysis. His research is funded by the National Science Foundation, the National Institutes of Health, the Department of Transportation, and the Susan G. Komen for the Cure Foundation. He has published over 50 peer-reviewed articles in leading conferences and journals. He received the Best Application Paper Award at the ACM SIGKDD conference in 2010 and was a finalist of the INFORMS Franz Edelman Award Competition in 2011. He is a member of IEEE, ACM, and SIAM.

References

- [1]. Aggarwal, Charu C.; Reddy, Chandan K., editors. Data Clustering: Algorithms and Applications. Chapman and Hall/CRC; 2013.

- [2]. Alon U, Barkai N, Notterman DA, Gish K, Ybarra S, Mack D, Levine AJ. Broad patterns of gene expression revealed by clustering analysis of tumor and normal colon tissues probed by oligonucleotide arrays. *Proceedings of the National Academy of Sciences of the United States of America*. 1999; 96(12):6745–6750. [PubMed: 10359783]
- [3]. Alqadah, Faris; Bader, Joel S.; Anand, Rajul; Reddy, Chandan K. Query-based biclustering using formal concept analysis; *SIAM International Conference on Data Mining*; 2012; p. 648-659.
- [4]. Anagnostopoulos, Aris; Dasgupta, Anirban; Kumar, Ravi. Approximation algorithms for co-clustering; *PODS '08: Proceedings of the twenty-seventh ACM SIGMOD-SIGACT-SIGART symposium on Principles of database systems*; NY, USA. 2008; p. 201-210.
- [5]. Aziz, Mohammad S.; Reddy, Chandan K. *Advances in Knowledge Discovery and Data Mining*. Springer; 2010. A robust seedless algorithm for correlation clustering; p. 28-37.
- [6]. Banerjee, Arindam; Dhillon, Inderjit; Ghosh, Joydeep; Merugu, Srujana; Modha, Dharmendra S. A generalized maximum entropy approach to bregman co-clustering and matrix approximation. *J. Mach. Learn. Res.* 2007; 8:1919–1986.
- [7]. Ben-Dor, Amir; Chor, Benny; Karp, Richard; Yakhini, Zohar. Discovering local structure in gene expression data: the order-preserving submatrix problem. *Journal of computational biology*. 2003; 10(3-4):373–384. [PubMed: 12935334]
- [8]. Burdick, Doug; Calimlim, Manuel; Gehrke, Johannes. Mafia: A maximal frequent itemset algorithm for transactional databases. *ICDE*. 2001:443–452.
- [9]. Causton HC, Ren B, Koh SS, Harbison CT, Kanin E, Jennings EG, Lee TI, True HL, Lander ES, Young RA. Remodeling of yeast genome expression in response to environmental changes. *Mol Biol Cell*. 2001; 12(2):323–337. [PubMed: 11179418]
- [10]. Cheng, Yizong; Church, George M. Biclustering of expression data; *Proceedings of the Eighth International Conference on Intelligent Systems for Molecular Biology*; 2000; p. 93-103.
- [11]. Cho, Hyuk; Dhillon, Inderjit S. Cocustering of human cancer microarrays using minimum sum-squared residue coclustering. *IEEE/ACM Trans. Comput. Biol. Bioinformatics*. 2008; 5(3):385–400.
- [12]. Cho RJ, Campbell MJ, Winzeler EA, Steinmetz L, Conway A, Wodicka L, Wolfsberg TG, Gabrielian AE, Landsman D, Lockhart DJ, Davis RW. A genome-wide transcriptional analysis of the mitotic cell cycle. *Molecular cell*. 1998; 2(1):65–73. [PubMed: 9702192]
- [13]. de la Fuente, Alberto. From ‘differential expression’ to ‘differential networking’ identification of dysfunctional regulatory networks in diseases. *Trends in Genetics*. 2010; 26(7):326–333. [PubMed: 20570387]
- [14]. Deodhar, Meghana; Ghosh, Joydeep. SCOAL: A framework for simultaneous co-clustering and learning from complex data. *ACM Trans. Knowl. Discov. Data*. 2010; 4:11:1–11:31.
- [15]. Deodhar, Meghana; Gupta, Gunjan; Ghosh, Joydeep; Cho, Hyuk; Dhillon, Inderjit. A scalable framework for discovering coherent co-clusters in noisy data; *ICML '09: Proceedings of the 26th Annual International Conference on Machine Learning*; 2009; p. 241-248.
- [16]. Dhillon, Inderjit S.; Mallela, Subramanyam; Modha, Dharmendra S. Information-theoretic co-clustering; *KDD '03: Proceedings of the ninth ACM SIGKDD international conference on Knowledge discovery and data mining*; New York, NY, USA. 2003; p. 89-98.
- [17]. Fan, Hongjian; Ramamohanarao, Kotagiri. Fast discovery and the generalization of strong jumping emerging patterns for building compact and accurate classifiers. *Knowledge and Data Engineering, IEEE Transactions on*. 2006; 18(6):721–737.
- [18]. Fang, Gang; Kuang, Rui; Pandey, Gaurav; Steinbach, Michael; Myers, Chad L.; Kumar, Vipin. Subspace differential coexpression analysis: problem definition and a general approach; *Pacific Symposium on Biocomputing*; 2010; p. 145-156.
- [19]. Fang, Gang; Pandey, Gaurav; Wang, Wen; Gupta, Manish; Steinbach, Michael; Kumar, Vipin. Mining low-support discriminative patterns from dense and high-dimensional data. *IEEE Transactions on Knowledge and Data Engineering*. 2010; 99 PrePrints.
- [20]. Golub TR, Slonim DK, Tamayo P, Huard C, Gaasenbeek M, Mesirov JP, Coller H, Loh ML, Downing JR, Caligiuri MA, Bloomfield CD, Lander ES. Molecular classification of cancer: class discovery and class prediction by gene expression monitoring. *Science*. Oct; 1999 286(5439): 531–537. [PubMed: 10521349]

- [21]. Hussain, Syed Fawad; Bisson, Gilles. Text categorization using word similarities based on higher order co-occurrences. *SDM*. 2010:1–12.
- [22]. Ihmels, Jan; Bergmann, Sven; Barkai, Naama. Defining transcription modules using large-scale gene expression data. *Bioinformatics*. 2004; 20(13):1993–2003. [PubMed: 15044247]
- [23]. Jiang, Daxin; Tang, Chun; Zhang, Aidong. Cluster analysis for gene expression data: A survey. *IEEE Trans. on Knowl. and Data Eng.* 2004; 16:1370–1386.
- [24]. Liu, Jinze; Yang, Jiong; Wang, Wei. Biclustering in gene expression data by tendency; Proceedings of the 2004 IEEE Computational Systems Bioinformatics Conference, CSB '04; Washington, DC, USA. 2004; p. 182-193.
- [25]. Macdonald, Tobey J.; Brown, Kevin M.; Lafleur, Bonnie; Peterson, Katia; Lawlor, Christopher; Chen, Yidong; Packer, Roger J.; Cogen, Philip; Stephan, Dietrich A. Expression profiling of medulloblastoma: PDGFRA and the RAS/MAPK pathway as therapeutic targets for metastatic disease. *Nature Genetics*. Oct; 2001 29(2):143–152. [PubMed: 11544480]
- [26]. Madeira, Sara C.; Oliveira, Arlindo L. Biclustering algorithms for biological data analysis: a survey. *IEEE/ACM transactions on computational biology and bioinformatics*. 2004; 1(1):24–45. [PubMed: 17048406]
- [27]. Odibat, Omar; Reddy, Chandan K. A generalized framework for mining arbitrarily positioned overlapping co-clusters; Proceedings of the SIAM International Conference on Data Mining (SDM); 2011. p. 343-354.
- [28]. Odibat, Omar; Reddy, Chandan K.; Giroux, Craig N. Differential biclustering for gene expression analysis; Proceedings of the ACM Conference on Bioinformatics and Computational Biology (BCB); 2010; p. 275-284.
- [29]. Okada, Yoshifumi; Inoue, Terufumi. Identification of differentially expressed gene modules between two-class DNA microarray data. *Bioinformation*. 2009; 4(4)
- [30]. Pensa, Ruggero G.; Boulicaut, Jean-François. Constrained co-clustering of gene expression data. *SDM*. 2008:25–36.
- [31]. Prelic, Amela; Bleuler, Stefan; Zimmermann, Philip; Wille, Anja; Buhlmann, Peter; Gruissem, Wilhelm; Hennig, Lars; Thiele, Lothar; Zitzler, Eckart. A systematic comparison and evaluation of biclustering methods for gene expression data. *Bioinformatics*. 2006; 22(9):1122–1129. [PubMed: 16500941]
- [32]. Reddy, Chandan K.; Chiang, Hsiao-Dong; Rajaratnam, Bala. Trust-tech-based expectation maximization for learning finite mixture models. *Pattern Analysis and Machine Intelligence. IEEE Transactions on*. 2008; 30(7):1146–1157.
- [33]. Serin, Akdes; Vingron, Martin. Debi: Discovering differentially expressed biclusters using a frequent itemset approach. *Algorithms for Molecular Biology*. 2011; 6(1):18. [PubMed: 21699691]
- [34]. Shan, Hanhuai; Banerjee, Arindam. Residual bayesian co-clustering for matrix approximation; Proc. SIAM International Conference on Data Mining; 2010; p. 223-234.
- [35]. Shi, Xiaoxiao; Fan, Wei; Yu, Philip S. Efficient semi-supervised spectral co-clustering with constraints; IEEE International Conference on Data Mining; 2010; p. 1043-1048.
- [36]. Song, Yangqiu; Pan, Shimei; Liu, Shixia; Wei, Furu; Zhou, Michelle X.; Qian, Weihong. Constrained coclustering for textual documents. *AAAI*. 2010
- [37]. Steinbach, Michael; Karypis, George; Kumar, Vipin. In: Grobelnik, Marko; Mladenic, Dunja; Milic-Frayling, Natasa, editors. A comparison of document clustering techniques; KDD-2000 Workshop on Text Mining, August 20; 2000; p. 109-111.
- [38]. Whitfield, Michael L.; Finlay, Deborah R.; Murray, John Isaac; Troyanskaya, Olga G.; Chi, Jen-Tsan; Pergamenschikov, Alexander; McCalmont, Timothy H.; Brown, Patrick O.; Botstein, David; Connolly, M. Kari Systemic and cell type-specific gene expression patterns in scleroderma skin. *Proceedings of the National Academy of Sciences*. 2003; 100(21):12319–12324.
- [39]. Xin, Xu; Ying, Lu; Tung, Anthony K. H.; Wang, Wei. Mining shifting-and-scaling co-regulation patterns on gene expression profiles; Proceedings of the 22nd International Conference on Data Engineering, ICDE '06; 2006. p. 89

- [40]. Zhang, Lijun; Chen, Chun; Bu, Jiajun; Chen, Zhengguang; Cai, Deng; Han, Jiawei. Locally discriminative coclustering. Knowledge and Data Engineering, IEEE Transactions on. 2012; 24(6):1025–1035.

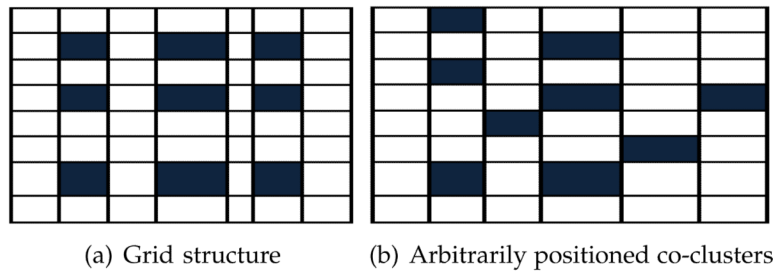


Fig. 1.
Types of co-cluster structures.

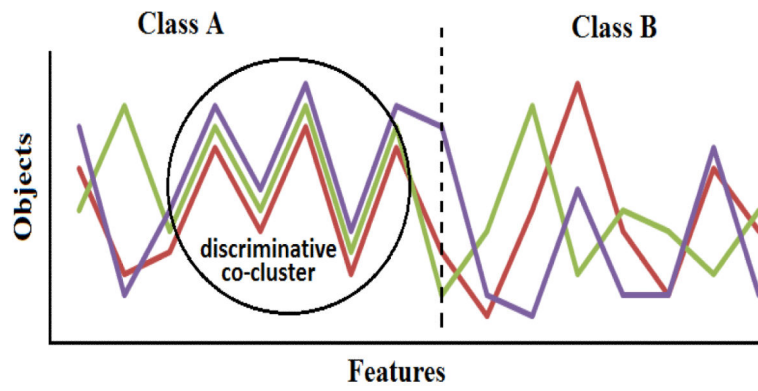


Fig. 2.

A set of three objects that are highly correlated in a subset of the features in class A, but they are not correlated in class B. Hence, these objects are considered as a discriminative co-cluster.

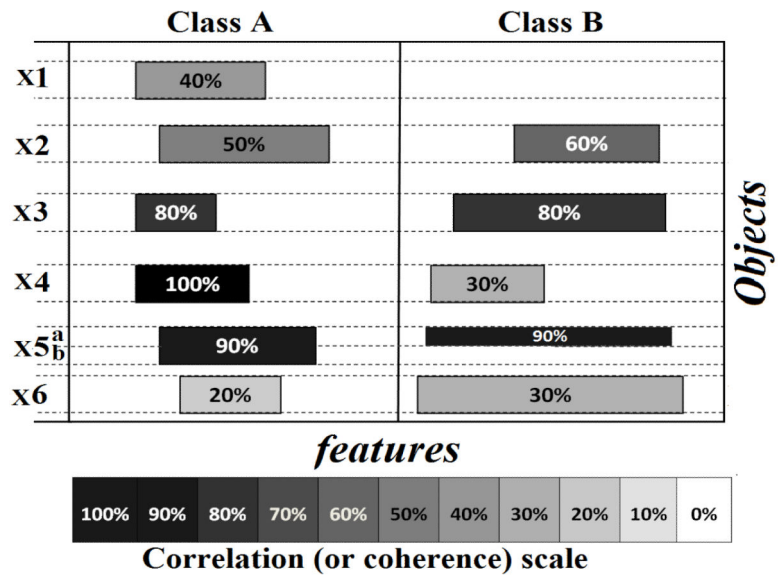


Fig. 3.
Example of discriminative co-clusters.

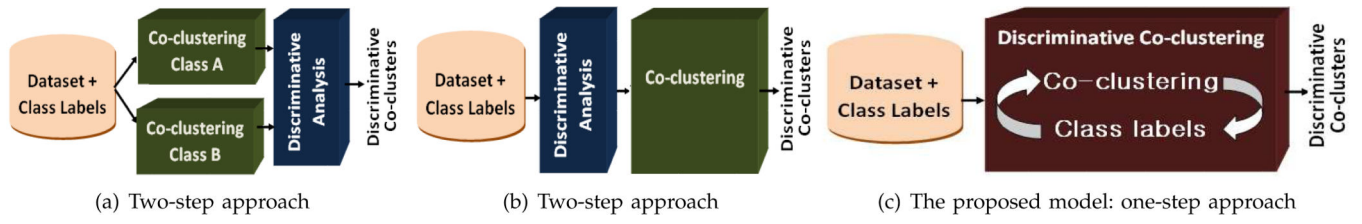


Fig. 4. Different approaches to obtain discriminative co-clusters.

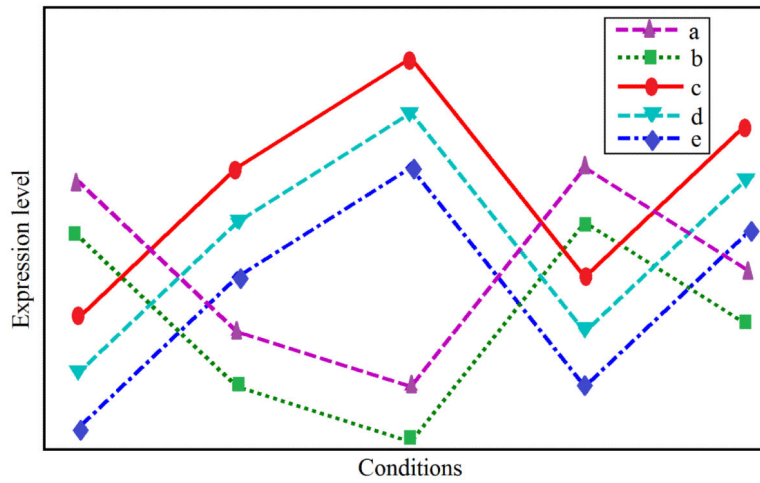


Fig. 5. Different types of relationships between genes in one co-cluster. The genes $\{a, b\}$ are positively correlated with each other, and the genes $\{c, d, e\}$ are positively correlated with each other. However, the genes $\{a, b\}$ are negatively correlated with the genes $\{c, d, e\}$.

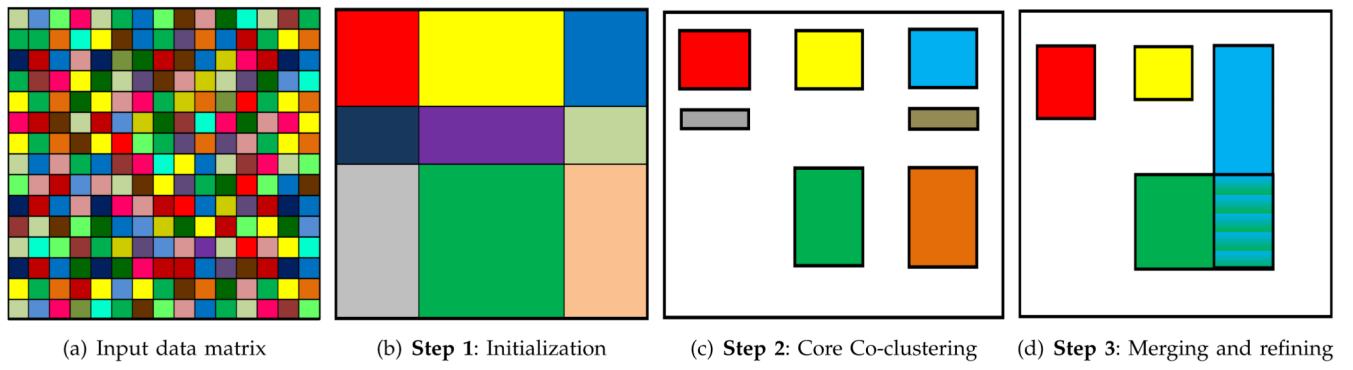


Fig. 6.
The main steps of the proposed *RAPOCC* algorithm.

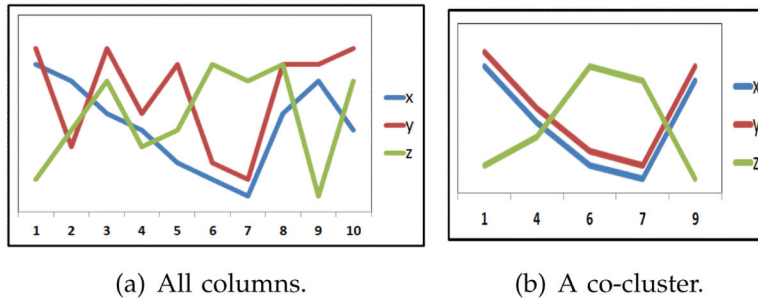


Fig. 7. (a) a plot for the entire running datasets. (b) a plot for the co-cluster extracted from the running dataset.

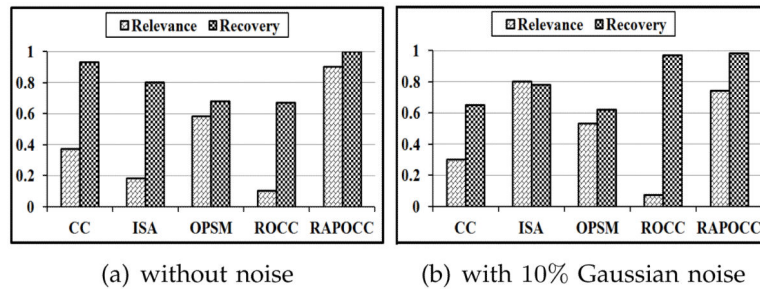


Fig. 8.
The co-clustering results on the synthetic datasets.

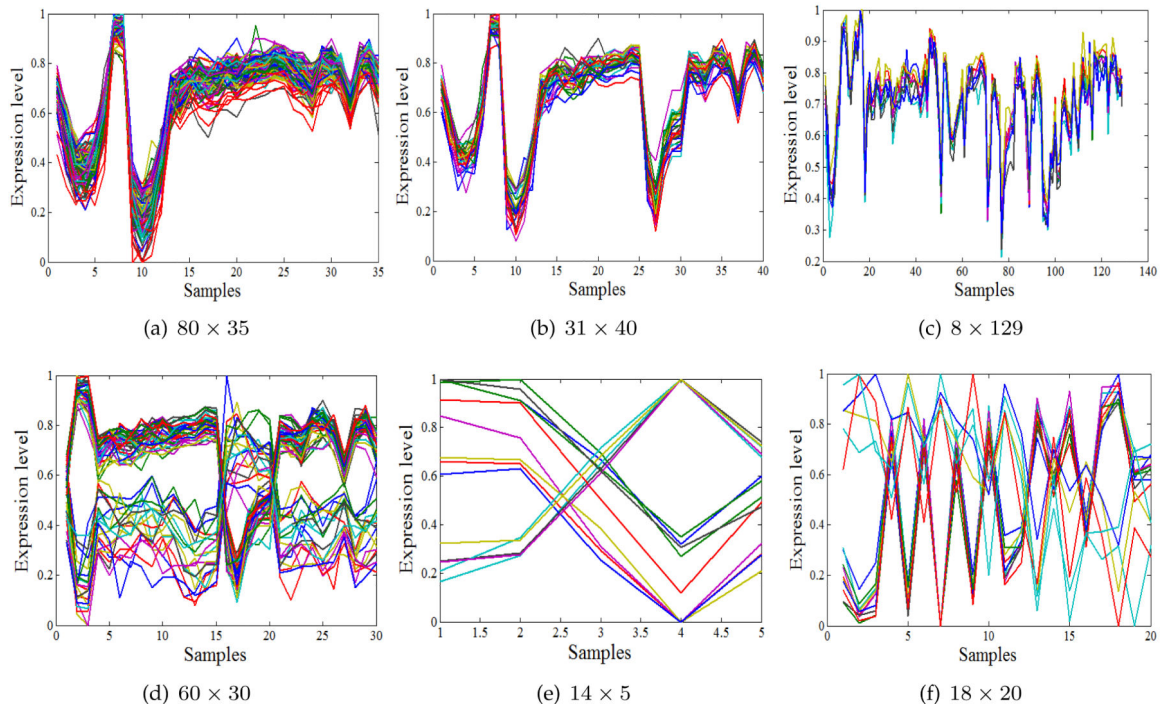


Fig. 9.

Examples of the co-clusters identified by the RAPOCC algorithm. The three co-clusters in the first row contain only the positively correlated genes which show similar patterns. These co-clusters were obtained from the Gasch yeast dataset. The three co-clusters in the second row contain positively and negatively correlated genes which show opposite patterns. These co-clusters were obtained from (d) Gasch yeast, (e) Scleroderma and (f) Causton yeast datasets.

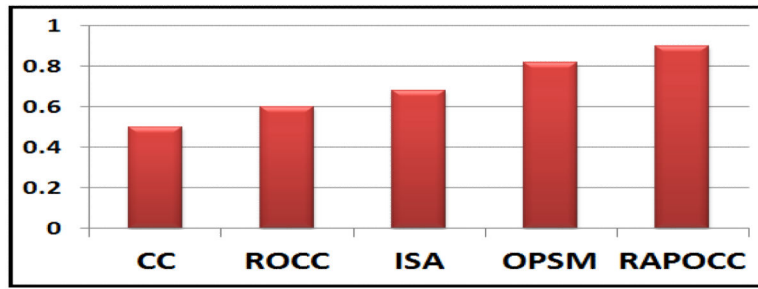


Fig. 10. Proportion of co-clusters that are significantly enriched (average of the 8 datasets).

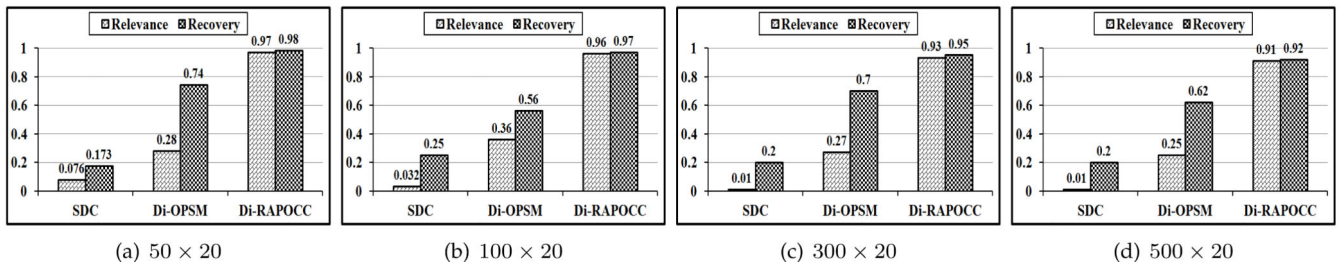


Fig. 11. Relevance and Recovery for SDC, Di-OPSM and Di-RAPOCC obtained from synthetic datasets.

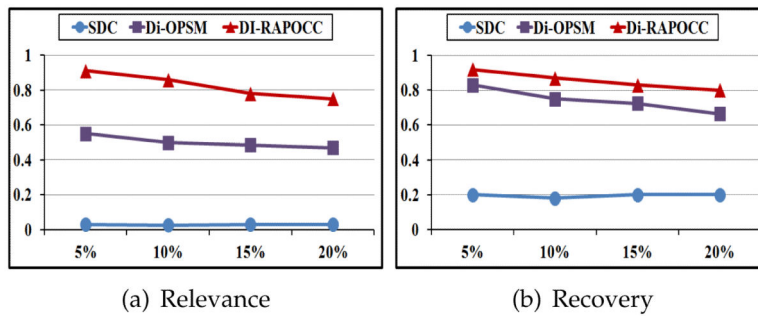


Fig. 12. Relevance and recovery obtained with noise levels of 5%, 10%, 15% and 20%, respectively.

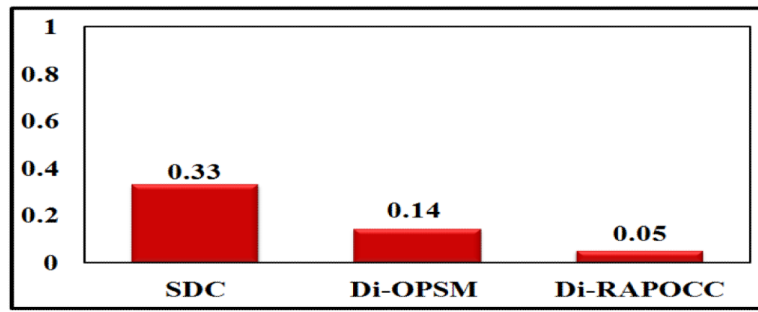


Fig. 13.
The inter-class overlapping on synthetic datasets.

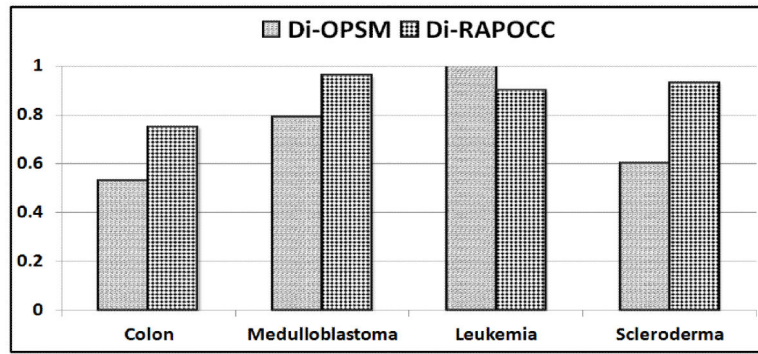


Fig. 14. Proportion of the co-clusters that are significantly enriched in each dataset (significance level = 5%).

TABLE 1

Notations used in this paper.

Notation	Description
D	input data matrix of M rows and N columns
κ	number of row clusters
ℓ	number of column clusters
ρ	mapping of row clusters
γ	mapping of column clusters
K	number of optimized co-clusters
X	Co-cluster of $ I $ rows and $ J $ columns
I	set of rows in co-cluster X
J	set of columns in co-cluster X
x_j	the j^{th} column in row x
$ \cdot $	the cardinality function
Notations for discriminative co-clustering	
N^A	No. of columns in class A
K^A	number of optimized co-clusters in class A
c_j^A	j^{th} column in class A , $1 \leq j \leq N^A $
$X_k^A.r(i)$	i^{th} row of the k^{th} co-cluster in class A
$X_k^B.c(j)$	j^{th} column of the k^{th} co-cluster in class B

TABLE 2

An illustration using a running example.

Row	c_1	c_2	c_3	c_4	c_5	c_6	c_7	c_8	c_9	c_{10}
x	9	8	6	5	3	2	1	6	8	5
y	10	4	10	6	9	3	2	9	9	10
z	2	5	8	4	5	9	8	9	1	8

TABLE 3

Results of h_G on the x and y rows in Table 2.

(i)	Columns $\{J^m\}$	$h_+(x, y, J^i)$
1	$J^1 = \{c_1, c_2, c_3, c_4, c_5, c_6, c_7, c_8, c_9, c_{10}\}$	0.9723
2	$J^2 = \{c_1, c_3, c_4, c_5, c_6, c_7, c_8, c_9, c_{10}\}$	0.9860
3	$J^3 = \{c_1, c_3, c_4, c_6, c_7, c_8, c_9, c_{10}\}$	0.9908
4	$J^4 = \{c_1, c_3, c_4, c_6, c_7, c_8, c_9\}$	0.9947
5	$J^5 = \{c_1, c_4, c_6, c_7, c_8, c_9\}$	0.9978
6	$J^6 = \{c_1, c_4, c_6, c_7, c_9\}$	1.0
7	$J^7 = \{c_4, c_6, c_7, c_9\}$	1.0
8	$J^8 = \{c_6, c_7, c_9\}$	1.0
9	$J^9 = \{c_6, c_9\}$	1.0
10	$J^{10} = \{c_9\}$	1.0
$h_G(x, y, J)$ (weighted average)		0.994

TABLE 4

Clustering of the running example dataset.

Row	c_1	c_2	c_3	c_4	c_5	c_6	c_7	c_8	c_9	c_{10}
$\zeta(x)$	3	3	2	2	1	1	1	2	3	2
$\zeta(y)$	3	1	3	2	3	1	1	3	3	3
$\zeta(z)$	1	2	3	2	2	3	3	3	1	3

TABLE 5

Description of the real-world gene expression datasets used in our experiments.

Dataset	Genes	Total samples	class A		class B	
			Description	samples	Description	samples
Leukemia [20]	5000	38	Acute lymphoblastic leukemia	11	Acute myeloid leukemia	27
Colon cancer [2]	2000	62	Normal	22	Tumor	40
Medulloblastoma [25]	2059	23	Metastatic	10	Non-metastatic	13
Scleroderma [38]	2773	27	Male	12	Female	15
Arabidopsis thaliana [31]	734	69	69 Different biological samples and pathways			
Gasch yeast [31]	2993	173	173 Time series gene expression data			
Cho yeast [12]	6240	14	17 Time points (<i>cell cycle</i>)			
Causton yeast [9]	4960	11	Response to environmental changes			

TABLE 6

Results of the five co-clustering methods on the eight gene expression datasets.

Dataset	Average coherence of co-clusters (No. of co-clusters)					Average volume of co-clusters (Avg No. of rows, Avg No. of columns)				
	CC	ISA	OPSM	ROCC	RAPOCC	CC	ISA	OPSM	ROCC	RAPOCC
Leukemia	0.9715 (20)	- (0)	0.9963 (37)	0.9775 (44)	0.9984 (25)	7611.2 (310, 15)	-	8475 (708, 20)	2544 (190, 10)	3543.7 (219, 13)
Colon	0.9884 (10)	0.9902 (21)	0.9810 (13)	0.9946 (62)	0.9986 (11)	15.5 (5.9, 3.6)	376 (148.3, 5.6)	2435 (619.2, 8.1)	881.2 (88, 10.6)	1437 (230.3, 7.8)
Medulloblastoma	0.9996 (10)	0.9906 (1)	0.9891 (10)	0.9892 (93)	0.9997 (15)	16.6 (5.9, 6.5)	10 (5, 2)	639 (225, 6)	258 (80.6, 3.2)	409.3 (82, 5)
Scleroderma	0.9838 (20)	0.9813 (2)	0.9862 (12)	0.9895 (47)	0.9950 (20)	2273.8 (110, 16)	15 (8, 2)	1303.4 (403, 8)	426 (63, 10)	1949 (380, 7)
Arabidopsis	0.9996 (20)	0.9569 (27)	0.9969 (12)	0.9952 (36)	0.9998 (20)	146.2 (19, 8)	40.6 (20, 2)	330.7 (98.8)	534.1 (41, 28)	2282.1 (191, 12)
Gasch yeast	0.9844 (20)	0.9907 (63)	0.9966 (14)	0.9945 (87)	0.9987 (25)	2424 (304, 43)	572.1 (67, 9)	2019.6 (522, 9)	2320.7 (115, 25)	2582.5 (272, 29)
Cho yeast	0.9322 (20)	- (0)	0.9923 (11)	0.9854 (33)	0.9960 (30)	950.5 (80, 12)	-	2015 (682, 7)	757.9 (152, 6)	1958 (392, 5)
Causton yeast	0.9220 (17)	- (0)	0.9907 (9)	0.9831 (20)	0.9965 (20)	2202.9 (219, 10)	-	2656.3 (941, 6)	800 (200, 4)	3897.5 (780, 5)

TABLE 7

Number of co-clusters from synthetic datasets.

Synthetic dataset	SDC	Di-OPSM	Di-RAPOCC
s=50	256	15	2
s=100	990	16	2
s=300	4, 451	16	3
s=500	10, 210	22	3

TABLE 8

Discriminative measures (synthetic datasets).

Synthetic dataset	<i>G</i>			<i>c</i>		
	SDC	OPM	RPC	SDC	OPM	RPC
$s=50, \eta=0$	0.51	0.54	0.69	0.51	0.55	0.72
$s=100, \eta=0$	0.50	0.68	0.71	0.54	0.0.54	0.70
$s=200, \eta=0$	0.49	0.63	0.70	0.54	0.66	0.71
$s=300, \eta=0$	0.52	0.51	0.67	0.51	0.64	0.70
$s=500, \eta=0$	0.51	0.64	0.71	0.52	0.63	0.72
$s=100, \eta=5\%$	0.53	0.57	0.71	0.51	0.60	0.70
$s=100, \eta=10\%$	0.52	0.65	0.67	0.53	0.61	0.65
$s=100, \eta=15\%$	0.51	0.63	0.76	0.49	0.63	0.70
$s=100, \eta=20\%$	0.52	0.64	0.72	0.50	0.61	0.65

TABLE 9

Discriminative measures (expression datasets)

Dataset	<i>G</i>			<i>C</i>		
	SDC	OPM	RPC	SDC	OPM	RPC
Colon	0.60	0.58	0.62	0.50	0.53	0.56
Medulloblastoma	0.49	0.54	0.59	0.51	0.53	0.55
Leukemia	-	0.57	0.59	-	0.56	0.58
Scleroderma	0.57	0.54	0.60	0.54	0.55	0.60

TABLE 10

The average No. of co-clusters, the average overlap and the average coherence

Dataset	No. of co-clusters in A			No. of co-clusters in B			Overlap			Average coherence (H)	
	SDC	OPM	RPC	SDC	OPM	RPC	SDC	OPM	RPC	OPM	RPC
Colon	155	10	15	1	3	13	0.0	0.01	0.04	0.992	0.997
Medulloblastoma	74,957	8	14	7,597	9	14	0.2	0.12	0.01	0.988	0.994
Leukemia	-	21	35	-	5	22	-	0.40	0.09	0.990	0.995
Scleroderma	48,623	12	10	469	10	9	0.04	0.17	0.0	0.986	0.998

TABLE 11

Comparisons between the three discriminative co-clustering algorithms.

Measure	SDC	OPM	RPC
No. of the co-clusters	High	Low	Medium
Size of the co-clusters	Small	Large	Medium
Coherence	-	Low	High
Discriminative coherence	Low	Medium	High
Inter-class overlap	High	Medium	Low
Recovery	Low	Medium	High
Relevance	Low	Medium	High