

# Package ‘BicARE’

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**Date** 2008-06-05

**Title** Biclustering Analysis and Results Exploration

**Depends** R (>= 1.8.0), Biobase (>= 2.5.5), multtest, GSEABase, GO.db

**Imports** methods

**Suggests** hgu95av2

**Author** Pierre Gestraud

**Maintainer** Pierre Gestraud <pierre.gestraud@curie.fr>

**Description** Biclustering Analysis and Results Exploration.

**License** GPL-2

**URL** <http://bioinfo.curie.fr>

**biocViews** Microarray, Transcription, Clustering

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BicARE-package

*BicARE***Description**

Biclustering Analysis and Results Exploration

**Details**

Package: BicARE  
 Version: 0.1.0  
 Date: 2008-06-05  
 Depends: R (>= 1.8.0), Biobase, multtest, GSEABase  
 License: GPL  
 biocViews: Microarray, Transcription, Statistics, Clustering  
 URL: <http://bioinfo.curie.fr>  
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**Index:**

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makeReport	Export the results as html files
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sample.biclustering	Example biclustering object
testAnnot	Find samples annotations over-represented in biclusters
testSet	Find gene sets that are enriched in a bicluster

Further information is available in the following vignettes:

BicARE [BicARE \(source, pdf\)](#)

**Author(s)**

Pierre Gestraud

Maintainer: Pierre Gestraud , <[pierre.gestraud@curie.fr](mailto:pierre.gestraud@curie.fr)>

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bicluster	<i>Extract a bicluster</i>
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**Description**

Extract a bicluster from an object of class biclustering

**Usage**

```
bicluster(biclustering, k, graph=TRUE)
```

**Arguments**

biclustering	an object of class "biclustering" created by function <a href="#">FLOC</a>
k	the number of the bicluster considered in the "biclustering" object
graph	boolean, indicating whether the graph should be plotted or not

**Value**

Returns the bicluster as a matrix with the genes on rows and the samples on columns. Result matrix is of class "bicluster". The "graph" option allows to plot the expression profiles of the genes across the conditions in the bicluster.

**Author(s)**

Pierre Gestraud

**Examples**

```
### extract the first bicluster
data(sample.biclustering)
sample.biclustering
bic <- bicluster(sample.biclustering, 1, graph=TRUE)
plot(bic)
```

---

FLOC	<i>Performs the FLOC algorithm</i>
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**Description**

Find a given number of biclusters using the a modified version of the FLOC algorithm.

**Usage**

```
FLOC(Data, k = 20, pGene = 0.5, pSample=pGene, r = NULL, N = 8, M
= 6, t = 500, blocGene = NULL, blocSample = NULL)
```

**Arguments**

Data	an <a href="#">ExpressionSet</a> or a matrix (with genes on rows and conditions on columns)
k	the number of biclusters searched
pGene	genes initial probability of membership to the biclusters
pSample	samples initial probability of membership to the biclusters
r	the residue threshold
N	minimal number of gene per bicluster
M	minimal number of conditions per bicluster
t	number of iterations
blocGene	a matrix indicating the directed initialisation for the genes (see details)
blocSample	a matrix indicating the directed initialisation for the conditions (see details)

**Details**

This biclustering algorithm is based on the FLOC algorithm (FLexible Overlapped biClustering) defined by Yang et al. (see references). It can discover a set of  $k$ , possibly overlapping, biclusters. If  $r$  is set to NULL, the residue threshold used in the analysis is the residue of `Data` divided by 10.

`blocGene` and `blocSample` are matrix of 0 and 1 with the rows representing the features (gene or samples) and the columns the biclusters. A 1 on line  $i$  and column  $j$  indicates that the feature  $i$  (gene or sample) will be include in the bicluster  $j$  during the initialisation step and will not be removed from it during the analysis. If the number of columns in these matrices is different from the number of bicluster searched,  $k$  is set to the maximal value of these two.

See [bicluster](#) to extract a bicluster from the biclustering result.

**Value**

Returns an object of class 'biclustering', a list containing at least :

Call	the matched call.
ExpressionSet	the data used
param	a data.frame with the algorithm parameters
bicRow	a matrix of boolean indicating the belonging of the genes to the biclusters
bicCol	the same as for <code>bicRow</code> but for the conditions
mat.resvol.bic	a matrix describing the biclusters

**Author(s)**

Pierre Gestraud (<pierre.gestraud@curie.fr>)

**References**

J. Yang, H. Wang, W. Wang, and P.S. Yu. An improved biclustering method for analyzing gene expression. *International Journal on Artificial Intelligence Tools*, 14(5):771-789, 2005

## Examples

```
data(sample.bicData)    ## subset of sample.ExpressionSet from Biobase
residue(sample.bicData) ## 0.3401921
resBic <- FLOC(sample.bicData, k=10, pGene=0.5, r=0.05, N=8, M=10, t=500)
resBic

## initialising samples of 2 biclusters
iniSample <- matrix(0, ncol=2, nrow=26)
## first bicluster initialised around Female cases
iniSample[pData(sample.bicData)$sex=="Female",1] <- 1
## second bicluster initialised around control cases
iniSample[pData(sample.bicData)$type=="Control",2] <- 1
resBic <- FLOC(sample.bicData, k=10, pGene=0.5, r=0.05, N=8, M=10, t=500, blocSample=iniSample)
resBic
```

---

makeReport

*Export the results as html files*

---

## Description

Creates a directory with html files containing the biclustering results.

## Usage

```
makeReport(dirPath, dirName, resBic, browse=TRUE)
```

## Arguments

dirPath	path to the directory
dirName	the name of the directory where the report will be created
resBic	a biclustering result
browse	logical. If TRUE the web browser will be opened

## Details

makeReport produces a html report of biclustering results in a new directory named dirName. If the browse argument is set to TRUE the web browser will be opened on the "home.html" file.

Make sure to have rights to create the result directory.

## Author(s)

Pierre Gestraud <pierre.gestraud@curie.fr>

## Examples

```
data(sample.biclustering)
dirPath <- getwd() ## report created in the current working directory
dirName <- "test"
makeReport(dirPath, dirName, sample.biclustering, browse=FALSE)
```

residue *Residue of a matrix*

---

**Description**

Returns the residue of a matrix.

**Usage**

```
residue(Data)
```

**Arguments**

Data            an [ExpressionSet-class](#) or a matrix

**Details**

This function computes the residue of a matrix as defined by Yang et al (see references).

**Author(s)**

Pierre Gestraud

**References**

J. Yang, H. Wang, W. Wang, and P.S. Yu. An improved biclustering method for analyzing gene expression. *International Journal on Artificial Intelligence Tools*, 14(5):771-789, 2005

**See Also**

[FLOC](#)

**Examples**

```
data(sample.bicData)
residue(sample.bicData)
```

---

sample.bicData            *Example data set for BicARE*

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**Description**

A subset of sample.ExpressionSet from package Biobase. The data for 26 cases, labeled A to Z and 350 genes. Each case has three covariates: sex (male/female), type (case/control) and score (testing score).

**Usage**

```
sample.bicData
```

**Format**

An ExpressionSet

---

sample.biclustering     *Example biclustering object*

---

### Description

A biclustering object created by the [FLOC](#) function on the sample.bicData with the following options : k=10, pGene = 0.3, pSample = 0.5, r = 0.025, N = 8, M = 8, t = 1000.

### Usage

```
sample.biclustering
```

### Format

a biclustering object

---

testAnnot     *Find samples annotations over-represented covariates in biclusters*

---

### Description

Characterisation of the biclusters in term of over-representation of sample covariates.

### Usage

```
testAnnot(resBic, annot=NULL, covariates="all")
```

### Arguments

resBic	a biclustering result from <a href="#">FLOC</a>
annot	annotation matrix, default value is set to NULL, then phenoData of the ExpressionSet is used
covariates	the names of the covariates that should be tested, default value is set to "all"

### Details

For each bicluster and each covariate a chi-squared test is performed to test the adequation between the distribution of the levels of the covariates in the bicluster and in the original dataset.

Multiple testing correction is performed by the Benjamini-Yekutieli procedure. The residuals of the tests indicate if the level is over or down represented in the bicluster.

Due to the amount of results it is advised to use the [makeReport](#) function to get a html report.

**Value**

A biclustering object containing `resBic` and updated with the results of the tests in `resBic$covar`. The results are presented as a list with :

<code>covar</code>	the samples covariates tested
<code>pvalues</code>	a matrix with the p-values of the tests
<code>adjpvalues</code>	a matrix with the p-values adjusted by the Benjamini Yekutieli procedure
<code>index</code>	a list of matrices with the numbers of each level in each bicluster
<code>residuals</code>	a list of matrices with the residuals of the tests for each modality in each bicluster

**Author(s)**

Pierre Gestraud

**Examples**

```
data(sample.biclustering)
resBic <- testAnnot(sample.biclustering, annot=NULL, covariates=c("sex", "type"))
```

---

<code>testSet</code>	<i>Find gene sets that are enriched in a bicluster</i>
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---

**Description**

Test of the over-representation of gene sets in the biclusters

**Usage**

```
testSet(resBic, geneSetCol)
```

**Arguments**

<code>resBic</code>	a biclustering object created by <a href="#">FLOC</a>
<code>geneSetCol</code>	a <a href="#">GeneSetCollection-class</a>

**Details**

The over-representation of a gene set in a bicluster is evaluated by an hypergeometric test.

The genes identifiers of the gene sets will automatically be mapped to the same as those used in the data.

Due to the amount of results it is advised to use the [makeReport](#) function to get a html report.

**Value**

A biclustering object containing `resBic` and updated with the results of the tests in `resBic$geneSet`. The results are presented as a list with :

<code>GeneSetCollection</code>	the <code>GeneSetCollection</code> used
<code>pvalues</code>	a matrix containing the pvalues of the tests for each <code>geneSet</code> and each bicluster
<code>adjpvalue</code>	a matrix containing the p-values adjusted by the Benjamini Yekutieli procedure



**Author(s)**

Pierre Gestraud <pierre.gestraud@curie.fr>

**Examples**

```
data(sample.biclustering)
gss <- GeneSetCollection(sample.biclustering$ExpressionSet[1:50,], setType=GOCollection())
resBic <- testSet(sample.biclustering, gss)
```

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