Package ‘CCPROMISE’

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Type Package

Title PROMISE analysis with Canonical Correlation for Two Forms of High Dimensional Genetic Data

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Description Perform Canonical correlation between two forms of high dimensional genetic data, and associate the first component of each form of data with a specific biologically interesting pattern of associations with multiple endpoints. A probe level analysis is also implemented.

Depends R (>= 3.3.0), stats, methods, CCP, PROMISE, Biobase, GSEABase, utils

License GPL (>= 2)

biocViews Microarray, GeneExpression

LazyLoad yes

NeedsCompilation no

R topics documented:

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Description

a tool to identify genes that are correlated between two set of genomic variables and are associated with a predefined pattern of associations with multiple endpoint variables.

Details

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The CCPROMISE (Canonical correlation with PROMISE analysis) is performed by calling function CCPROMISE. The two forms of genomic data such as gene expression and methylation are passed through minimal ExpressionSet; the gene annotation (defining relationship between a gene and the two forms of genomic data), phenotypic data and definition of R routines for calculating association statistics with individual endpoint variable are same as in PROMISE package. Please refer to PROMISE package for writing user defined routines.

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References


Examples

```r
## load data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)
data(exmplPat)
```
## Perform CCPROMISE test

test<- CCPROMISE(geneSet=exmplGeneSet,
                 ESet=exmplESet,
                 MSet=exmplMSet,
                 promise.pattern=exmplPat,
                 strat.var=NULL,
                 prlbl=NULL,
                 EMlbl=c("Expr", "Methyl"),
                 nbperm=TRUE,
                 max.ntail=10,
                 nperms=100,
                 seed=13)

---

CANN Canonical Correlation of Two Sets of Genomic Data

**Description**

Compute canonical correlation between two sets of genomic data.

**Usage**

CANN (geneSet, Edat, Mdat, EMlbl = c("Expr", "Methyl"), phdat)

**Arguments**

- **geneSet**: a gene set collection to annotate probes to gene
- **Edat**: data frame of the first form of genomic data, such as gene expression data with row being probes and column being subjects. The column names should match the row names phdat
- **Mdat**: data frame of the second form of genomic data, such as methylation data with row being probes and column being subjects. The column names should match the row names phdat
- **EMlbl**: label of the genomic data, default=c("Expr", "Methyl") for Edat and Mdat
- **phdat**: phenotype data with row being subjects and column being phenotype variables. The column names should match the column names of Edat and Mdat

**Details**

The function performs Canonical correlation between two forms genomic data for each gene (Edat and Mdat) defined by *gann*. If a gene only has one form of genomic data, the first principal component is used; If one form of data has numberof probesets exceeding the number of subjects, the first number of subjects probesets are used. The function return a list of three components. See *value* for details.
Value

The output of the function is a list of length 3 with three components:

- **CCres**: canonical correlation result: a data frame with row for each gene and six columns (Gene: gene names; n.EMlbl[1]: number of probes of first form genomic data; n.EMlbl[2]: number of probes of second form genomic data; CanonicalCR: Canonical correlation of first components; WilksPermPval: permutation p value of Wilks’ Lambda; WilksAsymPval: p value of F-approximations of Wilks’ Lambda).

- **FSTccscore**: the first component of canonical correlation: a data frame with row for each gene, first half of columns for first component of first form genomic data and second half of columns for first component of second form genomic data.

- **CCload**: a data frame of loading (each row is for a gene, first column is gene names, second column is the probeset ids of first form genomic data separated by '1', third column is the load for each probeset in first form genomic data separated by '1', fourth column is the probeset ids of second form genomic data separated by '1', fifth column is the load for each probeset in second form genomic data separated by '1')

Author(s)

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References


See Also

CCPROMISE

Examples

```r
## load example data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)
## Perform canonical correlation test
test1 <- CANN(geneSet=exmplGeneSet,
              Edat=exprs(exmplESet),
              Mdat=exprs(exmplMSet),
              EMlbl=c("Expr", "Methyl"),
              phdat=pData(exmplESet))
```

Description

PROMISE analysis of two genomic sets with multiple phenotypes under a predefined association pattern at gene level.
Usage

CCPROMISE (geneSet, ESet, MSet, promise.pattern, strat.var = NULL, prlbl = NULL, EMlbl = c("Expr", "Mthyl"), nbperm = FALSE, max.ntail = 100, nperms = 10000, seed = 13)

Arguments

geneSet a gene set collection to annotate probes to gene
ESet an ExpressionSet class contains minimum of exprs (expression matrix) of first form of genomic data such as gene expression and phenoData (AnnotatedDataFrame of end point data). Please refer to Biobase for details on how to create such an ExpressionSet expression set.
MSet an ExpressionSet class of second form of genomic data such as methylation levels, the subject id of MSet and ESet should be exactly same
promise.pattern PROMISE pattern
strat.var stratum variable
prlbl label of the genomic data, default=c('Expr', 'Methyl') for ESet and MSet
nbperm indicator of fast permutation using negative binomial strategy, taking two valid values: FALSE or TRUE. The default is FALSE.
max.ntail number of sucess if nbperm = T. Further permutation will not be performed for gene(s) or gene set(s) which max.ntail permuted statistics are greater or equal to the observed statistics, The default is 100.
nperms number of permutation, default = 10,000
seed initial seed of random number generator. The default is 13.

Details

The function performs PROMISE analysis for two forms of genomic data in minimal expression set format with a prefined phenotypic pattern. It calls two external function CANN and PROMISE2

Value

The output is a list of length 4. The 4 components are as following:

PRres PROMISE result for the first component of canonical correlation between two forms of genomic data. individual genes’ test statistics and p-values for each individual endpoint and PROMISE analysis
CCres result of canonical correlation analysis with six columns: Gene: Gene names; n.EMlbl[1]: number of probe set in the first form data; n.EMlbl[2]: number of probe set in the second form data; CanonicalCR: Canonical correlation of first components; WilksPermPval: permutation p value of Wilks’ Lambda; WilksAsymPval: p value of F-approximations of Wilks’ Lambda.
FSTccscore loads of first component of canonical correlation: a data frame of loading (each row is for a gene, first column is gene names, second column is the probe set ids of first form genomic data seperated by 'l', third column is the load for each probe set in first form genomic data seperated by 'l', fourth column is the probe set ids of second form genomic data seperated by 'l', fifth column is the load for each probe set in second form genomic data seperated by 'l')
CCload  a data frame of loading (each row is for a gene, first column is gene names, second column is the probeset ids of first form genomic data seperated by '|', third column is the load for each probeset in first form genomic data seperated by '|', fourth column is the probeset ids of second form genomic data seperated by '|', fifth column is the load for each probeset in second form genomic data seperated by '|')

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References

See Also
CANN PROMISE2

Examples
```r
## load data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)
data(exmplPat)
## Perform canonical correlation test
test<- CCPROMISE(geneSet=exmplGeneSet,
    ESet=exmplESet,
    MSet=exmplMSet,
    promise.pattern=exmplPat,
    strat.var=NULL,
    prlbl=NULL,
    EMlbl=c("Expr", "Methyl"),
    nbperm=FALSE,
    max.ntail=10,
    nperms=100,
    seed=13)
```
**exmplGeneSet**

**Value**

an example ExpressionSet contains conceptual data of 105 expression features measured by U133A array for 151 subjects. The phenotype data has 8 columns for the same 151 subjects.

**Description**

An conceptual example of gene set collection to annotate both form of genomic data to genes. The gene names can be extracted by method of setName() and probe ids can be extracted by method of geneIds().

**Usage**

data(exmplGeneSet)

**Value**

a conceptual gene set collection of 10 genes with 319 unique U133A expression probe ids or Infinium HumanMethylation450 probe ids.

---

**exmplMSet**

**Example of Conceptual Methylation Set**

**Description**

An conceptual ExpressionSet class contains minimum of exprs (matrix) of DNA methylation and phenoData (AnnotatedDataFrame of end point data).

**Usage**

data(exmplMSet)

**Value**

an conceptual example ExpressionSet of 735 DNA methylation probe ids for 151 subjects. The phenotype data has 8 columns for the same 151 subjects.
### exmplPat

**Example of Conceptual Phenotype Pattern Definition Set**

**Description**

An conceptual example of phenotype pattern definition set with three columns: stat.coef, stat.func, and endpt.vars; It defines an association pattern for three phenotypes.

**Usage**

data(exmplPat)

**Value**

a data frame

---

### PrbCor

**Probe Level Correlation of Two Sets of Genomic Data**

**Description**

Compute Spearman correlation of all probe combination between two sets of genomic data within a gene.

**Usage**

PrbCor (geneSet, Edat, Mdat, EMLbl = c("Expr", "Methyl"), phdat, pcut = 0.05)

**Arguments**

- **geneSet**: a gene set collection to annotate probes to gene
- **Edat**: data frame of the first form of genomic data, such as gene expression data with row being probes and column being subjects. The column names should match the row names phdat
- **Mdat**: data frame of the second form of genomic data, such as methylation data with row being probes and column being subjects. The column names should match the row names phdat
- **EMLbl**: label of the genomic data, default=c("Expr", "Methyl") for Edat and Mdat
- **phdat**: phenotype data with row being subjects and column being phenotype variables. The row names should match the column names of Edat and Mdat
- **pcut**: p value cutoff to eliminate probe pairs that are not significantly correlated. Default is 0.05

**Details**

The function performs Spearman correlation for all probe pairs between two forms genomic data within each gene (Edat and Mdat) defined by gen. If a gene only has one form of genomic data, the other form is coded as NA. The function return a list of two components. See value for details.
PrbPROMISE

Value

The output of the function is a list of length 2. The 2 components are as following:

- **res**: spearman correlation result: a data frame with row for each probe pair with correlation p value < pcut and five columns; Gene: Gene names; EMlbl[1]: probe id in the first form data; EMlbl[2]: probe id in the second form data; Spearman.rstat: Spearman r statistics; Spearman.p: Spearman p value.

- **gen**: Probe level data: a data frame with row for each probe pairs, first half of columns for first form genomic data and second half of columns for second form genomic data with sign reflecting the correlation of the probe pair.

Author(s)

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See Also

CCPROMISE

Examples

```r
## load example data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)

## Perform canonical correlation test

test1 <- PrbCor(geneSet = exmplGeneSet,
                  Edat = exprs(exmplESet),
                  Mdat = exprs(exmplMSet),
                  EMlbl = c("Expr", "Mthyl"),
                  phdat = pData(exmplESet))
```

PROMISE Analysis with Two Forms of Genomic Data at Probe Level

Description

PROMISE analysis of two genomic sets with multiple phenotypes under a predefined association pattern at probe level.

Usage

```r
PrbPROMISE (geneSet, ESet, MSet, promise.pattern, strat.var = NULL,
             prlbl = NULL, EMlbl = c("Expr", "Mthyl"), pcut = 0.05, nbperm = FALSE,
             max.ntail = 100, nperms = 10000, seed = 13)
```
Arguments

geneSet
  a gene set collection to annotate probes to gene

ESet
  an ExpressionSet class contains minimum of exprs (expression matrix) of first
  form of genomic data such as gene expression and phenoData (AnnotatedDataFrame
  of end point data). Please refer to Biobase for details on how to create such an
  ExpressionSet expression set.

MSet
  an ExpressionSet class of second form of genomic data such as methylation
  levels, the subject id of MSet and ESet should be exactly same

promise.pattern
  PROMISE pattern

strat.var
  stratum variable

prbl
  labels

EMlbl
  label of the genomic data, default=c(‘Expr’, ’Methyl’) for ESet and MSet

pcut
  p value cutoff to eliminate probe pairs that are not significantly correlated. De-
  fault is 0.05

nbperm
  indicator of fast permutation using negative binomial strategy, taking two valid
  values: FALSE or TRUE. The default is FALSE.

max.ntail
  number of success if nbperm = T. Further permutation will not be performed for
  gene(s) or gene set(s) which max.ntail permuted statistics are greater or equal
  to the observed statistics, The default is 100.

nperms
  number of permutation, default = 10,000

seed
  initial seed of random number generator. The default is 13.

Details

The function performs PROMISE analysis for two forms of genomic data in minimal expression set
format with a predefined phenotypic pattern. It calls two external function PrbCor and PROMISE2

Value

The output of the function is a list of length 2. The 2 components are as following:

PRres
  PROMISE result for the first component of canonical correlation between two
  forms of genomic data. individual genes’ test statistics and p-values for each
  individual endpoint and PROMISE analysis

CORres
  result of spearman correlation analysis of probe pairs within a gene with five
  columns: Gene: Gene names; EMlbl[1]: probe id in the first form data; EMlbl[2]:
  probe id in the second form data; Spearman.rstat: Spearman r statistics; Spear-
  man.p: Spearman p value.

Author(s)

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See Also

PrbCor PROMISE2
## load data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)
data(exmplPat)
## Perform probe level PROMISE analysis
test<-

```r
PROMISE2
```

### Examples

```r
## load data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)
data(exmplPat)
## Perform probe level PROMISE analysis
test<-
```

```r
test<-PrbPROMISE(geneSet=exmplGeneSet,
 ESet=exmplESet,
 MSet=exmplMSet,
 promise.pattern=exmplPat,
 strat.var=NULL,
 prlbl=c('LC50', 'MRD22', 'EFS', 'PR3'),
 EMlbl=c("Expr", "Methyl"),
 nbperm=TRUE,
 max.ntail=10,
 nperms=100,
 seed=13)
```

---

**PROMISE2**

**PROMISE Analysis of Two Genomic Sets**

---

**Description**

PROMISE analysis of two genomic sets with multiple phenotypes.

**Usage**

```r
PROMISE2 (exprSet, exprSet2, geneSet = NULL, promise.pattern,
 strat.var = NULL, nbperm = FALSE, max.ntail = 100, nperms = 10000,
 seed = 13)
```

**Arguments**

- `exprSet`: expression set of first genomic data
- `exprSet2`: expression set of second genomic data
- `geneSet`: geneSet should be NULL.
- `promise.pattern`: PROMISE pattern
- `strat.var`: stratum variable
- `nbperm`: indicator of fast permutation using negative binomial strategy, taking two valid values: FALSE or TRUE. The default is FALSE.
- `max.ntail`: number of sucess if nbperm = T. Further permutation will not be performed for gene(s) or gene set(s) which max.ntail permuted statistics are greater or equal to the observed statistics. The default is 100.
- `nperms`: number of permutation, default = 10,000
- `seed`: random seed, default = 13
Details

The function performs PROMISE analysis for two set genomic data with a predefined phenotypic pattern. It is an intermediate function called by CCPROMISE to perform PROMISE analysis with canonical correlation.

Value

The output of the function is a list of length 2. The 2 components are as following:

- `generes` individual genes’ test statistics and p-values for each individual endpoint and PROMISE analysis.
- `setres` Gene set level analysis is not implemented with value `NULL`.

Author(s)

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See Also

CCPROMISE

Examples

```r
## load data
data(exmplESet)
data(exmplMSet)
data(exmplGeneSet)
data(exmplPat)
## Perform canonical correlation test
test <- PROMISE2(exmplESet[1:10],
                 exmplMSet[1:10],
                 promise.pattern=exmplPat,
                 strat.var=NULL,
                 nbperm=FALSE,
                 max.ntail=10,
                 nperms=100,
                 seed=13)
```
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