

Package ‘nempi’

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Type Package

Title Inferring unobserved perturbations from gene expression data

Version 1.4.0

Depends R (>= 4.1), mnm

Description Takes as input an incomplete perturbation profile and differential gene expression in log odds and infers unobserved perturbations and augments observed ones. The inference is done by iteratively inferring a network from the perturbations and inferring perturbations from the network. The network inference is done by Nested Effects Models.

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Encoding UTF-8

LazyData true

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Imports e1071, nnet, randomForest, naturalsort, graphics, stats, utils, matrixStats, epiNEM

VignetteBuilder knitr

Suggests knitr, BiocGenerics, rmarkdown, RUnit

BugReports <https://github.com/cbg-ethz/nempi/issues>

URL <https://github.com/cbg-ethz/nempi/>

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R topics documented:

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| | |
|---------|-----------------------|
| classpi | <i>Classification</i> |
|---------|-----------------------|

Description

Builds and uses different classifiers to infer perturbation profiles

Usage

```
classpi(
  D,
  unknown = "",
  full = TRUE,
  method = "svm",
  size = NULL,
  MaxNWts = 10000,
  ...
)
```

Arguments

| | |
|---------|--|
| D | either a binary effects matrix or log odds matrix as for Nested Effects Models (see package 'nem') |
| unknown | colname of samples without mutation data, E.g. "" |
| full | if FALSE, does not change the known profiles |
| method | either one of svm, nn, rf |
| size | parameter for neural network (see package 'nnet') |
| MaxNWts | parameters for neural network (see package 'nnet') |
| ... | additional parameters for mnem::nem |

Value

plot

Author(s)

Martin Pirkl

Examples

```
D <- matrix(rnorm(1000*100), 1000, 100)
colnames(D) <- sample(seq_len(5), 100, replace = TRUE)
Gamma <- matrix(sample(c(0,1), 5*100, replace = TRUE, p = c(0.9, 0.1)), 5,
100)
Gamma <- apply(Gamma, 2, function(x) return(x/sum(x)))
Gamma[is.na(Gamma)] <- 0
rownames(Gamma) <- seq_len(5)
result <- classpi(D)
```

nempi

Main function for NEM based perturbation imputation.

Description

Infers perturbations profiles based on a sparse perturbation matrix and differential gene expression as log odds

Usage

```
nempi(
  D,
  unknown = "",
  Gamma = NULL,
  type = "null",
  full = TRUE,
  verbose = FALSE,
  logtype = 2,
  null = TRUE,
  soft = TRUE,
  combi = 1,
  converged = 0.1,
  complete = TRUE,
  mw = NULL,
  max_iter = 100,
  keepphi = TRUE,
  start = NULL,
  phi = NULL,
  ...
)
```

Arguments

| | |
|-----------|--|
| D | either a binary effects matrix or log odds matrix as for Nested Effects Models (see package 'nem') |
| unknown | colname of samples without mutation data, E.g. "" |
| Gamma | matrix with expectations of perturbations, e.g. if you have a binary mutation matrix, just normalize the columns to have sum 1 |
| type | "null": does not use the unknown samples for inference at the start, "random" uses them in a random fashion (not recommended) |
| full | if FALSE, does not change the known profiles |
| verbose | if TRUE gives more output during inference |
| logtype | log type for the log odds |
| null | if FALSE does not use a NULL node for uninformative samples |
| soft | if FALSE discretizes Gamma during the inference |
| combi | if combi > 1, uses a more complex algorithm to infer combinatorial perturbations (experimental) |
| converged | the absolute difference of log likelihood till convergence |
| complete | if TRUE uses the complete-data loglikelihood (recommended for many E-genes) |
| mw | if NULL infers mixture weights, otherwise keeps them fixed |
| max_iter | maximum iterations of the EM algorithm |
| keepphi | if TRUE, uses the previous phi for the next inference, if FALSE always starts with start network (and empty and full) |
| start | starting network as adjacency matrix |
| phi | if not NULL uses only this phi and does not infer a new one |
| ... | additional parameters for the nem function (see package mnem, function nem or mnem::nem) |

Value

nempi object

Author(s)

Martin Pirkl

Examples

```
D <- matrix(rnorm(1000*100), 1000, 100)
colnames(D) <- sample(seq_len(5), 100, replace = TRUE)
Gamma <- matrix(sample(c(0,1), 5*100, replace = TRUE, p = c(0.9, 0.1)), 5,
100)
Gamma <- apply(Gamma, 2, function(x) return(x/sum(x)))
Gamma[is.na(Gamma)] <- 0
rownames(Gamma) <- seq_len(5)
result <- nempi(D, Gamma = Gamma)
```

| | |
|---------|-------------------------------|
| nempibs | <i>Bootstrapping function</i> |
|---------|-------------------------------|

Description

Bootstrap algorithm to get a more stable result.

Usage

```
nempibs(D, bsruns = 100, bssize = 0.5, replace = TRUE, ...)
```

Arguments

| | |
|---------|--|
| D | either a binary effects matrix or log odds matrix as |
| bsruns | number of bootstraps |
| bssize | number of E-genes for each bootstrap |
| replace | if TRUE, actual bootstrap, if False sub-sampling |
| ... | additional parameters for the function nempi |

Value

list with aggregate Gamma and aggregate causal network phi

Author(s)

Martin Pirkl

Examples

```
D <- matrix(rnorm(1000*100), 1000, 100)
colnames(D) <- sample(seq_len(5), 100, replace = TRUE)
Gamma <- matrix(sample(c(0,1), 5*100, replace = TRUE, p = c(0.9, 0.1)), 5,
100)
Gamma <- apply(Gamma, 2, function(x) return(x/sum(x)))
Gamma[is.na(Gamma)] <- 0
rownames(Gamma) <- seq_len(5)
result <- nempibs(D, bsruns = 3, Gamma = Gamma)
```

pifit

Accuracy computation

Description

Compares the ground truth of a perturbation profile with the inferred profile

Usage

```
pifit(x, y, D, unknown = "", balanced = FALSE, propagate = TRUE, knowns = NULL)
```

Arguments

| | |
|-----------|--|
| x | object of class nempi |
| y | object of class mnemsim |
| D | data matrix |
| unknown | label for the unlabelled samples |
| balanced | if TRUE, computes balanced accuracy |
| propagate | if TRUE, propagates the perturbation through the network |
| knowns | subset of P-genes that are known to be perturbed (the other are neglected) |

Value

list of different accuracy measures: true/false positives/negatives, correlation, area under the precision recall curve, (balanced) accuracy

Author(s)

Martin Pirkl

Examples

```
library(mnem)
seed <- 42
Pgenes <- 10
Egenes <- 10
samples <- 100
uniform <- floor((Pgenes*Egenes)*0.1)
Nems <- mw <- 1
noise <- 1
multi <- c(0.2, 0.1)
set.seed(seed)
simmini <- simData(Sgenes = Pgenes, Egenes = Egenes,
Nems = Nems, mw = mw, nCells = samples,
uniform = uniform, multi = multi,
badCells = floor(samples*0.1))
data <- simmini$data
```

```
ones <- which(data == 1)
zeros <- which(data == 0)
data[ones] <- rnorm(length(ones), 1, noise)
data[zeros] <- rnorm(length(zeros), -1, noise)
lost <- sample(1:ncol(data), floor(ncol(data)*0.5))
colnames(data)[lost] <- ""
res <- nempi(data)
fit <- pifit(res, simmini, data)
```

plot.nempi

Plotting nempi

Description

Plot function for an object of class 'nempi'.

Usage

```
## S3 method for class 'nempi'
plot(x, barlist = list(), heatlist = list(), ...)
```

Arguments

| | |
|----------|---|
| x | object of class 'nempi' |
| barlist | additional arguments for function 'barplot' from package 'graphics' |
| heatlist | additional arguments for function 'HeatmapOP' from package 'epiNEM' |
| ... | additional arguments for function 'plotDnf' from package 'mnem' |

Value

Plots of the optimal network phi and perturbation matrix.

Author(s)

Martin Pirkl

Examples

```
D <- matrix(rnorm(1000*100), 1000, 100)
colnames(D) <- sample(seq_len(5), 100, replace = TRUE)
result <- nempi(D)
plot(result)
```

plotConvergence.nempi *Plot convergence of EM*

Description

Produces different convergence plots based on a nempi object

Usage

```
## S3 method for class 'nempi'  
plotConvergence(x, type = "b", ...)
```

Arguments

| | |
|------|--------------------------------|
| x | nempi object |
| type | see ?plot.default |
| ... | additional parameters for plot |

Value

plot

Author(s)

Martin Pirkel

Examples

```
D <- matrix(rnorm(1000*100), 1000, 100)  
colnames(D) <- sample(seq_len(5), 100, replace = TRUE)  
Gamma <- matrix(sample(c(0,1), 5*100, replace = TRUE, p = c(0.9, 0.1)), 5,  
100)  
Gamma <- apply(Gamma, 2, function(x) return(x/sum(x)))  
Gamma[is.na(Gamma)] <- 0  
rownames(Gamma) <- seq_len(5)  
result <- nempi(D, Gamma = Gamma)  
par(mfrow=c(2,3))  
plotConvergence(result)
```


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