# Package 'diffGeneAnalysis'

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Title Performs differential gene expression Analysis

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Description Analyze microarray data
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# **Description**

assocAnalysis performs the Associative-T method of differential gene analysis.

# Usage

```
assocAnalysis(bAdjusted, numSlides, ctrl, exp, sdrefgrp,
pctrl, pexpm,ctrlavjs, expmavjs, ctrlsds, expmsds, rgrplen)
```

# Arguments

numSlidesnumSlides represents the total number of chips in the microarray experiment.ctrlctrl is the number of control chips in the microarray experiment.expexp is the number of control chips in the microarray experiment.sdrefgrpstandard deviation of the computed refernce group.
exp exp is the number of control chips in the microarray experiment.
sdrefgrp standard deviation of the computed refernce group.
pctrl p values of the control chips on performin ttests.
pexpm p values of the experiment chips on performin ttests.
ctrlavjs averages of genes across all control chips.
expmavjs averages of genes across all experiment chips.
ctrlsds standard deviations of genes across all control chips.
expmsds standard deviations of genes across all experiment chips.
rgrplen total number of genes of the computed refernce group.

# Details

AssociativeAnalysis performs the Associative-T method of differential gene analysis. The results are displayed in a 10 column matrix as follows

Column Representation.

- 1 Gene Bank Id
- 2 Average Signal of the Control Chips/Channels.
- 3 SD of Control Chips/Channels.
- 4 Probability that a given gene in the Control Chips/Channels belongs or does not belong to background.
- 5 Average Signal of the Experimental Chips/Channels.
- 6 SD of Experimental Chips/Channels.
- 7 Probability that a given gene in the Experimental Chips/Channels belongs or does not belong to background.

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- 8 P-value from a Student T-test.
- 9 P-value from an Associative T-test.
- 10 Ratio of mean expression values (Control/Experimental).
- 11 Group Number.

Group Numbers are defined as follows:.

- A1 Expressed above background in both sample types, but over-expressed on the Experimental Chips/Channels.
- A2 Expressed above background in both sample types, but over-expressed on the Control Chips/Channels.
- A3 Expressed above background only on the Experimental Chips/Channel.
- A4 Expressed above background only in the Control Chips/Channel.
- 0 None of the above.

#### Value

A matrix of 11 columns as described in the details and n rows where n stands for the number of genes in the microarray dataset.

#### Author(s)

Choudary L Jagarlamudi

#### References

Dozmorov I,Centola,M. An associative analysis of gene expression array data.Bioinformatics.2003 Jan22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization,adjustment and analysis of gene expression data. Bioonformatics. 2004 Dec 12;20(18):3687-90

# **Examples**

#see refGroup for more details

biasAdjust

baisAdjust performs bias adjustment of normalized data

# Description

biasAdjust takes a normalized dataset and applies a multiplicative scalar derived from the data to help account for expression biases. These expression biases can come from many sources including dye bias, hybridization efficacy, changes in personnel, etc. After bias adjustment the data is ready for differential gene analysis.

# Usage

biasAdjust(normalized, numSlides)

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# Arguments

normalized normalized is a matrix of normalized microarray data. The first row consists of

headers ans the first column consists of gene names.

numSlides is the total number of chips of the microarray dataset.

#### Value

Returns a matrix of bias adjusted normalized data.

# Note

biasAdjust takes 2-3 seconds to execute under optimal conditions of size of datasets and speed of the machine..Tested on a Pentium IV 1.6Ghz, 256Mb RAM with 22464 genes and 10 chips it took 12 seconds to execute.

# Author(s)

Choudary L Jagarlamudi

#### References

Dozmorov I, Centola, M. An associative analysis of gene expression array data. Bioinformatics. 2003 Jan 22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization, adjustment and analysis of gene expression data. Bioonformatics. 2004 Dec 12;20(18):3687-90

# **Examples**

#biasAdjust(normalized,7)

curveFit

CurveFit data to a Gaussian distribution

# **Description**

CurveFit takes a vector of chipdata from microarray slides and fits the data to a Gaussian distribution through a non-linear least-squares optimization algorithm. The results are graphically depicted in a series of histograms. Each histogram represents a different initial seed (left to right: 2 bins, 3 bins, 4 bins, 4.5 bins, 5 bins, and 5.5 bins) that is passed to the curve fitting algorithm. The resulting fit for each histogram is superimposed with a solid blue line. The user is then able to visually select the 'best' fit.

# Usage

```
curveFit(chipdata, plot)
```

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#### **Arguments**

chipdata a vector of chipdata from microarray slides.

plot plot can take values of 1 or 0. If plot is 1 then the histogram with the curve fit

will be shown graphically.

#### Value

an object res which is a list containing the following components. res1]: mean of the computed background. res[2]: standard deviation of the computed background.

# Author(s)

Choudary L Jagarlamudi

# References

Dozmorov I,Centola,M. An associative analysis of gene expression array data. Bioinformatics.2003 Jan22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization, adjustment and analysis of gene expression data. Bioonformatics. 2004 Dec 12;20(18):3687-90

#### **Examples**

#see normalize for details.

dataTrim

DataTrim using pseudo winsorization algorithm

# **Description**

DataTrim cleans the data through a pseudo winsorization algorithm. First, the mean and SD are calculated. Then, any values above 2SD are trimmed. The mean is recalculated and any values less then 2 SD are trimmed. This process of cutting 2 SD above the mean and then 2 SD below the mean is repeated until no further cuts are possible.

# Usage

dataTrim(chipdata)

# **Arguments**

chipdata chipdata is a vector of chipdata from microarray chips.

### Value

Returns a vector of trimmed chipdata representing background based on pseudo winsorization algorithm.

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# Author(s)

Choudary L Jagarlamudi

#### References

Dozmorov I,Centola,M. An associative analysis of gene expression array data. Bioinformatics.2003 Jan22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization,adjustment and analysis of gene expression data. Bioonformatics.2004 Dec 12;20(18):3687-90

# **Examples**

#see normalize for details

|--|

# Description

Normalization of data utilizing information obtained from background fluorescence. Background fluoresce intensity values are used to determine a Gaussian distribution of lowly expressed genes, yielding the background estimates (mean and standard deviation).

# Usage

normalize(rawdata, numSlides,ctrl,expm,ctrlbg,expmbg)

# **Arguments**

rawdata	rawdata is matrix of microarray data. The first column consists of gene Names and the first row consists of headers.
numSlides	numSlides represents the total number of chips/slides in the microarray dataset including control and experiment.Control slides are always followed by experiment slides from left to right in the matrix.
ctrl	ctrl represents the total number of control chips in the microarray dataset.
expm	expm represents the total number of experiment chips in the microarray dataset.
ctrlbg	ctrlbg represents the percent of data to pick for background computation of the control chips.30 percent is the default.
expmbg	expmbg represents the percent of data to pick for background computation of the experiment chips.30 percent is the default.

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#### **Details**

The normalization algorithm trims the data based on innitial emperical estimates of the mean and standard deviation. All data beyond +/-2SD of the mean are cut iteratively. This procedure is repeated until no more cuts can be made. The trimmed data is then subjected to a non linear curve fitting procedure. The user is presented with six different pictures obtained using bars 2,3,4,4.5,5,and5.5 as mean. The user is given the freedom to select the best visual estimate of background. The user selected parameters are used to perform a z-Trnasformation on the data. The percent of data selected to compute background depends on the data obtained. The default is 30 percent. A normal distributed histogram should confirm that, else the user is allowed to pick a percent and make changes until the user sees a normal distributed histogram. Upon running normalize the user is presented with a set of 6 histograms. If the user is not happy with the default 30 percent, the user should go ahead and select a mean and confirm curvefit, then select 'no' to confirm histogram distribution. The user will be presented with a new set of 6 histograms. This process is repeated until the user selects the best Histogram distribution. This process is repeated for each individual chip.

# Value

A matrix of normalized values of rawdata

#### Author(s)

Choudary L Jagarlamudi

#### References

Dozmorov I,Centola,M. An associative analysis of gene expression array data. Bioinformatics.2003 Jan22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization,adjustment and analysis of gene expression data. Bioonformatics.2004 Dec 12;20(18):3687-90

# **Examples**

```
#rawdata is loaded in the package. Run example as follows:
#Read the description file for best results.
#data(rawdata)
#normalize(rawdata,7,3,4,0.15,0.60)
```

rawdata

Micro array dataset

# **Description**

Microarray dataset consisting of 7 chips.3 control and 4 experiment with 2382 genes. The first column consists of gene names, hence the dataset has 8 columns in total.

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#### Usage

```
data(rawdata)
```

#### **Format**

A data frame with 2382 observations on the following 11 variables.

# GeneID Gene IDS or Gene names

- c1 a numeric vector of gene intensities for control chip 1
- c2 a numeric vector of gene intensities for control chip 2
- c3 a numeric vector of gene intensities for control chip 3
- e1 a numeric vector of gene intensities for experiment chip 1
- e2 a numeric vector of gene intensities for experiment chip 2
- e3 a numeric vector of gene intensities for experiment chip 3
- e4 a numeric vector of gene intensities for experiment chip 4

#### Source

Dozmorov I,Centola,M. An associative analysis of gene expression array data. Bioinformatics.2003 Jan22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization,adjustment and analysis of gene expression data. Bioonformatics.2004 Dec 12;20(18):3687-90

# **Examples**

data(rawdata)

refGroup Reference Group computes a robust estimate of inter-assay variability (Standard Error)

# **Description**

refGroup takes a normalized, bias adjusted dataset and computes a robust estimate of inter-assay variability (Standard Error). This value is used here to perform T-tests. It can also be used in sample size calculations. The associative analysis method is then applied to the computed reference group.

# Usage

```
refGroup(biasAdjusted, numSlides, ctrl, exp, pval)
```

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#### **Arguments**

biasAdjusted biasAdjusted is a matrix of bias adjusted normalized microarray data. The first row consists of headers and the first column must consist of Gene Bank Id's.

numSlides numSlides represents the total number of chips in a microarray dataset including

control and experiment.

ctrl ctrl represents the total number of ctrl chips in the microarray.

exp exp represents the total number of experiment chips in the microarray.

pval pval is the strigency value for computing the reference group. 0.05 is the nor-

mally suggested value.

#### **Details**

AssociativeAnalysis performs the Associative-T method of differential gene analysis. The user is asked to enter values for E and R.E stand for the increase in fold over background and R stands for the ratio of experimental chips average over control chips average. The higher these values the higher will be the stringency. Example dataset used here was run with an E values of 1 and R value of 1.5.

The results are displayed in a 10 column matrix as follows

Column Representation.

- 1 Gene Bank ID
- 2 Average Signal of the Control Chips/Channels.
- 3 SD of Control Chips/Channels.
- 4 Probability that a given gene in the Control Chips/Channels belongs or does not belong to background.
- 5 Average Signal of the Experimental Chips/Channels.
- 6 SD of Experimental Chips/Channels.
- 7 Probability that a given gene in the Experimental Chips/Channels belongs or does not belong to background.
- 8 P-value from a Student T-test.
- 9 P-value from an Associative T-test.
- 10 Ratio of mean expression values (Control/Experimental).
- 11 Group Number.

Group Numbers are defined as follows:.

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- A2 Expressed above background in both sample types, but over-expressed on the Control Chips/Channels.
- A3 Expressed above background only on the Experimental Chips/Channel.
- A4 Expressed above background only in the Control Chips/Channel.
- 0 None of the above.

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# Value

A matrix of 11 columns as described in the details and n rows where n stands for the number of genes in the microarray dataset.

# Note

refGroup takes 4-5 seconds to execute under optimal conditions of size of datasets and speed of the machine. Tested on Pentium IV 1.6Ghz, 256 MB RAM with 22464 genes and 10 chips it took 25 seconds to execute.

# Author(s)

Choudary L Jagarlamudi

# References

Dozmorov I,Centola,M. An associative analysis of gene expression array data.Bioinformatics. 2003 Jan22;19(2):204-11

Knowlton N,Dozmorov I, Centola M. Microarray data Analysis Tool box(MDAT): for normalization,adjustment and analysis of gene expression data. Bioonformatics. 2004 Dec 12;20(18):3687-90

# **Examples**

#refGroup(bAdjusted,7,3,4,0.05)

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