

Package ‘AMARETTO’

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Type Package

Title Regulatory Network Inference and Driver Gene Evaluation using Integrative Multi-Omics Analysis and Penalized Regression

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Depends R (>= 3.6), impute, doParallel, grDevices, dplyr, methods, ComplexHeatmap

Description Integrating an increasing number of available multi-omics cancer data remains one of the main challenges to improve our understanding of cancer. One of the main challenges is using multi-omics data for identifying novel cancer driver genes. We have developed an algorithm, called AMARETTO, that integrates copy number, DNA methylation and gene expression data to identify a set of driver genes by analyzing cancer samples and connects them to clusters of co-expressed genes, which we define as modules. We applied AMARETTO in a pancancer setting to identify cancer driver genes and their modules on multiple cancer sites. AMARETTO captures modules enriched in angiogenesis, cell cycle and EMT, and modules that accurately predict survival and molecular subtypes. This allows AMARETTO to identify novel cancer driver genes directing canonical cancer pathways.

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biocViews

StatisticalMethod,DifferentialMethylation,GeneRegulation,GeneExpression,MethylationArray,Transcription,Preproce

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AMARETTO_CreateModuleData
AMARETTO_CreateModuleData

Description

AMARETTO_CreateModuleData

Usage

AMARETTO_CreateModuleData(AMARETTOinit, AMARETTOresults)

Arguments

AMARETTOinit List output from AMARETTO_Initialize().
 AMARETTOresults
 List output from AMARETTO_Run()

Value

result

Examples

```
data('ProcessedDataLIHC')
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
                                   NrModules = 2, VarPercentage = 50)
AMARETTOresults <- AMARETTO_Run(AMARETTOinit)
AMARETTO_MD <- AMARETTO_CreateModuleData(AMARETTOinit, AMARETTOresults)
```

AMARETTO_CreateRegulatorPrograms
AMARETTO_CreateRegulatorPrograms

Description

AMARETTO_CreateRegulatorPrograms

Usage

```
AMARETTO_CreateRegulatorPrograms(AMARETTOinit, AMARETTOresults)
```

Arguments

AMARETTOinit List output from AMARETTO_Initialize().
AMARETTOresults List output from AMARETTO_Run()

Value

result

Examples

```
data('ProcessedDataLIHC')  
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,  
                                   NrModules = 2, VarPercentage = 50)  
AMARETTOresults <- AMARETTO_Run(AMARETTOinit)  
AMARETTO_RP <- AMARETTO_CreateRegulatorPrograms(AMARETTOinit,AMARETTOresults)
```

AMARETTO_Download *AMARETTO_Download*

Description

Downloading TCGA dataset for AMARETTO analysis

Usage

```
AMARETTO_Download(CancerSite = "CHOL",  
                  TargetDirectory = TargetDirectory)
```

Arguments

CancerSite TCGA cancer code for data download
TargetDirectory Directory path to download data

Value

result

Examples

```
TargetDirectory <- file.path(getwd(),"Downloads/");dir.create(TargetDirectory)
CancerSite <- 'CHOL'
DataSetDirectories <- AMARETTO_Download(CancerSite,TargetDirectory = TargetDirectory)
```

```
AMARETTO_EvaluateTestSet
      AMARETTO_EvaluateTestSet
```

Description

Code to evaluate AMARETTO on a new gene expression test set. Uses output from AMARETTO_Run() and CreateRegulatorData().

Usage

```
AMARETTO_EvaluateTestSet(AMARETTOresults = AMARETTOresults,
  MA_Data_TestSet = MA_Data_TestSet,
  RegulatorData_TestSet = RegulatorData_TestSet)
```

Arguments

AMARETTOresults
AMARETTO output from AMARETTO_Run().

MA_Data_TestSet
Gene expression matrix from a test set (that was not used in AMARETTO_Run()).

RegulatorData_TestSet
Test regulator data from CreateRegulatorData().

Value

result

Examples

```
data('ProcessedDataLIHC')
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
  NrModules = 2, VarPercentage = 50)

AMARETTOresults <- AMARETTO_Run(AMARETTOinit)
AMARETTOtestReport <- AMARETTO_EvaluateTestSet(AMARETTOresults = AMARETTOresults,
  MA_Data_TestSet = AMARETTOinit$MA_matrix_Var,
  RegulatorData_TestSet = AMARETTOinit$RegulatorData)
```

AMARETTO_ExportResults

AMARETTO_ExportResults

Description

Retrieve a download of all the data linked with the run (including heatmaps)

Usage

```
AMARETTO_ExportResults(AMARETTOinit, AMARETTOresults, data_address,
  Heatmaps = TRUE, CNV_matrix = NULL, MET_matrix = NULL)
```

Arguments

| | |
|-----------------|-------------------------------|
| AMARETTOinit | AMARETTO initialize output |
| AMARETTOresults | AMARETTO results output |
| data_address | Directory to save data folder |
| Heatmaps | Output heatmaps as pdf |
| CNV_matrix | CNV_matrix |
| MET_matrix | MET_matrix |

Value

result

Examples

```
data('ProcessedDataLIHC')
TargetDirectory <- file.path(getwd(),"Downloads/");dir.create(TargetDirectory)
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
  NrModules = 2, VarPercentage = 50)

AMARETTOresults <- AMARETTO_Run(AMARETTOinit)
AMARETTO_ExportResults(AMARETTOinit,AMARETTOresults,TargetDirectory,Heatmaps = FALSE)
```

AMARETTO_HTMLreport

AMARETTO_HTMLreport

Description

Retrieve an interactive html report, including gene set enrichment analysis if asked for.

Usage

```
AMARETTO_HTMLreport(AMARETTOinit, AMARETTOresults, ProcessedData,
  show_row_names = FALSE, SAMPLE_annotation = NULL, ID = NULL,
  hyper_geo_test_bool = FALSE, hyper_geo_reference = NULL,
  output_address = "./", MSIGDB = TRUE, driverGSEA = TRUE,
  phenotype_association_table = NULL)
```

Arguments

AMARETTOinit AMARETTO initialize output
 AMARETTOresults AMARETTO results output
 ProcessedData List of processed input data
 show_row_names if True, sample names will appear in the heatmap
 SAMPLE_annotation SAMPLE annotation will be added to heatmap
 ID ID column of the SAMPLE annotation data frame
 hyper_geo_test_bool Boolean if a hyper geometric test needs to be performed. If TRUE provide a GMT file in the hyper_geo_reference parameter.
 hyper_geo_reference GMT file with gene sets to compare with.
 output_address Output directory for the html files.
 MSIGDB TRUE if gene sets were retrieved from MSIGDB. Links will be created in the report.
 driverGSEA if TRUE, module drivers will also be included in the hypergeometric test.
 phenotype_association_table a Data Frame, containing all modules phenotype association data. Optional.

Value

result

Examples

```

## Not run:
data('ProcessedDataLIHC')
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
                                   NrModules = 2, VarPercentage = 50)

AMARETTOresults <- AMARETTO_Run(AMARETTOinit)

AMARETTO_HTMLreport(AMARETTOinit= AMARETTOinit,AMARETTOresults= AMARETTOresults,
                    ProcessedData = ProcessedDataLIHC,
                    hyper_geo_test_bool=FALSE,
                    output_address='./')

## End(Not run)

```

AMARETTO_Initialize *AMARETTO_Initialize (version: reorder and filter MA_Matrix)*

Description

Code used to initialize the seed clusters for an AMARETTO run. Requires processed gene expressions (rna-seq or microarray), CNV (usually from a GISTIC run), and methylation (from MethylMix, provided in this package) data. Uses the function CreateRegulatorData() and results are fed into the function AMARETTO_Run().

Usage

```
AMARETTO_Initialize(ProcessedData = ProcessedData, Driver_list = NULL,
  NrModules, VarPercentage, PvalueThreshold = 0.001,
  RsquareThreshold = 0.1, pmax = 10, NrCores = 1, OneRunStop = 0,
  method = "union", random_seeds = NULL, convergence_cutoff = 0.01)
```

Arguments

| | |
|--------------------|--|
| ProcessedData | List of Expression, CNV and MethylMix data matrices, with genes in rows and samples in columns. |
| Driver_list | Custom list of driver genes to be considered in analysis |
| NrModules | How many gene co-expression modules should AMARETTO search for? Usually around 100 is acceptable, given the large number of possible driver-passenger gene combinations. |
| VarPercentage | Minimum percentage by variance for filtering of genes; for example, 75% would indicate that the CreateRegulatorData() function only analyses genes that have a variance above the 75th percentile across all samples. |
| PvalueThreshold | Threshold used to find relevant driver genes with CNV alterations: maximal p-value. |
| RsquareThreshold | Threshold used to find relevant driver genes with CNV alterations: minimal R-square value between CNV and gene expression data. |
| pmax | 'pmax' variable for glmnet function from glmnet package; the maximum number of variables aver to be nonzero. Should not be changed by user unless she/he fully understands the AMARETTO algorithm and how its parameters choices affect model output. |
| NrCores | A numeric variable indicating the number of computer/server cores to use for parallelization. Default is 1, i.e. no parallelization. Please check your computer or server's computing capacities before increasing this number. Parallelization is done via the RParallel package. Mac vs. Windows environments may behave differently when using parallelization. |
| OneRunStop | OneRunStop |
| method | Perform union or intersection of the driver genes evaluated from the input data matrices and custom driver gene list provided. |
| random_seeds | A numeric vector of length 2, containing two seed numbers for randomization : 1st for kmeans and 2nd for glmnet |
| convergence_cutoff | A numeric value (E.g. 0.01) representing the fraction of the total number of genes, in which, The algorithm is considered reaching convergence and will stop, if Nr of Gene-replacements in an iteration falls below this threshold * total number of genes. |

Value

result

Examples

```
data('ProcessedDataLIHC')
data('Driver_Genes')
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
                                   NrModules = 2, VarPercentage = 50)

## Not run:
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
                                   Driver_list = Driver_Genes[['MSigDB']],
                                   NrModules = 2, VarPercentage = 50)

## End(Not run)
```

AMARETTO_LarsenBased *AMARETTO_LarsenBased*

Description

AMARETTO_LarsenBased

Usage

```
AMARETTO_LarsenBased(Data, Clusters, RegulatorData, Parameters, NrCores,
                     random_seeds, convergence_cutoff)
```

Arguments

Data
Clusters
RegulatorData
Parameters
NrCores
random_seeds
convergence_cutoff

Value

result

AMARETTO_LearnRegulatoryProgramsLarsen
AMARETTO_LearnRegulatoryProgramsLarsen

Description

AMARETTO_LearnRegulatoryProgramsLarsen

Usage

```
AMARETTO_LearnRegulatoryProgramsLarsen(Data, Clusters, RegulatorData,
  RegulatorSign, Lambda, AutoRegulation, alpha, pmax, random_seeds)
```

Value

result

AMARETTO_Preprocess *AMARETTO_Preprocess*

Description

Wrapper code that analyzes process TCGA GISTIC (CNV) and gene expression (rna-seq or microarray) data via one call

Usage

```
AMARETTO_Preprocess(DataSetDirectories = DataSetDirectories,
  BatchData = BatchData)
```

Arguments

```
DataSetDirectories    DataSetDirectories
BatchData            BatchData
```

Value

result

Examples

```
## Not run:
TargetDirectory <- "Downloads" # path to data download directory
CancerSite <- 'CHOL'
DataSetDirectories <- AMARETTO_Download(CancerSite,TargetDirectory)
ProcessedData <- AMARETTO_Preprocess(DataSetDirectories,BatchData)

## End(Not run)
```

AMARETTO_ReassignGenesToClusters
AMARETTO_ReassignGenesToClusters

Description

AMARETTO_ReassignGenesToClusters

Usage

```
AMARETTO_ReassignGenesToClusters(Data, RegulatorData, Beta, Clusters,  
  AutoRegulation)
```

Value

result

AMARETTO_Run *AMARETTO_Run Function to run AMARETTO, a statistical algorithm to identify cancer drivers by integrating a variety of omics data from cancer and normal tissue.*

Description

AMARETTO_Run Function to run AMARETTO, a statistical algorithm to identify cancer drivers by integrating a variety of omics data from cancer and normal tissue.

Usage

```
AMARETTO_Run(AMARETTOinit)
```

Arguments

AMARETTOinit List output from AMARETTO_Initialize().

Value

result

Examples

```
data('ProcessedDataLIHC')  
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,  
  NrModules = 2, VarPercentage = 50)  
AMARETTOresults <- AMARETTO_Run(AMARETTOinit)
```

AMARETTO_VisualizeModule

AMARETTO_VisualizeModule

Description

Function to visualize the gene modules

Usage

```
AMARETTO_VisualizeModule(AMARETTOinit, AMARETTOresults, ProcessedData,  
  ModuleNr, show_row_names = FALSE, SAMPLE_annotation = NULL,  
  ID = NULL, order_samples = NULL)
```

Arguments

AMARETTOinit List output from AMARETTO_Initialize().

AMARETTOresults List output from AMARETTO_Run().

ProcessedData List of processed input data

ModuleNr Module number to visualize

show_row_names If TRUE, row names will be shown on the plot.

SAMPLE_annotation Matrix or Dataframe with sample annotation

ID Column used as sample name

order_samples Order samples in heatmap by mean or by clustering

Value

result

Examples

```
data('ProcessedDataLIHC')  
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,  
  NrModules = 2, VarPercentage = 50)  
  
AMARETTOresults <- AMARETTO_Run(AMARETTOinit)  
  
AMARETTO_VisualizeModule(AMARETTOinit = AMARETTOinit, AMARETTOresults = AMARETTOresults,  
  ProcessedData = ProcessedDataLIHC, ModuleNr = 1)
```

| | |
|--------|---------------|
| aprior | <i>aprior</i> |
|--------|---------------|

Description

Following four find empirical hyper-prior values

Usage

```
aprior(gamma.hat)
```

Value

result

| | |
|-----------|------------------|
| BatchData | <i>BatchData</i> |
|-----------|------------------|

Description

A dataset for conducting batch corection in TCGA samples

Usage

```
BatchData
```

Format

A data frame with 23263 observations and 3 variables:

Source

AMARETTO

| | |
|---------|----------------|
| Beta.NA | <i>Beta.NA</i> |
|---------|----------------|

Description

Beta.NA

Usage

```
Beta.NA(y, X)
```

Value

result

| | |
|--------|---------------|
| bprior | <i>bprior</i> |
|--------|---------------|

Description

bprior

Usage

```
bprior(gamma.hat)
```

Value

result

| | |
|--------------|---------------------|
| build.design | <i>build.design</i> |
|--------------|---------------------|

Description

Next two functions make the design matrix (X) from the sample info file

Usage

```
build.design(vec, des = NULL, start = 2)
```

Value

result

| | |
|---------------|----------------------|
| cacheResource | <i>cacheResource</i> |
|---------------|----------------------|

Description

cacheResource

Usage

```
cacheResource(TargetDirectory = TargetDirectory, resource = resource)
```

Value

result

| | |
|----------------|-----------------------|
| ComBat_NoFiles | <i>ComBat_NoFiles</i> |
|----------------|-----------------------|

Description

ComBat_NoFiles

Usage

```
ComBat_NoFiles(dat, saminfo, type = "txt", write = FALSE,
  covariates = "all", par.prior = TRUE, filter = FALSE, skip = 0,
  prior.plots = FALSE)
```

Value

result

| | |
|------------------|-------------------------|
| computeGisticURL | <i>computeGisticURL</i> |
|------------------|-------------------------|

Description

computeGisticURL

Usage

```
computeGisticURL(url = NULL, acronym = "CHOL")
```

Value

result

| | |
|---------------------|----------------------------|
| CreateRegulatorData | <i>CreateRegulatorData</i> |
|---------------------|----------------------------|

Description

Determine potential regulator genes.

Usage

```
CreateRegulatorData(MA_matrix = MA_matrix, CNV_matrix = NULL,
  MET_matrix = NULL, Driver_list = NULL, PvalueThreshold = 0.001,
  RsquareThreshold = 0.1, method = "union")
```

Value

result

| | |
|------------|-------------------|
| design.mat | <i>design.mat</i> |
|------------|-------------------|

Description

design.mat

Usage

```
design.mat(saminfo)
```

Value

result

| | |
|--------------|---------------------|
| Driver_Genes | <i>Driver_Genes</i> |
|--------------|---------------------|

Description

A list of cancer driver genes described in literature.

Usage

```
Driver_Genes
```

Format

List

Source

AMARETTO

| | |
|---------------|----------------------|
| filter.absent | <i>filter.absent</i> |
|---------------|----------------------|

Description

filters data based on presence/absence call

Usage

```
## S3 method for class 'absent'  
filter(x, pct)
```

Value

result

```
FindTranscriptionallyPredictive_CNV
      FindTranscriptionallyPredictive_CNV
```

Description

Function to identify which genes CNV significantly predict expression of that gene.

Usage

```
FindTranscriptionallyPredictive_CNV(MA_matrix, CNV_matrix,
      PvalueThreshold = 0.001, RsquareThreshold = 0.1)
```

Value

result

```
geneFiltering      geneFiltering
```

Description

Function to filter gene expression matrix

Usage

```
geneFiltering(Type, MAdat, Percentage)
```

Value

result

```
GeneSetDescription      GeneSetDescription
```

Description

GeneSetDescription

Usage

```
GeneSetDescription(filename, MSIGDB)
```

Arguments

| | |
|----------|--|
| filename | The name of the gmt file. |
| MSIGDB | If True, the gene set description column will be provided from MSIGDB. |

Value

result

| | |
|------------------|-------------------------|
| get_firehoseData | <i>get_firehoseData</i> |
|------------------|-------------------------|

Description

Downloading TCGA dataset via firehose

Usage

```
get_firehoseData(TargetDirectory = "./",
  TCGA_acronym_uppercase = "LUAD", dataType = "stddata",
  dataFileTag = "mRNAseq_Preprocess.Level_3", FFPE = FALSE,
  fileType = "tar.gz",
  gdacURL = "http://gdac.broadinstitute.org/runs/", untarUngzip = TRUE,
  printDisease_abbr = FALSE)
```

Value

result

| | |
|----------------|-----------------------|
| GmtFromModules | <i>GmtFromModules</i> |
|----------------|-----------------------|

Description

GmtFromModules

Usage

```
GmtFromModules(AMARETTOinit, AMARETTOresults, driverGSEA)
```

Arguments

| | |
|-----------------|---|
| AMARETTOinit | List output from AMARETTO_Initialize(). |
| AMARETTOresults | List output from AMARETTO_Run(). |
| driverGSEA | if TRUE , module driver genes will also be added to module target genes for GSEA. |

Value

result

 HyperGTestGeneEnrichment

Hyper Geometric Geneset Enrichment Test

Description

Calculates the p-values for unranked gene set enrichment based on two gmt files as input and the hyper geometric test.

Usage

```
HyperGTestGeneEnrichment(gmtfile, testgmtfile, NrCores,
  ref.numb.genes = 45956)
```

Arguments

| | |
|----------------|--|
| gmtfile | The gmt file with reference gene set. |
| testgmtfile | The gmt file with gene sets to test. In our case, the gmt file of the modules. |
| NrCores | Number of cores used for parallelization. |
| ref.numb.genes | The total number of genes teste, standard equal to 45 956 (MSIGDB standard). |

Value

result

 int.eprior

int.eprior

Description

Monte Carlo integration function to find the nonparametric adjustments

Usage

```
int.eprior(sdat, g.hat, d.hat)
```

Value

result

| | |
|---------------------|---------------|
| <code>it.sol</code> | <i>it.sol</i> |
|---------------------|---------------|

Description

Pass in entire data set, the design matrix for the entire data, the batch means, the batch variances, priors (m, t2, a, b), columns of the data matrix for the batch. Uses the EM to find the parametric batch adjustments

Usage

```
it.sol(sdat, g.hat, d.hat, g.bar, t2, a, b, conv = 1e-04)
```

Value

result

| | |
|---|----------|
| L | <i>L</i> |
|---|----------|

Description

likelihood function

Usage

```
L(x, g.hat, d.hat)
```

Value

result

| | |
|-----------------|------------------------|
| Lambda_Sequence | <i>Lambda_Sequence</i> |
|-----------------|------------------------|

Description

Lambda_Sequence

Usage

```
Lambda_Sequence(sx, sy)
```

Value

result

| | |
|------------|-------------------|
| list.batch | <i>list.batch</i> |
|------------|-------------------|

Description

Makes a list with elements pointing to which array belongs to which batch

Usage

```
list.batch(saminfo)
```

Value

result

| | |
|---------------|----------------------|
| MsigdbMapping | <i>MsigdbMapping</i> |
|---------------|----------------------|

Description

A dataset containing all MSIGDB pathways and their descriptions. .

Usage

```
MsigdbMapping
```

Format

List

Source

AMARETTO

| | |
|------------------|-------------------------------|
| plot_run_history | <i>Title plot_run_history</i> |
|------------------|-------------------------------|

Description

Title plot_run_history

Usage

```
plot_run_history(AMARETTOinit, AMARETTOresults)
```

Arguments

AMARETTOinit AMARETTO initialize output
 AMARETTOresults AMARETTO results output

Value

plot

Examples

```
data('ProcessedDataLIHC')
AMARETTOinit <- AMARETTO_Initialize(ProcessedData = ProcessedDataLIHC,
                                   NrModules = 2, VarPercentage = 50)

AMARETTOresults <- AMARETTO_Run(AMARETTOinit)

plot_run_history(AMARETTOinit,AMARETTOresults)
```

postmean

postmean

Description

postmean

Usage

```
postmean(g.hat, g.bar, n, d.star, t2)
```

Value

result

postvar

postvar

Description

postvar

Usage

```
postvar(sum2, n, a, b)
```

Value

result

| | |
|-------------------|--------------------------|
| Preprocess_MAdata | <i>Preprocess_MAdata</i> |
|-------------------|--------------------------|

Description

Preprocess_MAdata

Usage

```
Preprocess_MAdata(CancerSite = CancerSite, MAEO_ge = MAEO_ge,  
BatchData = BatchData)
```

Value

result

| | |
|--------|---------------|
| printf | <i>printf</i> |
|--------|---------------|

Description

Wrapper function for C-style formatted output.

Usage

```
printf(...)
```

Value

result

| | |
|-------------------|--------------------------|
| ProcessedDataLIHC | <i>ProcessedDataLIHC</i> |
|-------------------|--------------------------|

Description

A list of dataframes of processed toy example dataset from TCGA-LIHC.

Usage

```
ProcessedDataLIHC
```

Format

List

Source

AMARETTO

readGMT

readGMT

Description

readGMT

Usage

readGMT(filename)

Arguments

filename

Value

result

read_gct

read_gct

Description

Function to turn a .gct data files into a matrix format

Usage

read_gct(file_address)

Arguments

file_address Address of the input gct file.

Value

result

Examples

```
data_matrix<-read_gct(file_address="")
```

| | |
|-----------------|------------------------|
| Save_CancerSite | <i>Save_CancerSite</i> |
|-----------------|------------------------|

Description

Save_CancerSite

Usage

```
Save_CancerSite(CancerSite, TargetDirectory, DataSetDirectories,  
ProcessedData)
```

Value

result

| | |
|------------------------------------|---|
| TCGA_BatchCorrection_MolecularData | <i>TCGA_BatchCorrection_MolecularData</i> |
|------------------------------------|---|

Description

TCGA_BatchCorrection_MolecularData

Usage

```
TCGA_BatchCorrection_MolecularData(GEN_Data = GEN_Data,  
BatchData = BatchData, MinInBatch = MinInBatch)
```

Value

result

| | |
|------------------------------|-------------------------------------|
| TCGA_GENERIC_BatchCorrection | <i>TCGA_GENERIC_BatchCorrection</i> |
|------------------------------|-------------------------------------|

Description

TCGA_GENERIC_BatchCorrection

Usage

```
TCGA_GENERIC_BatchCorrection(GEN_Data = GEN_Data,  
BatchData = BatchData)
```

Value

result

TCGA_GENERIC_CheckBatchEffect
TCGA_GENERIC_CheckBatchEffect

Description

TCGA_GENERIC_CheckBatchEffect

Usage

TCGA_GENERIC_CheckBatchEffect(GEN_Data, BatchData)

Value

result

TCGA_GENERIC_CleanUpSampleNames
TCGA_GENERIC_CleanUpSampleNames

Description

TCGA_GENERIC_CleanUpSampleNames

Usage

TCGA_GENERIC_CleanUpSampleNames(GEN_Data = GEN_Data, IDlength = 12)

Value

result

TCGA_GENERIC_GetSampleGroups
TCGA_GENERIC_GetSampleGroups

Description

TCGA_GENERIC_GetSampleGroups

Usage

TCGA_GENERIC_GetSampleGroups(SampleNames)

Value

result

TCGA_GENERIC_MergeData

TCGA_GENERIC_MergeData

Description

TCGA_GENERIC_MergeData

Usage

TCGA_GENERIC_MergeData(NewIDListUnique, DataMatrix, MergeMethod)

Value

result

TCGA_Load_GISTICdata *TCGA_Load_GISTICdata*

Description

TCGA_Load_GISTICdata

Usage

TCGA_Load_GISTICdata(GisticDirectory)

Value

result

TCGA_Load_MolecularData

TCGA_Load_MolecularData

Description

TCGA_Load_MolecularData

Usage

TCGA_Load_MolecularData(MAEO_ge)

Value

result

| | |
|----------|-----------------|
| trim.dat | <i>trim.dat</i> |
|----------|-----------------|

Description

Trims the data of extra columns, note your array names cannot be named 'X' or start with 'X.'

Usage

```
trim.dat(dat)
```

Value

result

| | |
|-----------|------------------|
| write_gct | <i>write_gct</i> |
|-----------|------------------|

Description

write_gct

Usage

```
write_gct(data_in, file_address)
```

Value

result

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