

# Package ‘chroGPS’

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**Type** Package

**Title** chroGPS2: Generation, visualization and differential analysis of epigenome maps

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**Author** Oscar Reina, David Rossell

**Maintainer** Oscar Reina <oscar.reina@irbbarcelona.org>

**Description** We provide intuitive maps to visualize, analyze and compare the association between genetic elements based on their epigenetic profiles. The approach is based on Multi-Dimensional Scaling, and includes a parallelized implementation for handling high dimensional datasets. We provide several sensible distance metrics, and adjustment procedures to remove systematic biases typically observed when merging data obtained under different technologies or genetic backgrounds. We also provide functions and methods to perform differential analysis of epigenome maps at factor and gene level.

**License** GPL (>=2.14)

**Depends** R (>= 3.2.0), GenomicRanges, IRanges, methods, Biobase, MASS, graphics, stats, changepoint

**#Imports** graphics, cluster, DPpackage, ICSNP

**Imports** cluster, DPpackage, ICSNP, ellipse, vegan

**Enhances** parallel, XML, rgl, gplots, pheatmap, ChIPpeakAnno, org.Dm.eg.db, caTools, plotly

**Collate** adjustPeaks.R distGPS.R domainDist.R mds-class.R mds.R procrustesAdj.R clusGPS.R geneSetGPS.R getmodEncode.R gff2RDList.R gps2xgmml.R diffFactors.R diffGenes.R mergeReplicates.R profileClusters.R rankFactorsbyDomain.R rankFactorsbyProfile.R

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## R topics documented:

addVar . . . . .	2
adjustPeaks . . . . .	3
bg3 . . . . .	4
boostMDS . . . . .	5
clusGPS . . . . .	6
clusGPS-class . . . . .	9
combineGenesMatrix . . . . .	10
diffFactors . . . . .	11
diffGenes . . . . .	12
distGPS . . . . .	15
distGPS-class . . . . .	17
domainDist . . . . .	18
geneSetGPS . . . . .	19
getURL . . . . .	20
gff2RDList . . . . .	20
gps2xgmml . . . . .	21
mds . . . . .	22
mds-class . . . . .	24
mergeClusters . . . . .	25
mergeReplicates . . . . .	26
procrustesAdj . . . . .	28
profileClusters . . . . .	29
rankFactorsbyDomain . . . . .	30
rankFactorsbyProfile . . . . .	32
s2 . . . . .	33
splitDistGPS-class . . . . .	34
<b>Index</b>	<b>35</b>

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addVar	<i>Plot vector of a quantitative variable over a MDS map.</i>
--------	---

---

### Description

Given a quantitative variable as a numeric vector with one element for each point on a MDS map, calculate and plot the weight vector corresponding to that variable.

### Usage

```
addVar(mds1, z, plot = TRUE, label = "z", pos = 3, ...)
```

### Arguments

mds1	An object of class mds with the MDS object.
z	Numeric vector with the quantitative variable, one element for each point.
plot	Set to TRUE to calculate and draw the resulting vector on the MDS.
label	Something to be printed on the tip of the vector arrow, usually the name of the given variable.
pos	Graphical position where the label is drawn respect to the vector arrow tip.
...	Additional parameters given to the generic function plot.

**Value**

A named list with the vector components.

**Examples**

```
# Not run
# See chroGPS-manual.pdf for examples.
```

---

adjustPeaks	<i>Adjust peak width so that samples obtained under different conditions become comparable.</i>
-------------	---

---

**Description**

Peaks obtained under different conditions (e.g. chip-chip, chip-seq, mnase-seq) are typically not comparable in terms of their width. `adjustPeaks` modifies the mean and SD of the peak width distribution for each condition, so that they become equivalent to the condition with widest peaks. See details.

**Usage**

```
adjustPeaks(x, adjust, sampleid, logscale = TRUE)
```

**Arguments**

<code>x</code>	<code>GRangesList</code> indicating the binding sites for each sample/experiment.
<code>adjust</code>	Vector indicating the adjustment factor, i.e. the condition under which each sample has been obtained.
<code>sampleid</code>	Vector containing the sample identifier. <code>sampleid</code> should take the same value for samples obtained under different conditions, as this is used to detect the samples to be used for Procrustes adjustment.
<code>logscale</code>	If set to <code>TRUE</code> the mean and SD are matched for log width, otherwise the original widths are used. Working in log scale can help reduce the effect of outliers (e.g. an usually long binding site).

**Details**

In a sense, the peak calling resolution is decreased so that they become comparable to the less precise technology (notice that there is no reliable way to increase the precision given by a low-resolution technology).

**Value**

`GRangesList` object with adjusted widths.

**Methods**

**signature(x='GRangesList')** Each element in `x` contains the binding sites for a different sample. The start, end and chromosome of each binding sites should be accessed via `start`, `end` and `space`.

**See Also**

[procrustesAdj](#) for an alternative, more general, adjustment based on Procrustes. [distGPS](#) for computing distances, [mds](#) to create MDS-oriented objects.

**Examples**

```
#See examples in help(procrustesAdj)
```

---

bg3	<i>Sample binding site and related data from BG3 cell line in Drosophila melanogaster.</i>
-----	--

---

**Description**

chroGPS example dataset including CHIP-CHIP (modEncode) and CHIP-Seq (NCBI GEO GSE19325) data for *Drosophila melanogaster* BG3 cell line. The object `toydist`s stores precomputed `distGPS` objects (called `d`, `d2`, `d3`) for the epigenetic factors used in the dynamic vignette that comes with the package.

**Usage**

```
data(bg3)
```

**Source**

```
http://www.modencode.org http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE19325
```

**References**

```
http://www.modencode.org http://www.ncbi.nlm.nih.gov/geo/
```

**See Also**

`s2` for the analog S2 cell line object. `toydist`s for precomputed distance matrixes.

**Examples**

```
data(bg3)
class(bg3)
bg3
bg3names$Factor
# See vignette examples for several uses of these datasets.
```

---

 boostMDS

*Improve goodness-of-fit of a given MDS solution in terms of R-square.*


---

### Description

Given a distance matrix and a valid MDS representation for it, improve the R-square correlation between observed and approximated distances until converged is reached for a given threshold.

### Usage

```
boostMDS(D, Y, rate = 0.01, maxit = 50, tol = 0.001, samplesize,
         verbose = TRUE, scale = FALSE, seed = 149, plt = FALSE, mc.cores = 1)
```

### Arguments

D	Distance matrix.
Y	Matrix with points from a valid MDS solution for the distances in D.
rate	Grid step rate, start with 0.1 which usually is a good compromise, try also 0.01, 1, 10.
maxit	Maximum number of iterations.
tol	Tolerance for R-square convergence.
samplesize	When there are over 100 points to represent, the gradient descent step size is determined using a fraction <code>samplesize</code> of the original data points. By default 0.01 with a minimum of 100 points, which typically gives very stable results. Setting large <code>samplesize</code> can significantly increase the computational cost.
verbose	Give details of the gains in R-square and step size.
scale	Whether to scale the MDS coordinates in the output MDS.
seed	A random seed to be used in the resampling process if <code>samplesize &lt; 1</code> .
plt	Whether to plot the intermediate solutions or not.
mc.cores	Number of cores to use in parallelized grid step size search.

### Value

The function returns a matrix with the coordinates of a valid MDS solution for distance matrix D where the R-square correlation has been improved. However, have in mind that an MDS solution with better R-square does not necessarily mean the solution is easier to interpret. As with any MDS approach, a balance must be found between pure 'technical' goodness-of-fit and usefulness of the delivered solution in terms of answering the original hypothesis.

### References

boostMDS is based on hitMDS (High-Throughput Multidimensional Scaling, see <http://dig.ipk-gatersleben.de/hitmids/hitmids.html> for details)

## Examples

```
# Not run, see also chroGPS-manual.pdf file for examples
#data(geneSample)
#d = distGPS(geneSample,uniqueRows=TRUE)
#m = mds(d,type='isoMDS')
#m
#plot(m)
#m = boostMDS(d@d,m@points)
#plot(m)
```

---

clusGPS

*Computation of cluster density estimates for cluster contour representation and correct-classification rates (cluster robustness). A pre-computed clustering of elements used in the map has to be given as an input, which is useful to explore results using different clustering algorithms and methodologies (top-down, bottom-up, etc).*

---

## Description

After performing a pre-merging step so that all clusters have a minimum size, semiparametric bayesian density is estimated using a Dirichlet process mixture of normals. This is used both to compute bayesian mis-classification posterior probabilities (correct classification rates) and to estimate probability contours which can be visualized on the MDS map.

The functions `contour2dDP` and `plotContour` functions can be used to compute bayesian density estimates for a given set of elements (points) from a pre-generated 2D MDS object. These functions are used internally by `clusGPS` to draw cluster contours but are also useful to visualize other type of contours over the map (ie genes from a given Gene Ontology term, having a specific epigenetic mark of interest, etc).

The S4 accessors `clusNames`, `tabClusters` and `clusterID` retrieve information stored within a `clusGPS` object.

## Usage

```
clusGPS(d, m, h, sel=NULL, id=NULL, grid, ngrid=1000, densgrid=FALSE, preMerge=TRUE, type = "hclust",
"average", samplesize = 1, p.adjust = TRUE, k, mc.cores = 1,
set.seed = 149, verbose=TRUE, minpoints=70,...)
contour2dDP(x, ngrid, grid = NULL, probContour = 0.5, xlim, ylim,
labels = "", labcex = 0.01, col = colors()[393], lwd = 4,
lty = 1, contour.type = "single", contour.fill = FALSE,
minpoints=100, ...)
clusNames(clus)
tabClusters(clus,name)
clusterID(clus,name)
```

## Arguments

**d** Object of class `distGPS` with the pairwise observed dissimilarities between elements.

<code>m</code>	(Optional). Object of class <code>mds</code> with a MDS object generated from the distances in <code>d</code> . Only MDS type "boostMDS" is available. The <code>mds</code> function performs an optimization of the approximated distances in <code>m</code> in order to improve R-square correlation between them and the observed dissimilarities in <code>d</code> , maximizing goodness of fit.
<code>h</code>	(Optional). Object of class <code>hclust</code> with a pre-calculated clustering for the elements in <code>d</code> .
<code>sel</code>	(Optional). Logical vector indicating which elements from <code>d</code> will be used for performing hierarchical clustering with average linkage. This is useful if we want to focus on a given set of points only (i.e. those from a big cluster which we want to study in more detail).
<code>id</code>	(Optional). Label of the cluster which we want to further subdivide, ignoring points from all other clusters. Deprecated, use parameter <code>sel</code> specified above.
<code>grid</code>	Matrix of dimension <code>ngrid*nvar</code> giving the diagonal points of the grid where the density estimate is evaluated. The default value is <code>NULL</code> : grid dimensions are chosen according to the range of the data, and granularity is automatically determined according to data density, in order to provide a more accurate estimation in high density areas, where more resolution is needed.
<code>ngrid</code>	Number of grid points where the density estimate is evaluated. This argument is ignored if a grid is specified. The default value is 1000. Higher values are recommended if data presents very high density areas.
<code>densgrid</code>	Set to <code>true</code> to generate grid points from the quantile distribution of the data using the grid size defined by <code>ngrid</code> . This is useful if the data presents areas of very different density, ranging from very sparse to extremely dense areas, optimizing grid granularity where is necessary, therefore improving resolution of density estimation and reducing computation time.
<code>preMerge</code>	If <code>TRUE</code> will perform a first pre-merging step so that any cluster smaller than <code>minpoints</code> gets merged with its closest cluster based on their centroid distances. This is performed until no clusters <code>&lt; minpoints</code> exist.
<code>type</code>	Type of clustering to be performed. Currently only "hclust" (Agglomerative Nesting) is supported, but any other clustering type can be used by providing a pre-calculated object <code>h</code> . This variable is to become deprecated, since <code>clusGPS</code> will only work with a precomputed clustering.
<code>method</code>	Clustering method. See <code>hclust</code> for details. This variable is to become deprecated, since <code>clusGPS</code> will only work with a precomputed clustering.
<code>samplesize</code>	Proportion of elements to sample for computing clustering and density estimation. This is useful to generate density contours from a subset of the data, speeding up computation.
<code>p.adjust</code>	Set to <code>TRUE</code> to adjust the bayesian posterior probabilities of mis-classification.
<code>k</code>	Integer vector indicating the number of clusters on which density estimation will be computed for mis-classification or contour calculation.
<code>mc.cores</code>	Number of cores to be used for parallel computation with the <code>parallel</code> package.
<code>set.seed</code>	If <code>samplesize&lt;1</code> , random seed to be used to perform random sampling of the data.
<code>verbose</code>	Set to <code>TRUE</code> to output clustering process information.
<code>minpoints</code>	If <code>preMerge</code> is <code>FALSE</code> , then the algorithm will ignore clusters with fewer than <code>minpoints</code> elements. This is useful if the clustering method used tends to generate many very small clusters of limited use and difficult interpretation and for

	which density estimates may not be correctly computed. The default method is to preMerge clusters since this ensures density estimation is available for all clusters and helps interpreting the map, since no elements are ignored.
x	Numeric matrix indicating coordinates of the points for which a probability contour is calculated in contour2dDP.
probContour	Numeric matrix indicating coordinates of the points for which a probability contour is calculated in contour2dDP.
contour.type	For contour2dDP, type of contour, either 'single' (surrounding the points within the given probContour probability) or 'multiple' to generate terrain-like density contour lines.
contour.fill	Deprecated.
xlim,ylim,labels,labcex,col,lwd,lty	Graphical parameters given to contour2dDP.
clus	A valid clusGPS object from which we want to extract information.
name	Character indicating a valid name within a clusGPS object, from which we want to extract information.
...	Additional parameters.

### Value

The function clusGPS returns an object of class clusGPS. See help for clusGPS-methods for details. contour2dDP returns a DPdensity object with density contour information which can be plotted as 2D contours with our plotContour function, as well as with the plot function from the DPpackage package.

### Methods

**signature(d='distGPS',m='mds')** Hierarchical clustering is performed for the elements whose pairwise distances are given in d. For each cluster partition given in k, cluster identity for each element is returned, and semiparametric bayesian density estimation is computed using the point density information from m.

**plot** signature(m = "clusGPS"): S4 plot method for clusGPS objects.

**clusNames** signature(m = "clusGPS"): Retrieves names of the clustering configurations stored in clusGPS objects, one for each distance threshold indicated in k, that get automatically named accordingly.

**tabClusters** signature(m = "clusGPS"): Returns a table with the number of elements in each of the clusters found for an existing clustering configuration with name name within the clusGPS object.

**clusterID** signature(m = "clusGPS"): Returns a vector of cluster assignments for all the elements in an existing clustering configuration name within the clusGPS object.

### Author(s)

Oscar Reina

### Examples

```
# Not run
# data(s2)
# # Computing distances
```



```

# d <- distGPS(s2.tab,metric='tanimoto',uniqueRows=TRUE)
# # Creating MDS object
# mds1 <- mds(d,type='isoMDS')
# mds1
# plot(mds1)
# Precomputing clustering
# h <- hclust(as.dist(d@d),method='average')
# # Calculating densities (contours and probabilities), takes a while
# clus <- clusGPS(d,mds1,preMerge=TRUE,k=max(cutree(h,h=0.5)))
# # clus contains information for contours and probabilities
# plot(clus,type='contours',k=125,lwd=3,probContour=.75)
# plot(clus,type='stats',k=125,ylim=c(0,1))
# plot(clus,type='avgstat')
# plot(clus,type='density',k=3,ask=TRUE,xlim=range(mds1@points),ylim=range(mds1@points))

```

clusGPS-class

*Class "clusGPS"***Description**

Agglomerative Nesting for a distGPS object. Contains probability contours and bayesian posterior probability of mis-classification for the clusters evaluated.

**Details**

Parameters for the S4 plot method for mds objects.

Object of class "mds" with a 2D or 3D Multidimensional Scaling to be plotted.

**drawlabels:** TRUE to use rownames of the MDS points as text labels.

**labels:** Alternative character vector giving the text labels for the MDS points.

**plantar:** If a 3D MDS is used, set plantar to TRUE to plot projected views of the MDS using XY, YZ and XZ axis decomposition.

**point.cex:** Size of the points / spheres for the MDS plot.

**text.cex:** Size of text labels for the MDS points.

**text.pos:** Alignment position of the text labels respective to its points (1,2,3,4).

**point.col:** Color for the MDS points / spheres.

**text.col:** Color for the MDS text labels.

**point.pch:** PCH type for the MDS points.

**type.3d:** Use 'p' for points, 's' for spheres.

**radius:** Radius for the spheres on a 3D MDS plot. Automatically generated from point.cex and the number of points in the MDS.

**app:** Appearance of the 3D spheres on a 3D MDS plot, can be 'fill', 'lines', 'grid'.

**alpha:** Number between 0 and 1 with the level of transparency to be used on spheres on a 3D MDS.

**scalecol:** Set to TRUE to use a color scale for points, for instance to color points (genes) according to their expression level on a chroGPS-genes MDS plot.

**scale:** Scale to use to generate scale colors (for instance normalized gene expression for each element (gene) on chroGPS-genes MDS).

**palette:** Color palette to be used for scale colors.

**Objects from the Class**

Objects can be created by calls of the form `new("clusGPS", ...)`.

**Slots**

**h:** Object of class "hclust" with Agglomerative Nesting or user-provided cluster object.

**clus:** Object of class "list" with probability contour and bayesian posterior probability of mis-classification information for the clusters evaluated.

**adjusted:** Object of class "logical" indicating if bayesian posterior probabilities of mis-classification are adjusted for multiple testing.

**Author(s)**

Oscar Reina

**Examples**

```
showClass("clusGPS")
```

---

combineGenesMatrix	<i>Combine two datasets with epigenetic factor profiles at gene level in order to produce a differential chroGPS-genes map.</i>
--------------------	---

---

**Description**

Combine two datasets with epigenetic factor profiles at gene level in order to produce a differential chroGPS-genes map.

**Usage**

```
combineGenesMatrix(x, y, label.x, label.y, minFactors = 10, minGenes = 1000)
```

**Arguments**

x	Data frame or matrix with genes (rows) and epigenetic factors (columns), and values of 0/1 to indicate binding of a given epigenetic factor over genes.
y	Same as x, containing the second dataset to compare. Column names have to match between both datasets, with at least a minimum number of common ones.
label.x	Name for the x dataset.
label.y	Name for the y dataset.
minFactors	Minimum number of common factors between both datasets.
minGenes	Minimum number of genes in each dataset with epigenetic information (that is, minimum number of rows where all values are not zeros).

**Value**

A matrix with the combined epigenetic profile dataset to perform differential analysis.

**Author(s)**

Oscar Reina.

**See Also**

See also `distGPS`, `mds`, `diffGenes`.

**Examples**

```
## Not run
## See example in diffGenes function.
```

---

diffFactors	<i>Performs differential analysis of chroGPS-factors maps based on Procrustes analysis.</i>
-------------	---

---

**Description**

The function uses Procrustes analysis to compare two chroGPS-factors MDS solutions, providing a visual map highlighting differences and a ranked list of Procrustes squared errors between analog replicated factors in both maps.

**Usage**

```
diffFactors(m1,m2,name1='mds1',name2='mds2',minPoints=10,poslegend='topleft',plot=TRUE,pointcol.
```

**Arguments**

<code>m1</code>	A mds object, containing a chroGPS-factors MDS map.
<code>m2</code>	A mds object, containing a chroGPS-factors MDS map, must be conformable with <code>m1</code> (same number of elements and in the same order in the MDS points).
<code>name1</code>	Name for <code>m1</code> , by default 'mds1'
<code>name2</code>	Name for <code>m2</code> , by default 'mds2'
<code>minPoints</code>	Minimum number of points in each of the MDS objects.
<code>poslegend</code>	Position for the map legend, passed on to <code>legend</code> function.
<code>plot</code>	Wether to plot the differential map or not (in case we only want to retrieve Procrustes errors).
<code>pointcol.m1</code>	Color vector for points in MDS1.
<code>pointcol.m2</code>	Color vector for points in MDS2.
<code>textcol.m1</code>	Color vector for labels in MDS1. Labels are taken from MDS points rownames.
<code>textcol.m2</code>	Color vector for labels in MDS2. Labels are taken from MDS points rownames.
<code>pch.m1</code>	Pch for MDS1 points.
<code>pch.m2</code>	Pch for MDS2 points.
<code>cex.m1</code>	Cex for MDS1 points.
<code>cex.m2</code>	Cex for MDS2 points.
<code>segcol</code>	Color for segments joining replicates from both MDS.
<code>...</code>	Further arguments passed on to the <code>plot</code> function.

**Value**

The function returns a data frame with Procrustes errors between paired replicates in both chroGPS-factors MDS maps.

**Author(s)**

Oscar Reina.

**See Also**

See functions `distGPS`, `mds` for generating chroGPS-factors maps.

**Examples**

```
## Not run

data(s2)
data(repliSeq)
library(gplots)

# Modify colors and add some transparency
fnames <- s2names$Factor
s2names$Color[s2names$Color=='grey'] <- 'orange'
fcolors <- paste(col2hex(s2names$Color), 'BB', sep='')
bcolors <- paste(col2hex(s2names$Color), 'FF', sep='')

# Select time points to compare
m1 <- m.origs[['Early.Mid']]
m2 <- m.origs[['Late']]

## Perform differential Procrustes analysis
pp <- diffFactors(m1,m2)

## Plot both maps before and after adjustment
m3 <- pp$mds3
plot(0,xlim=c(-1,1),ylim=c(-1,1),xlab='',ylab='',xaxt='n',yaxt='n',col='NA')
segments(m1@points[,1],m1@points[,2],m3@points[,1],m3@points[,2],col='red')
par(new=TRUE)
plot(m1,drawlabels=TRUE,labels=s2names$Factor,point.pch=19,point.cex=4,text.cex=0.75,point.col=s2names$Color)
par(new=TRUE)
plot(m3,drawlabels=TRUE,labels=s2names$Factor,point.pch=19,point.cex=4,text.cex=0.75,point.col=s2names$Color)

## Plot Procrustes errors
pp <- pp$procrustes
par(las=1,mar=c(4,12,4,4)); barplot(sort(residuals(pp),decr=TRUE),horiz=TRUE,xlim=c(0,max(residuals(pp))+.1))
hist(residuals(pp),breaks=50)
```

---

diffGenes

*Performs differential analysis of chroGPS-genes maps based on Bayesian Correct Classification Rates.*

---

**Description**

The function uses Bayesian posterior probability of correct classification (CCR) of chroGPS-genes maps as computed by the `clusGPS` function to identify statistically significant epigenetic changes between two different backgrounds (technical and/or biological). The functions `find.fdr` and `find.threshold` are used internally to compute approximated FDRs from a vector of input posterior probabilities of correct classification (CCRs).

**Usage**

```
diffGenes(xy,m,clus,label.x,label.y,clusName=NULL,fdr=TRUE,mc.cores=1)
```

**Arguments**

<code>xy</code>	A matrix or data.frame object, containing a combined matrix with whole-genome epigenetic information for backgrounds X and Y, as obtained by the <code>combineGenesMatrix</code> function.
<code>m</code>	A mds object, containing a chroGPS-genes MDS map, obtained for the combined matrix <code>xy</code> .
<code>clus</code>	A <code>clusGPS</code> object, containing a chroGPS-genes cluster solution, obtained for the provided background via the <code>clusGPS</code> and <code>mergeClusters</code> functions.
<code>label.x</code>	Name for background X, by default 'mds1'
<code>label.y</code>	Name for background Y, by default 'mds2'
<code>clusName</code>	If <code>clus</code> contains more than one clustering solution (i.e. cutree dendrogram height cut), the name of the selected one. If no name is provided, the first element is used.
<code>fdr</code>	Set to TRUE (default) to compute False Discovery Rate of cluster correct classification from posterior probabilities.
<code>mc.cores</code>	Number of cores to use for parallel computations.

**Value**

The function returns a data frame with posterior probabilities and FDRs of cluster classifications for all genes in backgrounds X and Y, so that significant changes can be traced and reported easily.

**Author(s)**

Oscar Reina.

**See Also**

See functions `distGPS`, `mds`, `clusGPS`, `profileClusters` for generating chroGPS-genes maps. See function `plotTransitions` for examples on how to visualize differential gene maps results.

**Examples**

```
# Summarize factor replicates with method 'any' so that 1 replicate
# having the mark is enough
# Assuming s2.tab and bg3.tab contain the full datasets for dm3 genome
# and all factors used in Font-Burgada et al.
# s2.tab <- mergeReplicates(s2.tab,s2names$Factor,'any')
# bg3.tab <- mergeReplicates(bg3.tab,bg3names$Factor,'any')
```

```

# Join, use common factors. Then use common genes only from those that
# have at least one mark in both s2 and bg3
# x <- combineGenesMatrix(s2.tab,bg3.tab,'S2','BG3')

# Build map and cluster as always
# d <- distGPS(x,metric='tanimoto',uniqueRows=TRUE)

# Not run
# m <- mds(d,type='classic',splitMDS=TRUE,split=0.16,mc.cores=4)
# mm <- mds(d,m,type='boostMDS',samplesize=0.005,mc.cores=6)
# unique(rownames(m@points)==rownames(mm@points)) # sanity check
# This should be incorporated in the function code...
#m@points <- m@points[rownames(d@d),]
#mm@points <- mm@points[rownames(d@d),]

# Cluster
# h <- hclust(as.dist(d@d),method='average')
# clus <-
# clusGPS(d,mm,h,k=max(cutree(h,h=0.5)),ngrid=10000,mc.cores=8,recalcDist=FALSE,verbose=FALSE)
# clus.merged <- mergeClusters(clus,brake=0,mc.cores=8)
# clus
# clus.merged

# Use new function to profile clusters
# pc <- profileClusters2(x,clus.merged,normalize=TRUE)
# pheatmap(pc,trace='none',scale='none',col=bluered(100))

# Perform differential analysis
# x.diff <- res <-
# diffGPS.clus(x,mm,clus.merged,label.x='S2',label.y='BG3',clusName=clusNames(clus.merged)[1],fdr=TRUE,mc.cores=8)
# write.csv(x.diff,'diffgenes_fdrest.csv')

# Select genes changing clusters with FDR 0.05
# xx.diff <- x.diff[x.diff$ClusID.S2!=x.diff$ClusID.BG3 &
# x.diff$FDR.S2<0.25 & x.diff$FDR.BG3<0.25,]
# xx.diff$CC <- paste(xx.diff$ClusID.S2,xx.diff$ClusID.BG3,sep='.')
# head(sort(table(xx.diff$CC),decreasing=TRUE))
# write.csv(xx.diff,'kk_fdrest2.csv')

# Perform enrichment test using getEnrichedGO from chippeakanno package
# library(ChIPpeakAnno)
# library(org.Dm.eg.db)

# enriched.GO <- lapply(c('2.9','5.2'),function(cc) {
#   fbid <- as.character(xx.diff$geneid[xx.diff$CC==cc])
#   if (length(fbid)>=25)
#     ans <-
#   getEnrichedGO(annotatedPeak=fbid,orgAnn='org.Dm.eg.db',maxP=0.05,multiAdjMethod='BH')
#   else ans <- NULL
#   return(ans)
# })

# names(enriched.GO) <- c('2.9','5.2')
# enriched.GO <- enriched.GO[unlist(lapply(enriched.GO,length))>0]
# enriched.GO <- lapply(enriched.GO,function(x) lapply(x,function(y)

```

```
# unique(y[,-ncol(y)]))
# lapply(enriched.GO,head)

# Plot results with diffGPS.plot function
# res.sel <- res[res$ClusID.S2!=res$ClusID.BG3,]

# Plot
# diffGenes.plot(x,mm,clus.merged,res.sel,transitions='10.2',label.x='S2',label.y='BG3',fdr1=0.25,fdr2=0.25)
```

---

distGPS	<i>Compute matrix with pairwise distances between objects. Several GPS metrics are available.</i>
---------	---

---

## Description

The function computes pairwise distances between individuals (e.g. samples or genes) according to a user-specified metric. Several metrics are available. The precise definition of each metric depends on the class of the first argument (see details section).

## Usage

```
distGPS(x, metric='tanimoto', weights, uniqueRows=FALSE, genomelength=NULL, mc.cores=1)
```

## Arguments

x	Object for which we want to compute distances
metric	Desired distance metric. Valid options for chroGPS-factors map are 'tanimoto', 'avgdist', 'chisquare' and 'chi' (see details). For chroGPS-genes maps, metrics 'wtanimoto', 'euclidean' and 'manhattan' are also available.
weights	For signature(x='matrix'), an unnamed numeric vector with weights applied to every sample (column) in the original data. The typical example is when we have a sample (epigenetic factor) with several replicates available (biological or technical replicate, different antibody, etc.), and we want to treat them together (for instance giving a 1/nreplicates weight to each one). If not supplied, each replicate is considered as an individual sample (using 1 as weight for every sample).
uniqueRows	If set to TRUE and x is a matrix or data.frame, duplicated rows are removed prior to distance calculation. This can save substantial computing time and memory. Notice however that the dimension of the distance matrix is equal to the number of unique rows in x, instead of nrow. . (x).
genomelength	For 'chi' and 'chisquare' metrics, numeric value indicating the length of the genome. If not given the function uses the minimum length necessary to fit the total length of the result.
mc.cores	If mc.cores>1 and parallel package is loaded, computations are performed in parallel with mc.cores processors when possible.

## Details

For `GRangesList` objects, distances are defined as follows.

Let `a1` and `a2` be two `GRanges` objects. Define as `n1` the number of `a1` intervals overlapping with some interval in `a2`. Define `n2` analogously. The Tanimoto distance between `a1` and `a2` is defined as  $(n1+n2)/(nrow(z1)+nrow(z2))$ . The average distance between `a1` and `a2` is defined as  $.5*(n1/nrow(z1) + n2/nrow(z2))$ . The `wTanimoto` distance in `chroGPS-genes` weights each epigenetic factor (table columns) according to its frequency (table rows). The chi-square distance is defined as the usual chi-square distance on a binary matrix `B` which is automatically computed by `distGPS`. The binary matrix `B` is the matrix with `length(x)` rows and number of columns equal to the genome length, where `B[i, j]==1` indicates that element `i` has a binding site at base pair `j`. The chi distance is simply defined as the square root of the chi-square distance. Finally, euclidean and manhattan metrics have the same definition than in the base R function `dist`.

When choosing a metric one should consider the effect of outliers, i.e. samples with large distance to all other samples. Tanimoto and Average Distance take values between 0 and 1, and therefore outlying distances have a limited effect. Chi-square and Chi distances are not limited between 0 and 1, i.e. some distances may be much larger than others. The Chi metric is slightly more robust to outliers than the Chi-square metric.

For `matrix` or `data.frame` objects, `x` must be a matrix with 0's and 1's (or `FALSE` and `TRUE`). The usual definitions are used for Tanimoto (which is equivalent to Jaccard's index), Chi-square and Chi. Average overlap between rows `i` and `j` is simply the average between the proportion of elements in `i` also in `j` and the proportion of elements in `j` also in `i`.

## Value

Object of class `distGPS`, with matrix of pairwise dissimilarities (distances) between objects.

## Methods

`distGPS`:

Each element in `x` is assumed to indicate the binding sites for a different sample, e.g. epigenetic factor. Typically `space(x)` indicates the chromosome, `start(x)` the start position and `end(x)` the end position (in bp). Strand information is ignored.

**signature(x='GRangesList')** Rows in `x` contain individuals for which we want to compute distances. Columns in `x` contain the variables, and should only contain either 0's and 1's or `FALSE` and `TRUE`.

`splitDistGPS`:

This is a set of internal classes and functions to be used in the parallel computation of Multi-dimensional Scaling.

`uniqueCount`:

This function collapses a `chroGPS-genes` matrix or data frame so that elements with the same combination of variables are aggregated into a single entry. Elements become then identified by their unique pattern and a frequency count is also returned.

`as.matrix`:

Returns the raw distance matrix within the object.

## See Also

`mDS` to create MDS-oriented objects, `procrustesAdj` for Procrustes adjustment.



**Examples**

```
x <- rbind(c(rep(0,15),rep(1,5)),c(rep(0,15),rep(1,5)),c(rep(0,19),1),c(rep(1,5),rep(0,15)))
rownames(x) <- letters[1:4]
d <- distGPS(x,metric='tanimoto')
du <- distGPS(x,metric='tanimoto',uniqueRows=TRUE)
mds1 <- mds(d)
mds1
plot(mds1)
d <- distGPS(x,metric='chisquare')
mds1 <- mds(d)
mds1
plot(mds1)
```

---

distGPS-class	<i>Class "distGPS"</i>
---------------	------------------------

---

**Description**

Pairwise distances between elements. Function `distGPS` creates objects of this class. `splitDistGPS` in an private class used internally for parallel Multidimensional Scaling.

**Objects from the Class**

Objects can be created by calls of the form `new("distGPS",...)` to generate chroGPS-compliant distance matrices with user-defined metrics. The internal distance matrix can be extracted with the function `"as.matrix"`

**Slots**

**d:** Object of class `"matrix"` with pairwise dissimilarities (distances) between elements.

**metric:** Object of class `"character"` indicating the metric type used for calculating distances. See function `"distGPS"`.

**type:** Object of class `"character"`, deprecated.

**Author(s)**

Oscar Reina

**Examples**

```
showClass("distGPS")
data(s2)
data(toydists)
d
class(as.matrix(d))
as.matrix(d)[1:5,1:5]
```

---

domainDist

*Overview of intra and inter-domain distances.*


---

## Description

Given a distance of pairwise distances or dissimilarities between elements, return intra and inter-group sets of distances based on a given group definition. This is useful to get an insight on domain robustness for functional related genes or factors.

## Usage

```
domainDist(d, gps='factors', domain, type='intra', col='white', avg=FALSE,
plot=TRUE, ...)
```

## Arguments

d	Distance/Dissimilarities matrix, usually the slot d on a distGPS object, but any distance matrix can be given as input.
gps	'factors' for a chroGPS-factors distance matrix, 'genes' for a chroGPS-genes one.
domain	Character vector with group identity for each element d. It can be a functional domain classification (i.e. 'Activation', 'Repression', etc), given for each factor on a chroGPS-factors map or for each gene in a chroGPS-genes map. However, any classification of interest can be used (pathways, gene ontology, etc.)
type	Intradomain ('intra') or Interdomain ('inter') distance overview.
col	Character vector with colors to be passed to plot.
avg	TRUE to return also the average inter or intra domain distance.
plot	TRUE to generate inter/intra domain boxplots.
...	Additional parameters given to the generic function plot.

## Value

List of inter or intra domain distances.

## Examples

```
# Not run
# data(s2)
# d <- distGPS(s2,metric='avgdist',mc.cores=1)
# d.intra <- domainDist(as.matrix(d),domain=s2names$Color,type='intra',plot=TRUE)
# d.inter <- domainDist(as.matrix(d),domain=s2names$Color,type='inter',plot=TRUE)
```

---

geneSetGPS	<i>Highlight point (gene) position over a Multi-dimensional Scaling plot.</i>
------------	---

---

### Description

Given a list of genes of interest, the function highlights their position over a Multi-dimensional Scaling plot.

### Usage

```
geneSetGPS(x, m, genes, uniqueCount = TRUE, ...)
```

### Arguments

x	Matrix or data frame of observations x variables (typically genes x epigenetic factors), with gene identifiers as rownames.
m	Object of class mds with a valid Multidimensional Scaling representation for the elements in x.
genes	Character vector containing gene identifiers, matching those on rownames(x).
uniqueCount	Set to FALSE if the MDS has been generated directly from the data in x, otherwise set to TRUE to match gene identifiers with their unique pattern of observed variables.
...	Additional parameters given to the generic function plot.

### Value

Matrix with coordinates on the given input MDS object for the genes selected.

### Author(s)

Oscar Reina

### Examples

```
# Not run
# data(s2)
# d <- distGPS(s2.tab,metric='tanimoto',uniqueRows=TRUE)
# mds1 <- mds(d)
# set.seed(149)
# sampleGenes <- rownames(s2.tab)[sample(1:nrow(s2.tab),10,rep=FALSE)]
# pts <- geneSetGPS(s2.tab,mds1,genes=sampleGenes,uniqueCount=TRUE)
# plot(mds1)
# points(getPoints(pts),col='red',cex=3)
```

---

getUrl	<i>Retrieve file from URL.</i>
--------	--------------------------------

---

### Description

A function that can be used to retrieve any file of interest from the internet, in our case, mod-Encode binding site information GFF files into the working directory. See also help for function gff2RDList.

### Usage

```
getUrl(urls, filenames, extension='.gff3', method='internal')
```

### Arguments

urls	Character vector with one or more target URLs to download.
filenames	Character vector with the filename for each URL target.
extension	If desired, an extension to append to filenames.
method	Either 'internal' to use the system's default or 'wget' if it is installed.

### Value

Message indicating the path to downloaded file(s).

### Examples

```
# Not run
#getUrl('http://www.google.com/index.html','index','.html')
```

---

gff2RDList	<i>Retrieve binding site information from GFF3 files.</i>
------------	---

---

### Description

An auxiliary function to retrieve binding site information from GFF3 format files (for instance those downloaded from modEncode, see function getUrl).

### Usage

```
gff2RDList(filenames,listnames,dir,quote=NULL,chrprefix='')
```

### Arguments

filenames	GFF3 filenames to read.
listnames	Names for each read filename, will be used as names of the returned GRangesList. If not given, filenames will be used as listnames.
dir	Directory where the GFF3 files are located.
quote	Quote character used in the GFF3 files around fields, if any.
chrprefix	Prefix to be placed before the chromosome names if desired, for instance 'chr'.

**Value**

A list with Enriched and Depleted binding sites, each one is an object of class GRangesList with the GRanges objects containing the respective enriched or depleted binding sites from each GFF3 file.

**Examples**

```
# Not run
#getUrl('http://intermine.modencode.org/release-30/features.do?type=submission&action=export&format=gff3&su
#test <- gff2RDLlist('test.gff3',dir=getwd())
#test
#test$Enriched[[1]]
#test$Depleted[[1]]
```

gps2xgmm1

*Export an 'mds' object to Cytoscape .xgmm1 format***Description**

gps2xgmm1 creates a .xgmm1 file for visualizing MDS results in Cytoscape. Two-dimensional MDS maps can be visualized in Cytoscape as usual. For three-dimensional maps Cytoscape's 3D Renderer ([http://wiki.cytoscape.org/Cytoscape\\_3/3D\\_Renderer](http://wiki.cytoscape.org/Cytoscape_3/3D_Renderer)) is required.

**Usage**

```
gps2xgmm1(x, fname='out.xgmm1', names.arg, fontSize=4, col=gplots::col2hex('steelblue'), cex)
```

**Arguments**

x	Object of class mds
fname	Name of output file
names.arg	Names for each point. If missing, they're taken from x.
fontSize	Font size
col	Fill colour(s) for the plotting symbols. Should be given in hexadecimal, e.g. as returned by function col2hex from gplots. Tips: col2hex('steelblue') looks nice in 2D/3D plots, col2hex('steelblue') looks nice in 2D plots and a bit faded on 3D plots.
cex	Expansion factor for plotting symbols. By default, cex=12 for 2D plots and cex=100 for 3D plots.

**Details**

The .xgmm1 file contains the map co-ordinates in 2 or 3 dimensions, depending on the number of dimensions stored in the input mds object. To visualize properly a file with 3D co-ordinates, you need to install Cytoscape's 3D Renderer ([http://wiki.cytoscape.org/Cytoscape\\_3/3D\\_Renderer](http://wiki.cytoscape.org/Cytoscape_3/3D_Renderer)) and start Cytoscape following the instructions provided therein.

An .xgmm1 file with 3D co-ordinates can still be visualized in regular Cytoscape but the z-axis will be ignored.

**Value**

Generates an .xgmml file that can be opened in Cytoscape (File -> Import -> Network).

**Examples**

```
#See help(mds) for an example
```

---

 mds

*Metric and non-metric Multidimensional Scaling for a distGPS object.*

---

**Description**

Generation of Multidimensional Scaling objects for the dissimilarities between elements given as an input in a `distGPS` object. Metric and non-metric algorithms are available, as well as an optimization algorithm for improving r-square correlation between observed and approximated distances. The MDS calculation for a given distance matrix can be splitted into smaller individual tasks and run in parallel, greatly improving CPU time and system memory usage. The S4 accessor functions `getR2`, `getStress`, `getPoints` retrieve R-square correlation, stress and points stored within a `mds` object respectively. The function `is.adj` is useful to know if a certain `chroGPS` MDS map has been adjusted by Procrustes or not (see `help` for `procrustesAdj` for details.)

**Usage**

```
mds(d, m = NULL, k = 2, type = "classic", add = FALSE, cor.method = "pearson", splitMDS = FALSE, split
getR2(m)
getStress(m)
getPoints(m)
```

**Arguments**

<code>d</code>	Object of class <code>distGPS</code> with the pairwise observed dissimilarities between elements, a distance matrix.
<code>m</code>	(Optional). Object of class <code>mds</code> with a MDS object generated from the distances in <code>d</code> . Only MDS type "boostMDS" is available. The <code>mds</code> function performs an optimization of the approximated distances in <code>m</code> in order to improve r-square correlation between them and the observed dissimilarities in <code>d</code> , maximizing goodness of fit.
<code>k</code>	Dimensionality of the reconstructed space, typically set to 2 or 3.
<code>type</code>	Set to "classic" to perform classical MDS (uses function <code>cmdscale</code> from package <code>stats</code> ). Set to "isoMDS" to use Kruskal's non-metric MDS (uses function <code>isoMDS</code> from package <code>MASS</code> ) Set to "boostMDS" to perform r-square optimization of a pre-computed input MDS for that distance matrix.
<code>add</code>	Logical indicating if an additive constant $c^*$ should be computed, and added to the non-diagonal dissimilarities such that all $n-1$ eigenvalues are non-negative in <code>cmdscale</code> .
<code>cor.method</code>	A character string indicating which correlation coefficient (or covariance) is to be computed. One of "pearson" (default), "kendall", or "spearman", can be abbreviated.

<code>splitMDS</code>	Set to TRUE to perform computation of the MDS in parallel (see parameters below).
<code>split</code>	Proportion of elements to include in each (but last) distance submatrix.
<code>overlap</code>	Proportion of elements to be used as common anchor points between two adjacent distance submatrixes. These points will be used as spatial references to stitch each two adjacent MDS objects by Procrustes.
<code>stepSize</code>	Size for the quadratic search step to be used for R-square optimization if <code>boostMDS</code> is called, see specific help function for details.
<code>reshuffle</code>	Set to TRUE to perform random resampling of the input distance matrix before splitting it for parallel computation. This is often necessary to sufficiently capture the inherent variability of the data in each distance submatrix so that the stitching process can work properly, as the original data may present an arbitrary sorting of some kind. If a previous resampling of the data has been performed, this is not necessary.
<code>set.seed</code>	Random seed to perform the resampling.
<code>mc.cores</code>	Number of cores to be passed to the <code>mclapply</code> function from the <code>parallel</code> package, used to perform the parallel MDS computations.
<code>...</code>	Additional parameters passed to <code>cmdscale</code> , <code>isoMDS</code> or <code>boostMDS</code> , see each individual help file for details.

### Value

The function returns a `mds` object. See `help("mds-Class")` for details.

### Methods

- mds** `signature(d = "distGPS", m = "missing")`: Creates a `mds` object with points in a  $k$ -dimensional space approximating the pairwise distances in `d`.
- mds** `signature(d = "distGPS", m = "mds")`: For the observed dissimilarities in `d` and a valid spatial representation of them in `m`, the function returns a `mds` object with an optimized representation of `d` in terms of R-square. The MDS stress measure is also returned. See `help` for `boostMDS` for details.
- plot** `signature(m = "mds")`: S4 plot method for `mds` objects.

### Author(s)

Oscar Reina

### See Also

See functions `cmdscale`, `isoMDS` from package `MASS`.

### Examples

```
x <- rbind(c(rep(0,15),rep(1,5)),c(rep(0,15),rep(1,5)),c(rep(0,19),1),c(rep(1,5),rep(0,15)))
rownames(x) <- letters[1:4]
d <- distGPS(x,metric='tanimoto',uniqueRows=TRUE)
mds1 <- mds(d)
mds1
plot(mds1)
#gps2xgmm1(mds1, fname='chroGPS_factors.xgmm1', fontSize=4,col=col2hex('red'), cex=8)
```

---

 mds-class

 Class "mds"
 

---

## Description

Multidimensional Scaling. Function `mds` creates object of this class.

## Details

Parameters for the S4 plot method for `mds` objects.

Object of class "mds" with a 2D or 3D Multidimensional Scaling to be plotted.

`drawlabels`: TRUE to use rownames of the MDS points as text labels.

`labels`: Alternative character vector giving the text labels for the MDS points.

`planar`: If a 3D MDS is used, set `planar` to TRUE to plot projected views of the MDS using XY, YZ and XZ axis decomposition.

`point.cex`: Size of the points / spheres for the MDS plot.

`text.cex`: Size of text labels for the MDS points.

`text.pos`: Alignment position of the text labels respective to its points (1,2,3,4).

`point.col`: Color for the MDS points / spheres.

`text.col`: Color for the MDS text labels.

`point.pch`: PCH type for the MDS points.

`type.3d`: Use 'p' for points, 's' for spheres.

`radius`: Radius for the spheres on a 3D MDS plot. Automatically generated from `point.cex` and the number of points in the MDS.

`app`: Appearance of the 3D spheres on a 3D MDS plot, can be 'fill', 'lines', 'grid'.

`alpha`: Number between 0 and 1 with the level of transparency to be used on spheres on a 3D MDS.

`scalecol`: Set to TRUE to use a color scale for points, for instance to color points (genes) according to their expression level on a chroGPS-genes MDS plot.

`scale`: Scale to use to generate scale colors (for instance normalized gene expression for each element (gene) on chroGPS-genes MDS).

`palette`: Color palette to be used for scale colors.

`xlim`: Graphical limit for the X axis.

`ylim`: Graphical limit for the Y axis.

`zlim`: Graphical limit for the Z axis for 3D plots.

## Objects from the Class

Objects can be created by calls of the form `new("mds", ...)`.



**Slots**

**points:** Object of class "matrix" with coordinates in the approximated space.

**Type:** Object of class "character" with the type of MDS (classicMDS, isoMDS).

**Adj:** Object of class "logical" indicating if the MDS object has been adjusted by Procrustes or not.

**R. square:** Object of class "numeric" with the percentage of variability from the original dissimilarities captured in the approximated distances.

**stress:** Object of class "numeric" with the stress for the returned MDS configuration.

**Author(s)**

Oscar Reina

**See Also**

cmdscale from package base. isoMDS from package MASS.

**Examples**

```
showClass("mds")
```

---

mergeClusters	<i>Unsupervised cluster merging based on their observed overlap with automatic changepoint detection.</i>
---------------	---

---

**Description**

The function uses contour density estimation as computed by the `clusGPS` function to merge significantly overlapping clusters in an unsupervised manner. In each step, clusters with highest overlap are merged, their individual density estimates are updated in a computational feasible manner, and the process continues until the maximum overlap between any given pair of clusters drops swiftly, as detected by the `cpt.mean` function in the `changepoint` package.

**Usage**

```
mergeClusters(clus, clus.method = "unweighted", cpt.method = "mean", logscale = TRUE, brake = rep(1,
```

**Arguments**

<code>clus</code>	A <code>clusGPS</code> object from which we want to merge clusters with a significant overlap when visualized on a <code>chroGPS</code> MDS map. This is quite useful when a clustering method (i.e. hierarchical clustering with average linkage) tends to return a high number of overlapping clusters.
<code>clus.method</code>	Currently only 'unweighted' method is supported, that is, cluster overlap is computed based on spatial location of contours, but the computed overlaps are not weighted for cluster size.
<code>cpt.method</code>	Use 'mean' for using <code>cpt.mean</code> function in <code>changepoint</code> package for computing overlap changepoint. Use 'var' for <code>cpt.var</code> . See specific function help for details.

logscale	Defaults to TRUE. Whether to use decimal or log scale values for computing overlap changepoint.
brake	(Optional). By default, the function returns the clusters from the optimal merging step as detected by the changepoint functions (brake=1). By using smaller values (0, -1, -2, ...) or bigger ones (2, 3, 4, ...) the algorithm can be forced to return the result from any previous or later merging step respectively.
plt	Set to TRUE to visualize maximum cluster overlap for each merging step and changepoint detection (optimal merging step).
mc.cores	Numbers of cores to use in parallel computation.

**Value**

A clusGPS object where significantly overlapping clusters are merged, highly improving visualization, cluster robustness and further study of the epigenetic configuration of the chroGPS map.

**Author(s)**

Oscar Reina.

**References**

Changepoint package from Killick et al, 2012.

**See Also**

See documentation for package changepoint, clusGPS for epigenetic cluster generation.

**Examples**

```
# Not run
# data(s2)
# # Computing distances
# d <- distGPS(s2.tab,metric='tanimoto',uniqueRows=TRUE)
# # Creating MDS object
# mds1 <- mds(d,type='isoMDS')
# mds1
# plot(mds1)
# Precomputing clustering
# h <- hclust(as.dist(d@d),method='average')
# # Calculating densities (contours and probabilities), takes a while
# clus <- clusGPS(d,mds1,preMerge=TRUE,k=300) # Generating a high number of clusters
# clus <- mergeClusters(clus)
```

---

mergeReplicates	<i>Merges information from epigenetic replicates at factor, gene and chroGPS MDS map level.</i>
-----------------	---

---

**Description**

The function performs effectively merging of epigenetic replicate information, either as genomic intervals or as gene epigenetic profiles. It also allows merging of final chroGPS MDS factor maps so that multiple points for the same epigenetic factor (i.e. different antibodies or experimental sources) are returned as a single location in the map (centroid).

**Usage**

```
mergeReplicates(x, id, mergeBy='any', mc.cores=1)
```

**Arguments**

<code>x</code>	Object containing the epigenetic information used to generate either factors or gene maps. Can also be a final chroGPS-factors MDS map object.
<code>id</code>	Identifier for epigenetic factors (list elements for <code>x='GRangesList'</code> , column names for <code>x='matrix'</code> or <code>'data.frame'</code> , row names for <code>x='mds'</code> ). This information will be used to identify valid replicates to be merged.
<code>mergeBy</code>	Either a character vector with possible values <code>'any'</code> or <code>'all'</code> to merge replicate genomic intervals or gene epigenetic profiles based on a logical <code>'OR'</code> or <code>'AND'</code> fashion, or a decimal number between (0,1] indicating the minimum proportion of replicates needed to consider.
<code>mc.cores</code>	Number of cores to use in calls to <code>parallel::mclapply</code>

**Value**

An object from the same class as `x`, containing a data set with merged replicates as specified by the `'mergeBy'` argument.

**Methods**

**signature(x='GRangesList')** Each element in `x` is assumed to indicate the binding sites for a different sample, e.g. epigenetic factor. Typically `space(x)` indicates the chromosome, `start(x)` the start position and `end(x)` the end position (in bp). Strand information is ignored.

**signature(x='matrix')** Rows in `x` contain individuals for which we want to compute distances. Columns in `x` contain the variables, and should only contain either 0's and 1's or FALSE and TRUE.

**signature(x='mds')** A MDS object containing a chroGPS-factors map.

**See Also**

[mds](#) to create MDS-oriented objects

**Examples**

```
## Not run

#data(s2)
#data(bg3)

#names(s2)
#names(bg3)

# Unify replicates
#mnames <- sort(unique(intersect(s2names$Factor, bg3names$Factor)))
#sel <- s2names$Factor %in% mnames
#s2.repset <- mergeReplicateList(s2[sel], id=s2names$Factor[sel], mergeBy='any')
#sel <- bg3names$Factor %in% mnames
#bg3.repset <- mergeReplicateList(bg3[sel], id=bg3names$Factor[sel], mergeBy='any')
```

```
#names(s2.repset)
#names(bg3.repset)
```

---

procrustesAdj      *Use Procrustes to adjust an MDS map containing samples obtained under different conditions, e.g. technology or genetic backgrounds.*

---

### Description

The function adjusts a previous mds to take into account that samples were obtained under different conditions, e.g. technological or genetic. Pairwise adjustments are performed by identifying samples present in both conditions and using Procrustes. When there are more than two conditions, sequential pairwise adjustments are applied (in the order that maximizes the number of common samples in each pairwise adjustment).

### Usage

```
procrustesAdj(mds1, d, adjust, sampleid)
```

### Arguments

mds1	Object of class mds with a Multi-dimensional scaling analysis on a distance matrix, typically obtained by a previous call to mds.
d	Object of class distGPS with the matrix used to create the Multidimensional Scaling object usually through a call to mds.
adjust	Vector indicating the adjustment factor, i.e. the condition under which each sample has been obtained.
sampleid	Vector containing the sample identifier. sampleid should take the same value for samples obtained under different conditions, as this is used to detect the samples to be used for Procrustes adjustment.

### Details

We implement the Procrustes adjustment as follows. First we identify common samples, i.e. those obtained both under conditions A and B. Second, we use Procrustes to estimate the shift, scale and rotation that best matches the position of the samples in B to those in A. If only 1 sample was obtained under both conditions, only the shift is estimated. Last, we apply the estimated shift, scale and rotation to all B samples. That is, the Procrustes parameters are estimated using common samples only, which are then applied to all samples to perform the adjustment.

Notice that the R square of the adjusted mds is typically improved after Procrustes adjustment, since distances between samples obtained under different conditions are set to NA and therefore MDS needs to approximate distances between less points.

When several replicates are available for a given sampleid under the same condition (adjust), the average position of all replicates is used.

### Value

Adjusted mds object. Have in mind that only original distances between samples obtained under the same condition should be conserved, as the adjusted distances manipulated by Procrustes no longer correlate with the distances between their points in the adjusted MDS.

## Methods

**signature(x='mds')** x is a mds object with the results of an MDS analysis.

## See Also

[distGPS](#) for computing distances, [mds](#) to create MDS-oriented objects.

## Examples

```
# Unadjusted map
data(s2)
data(s2Seq)
data(toydist) # precomputed distances
# d2 <- distGPS(c(reduce(s2),reduce(s2Seq)),metric='avgdist') # not run
mds2 <- mds(d2,k=2,type='isoMDS')
cols <- c(as.character(s2names$Color),as.character(s2SeqNames$Color))
sampleid <-
  c(as.character(s2names$Factor),as.character(s2SeqNames$Factor))
pchs <- rep(c(20,17),c(length(s2),length(s2Seq)))
point.cex <- rep(c(8,5),c(length(s2),length(s2Seq)))
par(mar=c(2,2,2,2))
plot(mds2,drawlabels=TRUE,point.pch=pchs,point.cex=point.cex,text.cex=.7,
point.col=cols,text.col='black',labels=sampleid,font=2)
#legend('topleft',legend=sprintf('R2=%.3f - %stress=%.3f',getR2(mds2),getStress(mds2)),bty='n',cex=1)
legend('topright',legend=c('ChIP-Chip','ChIP-Seq'),pch=c(20,17),pt.cex=c(1.5,1))

# Procrustes Adjusted map
adjust <- rep(c('chip','seq'),c(length(s2),length(s2Seq)))
sampleid <-
  c(as.character(s2names$Factor),as.character(s2SeqNames$Factor))
mds3 <- procrustesAdj(mds2,d2,adjust=adjust,sampleid=sampleid)
par(mar=c(0,0,0,0),xaxt='n',yaxt='n')
plot(mds3,drawlabels=TRUE,point.pch=pchs,point.cex=point.cex,text.cex=.7,
point.col=cols,text.col='black',labels=sampleid,font=2)
#legend('topleft',legend=sprintf('R2=%.3f - %stress=%.3f',getR2(mds3),getStress(mds3)),bty='n',cex=1)
legend('topright',legend=c('ChIP-Chip','ChIP-Seq'),pch=c(20,17),pt.cex=c(1.5,1))

# Peak Width Adjusted map
s2.pAdj <-
adjustPeaks(c(reduce(s2),reduce(s2Seq)),adjust=adjust,sampleid=sampleid,logscale=TRUE)
# d3 <- distGPS(s2.pAdj,metric='avgdist')
mds4 <- mds(d3,k=2,type='isoMDS')
par(mar=c(0,0,0,0),xaxt='n',yaxt='n')
plot(mds4,drawlabels=TRUE,point.pch=pchs,point.cex=point.cex,text.cex=.7,
point.col=cols,text.col='black',labels=sampleid,font=2)
#legend('topleft',legend=sprintf('R2=%.3f - %s=%.3f',getR2(mds4),getStress(mds4)),bty='n',cex=1)
legend('topright',legend=c('ChIP-Chip','ChIP-Seq'),pch=c(20,17),pt.cex=c(1.5,1))
```

**Description**

Assess epigenetic profiles for genes present in each cluster as obtained by the `clusGPS` function.

**Usage**

```
profileClusters(x, clus, clusName = NULL, normalize = FALSE, mc.cores = 1, ...)
```

**Arguments**

<code>x</code>	Matrix or data.frame with epigenetic profiles used to generate chroGPS-genes map. See <code>distGPS</code> .
<code>clus</code>	A valid <code>clusGPS</code> object from the chroGPS-genes map generated from <code>x</code> .
<code>clusName</code>	If desired, the name of a clustering solution within the <code>clusGPS</code> object. See <code>clusNames</code> .
<code>normalize</code>	Normalize epigenetic profile information with respect to whole genome (to assess enrichments/depletions).
<code>mc.cores</code>	Cores to use in call to <code>parallel::mclapply</code> .
<code>...</code>	Additional arguments.

**Value**

Data frame with epigenetic enrichment profiles for all clusters (rows) and epigenetic factors (columns), to be used for downstream assessments or visualization (heatmaps).

**Author(s)**

Oscar Reina.

**See Also**

`distGPS`, `clusGPS`.

**Examples**

```
## Not run
```

---

`rankFactorsbyDomain`     *Function to help selecting candidate epigenetic factors based on Epigenetic Domain cohesion/separation.*

---

**Description**

Function to help selecting candidate epigenetic factors based on Epigenetic Domain cohesion/separation. This function ranks all combinations of a certain size of epigenetic factors in the selected domain based on how much they contribute to provide a good cohesion and separation of the points of their epigenetic domain against the rest. This allows selection of the 'best' combination of epigenetic factors in a certain domain to generate informative epigenetic maps.

**Usage**

```
rankFactorsbyDomain(d, sampleinfo, ranktype = "domainDist", selName = "Color", selValue, k = NULL, m
```

**Arguments**

d	A distGPS object with the epigenetic distances, see function distGPS.
sampleinfo	A data.frame object with at least the domain information for the epigenetic factors in the provided distance matrix. Sampleinfo rownames need to match row and colnames in the distance matrix.
ranktype	Character vector indicating the method to use. Currently only supporting 'domainDist' method.
selName	Name of the column containing the epigenetic domain information in the sampleinfo dataframe.
selValue	Character indicating the name of the domain to be evaluated.
k	Size of the factor combination to be evaluated.
mc.cores	Cores to use in calls to parallel::mclapply.

**Value**

Named list with each evaluated combination of factors and:

inter	Their resulting average inter-domain distance.
intra	Their resulting average intra-domain distance.

**Author(s)**

Oscar Reina.

**See Also**

[distGPS](#), [domainDist](#)

**Examples**

```
## Rank Factors by Domain, using intra/inter domain distance

data(s2)
data(toydists)
#d <- distGPS(s2,metric='avgdist',mc.cores=8) # Compute distances
rownames(s2names) <- s2names$ExperimentName

# Known domains
# Call rankFactorsbyDomain for HP1a repression domain, select a combination of 4
# factors
library(caTools)
rank.factors.4 <- rankFactorsbyDomain(d,s2names,ranktype='domainDist',selName='Color',selValue='lightblue',
ddd <- as.data.frame(do.call(rbind,lapply(rank.factors.4,unlist)))
ddd <- ddd[order(ddd$intra,decreasing=FALSE),]
head(ddd)
```

---

rankFactorsbyProfile *Function to help selecting candidate epigenetic factors based on their predictive capabilities.*

---

### Description

Function to help selecting candidate epigenetic factors based on their predictive capabilities. This function evaluates how information from a certain set of epigenetic factors can be used to accurately predict information from the rest using linear and logistic regression.

### Usage

```
rankFactorsbyProfile(x,minFactors=5,ranktype='glm',glm.threshold=0.5,verbose=TRUE,maxIter=ncol(x))
```

### Arguments

x	A genes x factors table as used in the distGPS function for generation of epigenetic gene maps.
minFactors	Minimum set of 'core' factors to retain. Factors are removed based on how accurately they can be predicted by others.
ranktype	Either 'lm' for using linear regression, or 'glm' for logistic.
glm.threshold	For 'glm', threshold to round values obtained from the predict function so that they are compared with the real values. Defaults to 0.5.
verbose	Informs about progression of the linear or logistic regression steps.
maxIter	Deprecated.
mc.cores	Cores to use in calls to parallel::mclapply.

### Value

A data frame with the epigenetic factors in the order they are removed and their prediction accuracy.

### Author(s)

Oscar Reina.

### See Also

[distGPS](#), [domainDist](#), [rankFactorsbyDomain](#)

### Examples

```
## Not run

data(s2)

# Unknown domains
# Perform computation
glm.rank <- rankFactorsbyProfile(s2.tab,ranktype='glm',glm.threshold=0.75,mc.cores=1)

# Returned objects are lists named by the factor with highest prediction accuracy in each iteration
names(glm.rank)
```



---

s2

*Sample binding site and related data from S2 and BG3 cell lines in Drosophila melanogaster.*

---

## Description

chroGPS example dataset including ChIP-CHIP (modEncode) and ChIP-Seq (NCBI GEO GSE19325) data for *Drosophila melanogaster* S2 and BG3 cell lines as well as S2 wildtype gene expression values coming from Affymetrix *Drosophila2* arrays. The object `toydist` stores precomputed `distGPS` objects (called `d`, `d2`, `d3`) for the epigenetic factors used in the dynamic vignette that comes with the package. The objects `d.origs`, `m.origs`, contain the `distGPS` and `mds` objects for the 76 S2 epigenetic factors used in Font-Burgada et al. 2014, computed over modENCODE Origins of Replication at four different replication time points.

## Usage

```
data(s2)
data(s2Seq)
data(bg3)
data(repliSeq)
```

## Source

<http://www.modencode.org> <http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE19325>

## References

<http://www.modencode.org> <http://www.ncbi.nlm.nih.gov/geo/>

## Examples

```
data(s2)
class(s2)
s2
s2names$Factor

data(s2Seq)
s2Seq

data(bg3)
names(bg3)

data(repliSeq)
class(d.origs)
class(m.origs)
names(d.origs)
d.origs[[1]]
m.origs[[1]]
```

# See vignette examples for several uses of these datasets.

---

splitDistGPS-class      *Class "splitDistGPS"*

---

**Description**

Set of pairwise distances between elements. This is an internal class to be used with the parallel version of mds, and should not be used on its own.

**Objects from the Class**

Objects from this class are used internally for parallel Multidimensional Scaling. See mds for details.

**Slots**

**d:** List of distGPS objects.

**size:** Object of class "numeric" indicating the size of the individual distGPS objects in the list. See function mds.

**o:** Object of class "numeric" with the overlap (anchor points) between adjacent distGPS objects. See function mds.

**shuffle:** Object of class "numeric", deprecated.

**Author(s)**

Oscar Reina

**Examples**

```
showClass("splitDistGPS")
```

# Index

- \*Topic **~changepoint**
  - mergeClusters, 25
- \*Topic **~clustering**
  - diffGenes, 12
  - mergeClusters, 25
- \*Topic **~mds**
  - diffFactors, 11
  - diffGenes, 12
- \*Topic **classes**
  - clusGPS-class, 9
  - distGPS-class, 17
  - mds-class, 24
  - splitDistGPS-class, 34
- \*Topic **clustering**
  - distGPS, 15
  - mergeReplicates, 26
- \*Topic **cluster**
  - clusGPS, 6
- \*Topic **datasets**
  - bg3, 4
  - s2, 33
- \*Topic **graphics**
  - geneSetGPS, 19
- \*Topic **graphs**
  - mds, 22
- \*Topic **manip**
  - gps2xgmml, 21
- \*Topic **mds**
  - mds, 22
- \*Topic **modEncode**
  - bg3, 4
  - s2, 33
- \*Topic **multivariate,cluster**
  - addVar, 2
  - adjustPeaks, 3
  - boostMDS, 5
  - domainDist, 18
  - getURL, 20
  - gff2RDList, 20
  - procrustesAdj, 28
- \*Topic **multivariate**
  - distGPS, 15
  - mergeReplicates, 26
- addVar, 2
- adjustPeaks, 3
- adjustPeaks,GRangesList-method (adjustPeaks), 3
- adjustPeaks-methods (adjustPeaks), 3
- as.matrix,distGPS-method (distGPS-class), 17
- bg3, 4
- bg3names (bg3), 4
- boostMDS, 5
- clusGPS, 6
- clusGPS,distGPS,mds-method (clusGPS), 6
- clusGPS-class, 9
- clusGPS-method (clusGPS-class), 9
- clusGPS-methods (clusGPS), 6
- clusNames (clusGPS), 6
- clusNames,clusGPS-method (clusGPS), 6
- clusterID (clusGPS), 6
- clusterID,clusGPS-method (clusGPS), 6
- combineGenesMatrix, 10
- contour2dDP (clusGPS), 6
- d (s2), 33
- d2 (s2), 33
- d3 (s2), 33
- diffFactors, 11
- diffFactors-methods (diffFactors), 11
- diffGenes, 12
- diffGenes-methods (diffGenes), 12
- distGPS, 4, 15, 29, 31, 32
- distGPS,data.frame-method (distGPS), 15
- distGPS,GRangesList-method (distGPS), 15
- distGPS,matrix-method (distGPS), 15
- distGPS-class, 17
- distGPS-methods (distGPS), 15
- domainDist, 18, 31, 32
- find.fdr (diffGenes), 12
- find.threshold (diffGenes), 12
- geneSetGPS, 19
- geneSetGPS,data.frame,mds,character-method (geneSetGPS), 19

- geneSetGPS, matrix, mds, character-method (geneSetGPS), 19
- geneSetGPS-methods (geneSetGPS), 19
- getPoints (mds), 22
- getPoints, mds-method (mds), 22
- getR2 (mds), 22
- getR2, mds-method (mds), 22
- getStress (mds), 22
- getStress, mds-method (mds), 22
- getURL, 20
- gff2RDList, 20
- gps2xgmm1, 21
- gps2xgmm1, mds, ANY-method (gps2xgmm1), 21
- gps2xgmm1, mds-method (gps2xgmm1), 21
  
- hclust, 7
- hclust-class (clusGPS-class), 9
  
- is.adj (mds), 22
- is.adj, mds-method (mds), 22
  
- m.origs (s2), 33
- mds, 4, 16, 22, 27, 29
- mds, distGPS, mds-method (mds-class), 24
- mds, distGPS, missing-method (mds), 22
- mds, splitDistGPS, missing-method (mds), 22
- mds-class, 24
- mds-methods (mds), 22
- mergeClusters, 25
- mergeClusters, clusGPS-method (mergeClusters), 25
- mergeClusters, list-method (mergeClusters), 25
- mergeClusters-methods (mergeClusters), 25
- mergeReplicateList (mergeReplicates), 26
- mergeReplicateMatrix (mergeReplicates), 26
- mergeReplicateMDS (mergeReplicates), 26
- mergeReplicates, 26
- mergeReplicates, data.frame-method (mergeReplicates), 26
- mergeReplicates, GRangesList-method (mergeReplicates), 26
- mergeReplicates, list-method (mergeReplicates), 26
- mergeReplicates, matrix-method (mergeReplicates), 26
- mergeReplicates, mds-method (mergeReplicates), 26
- mergeReplicates-methods (mergeReplicates), 26
  
- plot, clusGPS, ANY-method (clusGPS-class), 9
- plot, clusGPS-method (clusGPS-class), 9
- plot, mds, ANY-method (mds), 22
- plot, mds-method (mds-class), 24
- plotContour (clusGPS), 6
- plotDiffGenes (diffGenes), 12
- procrustesAdj, 4, 16, 28
- procrustesAdj, mds, distGPS-method (procrustesAdj), 28
- procrustesAdj-methods (procrustesAdj), 28
- profileClusters, 29
  
- rankFactorsbyDomain, 30, 32
- rankFactorsbyDomain, data.frame-method (rankFactorsbyDomain), 30
- rankFactorsbyDomain, distGPS-method (rankFactorsbyDomain), 30
- rankFactorsbyDomain, matrix-method (rankFactorsbyDomain), 30
- rankFactorsbyDomain-methods (rankFactorsbyDomain), 30
- rankFactorsbyProfile, 32
- repliSeq (s2), 33
  
- s2, 33
- s2names (s2), 33
- s2Seq (s2), 33
- s2SeqNames (s2), 33
- show, clusGPS-method (clusGPS-class), 9
- show, distGPS-method (distGPS-class), 17
- show, mds-method (mds-class), 24
- show, splitDistGPS-method (splitDistGPS-class), 34
- splidDistGPS-class (splitDistGPS-class), 34
- splitDistGPS, data.frame-method (distGPS), 15
- splitDistGPS, distGPS-method (splitDistGPS-class), 34
- splitDistGPS, matrix-method (distGPS), 15
- splitDistGPS-class, 34
- splitDistGPS-class (distGPS-class), 17
  
- tabClusters (clusGPS), 6
- tabClusters, clusGPS-method (clusGPS), 6
- toydists (s2), 33
  
- uniqueCount (distGPS), 15