# Package 'caOmicsV'

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Title Visualization of multi-dimentional cancer genomics data
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<b>Description</b> caOmicsV package provides methods to visualize multi-dimentional cancer genomics data including of patient information, gene expressions, DNA methylations, DNA copy number variations, and SNP/mutations in matrix layout or network layout
License GPL (>=2.0)
<b>Depends</b> R ( $>= 3.1.1$ ), igraph ( $>= 0.7.1$ ), bc3net ( $>= 1.0.2$ )
biocViews Visualization, Network

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caOmicsV-package

caOmicsV bioMatrix and bioNetCircos Layout Plot

#### **Description**

This package provides methods to display genomics data with two layouts:

bioMatrix layout: sample info and multiple genomics data are displayed as matrix with rows for features(phenotypes and genes, and columns for each sample. Omics data will be plotted with different layers when necessary.

bioNetCircos layout: sample info and multiple genomics data are displayed on an biological network with node for features (e.g., genes) and on each node, sample info and genomics data are plotted with circular layout.

#### **Details**

Package: caOmicsV Type: Package Version: 0.99.0 Date: 2015-03-06 License: GPL >= 2.0

#### Author(s)

Henry Zhang Maintainer: Henry Zhang <a href="mail.nih.gov">hzhang@mail.nih.gov</a>

#### **Examples**

library(caOmicsV)

bioMatrixLegend

Plot Legend on caOmicsV bioMatrix Layout

## **Description**

Draw legend including of heatmap color scale and data categories. Graphic device must be initialized first.

# Usage

bioMatrixLegend(heatmapNames=NULL, categoryNames=NULL, binaryNames=NULL, heatmapMin=NULL, heatmapMax=NULL, colorType="BlueWhiteRed") 4 bioNetCircosPlot

#### **Arguments**

categoryNames character vector of length 2, name(s) of dataset for heatmap, e.g., "RNASeq" and/or "miRNASeq" character vector of length 2 or more, names of categories, e.g., "Methylation High", "Methylation Low", ...

binaryNames character vector of length 2, names of binary data, e.g., "DNA Amplification" and "DNA Deletion".

heatmapMin numeric, minimum values of heatmap plot data, default -3 (z-scores)

heatmapMax numeric, maximum values of heatmap plot data, default 3 (z-scores)

colorType characte vector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed",

"GreenBlackRed", or "YellowToRed"

#### **Details**

This function will plot legend on the bottom of matrix layout if any argument is defined. The order of legend items (from left to right) is heatmap color scale followed by colored boxes for category data legend then colored points for binary data legend.

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
data(caOmicsV.biomatrix.eset)
plotBioMatrix(caOmicsV.biomatrix.eset, summaryType="text")
bioMatrixLegend(heatmapNames=c("RNASeq", "miRNASeq"),
    categoryNames=c("Methyl H", "Methyl ZL"),
    binaryNames=c("CN LOSS", "CN Gain"),
    heatmapMin=-3, heatmapMax=3, colorType="BlueWhiteRed")
```

bioNetCircosPlot

caOmicsV bioNetCircos Layout Plot

#### **Description**

Plot one track of caOmics data on each node of a biological network. Supported plot types include polygon, bar, points, heatmap, and line plots.

# Usage

```
bioNetCircosPlot(dataValues=NULL, plotType="polygon", outer, inner,
    plotColors=NULL, maxValue=NULL, minValue=NULL)
```

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#### **Arguments**

dataValues an numeric matrix with columns for samples and rows for genes plotType character vector for plot type, supporting polygon, points, and lines.

outer non-negative numeric, outer location of a data track inner non-negative numeroc, inner location of a data track plotColors character vector specifying Colors for plot items

maxValue numeric, the biggest value of plot data or user defined top threshold. Set to

NULL to use the biggest value in dataset.

minValue numeric, the smallest value of plot data or user defined bottom threshol. Set to

NULL to use the smallest value in dataset.

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)
initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout()

binaryData <- caOmicsV.bionet.eset$binaryData[[1]]
sampleColors <- c(rep("blue", ncol(binaryData)))
plotType <- "points"
inner <- 2
outer <- 3
bioNetCircosPlot(binaryData, plotType, outer, inner, sampleColors)</pre>
```

bioNetLegend

Draw Legend for caOmicsV bioNet Plot

# Description

The bioNet legend includes a heatmap color scale and names for each track, such as "1. Tissue: T(red), N(blue)", "2. Methylation", "3. miRNA hsa-mir-424", "4. Gene Expression", ... A graphic device and igraph object must be initialized first

#### Usage

```
bioNetLegend(dataNames, textCoor=NULL, heatmapCoor=NULL,
    scaleWidth, scaleHeight, heatmapMin=NULL, heatmapMax=NULL,
    colorType="BlueWhiteRed", direction="h")
```

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#### **Arguments**

dataNames character vector, names of data on each circular track

textCoor numeric vector of length 2, x and y coordinates for legend text.

heatmapCoor numeric vector of length 2, x and y coordinates for heatmap colour scale

scaleWidth non-negative numeric, length (width) of heatmap color scale

scaleHeight non-negative numeric, height of heatmap color scale
heatmapMin numeric, minimum value of heatmap color scale
heatmapMax numeric, maximum value of heatmap color scale

colorType characte veector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed",

"GreenBlackRed", or "YellowToRed"

direction character, direction of heatmap color scale, either 'h' for horizontal or 'v' for

vertical.

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
data(caOmicsV.bionet.eset)
plotBioNetCircos(caOmicsV.bionet.eset)
dataNames <- c("Tissue Type", "RNASeq", "miRNASeq", "Methylation", "CNV")
bioNetLegend(dataNames, heatmapMin=-3, heatmapMax=3)</pre>
```

caOmicsV.biomatrix.eset

Demo Dataset for caOmicsV bioMatrix Plot

# Description

eset for demo of caOmicsV bioMatrix Plot

# Usage

```
data("caOmicsV.biomatrix.eset")
```

#### **Format**

The format is: List of 8

\$ sampleNames : chr [1:60] "BC.A216.Normal" "BD.A2L6.Normal"

"BD.A3EP.Normal" "DD.A113.Normal" ...

\$ geneNames : chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...

\$ secondGeneNames: chr [1:26] "hsa-mir-10b" "hsa-mir-139"

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```
"hsa-mir-10b" "hsa-mir-10b" ...
$ sampleInfo : chr [1:2, 1:60] "TCGA.BC.A216.Normal"
"Solid Tissue Normal" "TCGA.BD.A2L6.Normal" "Solid Tissue Normal" ...
..- attr(*, "dimnames")=List of 2
.. ..$: chr [1:2] "sampleID" "sample_type"
.. ..$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
$ heatmapData :List of 2
..$: num [1:26, 1:60] 1.157 -0.623 0.667 0.976 0.868 ...
....- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
..$: num [1:26, 1:60] -1.5 1.08 -1.5 -1.5 -1.5 ...
....- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
$ categoryData :List of 1
..$: num [1:26, 1:60] 1 0 1 1 1 0 0 0 1 0 ...
....- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
$ binaryData :List of 2
..$: num [1:26, 1:60] 0 0 0 0 0 0 0 0 0 0 ...
...- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
..$: num [1:26, 1:60] 0 0 0 0 0 0 0 0 0 ...
....- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
$ summaryInfo :List of 1
..$: chr [1:26, 1:2] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
....- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:2] "GeneSymbol" "logFC"
```

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#### Value

None

#### **Examples**

```
data("caOmicsV.biomatrix.eset")
```

#### **Description**

eset for demo of caOmicsV bionetCircos Plot

## Usage

```
data("caOmicsV.bionet.eset")
```

\$ categoryData :List of 1

#### **Format**

```
The format is: List of 8
$ sampleNames : chr [1:60] "BC.A216.Normal" "BD.A2L6.Normal"
"BD.A3EP.Normal" "DD.A113.Normal" ...
$ geneNames : chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
$ secondGeneNames: NULL
$ sampleInfo: chr [1:2, 1:60] "TCGA.BC.A216.Normal"
"Solid Tissue Normal" "TCGA.BD.A2L6.Normal" "Solid Tissue Normal" ...
..- attr(*, "dimnames")=List of 2
.. ..$: chr [1:2] "sampleID" "sample_type"
.. ..$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
$ heatmapData :List of 2
..$: num [1:26, 1:60] 1.157 -0.623 0.667 0.976 0.868 ...
...- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
..$: num [1:26, 1:60] -1.5 1.08 -1.5 -1.5 -1.5 ...
....- attr(*, "dimnames")=List of 2
.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...
.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"
"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...
```

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```
..$: num [1:26, 1:60] 0.825 0.364 0.545 0.798 0.606 ...

...- attr(*, "dimnames")=List of 2

.....$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...

.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal"

"TCGA.BD.A3EP.Normal" "TCGA.DD.A113.Normal" ...

$ binaryData :List of 1

...$: num [1:26, 1:60] 0.0266 0.0069 0.0193 0.0117 0.0081 0.0242

0.0069 0.0079 0.0348 0.01 ...

...- attr(*, "dimnames")=List of 2

......$: chr [1:26] "ECM1" "SLC26A6" "ADAMTS13" "FCN3" ...

.....$: chr [1:60] "TCGA.BC.A216.Normal" "TCGA.BD.A2L6.Normal" "TCGA.BD.A3EP.Normal"

"TCGA.DD.A113.Normal" ...

$ summaryInfo: NULL
```

#### Value

None

# **Examples**

data("caOmicsV.bionet.eset")

CA\_OMICS\_ENV

caOmicsV Environment

## **Description**

The caOmicsV Environment holds and protects all parameters and objects used for caOmicsV plot.

# Usage

CA\_OMICS\_ENV

#### **Format**

The format is: <environment: 0x26b3e00>

#### Value

None

# **Examples**

is.environment(CA\_OMICS\_ENV)

CA\_OMICS\_NAME

The Name of caOmicsV Environment

# Description

"CA\_OMICS\_ENV" is used for caOmicsV environment name.

## Usage

CA\_OMICS\_NAME

#### **Format**

The format is: chr "CA\_OMICS\_ENV"

## Value

None

# **Examples**

caOmicsVEnvironment <- get(CA\_OMICS\_NAME, envir=globalenv())</pre>

CA\_OMICS\_NA\_STRING

The Default NA String Used by caOmicsV Package

# Description

The default NA string used by caOmicV package is NULL

# Usage

CA\_OMICS\_NA\_STRING

#### **Format**

The format is: NULL

## Value

None

# **Examples**

CA\_OMICS\_NA\_STRING

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CNVDemoData

Copy Number Variation Demo Data Set

#### **Description**

A data frame with copy number variation of 26 genes in 60 samples. Used for point plot demo on caOmicsV bioNetCircos layout, or bianry data plot on caOmicsV bioMatrix layout after transformed to bionary data.

#### Usage

data("CNVDemoData")

#### **Format**

A data frame with 26 observations on the following 61 variables.

Gene\_Symbol a factor with levels ACTN1 ADAMTS13 AMIGO3 ATP2A1 BCO2 CDKN3 CFP CNBP COL15A1 CSRNP1 CXCL12 DBH DDX55 ECM1 ELOVL1 ESM1 FAM81A FCN3 KCNQ1 LEPREL1 LIFR LILRA6 LILRB5 LOC222699 LOC283050 LRRC16A LYVE1 MAEL MAN1B1 MRPS25 MT1F NIPA2 NPHP4 NR5A2 OLFML3 PLVAP PROX1 PTH1R RBL2 RCAN1 RND3 SEMA3F SLC26A6 TOMM40L TSEN34 VDAC3

```
TCGA.BC.A216.Normal a numeric vector
```

TCGA.BD.A2L6.Normal a numeric vector

TCGA.BD.A3EP.Normal a numeric vector

TCGA.DD.A113.Normal a numeric vector

TCGA.DD.A114.Normal a numeric vector

TCGA.DD.A118.Normal a numeric vector

TCGA.DD.A119.Normal a numeric vector

TCGA.DD.A11A.Normal a numeric vector

TCGA.DD.A11B.Normal a numeric vector

TCGA.DD.A11C.Normal a numeric vector

TCGA.DD.A11D.Normal a numeric vector

TCGA.DD.A1EB.Normal a numeric vector

TCGA.DD.A1EC.Normal a numeric vector

TCGA.DD.A1EG.Normal a numeric vector TCGA.DD.A1EH.Normal a numeric vector

TCGA.DD.A1EI.Normal a numeric vector

TCGA.DD.A1EJ.Normal a numeric vector

TCGA.DD.A1EL.Normal a numeric vector

TCGA.DD.A39V.Normal a numeric vector

TCGA.DD.A39W.Normal a numeric vector

TCGA.DD.A39X.Normal a numeric vector

TCGA.DD.A39Z.Normal a numeric vector

TCGA.DD.A3A1.Normal a numeric vector

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```
TCGA.DD.A3A2.Normal a numeric vector
TCGA.DD.A3A3.Normal a numeric vector
TCGA.EP.A12J.Normal a numeric vector
TCGA.EP.A26S.Normal a numeric vector
TCGA.ES.A2HT.Normal a numeric vector
TCGA.FV.A23B.Normal a numeric vector
TCGA.FV.A2QR.Normal a numeric vector
TCGA.BC.A216.Tumor a numeric vector
TCGA.BD.A2L6.Tumor a numeric vector
TCGA.BD.A3EP.Tumor a numeric vector
TCGA.DD.A113.Tumor a numeric vector
TCGA.DD.A114.Tumor a numeric vector
TCGA.DD.A118.Tumor a numeric vector
TCGA.DD.A119.Tumor a numeric vector
TCGA.DD.A11A.Tumor a numeric vector
TCGA.DD.A11B.Tumor a numeric vector
TCGA.DD.A11C.Tumor a numeric vector
TCGA.DD.A11D.Tumor a numeric vector
TCGA.DD.A1EB.Tumor a numeric vector
TCGA.DD.A1EC.Tumor a numeric vector
TCGA.DD.A1EG.Tumor a numeric vector
TCGA.DD.A1EH.Tumor a numeric vector
TCGA.DD.A1EI.Tumor a numeric vector
TCGA.DD.A1EJ.Tumor a numeric vector
TCGA.DD.A1EL.Tumor a numeric vector
TCGA.DD.A39V.Tumor a numeric vector
TCGA.DD.A39W.Tumor a numeric vector
TCGA.DD.A39X.Tumor a numeric vector
TCGA.DD.A39Z.Tumor a numeric vector
TCGA.DD.A3A1.Tumor a numeric vector
TCGA.DD.A3A2.Tumor a numeric vector
TCGA.DD.A3A3.Tumor a numeric vector
TCGA.EP.A12J.Tumor a numeric vector
TCGA.EP.A26S.Tumor a numeric vector
TCGA.ES.A2HT.Tumor a numeric vector
TCGA.FV.A23B.Tumor a numeric vector
TCGA.FV.A2QR.Tumor a numeric vector
```

#### **Examples**

data(CNVDemoData)

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convertToZScores

Calculate z-scores for A Data Matrix

#### **Description**

Calculate z-scores for data values in a data frame. The row ID must be in the first column followed by data values.

## Usage

```
convertToZScores(exprData)
```

## **Arguments**

exprData

A data frame with first column as row IDs and others are numeric values.

#### Value

A data frame with z scores for each row. The first column is still row IDs.

#### Author(s)

Henry Zhang

# **Examples**

```
data(RNASeqDemoData)
exprZ <- convertToZScores(RNASeqDemoData)</pre>
```

drawBioNetNodeBackground

Draw Background for A Data Track On Nodes of caOmicsV bioNet Layout

# Description

Paint (with any color other than white) background for a circular track on each node on caOmicsV bioNetCircos layout. Graphic device and igraph object must exist.

#### Usage

```
drawBioNetNodeBackground(trackLocations, bgColor=gray(0.9, alpha=0.5))
```

## **Arguments**

```
trackLocations a list returned by getBioNetPlotLocations() function.

bgColor character vector or R color specification for background color
```

# Value

None

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#### Author(s)

Henry Zhang

## **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)
initializeBioNetCircos(bioNet)

bioNetGraph <- getBioNetGraph()
outer <- 1.5
inner <- 1.0

nodeCenter <- as.numeric(bioNetGraph$layout[1,])
plotLocations <- getBioNetPlotLocations(nodeCenter, outer, inner)
showBioNetNodesLayout()
eraseBioNetNode()
drawBioNetNodeBackground(plotLocations)</pre>
```

eraseBioNetNode

Erase Background of All Nodes on caOmicsV bioNetCircos Layout

## **Description**

Erase all nodes on a caOmicsV bioNetCircos layout except of the edges. Graphic device and igraph object must be initialized first.

## Usage

```
eraseBioNetNode()
```

#### Value

None

## Author(s)

Henry Zhang

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)
initializeBioNetCircos(bioNet)

showBioNetNodesLayout()
eraseBioNetNode()</pre>
```

getBezierCurve 15

getBezierCurve	Calculate x and y Coordinates for A Quandratic Bezier Curve
----------------	---

#### **Description**

Calculate x and y coordinates for a quandratic Bezier curve between two points with the equation: B(t) = (1-t)((1-t)P0 + tP1) + t((1-t)P1 + tP2) where P0 is the start point, P2 is the end point, and P1 is the control point. P1 will be adjusted based on the distance of two points.

## Usage

```
getBezierCurve(lineStart, lineEnd, totalPoints)
```

## **Arguments**

lineStart numeric vector, the coordinate of a point where Bezier line starts

lineEnd numeric vector, the coordinate of a point where Bezier line ends

totalPoints non-negative numeric, total number of points that form a Bezier line

#### Value

posX x coordinates of points that form Bezier line posY y coordinates of points that form Bezier line

#### Author(s)

Henry Zhang

# **Examples**

```
lineStart <- c(0, 1)
lineEnd <- c(1, 0)
totalPoints <- 2000
the_line <- getBezierCurve(lineStart, lineEnd, totalPoints)</pre>
```

```
getBioMatrixDataRowTop
```

Get y Coordinate for Top of A Row on bioMatrix Layout

#### **Description**

Caluclate the y coordinate of a row top on bioMatrix layout. The bioMatrix layout must be initialized first

## Usage

```
getBioMatrixDataRowTop(rowNumber, areaName=c("omicsData", "phenotype"))
```

#### **Arguments**

rowNumber non-negative integer, number of the row

areaName character vector, either "phenotype" or "omicsdata"

#### Value

non-negative numeric, the y coordinate of the row top.

#### Author(s)

Henry Zhang

#### **Examples**

```
initializeBioMatrixPlot()
yTop <- getBioMatrixDataRowTop(2, areaName="omicsData")</pre>
```

getBioMatrixParameters

Methods to Get caOmicsV BioMatrix Plot Parameters

## **Description**

Get methods to retrieve parameters for caOmicsV bioMatrix layout plot stored in caOmicsV environment. bioMatrix layout must be initialized first.

# Usage

```
getBioMatrixBasePositions()
getBioMatrixColumnPadding()
getBioMatrixDataAreaWidth()
getBioMatrixGeneLabelWidth()
getBioMatrixGeneNumber()
getBioMatrixLegendHeight()
getBioMatrixPhenotypeNumber()
getBioMatrixPlotAreaHeigth()
getBioMatrixRemarkWidth()
getBioMatrixRowPadding()
getBioMatrixSampleHeight()
getBioMatrixSampleIDHeight()
getBioMatrixSampleNumber()
getBioMatrixSampleNumber()
getBioMatrixSampleWidth()
```

## Value

getBioMatrixBasePositions() returns a numeric matrix of defult x and y coordinates of rectangles for each samples at a row.

getBioMatrixColumnPadding() returns a non-negative numeric value in inch for padding between two samples, default 0.025.

getBioMatrixDataAreaWidth() returns a non-negative numeric value in inch for width of data plot area, total samples times the sum of sample width and columnPadding.

getBioMatrixDataRowTop() returns y coordinate for the top of a sample row.

getBioMatrixGeneLabelWidth() returns a non-negative numeric value in inch for length of left labels (gene names).

getBioMatrixGeneNumber() returns total number of genes to be plotted.

getBioMatrixLegendHeight() returns a non-negative numeric value in inch for the height of legend area.

getBioMatrixPhenotypeNumber() returns total number of phenotypes.

getBioMatrixPlotAreaHeigth() returns a non-negative numeric value in inch height of all plot areas (sample name area, data plot area, and legend area).

getBioMatrixPlotAreaWidth() returns a non-negative numeric value in inch for width of all plot areas (left labels (gene names), data plot area, and right labels).

getBioMatrixRemarkWidth() returns a non-negative numeric value in inch for width on the right side of data plot, usually for second gene labels or summary data plot.

getBioMatrixRowPadding() returns a non-negative numeric value in inch for height of padding between two rows.

getBioMatrixSampleHeight() returns a non-negative numeric value in inch for height of a sample row.

getBioMatrixSampleIDHeight() returns a non-negative numeric value in inch for height of smaple labels (on the top of phenotype plot area).

getBioMatrixSampleNumber() returns the total number of samples in plot datasets.

getBioMatrixSampleWidth() returns a non-negative numeric value in inch for width of a rectangle (sample).

## Author(s)

Henry Zhang

#### **Examples**

#### initializeBioMatrixPlot()

```
<- getBioMatrixBasePositions()</pre>
positions
colPadding
             <- getBioMatrixColumnPadding()</pre>
dataAreaWidth <- getBioMatrixDataAreaWidth()</pre>
geneNameWidth <- getBioMatrixGeneLabelWidth()</pre>
numOfGenes
               <- getBioMatrixGeneNumber()</pre>
legendHeight <- getBioMatrixLegendHeight()</pre>
numOfFeatures <- getBioMatrixPhenotypeNumber()</pre>
dataAreaHeight <- getBioMatrixPlotAreaHeigth()</pre>
plotAreaWidth <- getBioMatrixPlotAreaWidth()</pre>
sumAreaWidth <- getBioMatrixRemarkWidth()</pre>
rowPadding <- getBioMatrixRowPadding()</pre>
sampleHeight <- getBioMatrixSampleHeight()</pre>
sampleIDHeight <- getBioMatrixSampleIDHeight()</pre>
numOfSamples <- getBioMatrixSampleNumber()</pre>
sampleWidth
               <- getBioMatrixSampleWidth()</pre>
```

getBioNetNodeLinkLine Get X and Y Coordinates for An Anrrow between Two Nodes

# Description

Caluculate x and y coordinates for an customized arrow head and tail with defined length to connect two nodes.

# Usage

```
getBioNetNodeLinkLine(lineX, lineY, arrowSize=1, lineLength)
```

## **Arguments**

lineX numeric vector, x coordinates of the link line
numeric vector, y coordinates of the link line

arrowSize non-negative numeric, scaling factor for arrow size, default 1

lineLength non negative integer, the length of link line

#### **Details**

An arrow is drawn as an polygon. By default, the arrow is in inside of a circle (radius 1) without tail and it points to radian 0. The tail, if any, will be added to the left.

#### Value

A two dimensional numeric matrix for x and y coordinates of the arrow.

#### Author(s)

Henry Zhang

```
from <- c(1, 1)
to <- c(2, 2)
lineX <- seq(from[1], to[1], 1000)
lineY <- seq(from[2], to[2], 1000)

lineLength <- sqrt((from[1]-to[1])^2 + (from[2]-to[2])^2)
positions <- getBioNetNodeLinkLine(lineX, lineY, arrowSize=1, lineLength)</pre>
```

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getBioNetParameters

Methods to Get caOmicsV BioNetCircos Plot Parameters

#### **Description**

Methods for retrieving plot parameters of caOmicsV bioNetCircos layout stored in caOmicsV environment. The bioNetCircos layout must be initialized first.

## Usage

```
getBioNetBasePositions()
getBioNetGraph()
getBioNetNodePaddingScale()
getBioNetNodeParameters()
getBioNetNodePlotAreaBoundary()
getBioNetNodeRadius()
getBioNetPlotAreaWidth()
getBioNetPlotSampleWidth()
getBioNetPlotTotalSample()
```

#### Value

getBioNetBasePositions() returns numeric matrix containing x and y coordinates of points on a circular line with radius of 1 and degrees of text rotation on each point.

getBioNetGraph() returns an igraph object which representing the biological network built with user's inputs.

getBioNetNodePaddingScale() returns a numeric value for padding between two nodes.

getBioNetNodeParameters() returns a list containing totalSamples, sampleWidth, nodeRadius, nodePadding, plotAreaWidth, inner, and outer boundary of plot area.

 $getBioNetNodePlotAreaBoundary()\ returns\ a\ numeric\ vector\ for\ outer\ and\ inner\ boundary\ of\ a\ node\ on\ caOmicsV\ bioNetCircos\ layout.$ 

getBioNetNodeRadius() returns the numeric value for radius of a node on caOmicsV bioNetCircos layout.

getBioNetPlotAreaWidth() returns the width of bioNetCircos layout plot area.

getBioNetPlotSampleWidth() returns total points a sample will need on a circular track.

getBioNetPlotTotalSample() returns the total number of samples to be plotted on each node.

# Author(s)

Henry Zhang

```
data(caOmicsV.bionet.eset)
eSet <- caOmicsV.bionet.eset
expr <- eSet$heatmapData[[1]]
bioNet <- bc3net(expr)
initializeBioNetCircos(bioNet)</pre>
```

```
getBioNetPlotLocations
```

Get bioNetCircos Plot Locations

## **Description**

Get plot loactions for a node on caOmicsC bioNetCircos layout

#### Usage

```
getBioNetPlotLocations(nodeCenter, outer, inner)
```

## Arguments

nodeCenter numeric, x and y coordinates of the node center

outer non-negative numeric, outer limit of plot track relative to node center non-negative numeric, inner limit of plot track relative to ode center

#### Value

nodeCenter numeric, x and y coordinates of the node center

outPositions two dimensional numeric matrix for x and y coordinates of outer boundary for

plot

plot

 $\verb"positionIndex"$ 

matrix with index of x and y coordinates of base plot position for each sample

#### Author(s)

Henry Zhang

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)
initializeBioNetCircos(bioNet)
bioNetGraph <- getBioNetGraph()
outer <- 1.5</pre>
```

```
inner <- 1.0

nodeCenter <- as.numeric(bioNetGraph$layout[1,])
plotLocations <- getBioNetPlotLocations(nodeCenter, outer, inner)</pre>
```

getBioNetSamplePlotPosition

Calculate x and y Coordinates for Each Sample on Default Node

## **Description**

Calculate x and y coordinates for each sample on default node. The output will be a three column matrix representing the left, center, and right position for each sample on circumference of default node. The center positions are for points plot and others are for polygon plot. bionetCircos layout must be initialized first.

#### Usage

```
getBioNetSamplePlotPosition(totalSamples)
```

#### **Arguments**

totalSamples non-negative integer, total numbe of samples to be plotted

#### Value

matrix with index of x and y coordinates for each sample

#### Author(s)

Henry Zhang

## **Examples**

```
totalSamples <- 100
samplePositions <- getBioNetSamplePlotPosition(totalSamples)</pre>
```

getCaOmicsVColors

Get Default Colors Used by caOmicsV Plot

# Description

Display default colors used by caOmicsV package

#### **Usage**

```
getCaOmicsVColors()
```

## Value

A character vector of length 8 for R colors including of "red", "blue", "black", "green", "cyan", "brown", "magenta", and "gold".

22 getDefaultNaStrings

#### Author(s)

Henry Zhang

## **Examples**

```
defaultColors <- getCaOmicsVColors()</pre>
```

```
getCaOmicsVPlotTypes Get Plot Types Supported by caOmicsV Package
```

## **Description**

Retrieve the plot types supported by current version of caOmicsV package

## Usage

```
getCaOmicsVPlotTypes()
```

#### Value

```
Character vector including "polygon", "bar", "points", "heatmap", "line", "category", "binary"
```

## Author(s)

Henry Zhang

## **Examples**

```
plotType <- getCaOmicsVPlotTypes()</pre>
```

```
{\tt getDefaultNaStrings} \qquad \textit{Default NA String}
```

# **Description**

Get the default NS strings used by caOmicsV package

# Usage

```
getDefaultNaStrings()
```

## Value

Returns na strings used by caOmicsV package. Default is "NULL".

# Author(s)

Henry Zhang

```
naStr <- getDefaultNaStrings()</pre>
```

getESet 23

getESet	Prepare Data Set for caOmicsV Plot	
---------	------------------------------------	--

#### **Description**

This function will validate each dataset then convert them to matrix and wrap all of them in one list object.

#### Usage

```
getESet(sampleNames, geneNames, sampleData, heatmapData=list(),
    categoryData=list(), binaryData=list(), summaryData=list(),
    secondGeneNames=NULL)
```

#### **Arguments**

sampleNames character vector, sample names, must be same or exist in every data set. character vector, gene names, must be same or exist in every data set. geneNames sampleData data frame with rows for samples and columns for features. heatmapData list of data frame(s) for heatmap plot. The first column of each data frame is row names and others are numeric values. The list could be empty, or having one or more data frame in a list object. Heatmap data should be log2 values or z-scores. list of data frame(s). The first column of each data frame is row names and categoryData others are numeric values. The list could be empty, or having one or more data frame in a list object binaryData list of data frame(s). The first column of each data frame is row names and others are binary values. The listcould be none, or one or more data frame in a list object summaryData list of data frames with summary information for samples (columns) or for genes (rows). The first column is for ID following by one or more columns of summary secondGeneNames

character vector, gene names that will be plot on right side of biomatrix plot

# Value

layout

sampleNames character verctor, sample names geneNames character verctor, gene names secondGeneNames character verctor, for example, miRNA names sampleInfo a data frame, sample information such as Tumor/Normal, age, diagnosis heatmapData list of data matrix(s), e.g., RNASeq read counts at gene level categoryData list of data matrix(s), such as SNP in a gene, homozygous, or heterozygous, or wildtype binaryData list of data matrix(s), e.g., mutation status of the gene list of data matrix(s), such as percentage of highly expressed miRNA in all samsummaryInfo ples

#### Author(s)

Henry Zhang

#### **Examples**

```
data(sampleDemoData)
data(RNA2miRNA)
data(RNASeqDemoData)
data(miRNADemoData)
data(methylDemoData)
data(CNVDemoData)
sampleNames <- as.character(sampleDemoData[,1])</pre>
geneNames <- as.character(RNA2miRNA[,1])</pre>
secondGeneNames <- as.character(RNA2miRNA[,2])</pre>
normals <- grep("Normal", colnames(RNASeqDemoData))</pre>
tumors <- grep("Tumor", colnames(RNASeqDemoData))</pre>
tumorExpr <- RNASeqDemoData[, tumors]</pre>
normalExpr <- RNASeqDemoData[, normals]</pre>
meanLog2Fold <- log2(rowMeans(tumorExpr/normalExpr))</pre>
summaryData <- data.frame(geneNames, meanLog2Fold)</pre>
eSet <- getESet(sampleNames, geneNames, sampleDemoData,</pre>
                 heatmapData=list(RNASeqDemoData, miRNADemoData),
                 categoryData=list(methylDemoData),
                 binaryData=list(CNVDemoData),
                 summaryData=list(summaryData),
                 secondGeneNames)
```

getHeatmapColorScales Get caOmicsV Heatmap Color Scales

#### **Description**

Generate a color map for heatmap color scales

#### Usage

```
getHeatmapColorScales(colorType)
```

## **Arguments**

```
colorType character vector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed", "GreenBlackRed", "YellowToRed", and "Black".
```

#### Value

An RGB color matrix with demension of 255 by 2.

#### Author(s)

Henry Zhang

getPlotOmicsData 25

## **Examples**

```
colorMap <- getHeatmapColorScales("BlueWhiteRed")</pre>
```

getPlotOmicsData

Extract Subset from A Data Frame

## **Description**

Extract a subset from the imput data for a set of samples and genes

# Usage

```
getPlotOmicsData(omicsData, sampleNames, geneNames)
```

## **Arguments**

omicsData a data frame with all samples and all genes

sampleNames character vector, names of samples to be extracted from dataset.

geneNames Character vector, names of genes to be extracted from dataset.

## Value

A data frame with subset of input data

#### Author(s)

Henry Zhang

## **Examples**

```
data(RNASeq)
data(RNASeqDemoData)
geneNames <- as.character(RNASeqDemoData[,1])
sampleNames <- colnames(RNASeqDemoData)[-1]
plotData <- getPlotOmicsData(RNASeq, sampleNames, geneNames)</pre>
```

getPlotSampleData

Extract Subset of Sample Information

# Description

Extract required rows and columns from a sample dataset

# Usage

```
getPlotSampleData(sampleData, sampleNames)
```

#### **Arguments**

sampleData Data frame with rows for samples and columns for features. Column 1 must be

sample names.

sampleNames character vector, names of samples to select.

#### Value

A data frame with subset of sample data and with the row order same as sampleNames.

#### Author(s)

Henry Zhang

## **Examples**

```
data(sampleDemoData)
sampleNames <- as.character(sampleDemoData[10:40,1])
sampleInfo <- getPlotSampleData(sampleDemoData, sampleNames)</pre>
```

getPlotSummaryData

Extract Summary Subset for Plotting

## **Description**

Extract required rows and columns from a summary data set.

#### Usage

```
getPlotSummaryData(summaryData, sampleNames=NULL, geneNames=NULL)
```

#### **Arguments**

summaryData a data frame with summary data for each gene (rows are for genes and columns

are summary values) or for each sample (rows are summary values and columns

are sample names)

sampleNames character vector, names of samples/columns to be extracted. Set to NULL when

summary data is for genes.

geneNames character vector, names of genes to be extracted. Set to NULL when summary

data is for samples

#### Value

A data frame with subset of input data and with the same orders as geneNames and sampleNames.

#### Author(s)

Henry Zhang

getRelatedPlotData 27

#### **Examples**

```
data(RNASeqDemoData)
exprValue <- as.matrix(RNASeqDemoData[, 2:ncol(RNASeqDemoData)])
summaryData <- data.frame(as.character(RNASeqDemoData[,1]),
    rowMeans(exprValue), log2(rowMeans(exprValue)))
geneSymbols <- c("ECM1", "SLC26A6", "ADAMTS13", "FCN3", "CFP")
sumByGene <- getPlotSummaryData(summaryData, geneNames=geneSymbols)
summaryData <- rbind(colMeans(exprValue), log2(colMeans(exprValue)))
summaryData <- data.frame(c("sampleMean", "log2mean"), summaryData)
colnames(summaryData) <- colnames(exprValue)
sampleID <- colnames(exprValue)[c(2:21, 32:41)]
sumBySample <- getPlotSummaryData(summaryData, sampleID)</pre>
```

getRelatedPlotData

Extract subset from A Data Frame Based on Relational Information

## **Description**

Extract a subset of plot data based on relational information, e.g., expression of miRNA or DNA copy number variationa that are corelated to differentially expressed genes

#### Usage

```
getRelatedPlotData(omicsData, linkData, geneNames)
```

#### **Arguments**

omicsData a data frame, the dataset from which subset is extracted

linkData a data frame, usually gene names and their related items. The first column must

be the items to which the second item is linked to.

geneNames character vector, subset of gene names for subset.

#### Value

A data frame with subset of the first input data.

#### Author(s)

Henry Zhang

```
data(miRNA)
data(RNA2miRNA)
geneNames <- as.character(RNA2miRNA[,1])
miRNAexpr <- getRelatedPlotData(miRNA, RNA2miRNA, geneNames)</pre>
```

28 initializeBioMatrixPlot

```
initializeBioMatrixPlot
```

Set Up Parameters for caOmicsV bioMatrix Plot Layout

#### **Description**

Initialize parameters for caOmicsV bioMatrix layout to set up plot area size and item sizes.

#### Usage

```
initializeBioMatrixPlot(numOfGenes=100, numOfSamples=100,
    numOfPhenotypes=1, sampleHeight=0.4, sampleWidth=0.1,
    columnPadding=0.025, rowPadding=0.1, geneNameWidth=1,
    remarkWidth=1, sampleNameHeight=1, legendHeight=1)
```

#### **Arguments**

numOfGenes non-negative numeric, total number of genes to be plotted numOfSamples non-negative numeric, total number of samples to be plotted numOfPhenotypes non-negative numeric, total number of phenotypes to be plotted non-negative numeric, height of rectangle area in inch for a sample plot sampleHeight non-negative numeric, width of rectangle area in inch for a sample plot sampleWidth columnPadding non-negative numeric, width of padding in inch between two rectangles (samples) non-negative numeric, height of padding in inch between two rows (genes) rowPadding  ${\tt geneNameWidth}$ non-negative numeric, width of plot area in inch for gene name plot non-negative numeric, width of plot area in inch for second set of gene names remarkWidth and summary data. sampleNameHeight non-negative numeric, height of plot area in inch for sample names (sample name are plotted vertically).

non-negative numeric, height of plot area in inch for legend.

#### Value

None

#### Author(s)

Henry Zhang

legendHeight

```
initializeBioMatrixPlot(numOfGenes=100, numOfSamples=100,
    numOfPhenotypes=1, sampleHeight=0.4, sampleWidth=0.1,
    columnPadding=0.025, rowPadding=0.1, geneNameWidth=1,
    remarkWidth=1, sampleNameHeight=1, legendHeight=1)
```

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#### initialize Bio Net Circos

Initialize caOmicsV bioNetCircos Layout Plot

## **Description**

Set up parameters for layout of caOmicsV bioNetCircos plot including total number of samples, default width of a sample on a circos track, node radius, padding width between two nodes, plot area of each node, default x and y coordinates of points on a circle with radius of 1, and node layout.

## Usage

## **Arguments**

bioNet	an igraph object
totalSamples	non-negative numeric, total number of samples to be plotted
sampleWidth	non-negative numeric, total number of points to represent a sample on a circular track.
nodeRadius	non-negative numeric, radius of a node on biological network.
nodePadding	non-negative numeric, padding width between two node on biological network.
plotAreaWidth	non-negative numeric, outside boundary of plot area of a node, relative to node radius, default 1 (same as nodeRadius)
layout	two dimentioanl numeric matrix, node layout of the igraph returned by igraph layout method

#### Value

None

#### Author(s)

Henry Zhang

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=100, sampleWidth=100, nodeRadius=1, nodePadding=1, plotAreaWidth=1, layout=layout.fruchterman.reingold(bioNet))</pre>
```

30 labelBioNetNodeNames

labelBioNetNodeNames Label Names for Each Node on Network Graph

## **Description**

Plot name for one or more nodes on an igraph network. igraph object and graphic device must be initialized first.

## Usage

## **Arguments**

nodeList non-negatice integer, index of node(s) on a bioNet layout

labelColor character vector, colord for text (labels)

labelLocation character vector, location relative to node center, either "bottom", "left", "top",

or "right"

labelOffset non-negative numeric, distance from node outside boundary

## Value

None

## Author(s)

Henry Zhang

linkBioNetNodes 31

linkBioNetNodes	Draw A Customized Arrow between Two Nodes		
-----------------	---	--	--

# Description

Plot a customized arrow between two nodes to replace the edge. bioNetCorcos layout must be initialized first.

# Usage

```
linkBioNetNodes(fromNode, toNode, lineColor = "black", arrowSize = 1)
```

## **Arguments**

fromNode non negative integer, the start node to be linked toNode non negative integer, the end node to be linked

lineColor character vector, color of the arrow

arrowSize non-negative numeric, scaling factor for arrow size, default 1

#### Value

None

#### Author(s)

Henry Zhang

# Examples

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)
initializeBioNetCircos(bioNet)
showBioNetNodesLayout()
linkBioNetNodes(fromNode=2, toNode=5, lineColor="red", arrowSize=1)</pre>
```

linkBioNetSamples

Link Two samples Inside a Node with Quandratic Bezier Curve

## **Description**

Draw a quandratic Bezier curve line between two samples inside of a node. bioNetCorcos layout must be initialized first.

# Usage

```
linkBioNetSamples(nodeIndex, fromSample, toSample, outer, plotColors)
```

32 methylDemoData

#### **Arguments**

nodeIndex non-negative integer, the node on which link line is drawn

fromSample non-negative integer, the first sample to be linked toSample non-negative integer, the second sample to be linked

outer non-negative numeric, the start and end of link line relative to node center

plotColors character vector, color for the link line (ribbon)

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

methylDemoData

Methylation Demo Data Set

## **Description**

A data frame with bea values of 26 genes in 60 samples. Used for bar plot on caOmicsV bioNet-Circos layout demo, and for categoray plot on caOmicsV bioNet layout demo after transforming to category data.

## Usage

```
data("methylDemoData")
```

#### **Format**

A data frame with 26 observations on the following 61 variables.

Gene\_Symbol a factor with levels ACTN1 ADAMTS13 AMIGO3 ATP2A1 BCO2 CDKN3 CFP CNBP COL15A1 CSRNP1 CXCL12 DBH DDX55 ECM1 ELOVL1 ESM1 FAM81A FCN3 KCNQ1 LEPREL1 LIFR LILRA6 LILRB5 LOC222699 LOC283050 LRRC16A LYVE1 MAEL MAN1B1 MRPS25 MT1F NIPA2 NPHP4 NR5A2 OLFML3 PLVAP PROX1 PTH1R RBL2 RCAN1 RND3 SEMA3F SLC26A6 TOMM40L TSEN34 VDAC3

```
TCGA.BC.A216.Normal a numeric vector
```

TCGA.BD.A2L6.Normal a numeric vector

methylDemoData 33

```
TCGA.BD.A3EP.Normal a numeric vector
TCGA.DD.A113.Normal a numeric vector
TCGA.DD.A114.Normal a numeric vector
TCGA.DD.A118.Normal a numeric vector
TCGA.DD.A119.Normal a numeric vector
TCGA.DD.A11A.Normal a numeric vector
TCGA.DD.A11B.Normal a numeric vector
TCGA.DD.A11C.Normal a numeric vector
TCGA.DD.A11D.Normal a numeric vector
TCGA.DD.A1EB.Normal a numeric vector
TCGA.DD.A1EC.Normal a numeric vector
TCGA.DD.A1EG.Normal a numeric vector
TCGA.DD.A1EH.Normal a numeric vector
TCGA.DD.A1EI.Normal a numeric vector
TCGA.DD.A1EJ.Normal a numeric vector
TCGA.DD.A1EL.Normal a numeric vector
TCGA.DD.A39V.Normal a numeric vector
TCGA.DD.A39W.Normal a numeric vector
TCGA.DD.A39X.Normal a numeric vector
TCGA.DD.A39Z.Normal a numeric vector
TCGA.DD.A3A1.Normal a numeric vector
TCGA.DD.A3A2.Normal a numeric vector
TCGA.DD.A3A3.Normal a numeric vector
TCGA.EP.A12J.Normal a numeric vector
TCGA.EP.A26S.Normal a numeric vector
TCGA.ES.A2HT.Normal a numeric vector
TCGA.FV.A23B.Normal a numeric vector
TCGA.FV.A2QR.Normal a numeric vector
TCGA.BC.A216.Tumor a numeric vector
TCGA.BD.A2L6.Tumor a numeric vector
TCGA.BD.A3EP.Tumor a numeric vector
TCGA.DD.A113.Tumor a numeric vector
TCGA.DD.A114.Tumor a numeric vector
TCGA.DD.A118.Tumor a numeric vector
TCGA.DD.A119.Tumor a numeric vector
TCGA.DD.A11A.Tumor a numeric vector
TCGA.DD.A11B.Tumor a numeric vector
TCGA.DD.A11C.Tumor a numeric vector
TCGA.DD.A11D.Tumor a numeric vector
TCGA.DD.A1EB.Tumor a numeric vector
```

miRNA miRNA

```
TCGA.DD.A1EC.Tumor a numeric vector
TCGA.DD.A1EG.Tumor a numeric vector
TCGA.DD.A1EH.Tumor a numeric vector
TCGA.DD.A1EI.Tumor a numeric vector
TCGA.DD.A1EJ.Tumor a numeric vector
TCGA.DD.A1EL.Tumor a numeric vector
TCGA.DD.A39V.Tumor a numeric vector
TCGA.DD.A39W.Tumor a numeric vector
TCGA.DD.A39X.Tumor a numeric vector
TCGA.DD.A39Z.Tumor a numeric vector
TCGA.DD.A3A1.Tumor a numeric vector
TCGA.DD.A3A2.Tumor a numeric vector
TCGA.DD.A3A3.Tumor a numeric vector
TCGA.EP.A12J.Tumor a numeric vector
TCGA.EP.A26S.Tumor a numeric vector
TCGA.ES.A2HT.Tumor a numeric vector
TCGA.FV.A23B.Tumor a numeric vector
TCGA.FV.A2QR.Tumor a numeric vector
```

#### **Examples**

data(methylDemoData)

miRNA

Selected miRNA Read Counts

# Description

A data frame with read counts of 14 miRNA in 86 samples.

## Usage

```
data("miRNA")
```

#### **Format**

A data frame with 14 observations on the following 86 variables.

TCGA.BC.A112.Tumor a numeric vector

```
miRNA_ID a factor with levels hsa.mir.101.1 hsa.mir.101.2 hsa.mir.10b hsa.mir.1180 hsa.mir.125b.2 hsa.mir.139 hsa.mir.142 hsa.mir.151 hsa.mir.183 hsa.mir.22 hsa.mir.25 hsa.mir.424 hsa.mir.450b hsa.mir.93

TCGA.2Y.A9H1.Tumor a numeric vector

TCGA.BC.4073.Tumor a numeric vector

TCGA.BC.A10W.Tumor a numeric vector
```

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```
TCGA.BC.A216.Tumor a numeric vector
TCGA.BC.A216.Normal a numeric vector
TCGA.BD.A2L6.Tumor a numeric vector
TCGA.BD.A2L6.Normal a numeric vector
TCGA.BD.A3EP.Tumor a numeric vector
TCGA.BD.A3EP.Normal a numeric vector
TCGA.BW.A5NP.Tumor a numeric vector
TCGA.CC.A1HT.Tumor a numeric vector
TCGA.CC.A7IG.Tumor a numeric vector
TCGA.CC.A9FS.Tumor a numeric vector
TCGA.DD.A113.Tumor a numeric vector
TCGA.DD.A113.Normal a numeric vector
TCGA.DD.A114.Tumor a numeric vector
TCGA.DD.A114.Normal a numeric vector
TCGA.DD.A118.Tumor a numeric vector
TCGA.DD.A118.Normal a numeric vector
TCGA.DD.A119.Tumor a numeric vector
TCGA.DD.A119.Normal a numeric vector
TCGA.DD.A11A.Tumor a numeric vector
TCGA.DD.A11A.Normal a numeric vector
TCGA.DD.A11B.Tumor a numeric vector
TCGA.DD.A11B.Normal a numeric vector
TCGA.DD.A11C.Tumor a numeric vector
TCGA.DD.A11C.Normal a numeric vector
TCGA.DD.A11D.Tumor a numeric vector
TCGA.DD.A11D.Normal a numeric vector
TCGA.DD.A1EB.Tumor a numeric vector
TCGA.DD.A1EB.Normal a numeric vector
TCGA.DD.A1EC.Tumor a numeric vector
TCGA.DD.A1EC.Normal a numeric vector
TCGA.DD.A1EE.Normal a numeric vector
TCGA.DD.A1EG.Tumor a numeric vector
TCGA.DD.A1EG.Normal a numeric vector
TCGA.DD.A1EH.Tumor a numeric vector
TCGA.DD.A1EH.Normal a numeric vector
TCGA.DD.A1EI.Tumor a numeric vector
TCGA.DD.A1EI.Normal a numeric vector
TCGA.DD.A1EJ.Tumor a numeric vector
TCGA.DD.A1EJ.Normal a numeric vector
TCGA.DD.A1EL.Tumor a numeric vector
```

36 miRNA

TCGA.DD.A1EL.Normal a numeric vector TCGA.DD.A39V.Tumor a numeric vector TCGA.DD.A39V.Normal a numeric vector TCGA.DD.A39W.Tumor a numeric vector TCGA.DD.A39W.Normal a numeric vector TCGA.DD.A39X.Tumor a numeric vector TCGA.DD.A39X.Normal a numeric vector TCGA.DD.A39Z.Tumor a numeric vector TCGA.DD.A39Z.Normal a numeric vector TCGA.DD.A3A1.Tumor a numeric vector TCGA.DD.A3A1.Normal a numeric vector TCGA.DD.A3A2.Tumor a numeric vector TCGA.DD.A3A2.Normal a numeric vector TCGA.DD.A3A3.Tumor a numeric vector TCGA.DD.A3A3.Normal a numeric vector TCGA.DD.A3A6.Normal a numeric vector TCGA.DD.A4NG.Tumor a numeric vector TCGA.DD.A4NV.Tumor a numeric vector TCGA.ED.A4XI.Tumor a numeric vector TCGA.ED.A82E.Tumor a numeric vector TCGA.EP.A12J.Tumor a numeric vector TCGA.EP.A12J.Normal a numeric vector TCGA.EP.A26S.Tumor a numeric vector TCGA.EP.A26S.Normal a numeric vector TCGA.ES.A2HS.Tumor a numeric vector TCGA.ES.A2HT.Tumor a numeric vector TCGA.ES.A2HT.Normal a numeric vector TCGA.FV.A23B.Tumor a numeric vector TCGA.FV.A23B.Normal a numeric vector TCGA.FV.A2QR.Tumor a numeric vector TCGA.FV.A2QR.Normal a numeric vector TCGA.FV.A4ZQ.Tumor a numeric vector TCGA.G3.A3CH.Tumor a numeric vector TCGA.G3.A6UC.Tumor a numeric vector TCGA.G3.AAV3.Tumor a numeric vector TCGA.K7.A5RF.Tumor a numeric vector TCGA.LG.A9QD.Tumor a numeric vector TCGA.08.A75V.Tumor a numeric vector TCGA.RC.A7SH.Tumor a numeric vector TCGA.UB.AA0U.Tumor a numeric vector TCGA.ZP.A9CZ.Tumor a numeric vector

#### **Examples**

data(miRNA)

miRNADemoData 37

miRNADemoData

miRNASeq Demo Data Set

## **Description**

A data frame with miRNA read counts for 26 genes in 60 samples. Used for heatmap plot demo on both caOmicsV bioNetCircos layout and bioMatrix layout after transformed to log2 values.

#### Usage

```
data("miRNADemoData")
```

#### **Format**

A data frame with 26 observations on the following 61 variables.

```
miRNA_ID a character vector
```

TCGA.BC.A216.Normal a numeric vector

TCGA.BD.A2L6.Normal a numeric vector

TCGA.BD.A3EP.Normal a numeric vector

TCGA.DD.A113.Normal a numeric vector

TCGA.DD.A114.Normal a numeric vector

TCGA.DD.A118.Normal a numeric vector

TCGA.DD.A119.Normal a numeric vector

TCGA.DD.A11A.Normal a numeric vector

TCGA.DD.A11B.Normal a numeric vector

TCGA.DD.A11C.Normal a numeric vector

TCGA.DD.A11D.Normal a numeric vector

TCGA.DD.A1EB.Normal a numeric vector

TCGA.DD.A1EC.Normal a numeric vector

TCGA.DD.A1EG.Normal a numeric vector

TCGA.DD.A1EH.Normal a numeric vector

TCGA.DD.A1EI.Normal a numeric vector

TCGA.DD.A1EJ.Normal a numeric vector

TCGA.DD.A1EL.Normal a numeric vector

TCGA.DD.A39V.Normal a numeric vector

TCGA.DD.A39W.Normal a numeric vector

TCGA.DD.A39X.Normal a numeric vector

TCGA.DD.A39Z.Normal a numeric vector

TCGA.DD.A3A1.Normal a numeric vector

TCGA.DD.A3A2.Normal a numeric vector

TCGA.DD.A3A3.Normal a numeric vector

TCGA.EP.A12J.Normal a numeric vector

38 miRNADemoData

```
TCGA.EP.A26S.Normal a numeric vector
TCGA.ES.A2HT.Normal a numeric vector
TCGA.FV.A23B.Normal a numeric vector
TCGA.FV.A2QR.Normal a numeric vector
TCGA.BC.A216.Tumor a numeric vector
TCGA.BD.A2L6.Tumor a numeric vector
TCGA.BD.A3EP.Tumor a numeric vector
TCGA.DD.A113.Tumor a numeric vector
TCGA.DD.A114.Tumor a numeric vector
TCGA.DD.A118.Tumor a numeric vector
TCGA.DD.A119.Tumor a numeric vector
TCGA.DD.A11A.Tumor a numeric vector
TCGA.DD.A11B.Tumor a numeric vector
TCGA.DD.A11C.Tumor a numeric vector
TCGA.DD.A11D.Tumor a numeric vector
TCGA.DD.A1EB.Tumor a numeric vector
TCGA.DD.A1EC.Tumor a numeric vector
TCGA.DD.A1EG.Tumor a numeric vector
TCGA.DD.A1EH.Tumor a numeric vector
TCGA.DD.A1EI.Tumor a numeric vector
TCGA.DD.A1EJ.Tumor a numeric vector
TCGA.DD.A1EL.Tumor a numeric vector
TCGA.DD.A39V.Tumor a numeric vector
TCGA.DD.A39W.Tumor a numeric vector
TCGA.DD.A39X.Tumor a numeric vector
TCGA.DD.A39Z.Tumor a numeric vector
TCGA.DD.A3A1.Tumor a numeric vector
TCGA.DD.A3A2.Tumor a numeric vector
TCGA.DD.A3A3.Tumor a numeric vector
TCGA.EP.A12J.Tumor a numeric vector
TCGA.EP.A26S.Tumor a numeric vector
TCGA.ES.A2HT.Tumor a numeric vector
TCGA.FV.A23B.Tumor a numeric vector
TCGA.FV.A2QR.Tumor a numeric vector
```

## **Examples**

data(miRNADemoData)

plotBioMatrix 39

## **Description**

A sample way to plot sample information and genomic data with default settings.

#### Usage

```
plotBioMatrix(eSet, summaryType=c("text", "bar"), summarybyRow=TRUE,
    heatmapMax=NULL, heatmapMin=NULL, heatmapColor="BlueWhiteRed")
```

## **Arguments**

eSet an object, returned from method of getESet(), with all plot data
summaryType character vector, either "text" or "bar", for plot type of summary data
summarybyRow logic, if the summary data is for each row or for each column
numeric, maximum value for heatmap plot, set to NULL to use the maximum value in input data.
heatmapMin numeric, minimum value for heatmap plot, set to NULL to use the minimum value in input data.
heatmapColor character vector,one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed",

## Value

None

## Author(s)

Henry Zhang

## **Examples**

```
data(caOmicsV.biomatrix.eset)
plotBioMatrix(caOmicsV.biomatrix.eset, summaryType="text", summarybyRow=TRUE,
    heatmapMax=NULL, heatmapMin=NULL, heatmapColor="BlueWhiteRed")
```

"GreenBlackRed", or "YellowToRed"

 $plot {\tt BioMatrixBars} \qquad \textit{Bar Plot on caOmicsV bioMatrix Layout}$ 

# Description

Bar plot method for caOmicsV bioMatrix layout with non-negative numeric matrix or vector with values in range of  $0 \sim 1$ . bioMatrix layout and graphic device must be initialized first. must be initialized first.

#### Usage

#### **Arguments**

barData non-negative numeric matrix or vector with values in range of  $0 \sim 1$ 

barColor character vector for color name or R color specification

areaName character vector, name of plot area, currentlt use "omicsData" only

byRow logic, whether plot bars for each row or not

skipPlotRow non-negative integer, how many row(s) to be skipped from first row

skipPlotColumns

non-negative integer, how many row(s) to be skipped from first column

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
showBioMatrixPlotLayout("Gene", paste("Sample", 1:50), "Diagnosis")
barData <- matrix(c(rep(0.25, 15), rep(0.75, 20), rep(0.5, 15)), byrow=TRUE)
plotBioMatrixBars(barData, barColor="red")</pre>
```

plotBioMatrixBinaryData

Binary Data Plot on caOmicsV bioMatrix Layout

## **Description**

Plot binary data as points in the inside of each rectangle(sample). This function plot all rows on omics data area and only the positive samples will be shown with colored points. For one row plot, pass data as vector and supply correct skipPlotRow parameter to define where to plot. bioMatrix layout and graphic device must be initialized first.

#### Usage

#### **Arguments**

binaryData vector or matrix with values of 0 and 1 only character vector, either "omicsData" or "phenotype" areaName non-negative integer, same as pch, default 19 scatterTypenon-negative numric, same as cex scatterSize totalSubRow non-negative integer, how many sub-rows in a sample area subRowIndex non-negative integer, which subrow will be plotted sampleColor character vector for color name(s) or R color specification skipPlotRow non-negative integer, total rows on plot area that should be skipped, default 0

#### Value

None

#### Author(s)

Henry Zhang

## **Examples**

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
showBioMatrixPlotLayout("Gene", paste("Sample", 1:50), "Diagnosis")
binaryData <- matrix(c(rep(1, 15), rep(0, 20), rep(1, 15)), nrow=1)
plotBioMatrixBinaryData(binaryData, scatterType=16)</pre>
```

 ${\tt plotBioMatrixCategoryData}$ 

Plot Category Data on caOmicsV bioMatrix Layout

## **Description**

Draw rectangle outline for one or more row(s) of samples to represent categorical values. This function highlights all samples on each row. bioMatrix layout and graphic device must be initialized first.

#### Usage

```
plotBioMatrixCategoryData(categoryData, areaName=c("omicsData", "phenotype"),
    sampleColors=palette(), lineWidth=1, skipPlotRow=0)
```

## **Arguments**

categoryData	vector or matrix of categorical values, such as 'High', 'low', and 'No'
areaName	character vector, either "omicsdata" or "phenotype"
sampleColors	character vector for color names or vector of R color specification
lineWidth	non-negatice integer, graphic parameter for lwd (line width), default 1
skipPlotRow	non-negative integer, total rows on plot area that should be skipped when only one row to plot, default 0.

#### Value

None

#### Author(s)

Henry Zhang

# Examples

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
showBioMatrixPlotLayout("Gene", paste("Sample", 1:50), "Diagnosis")
categoryData <- matrix(c(rep(1, 15), rep(0, 20), rep(1, 15)), nrow=1)
plotBioMatrixCategoryData(categoryData, areaName="omicsData")</pre>
```

plotBioMatrixHeatmap Heatma

Heatmap Plot on caOmicsV bioMatrix Layout

## **Description**

Headmap plot method for caOmicsV bioMatrix layout. This function will plot all rows of the input data. bioMatrix layout and graphic device must be initialized first.

#### Usage

```
plotBioMatrixHeatmap(exprData, topAdjust=0, bottomAdjust=0, maxValue=NULL,
    minValue=NULL, heatmapColor="BlueWhiteRed", skipPlotRow=0)
```

#### **Arguments**

exprData numeric matrix (log2 values) with row names topAdjust non-negative numeric, height of top y coordinate should be reduced to show different layers, default 0 bottomAdjust non-negative numeric, height of bottom y coordinate should be reduced for a small rectangle, default 0 maxValue numeric, value for highest color in heatmap, set to NULL to use the maximum value in expression dataset minValue numeric, value for lowest color in heatmap, set to NULL to use the minimum value in expression dataset heatmapColor character vector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed", "GreenBlackRed", or "YellowToRed" skipPlotRow non-negative integer, total rows on plot area that should be skipped, default 0

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
showBioMatrixPlotLayout("Gene", paste("Sample", 1:50), "Diagnosis")

data(caOmicsV.biomatrix.eset)
exprData <- caOmicsV.biomatrix.eset$heatmapData[[1]]
exprData <- matrix(exprData[1, 1:50], nrow=1)
plotBioMatrixHeatmap(exprData, maxValue=3, minValue=-3)</pre>
```

plotBioMatrixRowNames Plot Row Names on caOmicsV bioMatrix Layout

# Description

Plot row names on the left or right side of biomatrix plot area. Penotype names, gene names, and remark notes are all plotted with this function. bioMatrix layout an dgraphic device must be initialized first.

#### Usage

#### **Arguments**

geneNames character vector, row names to be plotted

areaName character vector, either "omicsData" or "phenotype"

colors character vector of color names or R color specification

side character vector, either "left" or "right"

skipPlotRows non-negative integer, total rows on plot area that should be skipped from the first

row, default 0

skipPlotColumns

non-negative integer, columns (sampleWidth) will be skipped when plotting

items on remark area

#### Value

None

# Author(s)

Henry Zhang

#### **Examples**

```
initializeBioMatrixPlot(numOfGenes=20, numOfSamples=50)
geneNames <- paste0("gene_", 1:20)</pre>
miRNANames <- paste0("miRNA_", 1:20)</pre>
showBioMatrixPlotLayout(geneNames, paste("Sample", 1:50), "Diagnosis")
plotBioMatrixRowNames(miRNANames, areaName="omicsData", colors="blue",
    side="right")
```

plotBioMatrixSampleData

Plot Sample Data on caOmicsV bioMatrix Layout

## **Description**

Plot colored polygons on phenotyp area of caOmicsV bioMatrix layout to show sample information such as diagnosis, tissue type, ... bioMatrix layout and graphic device must be initialized first.

## Usage

```
plotBioMatrixSampleData(rowNumber, areaName, fillColor=NA, borderColor=NA,
    topAdjust=0, bottomAdjust=0)
```

## **Arguments**

rowNumber	non-negative integer, number of the row where the data to be ploted
areaName	chracter vector, either "phenotype" or "omicsdata"
fillColor	chracter vector of color names or vector of R color specification
borderColor	chracter vector or a R colors specification for boarder color
topAdjust	non-negative numeric, height that will be reduced from top
bottomAdjust	non-negative numeric, height that will be reduced from bottom

# Value

None

#### Author(s)

Henry Zhang

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
showBioMatrixPlotLayout("Gene", paste("Sample", 1:50), "Diagnosis")
sampleColor <- c(rep("blue", 25), rep("red", 25))</pre>
plotBioMatrixSampleData(rowNumber=1, sampleColor, areaName="phenotype")
```

```
plotBioMatrixSampleNames
```

Label Sample Names on the Top of caOmicsV bioMatrix Layout

## **Description**

Text plot on the top of phenotyp area to show sample names. bioMatrix layout and graphic device must be initialized first.

# Usage

```
plotBioMatrixSampleNames(sampleNames, sampleColors)
```

# **Arguments**

```
sampleNames character vector, sample names to be plotted sampleColors character vector or R color name(s) for text color(s)
```

## Value

None

#### Author(s)

Henry Zhang

#### **Examples**

plotBioNetBars

Bar Plot on caOmicsV bioNetCircos Layout

# Description

Bar plot method for caOmicsV bioNetCircos layout. This will plot one track of bars for every node. bioNetCircos layout and graphic device must be initialized first.

# Usage

```
plotBioNetBars(dataValues, outer, inner, plotColors)
```

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#### **Arguments**

dataValues numeric matrix with range of  $0 \sim 1$  for bar height, total rows of the matrix must

be same as the number of nodes and row names must be same as the vertex

names in bioNetGraph

outer non-negative numeric, the outside boundary of plot area from node center non-negative numeric, the inside boundary of plot area from node center

plotColors character vector or vector of R color specification, color names for each sample,

pre-generated to control sample colors

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout()

methyl <- caOmicsV.bionet.eset$categoryData[[1]]
sampleColors <- c(rep("red", 20), rep("blue", 20), rep("cyan", 20))
plotBioNetBars(methyl, outer=1.6, inner=1.5, plotColors=sampleColors)</pre>
```

plotBioNetCircos

Default Plot Method for caOmicsV bioNetCircos Layout

## **Description**

A sample way to plot sample information and genomic data with default settings. This method is mainly for demo purpose. In most cases, users need make changes based on this display, specifically, the igraph layout.

#### Usage

```
plotBioNetCircos(eSet, graph=NULL, heatmapMax=NULL, heatmapMin=NULL,
    heatmapColor="BlueWhiteRed")
```

# Arguments

eSet an object returned from method of getESet() that contains all data for plot graph an igraph object generated with one plot dataset, e.g, gene expression value heatmapColor character vector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed",

"GreenBlackRed", "YellowToRed", and "Black".

plotBioNetHeatmap 47

heatmapMax numeric, maximum value for heatmap plot, set to NULL to use the biggest value

in input data set.

heatmapMin numeric, minimum value for heatmap plot, set to NULL to use the smallest value

in input data set.

#### Value

None

#### Author(s)

Henry Zhang

# **Examples**

```
library(caOmicsV)
data(caOmicsV.bionet.eset)
plotBioNetCircos(caOmicsV.bionet.eset)
```

plotBioNetHeatmap

Heatmap Plot for Each Node on caOmicsV bioNetCircos Layout

## **Description**

Headmap plot on caOmicsv bioNetCircos layout. This method plots one track of heatmap for every node. bioNetCircos layout and graphic device must be initialized first.

#### Usage

```
plotBioNetHeatmap(dataValues, maxValue=NULL, minValue=NULL, outer, inner, plotColors)
```

## **Arguments**

dataValues numeric matrix of log2 values for heatmap plot. Total rows of the matrix must be

same as the number of nodes and rownames must be same as the vertex names

in bioNetGraph

maxValue numeric, the biggest value of plot data minValue numeric, the smallest value of plot data

outer non-negative numeric, the outside boundary of plot area from node center non-negative numeric, the inside boundary of plot area close to node center plotColors character vector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed",

"GreenBlackRed", "YellowToRed", "BlackOnly".

#### Value

None

#### Author(s)

Henry Zhang

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#### **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout()
plotBioNetHeatmap(expr, outer=3, inner=2, plotColors="BlueWhiteRed")</pre>
```

plotBioNetLines

Line Plot on caOmicsV bioNetCircos Layout

## **Description**

Line plot on caOmicsv bioNetCircos layout. This method plots one track of line for every node. bionetCircos layout and graphic device must be initialized first.

## Usage

```
plotBioNetLines(dataValues, outer, inner, maxValue=NULL, minValue=NULL,
    plotColors=rep("black", ncol(dataValues)))
```

#### **Arguments**

dataValues numeric matrix of plot data

outer non-negative numeric, the outside boundary of plot area from node center non-negative numeric, the inside boundary of plot area from node center

maxValue numeric, the biggest value of plot data minValue numeric, the smallest value of plot data

plotColors character vector or vector of R specification, colours for each sample

#### Value

None

#### Author(s)

Henry Zhang

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout()
plotBioNetLines(expr, outer=13, inner=2)</pre>
```

plotBioNetPoints 49

plotBioNetPoints	Point Plot on caOmicsV bioNetCircos Layout	

# Description

Point plot on caOmicsv bioNetCircos layout. This method plots one track of points for every node. bioNetCircos layout and graphic device must be initialized first.

## Usage

# Arguments

dataValues	numeric matrix of plot data
maxValue	numeric, the biggest value of plot data
minValue	numeric, the smallest value of plot data
outer	non-negative numeric, the outer boundary of plot area from node center
inner	non-negative numeric, the inner boundary of plot area from node center
plotColors	character vector or vector of R color specification, colors for each sample
sizeByValue	logic, if true, the data value will be used for point size
pch	charcter for point type, same as the one in par()

## Value

None

# Author(s)

Henry Zhang

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout()
plotBioNetPoints(expr, outer=3, inner=2)</pre>
```

plotBioNetPolygons Polygon Plot on caOmicsV bioNetCircos Layout

## **Description**

Plot category data as polygons on node(s) of caOmicsV bioNetCircos layout. bioNetCircos layout and graphic device must be initialized first. Polygon colours are converted from category value with build in colour series. Use getcaOmicsVColors() and setcaOmicsVColors(colorList) to see or reset customized colours.

#### Usage

```
plotBioNetPolygons(dataValues, outer, inner)
```

#### **Arguments**

dataValues matrix of character or numeric for category data

outer non-negative numeric, the outer boundary of plot area from node center non-negative numeric, the innner boundary of plot area from node center

#### Value

None

## Author(s)

Henry Zhang

## **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout()
dataValues <- matrix(rep(c(1:3), each=20), nrow=1)
plotBioNetPolygons(dataValues, outer=3, inner=2)</pre>
```

 ${\it plot Heatmap Color Scale \ for \ both \ caOmics V \ bioMatrix \ and \ bioNew-Circos \ Layout}$ 

# Description

Draw heatmap color scale for legends of bioMatrix plot and bioNetCircos plot. Graphic device must be initialized.

#### Usage

#### **Arguments**

coorX	numeric, x coordinates for the top left location of color scale
coorY	numeric, y coordinates for the top left location of color scale
scaleWidth	non-negative numeric, width of color scale
scaleHeight	non-negative numeric, height of color scale
colorType	character vector, one of "BlueWhiteRed", "GreenWhiteRed", "GreenYellowRed", "GreenBlackRed", "YellowToRed", and "BlackOnly"
minValue	numeric, the smallest value associated with the lowest coluor
maxValue	numeric, the biggest value associated with the highest coluor

One character, either "h" for horizontal or "v" for vertical

#### Value

None

#### Author(s)

Henry Zhang

direction

## **Examples**

resetBioNetNodePlotAreaBoundary

Update Node Plot Area Boundary on caOmicsV bioNetCircos Layout

## **Description**

Record the plotted area boundary for all nodes on the igraph. These boundary may be needed for drawn customized arrows and label node names. bioNetCircos layout must be initialized first. This function is for internal use only.

## Usage

```
resetBioNetNodePlotAreaBoundary(inner=getBioNetNodeRadius(), outer)
```

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## **Arguments**

inner non-negative numeric, the inner boundary of area that has been plotted outer non-negative numeric, the outer boundary of area that has been plotted

#### Value

None

## Author(s)

Henry Zhang

#### **Examples**

resetBioNetNodePlotAreaBoundary(outer=1.5)

RNA2miRNA

Paired miRNA and Genes with Nagetive Correlation Coefficients

# **Description**

A data frame contains 26 genes and their most negative correlated miRNA. Used for demo of generation a new data frame with gene names and related miRNA read counts.

# Usage

data("RNA2miRNA")

## **Format**

A data frame with 26 observations on the following 2 variables.

GeneSymbol a factor with levels ADAMTS13 AMIGO3 BCO2 CDKN3 CFP COL15A1 CSRNP1 CXCL12 DBH ECM1 ESM1 FCN3 LEPREL1 LIFR LILRB5 LOC222699 LYVE1 MT1F OLFML3 PLVAP PTH1R RCAN1 RND3 SEMA3F SLC26A6 TOMM40L

miRNA\_ID a factor with levels hsa.mir.101.1 hsa.mir.101.2 hsa.mir.10b hsa.mir.139 hsa.mir.424 hsa.mir.450b

## **Examples**

data(RNA2miRNA)

RNASeq 53

RNASeq

An Sample of RNASeq Data Set

## **Description**

A data frame with RNASeq reads of 47 genes in 76 samples.

#### Usage

```
data("RNASeq")
```

#### **Format**

A data frame with 47 observations on the following 77 variables.

GeneSymbol a factor with levels A1BG ADAMTS13 AMIGO3 ARMCX6 BCO2 C17orf68 CBLN4 CDKN3 CFP COL13A1 COL15A1 CSRNP1 CXCL12 DBH DNA2 ECM1 ESM1 FAM20A FCN3 GJB2 HOXD9 KIF12 LEPREL1 LIFR LILRB5 LOC222699 LOC441455 LYVE1 MOBKL1A MT1F NPM3 OLFML3 PCDHB7 PLVAP PRICKLE4 PTH1R RCAN1 RND3 RNF125 SEMA3F SIRT5 SLC26A6 SPACA5 TKTL1 TOMM40L UBE2J2 ZNF273

```
TCGA.DD.A3A3.Normal a numeric vector
```

TCGA.DD.A1EI.Normal a numeric vector

TCGA.BC.A216.Normal a numeric vector

TCGA.DD.A11A.Normal a numeric vector

TCGA.EP.A26S.Normal a numeric vector

TCGA.DD.A118.Normal a numeric vector

TCGA.DD.A3A1.Normal a numeric vector

TCGA.BD.A3EP.Normal a numeric vector

TCGA.EP.A12J.Normal a numeric vector

TCGA.DD.A11B.Normal a numeric vector

TCGA.BD.A2L6.Normal a numeric vector

TCGA.DD.A3A2.Normal a numeric vector

TCGA.FV.A2QR.Normal a numeric vector

TCGA.DD.A1EJ.Normal a numeric vector

TCGA.DD.A11D.Normal a numeric vector

TCGA.DD.A39X.Normal a numeric vector

TCGA.DD.A39V.Normal a numeric vector

TCGA.FV.A23B.Normal a numeric vector

TCGA.DD.A1EH.Normal a numeric vector

TCGA.DD.A1EG.Normal a numeric vector

TCGA.DD.A11C.Normal a numeric vector

TCGA.DD.A113.Normal a numeric vector

TCGA.DD.A1EC.Normal a numeric vector

TCGA.ES.A2HT.Normal a numeric vector

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```
TCGA.DD.A119.Normal a numeric vector
TCGA.DD.A1EL.Normal a numeric vector
TCGA.DD.A1EB.Normal a numeric vector
TCGA.DD.A39W.Normal a numeric vector
TCGA.DD.A39Z.Normal a numeric vector
TCGA.DD.A114.Normal a numeric vector
TCGA.DD.A1EH.Tumor a numeric vector
TCGA.ED.A4XI.Tumor a numeric vector
TCGA.DD.A4NB.Tumor a numeric vector
TCGA.DD.A118.Tumor a numeric vector
TCGA.BD.A3EP.Tumor a numeric vector
TCGA.DD.A1ED.Tumor a numeric vector
TCGA.DD.A39X.Tumor a numeric vector
TCGA.DD.A39Z.Tumor a numeric vector
TCGA.BC.A216.Tumor a numeric vector
TCGA.DD.A11A.Tumor a numeric vector
TCGA.DD.A113.Tumor a numeric vector
TCGA.BC.A10Z.Tumor a numeric vector
TCGA.DD.A119.Tumor a numeric vector
TCGA.DD.A3A3.Tumor a numeric vector
TCGA.ED.A82E.Tumor a numeric vector
TCGA.FV.A2QR.Tumor a numeric vector
TCGA.DD.A1EL.Tumor a numeric vector
TCGA.FV.A23B.Tumor a numeric vector
TCGA.CC.5258.Tumor a numeric vector
TCGA.EP.A12J.Tumor a numeric vector
TCGA.DD.A3A1.Tumor a numeric vector
TCGA.DD.A11D.Tumor a numeric vector
TCGA.ED.A5KG.Tumor a numeric vector
TCGA.DD.A114.Tumor a numeric vector
TCGA.DD.A39W.Tumor a numeric vector
TCGA.DD.A1EC.Tumor a numeric vector
TCGA.EP.A26S.Tumor a numeric vector
TCGA.RC.A7S9.Tumor a numeric vector
TCGA.BD.A2L6.Tumor a numeric vector
TCGA.UB.A7MD.Tumor a numeric vector
TCGA.DD.A11B.Tumor a numeric vector
TCGA.ED.A7XP.Tumor a numeric vector
TCGA.DD.A1EI.Tumor a numeric vector
TCGA.UB.A7MA.Tumor a numeric vector
```

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```
TCGA.DD.A73A.Tumor a numeric vector TCGA.DD.A3A2.Tumor a numeric vector TCGA.DD.A1EB.Tumor a numeric vector TCGA.HP.A5MZ.Tumor a numeric vector TCGA.DD.A1EJ.Tumor a numeric vector TCGA.DD.A1EJ.Tumor a numeric vector TCGA.RC.A7SB.Tumor a numeric vector TCGA.RC.A6M4.Tumor a numeric vector TCGA.ES.A2HT.Tumor a numeric vector TCGA.DD.A11C.Tumor a numeric vector TCGA.DD.A39V.Tumor a numeric vector TCGA.MI.A75C.Tumor a numeric vector TCGA.MI.A75C.Tumor a numeric vector
```

#### **Examples**

data(RNASeq)

RNASeqDemoData

RNASeq Demo Data Set

#### **Description**

A data frame with RNASeq read counts of 26 genes in 60 samples. Used for heatmap plot on caOimcsV bioNetCircos layout and bioMatirx layout after transformed to log2 values

#### Usage

```
data("RNASeqDemoData")
```

#### **Format**

A data frame with 26 observations on the following 61 variables.

GeneSymbol a factor with levels A1BG ADAMTS13 AMIGO3 ARMCX6 BCO2 C17orf68 CBLN4 CDKN3 CFP COL13A1 COL15A1 CSRNP1 CXCL12 DBH DNA2 ECM1 ESM1 FAM20A FCN3 GJB2 HOXD9 KIF12 LEPREL1 LIFR LILRB5 LOC222699 LOC441455 LYVE1 MOBKL1A MT1F NPM3 OLFML3 PCDHB7 PLVAP PRICKLE4 PTH1R RCAN1 RND3 RNF125 SEMA3F SIRT5 SLC26A6 SPACA5 TKTL1 TOMM40L UBE2J2 ZNF273

```
TCGA.BC.A216.Normal a numeric vector TCGA.BD.A2L6.Normal a numeric vector TCGA.BD.A3EP.Normal a numeric vector TCGA.DD.A113.Normal a numeric vector TCGA.DD.A114.Normal a numeric vector TCGA.DD.A118.Normal a numeric vector TCGA.DD.A119.Normal a numeric vector
```

TCGA.DD.A11A.Normal a numeric vector

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```
TCGA.DD.A11B.Normal a numeric vector
TCGA.DD.A11C.Normal a numeric vector
TCGA.DD.A11D.Normal a numeric vector
TCGA.DD.A1EB.Normal a numeric vector
TCGA.DD.A1EC.Normal a numeric vector
TCGA.DD.A1EG.Normal a numeric vector
TCGA.DD.A1EH.Normal a numeric vector
TCGA.DD.A1EI.Normal a numeric vector
TCGA.DD.A1EJ.Normal a numeric vector
TCGA.DD.A1EL.Normal a numeric vector
TCGA.DD.A39V.Normal a numeric vector
TCGA.DD.A39W.Normal a numeric vector
TCGA.DD.A39X.Normal a numeric vector
TCGA.DD.A39Z.Normal a numeric vector
TCGA.DD.A3A1.Normal a numeric vector
TCGA.DD.A3A2.Normal a numeric vector
TCGA.DD.A3A3.Normal a numeric vector
TCGA.EP.A12J.Normal a numeric vector
TCGA.EP.A26S.Normal a numeric vector
TCGA.ES.A2HT.Normal a numeric vector
TCGA.FV.A23B.Normal a numeric vector
TCGA.FV.A2QR.Normal a numeric vector
TCGA.BC.A216.Tumor a numeric vector
TCGA.BD.A2L6.Tumor a numeric vector
TCGA.BD.A3EP.Tumor a numeric vector
TCGA.DD.A113.Tumor a numeric vector
TCGA.DD.A114.Tumor a numeric vector
TCGA.DD.A118.Tumor a numeric vector
TCGA.DD.A119.Tumor a numeric vector
TCGA.DD.A11A.Tumor a numeric vector
TCGA.DD.A11B.Tumor a numeric vector
TCGA.DD.A11C.Tumor a numeric vector
TCGA.DD.A11D.Tumor a numeric vector
TCGA.DD.A1EB.Tumor a numeric vector
TCGA.DD.A1EC.Tumor a numeric vector
TCGA.DD.A1EG.Tumor a numeric vector
TCGA.DD.A1EH.Tumor a numeric vector
TCGA.DD.A1EI.Tumor a numeric vector
TCGA.DD.A1EJ.Tumor a numeric vector
TCGA.DD.A1EL.Tumor a numeric vector
```

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```
TCGA.DD.A39V.Tumor a numeric vector TCGA.DD.A39W.Tumor a numeric vector TCGA.DD.A39X.Tumor a numeric vector TCGA.DD.A39Z.Tumor a numeric vector TCGA.DD.A3A1.Tumor a numeric vector TCGA.DD.A3A2.Tumor a numeric vector TCGA.DD.A3A3.Tumor a numeric vector TCGA.EP.A12J.Tumor a numeric vector TCGA.EP.A26S.Tumor a numeric vector TCGA.ES.A2HT.Tumor a numeric vector TCGA.FV.A23B.Tumor a numeric vector TCGA.FV.A20R.Tumor a numeric vector
```

#### **Examples**

data(RNASeqDemoData)

sampleDemoData

Sample Tissue Type Demo Data Set

## **Description**

A data frame with sample names and tissue type for 60 samples. Used to plot tissue type on both caOmicsV bioMatrix layout and bioNetCircos layout.

#### Usage

```
data("sampleDemoData")
```

## Format

A data frame with 60 observations on the following 2 variables.

```
sample_names a factor with levels TCGA.BC.A216.Normal TCGA.BC.A216.Tumor TCGA.BD.A2L6.Normal
    TCGA.BD.A2L6.Tumor TCGA.BD.A3EP.Normal TCGA.BD.A3EP.Tumor TCGA.DD.A113.Normal
    TCGA.DD.A113.Tumor TCGA.DD.A114.Normal TCGA.DD.A114.Tumor TCGA.DD.A118.Normal
    TCGA.DD.A118.Tumor TCGA.DD.A119.Normal TCGA.DD.A119.Tumor TCGA.DD.A11A.Normal
    TCGA.DD.A11A.TumorTCGA.DD.A11B.NormalTCGA.DD.A11B.TumorTCGA.DD.A11C.Normal
    TCGA.DD.A11C.Tumor TCGA.DD.A11D.Normal TCGA.DD.A11D.Tumor TCGA.DD.A1EB.Normal
    TCGA.DD.A1EB.Tumor TCGA.DD.A1EC.Normal TCGA.DD.A1EC.Tumor TCGA.DD.A1EG.Normal
    TCGA.DD.A1EG.Tumor TCGA.DD.A1EH.Normal TCGA.DD.A1EH.Tumor TCGA.DD.A1EI.Normal
    TCGA.DD.A1EI.Tumor TCGA.DD.A1EJ.Normal TCGA.DD.A1EJ.Tumor TCGA.DD.A1EL.Normal
    TCGA.DD.A1EL.Tumor TCGA.DD.A39V.Normal TCGA.DD.A39V.Tumor TCGA.DD.A39W.Normal
    TCGA.DD.A39W.Tumor TCGA.DD.A39X.Normal TCGA.DD.A39X.Tumor TCGA.DD.A39Z.Normal
    TCGA.DD.A39Z.Tumor TCGA.DD.A3A1.Normal TCGA.DD.A3A1.Tumor TCGA.DD.A3A2.Normal
    TCGA.DD.A3A2.Tumor TCGA.DD.A3A3.Normal TCGA.DD.A3A3.Tumor TCGA.EP.A12J.Normal
    TCGA.EP.A12J.Tumor TCGA.EP.A26S.Normal TCGA.EP.A26S.Tumor TCGA.ES.A2HT.Normal
    TCGA.ES.A2HT.TumorTCGA.FV.A23B.NormalTCGA.FV.A23B.TumorTCGA.FV.A2QR.Normal
    TCGA.FV.A2QR.Tumor
```

tissue\_type a factor with levels Normal Tumor

## **Examples**

data(sampleDemoData)

setBioMatrixBaseCoordinates

Setup Base Coordinates for caOmicsV bioMatrix Layout

# Description

Initialize x and y coordinates for each plot area on bioMatrix layout. This function is for internal use.

# Usage

# **Arguments**

numOfSamples	non-negative integer, number of samples to be plotted
sampleWidth	non-negative numeric, width of rectangle in inch for each sample, default 0.1
columnPadding	non-negative numeric, padding width in inch between two samples, default $0.025$
sampleHeight	non-negative numeric, height of rectangle in inchfor each sample, default 0.4
geneNameWidth	non-negative numeric, width of gene labeling area in inch, default 1
remarkWidth	non-negative numeric, width of remark area in inch, default 1

## Value

None

# Author(s)

Henry Zhang

setBioMatrixPlotArea 59

setBioMatrixPlotArea Setup Plot Area for caOimcsV bioMatrix Layout

## **Description**

Setup plot area including of sample name area (sampleHeight), phenotype area, gene label area (geneNameWidth), and remark area (remarkWidth, and other descriptions). bioMatrix layout muse be initialized first. This function is for internal use only.

#### Usage

```
setBioMatrixPlotArea()
```

#### Value

None

#### Author(s)

Henry Zhang

## **Examples**

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
setBioMatrixPlotArea()
```

setBioMatrixPlotParameters

Setup Plot Parameters for caOmicsV bioMatrix Layout

## **Description**

Put biomatrix plot parameters to CA\_OMICS\_ENV environment. This function is for internal use only and all arguments are validated outside in advance.

## Usage

# Arguments

numOfGenes non-negative integer, total number of genes to be plotted
numOfSamples non-negative integer, total number of samples to be plotted
numOfPhenotypes non-negative integer, total number of phenotypes to be plotted
sampleHeight non-negative numeric, height of polygon (sample) in inch on each row
non-negative numeric, width of polygon (sample) in inch on each row

columnPadding non-negative numeric, padding width in inch between two polygons (samples)

rowPadding non-negative numeric, padding height in inch between two rows

geneNameWidth non-negative numeric, width of left text plot area in inch for gene names

remarkWidth non-negative numeric, width of plot area in inch for right side text plot (second

set of gene names) and/or summary data plot

sampleNameHeight

non-negative numeric, height of plot area in inch for sample names

legendHeight non-negative numeric, height of legend area

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
setBioMatrixPlotParameters(numOfGenes=30, numOfSamples=50, numOfPhenotypes=1,
    sampleHeight=0.4, sampleWidth=0.2, columnPadding=0.05, rowPadding=0.5,
    geneNameWidth=1, remarkWidth=2, sampleNameHeight=2, legendHeight=2)
```

 ${\tt setBioNetCircosBasePlotPositions}$ 

Setup Default Plot Positions for a Node on caOmicsV bioNetCircor Layout

#### Description

This function calculates default x and y coordinates for points on a circular line. Degrees a character string should be rotated on each point are also calculated. THis function is for internal call only.

# Usage

```
setBioNetCircosBasePlotPositions(totalSamples=100, sampleWidth=100)
```

## **Arguments**

totalSamples non-negative numeric, total number of samples to be plotted

sampleWidth non-negative numeric, width (how many points) of a sample on the circular line

#### Value

None

#### Author(s)

Henry Zhang

setBioNetNodeLayout 61

#### **Examples**

 ${\tt setBioNetCircosBasePlotPositions(totalSamples=100, sampleWidth=100)}$ 

setBioNetNodeLayout

Setup Layout for the igraph Object

## **Description**

Set up node layout on a biological network. The layout is taken from an igraph layout and scaled to allocate circos plot area for each node.

# Usage

```
setBioNetNodeLayout(bioNet, layout=layout.auto(bioNet))
```

# Arguments

bioNet an igraph object for which layout will be set

layout object from layout method of igraph package

#### Value

None

## Author(s)

Henry Zhang

## **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)
setBioNetNodeLayout(bioNet, layout=layout.auto(bioNet))</pre>
```

setBioNetPlotAreaBackground

Setup Plot Area Background for Nodes of caOmicsV bioNetCircos Layout

# Description

Change the plot area background of igraph nodes. Use white color to erase background and grey to show the plot area boundary.

## Usage

```
setBioNetPlotAreaBackground(bgColor=grey(0.75, alpha=0.5))
```

62 setBioNetPlotParameters

#### **Arguments**

bgColor character vector of a colour name or a R colour specification

#### Value

None

#### Author(s)

Henry Zhang

#### **Examples**

```
setBioNetPlotAreaBackground(grey(0.75, alpha=0.5))
```

setBioNetPlotParameters

Initialize Plot Parameters for caOmicsV bioNetCircos Plot

#### **Description**

Set up plot parameters for bioNetCircos layout including of totalSamples, sampleWidth, nodeRadius, nodePadding, plotAreaWidth, outer and inner boundaries of plotArea, as well as default plot colors. This function is for internal use only.

## Usage

```
setBioNetPlotParameters(totalSamples, sampleWidth, nodeRadius, nodePadding,
    plotAreaWidth)
```

#### **Arguments**

totalSamples non-negative integer, total number of samples to be plotted

sampleWidth non-negative integer, width (number of points) of a sample on the circular line

nodeRadius non-negative numeric, radius of the node

nodePadding non-negative numeric, padding width in inch between two nodes

plotAreaWidth non-negative numeric, total width of plot area.

#### Value

None.

## Author(s)

Henry zhang

setCaOmicsVColors 63

setCaOmicsVColors

Setup Default Plot Colors for caOmicsV Plot

## **Description**

Set or change default colors used for caOmicsV Plot.

# Usage

```
setCaOmicsVColors(colorList=NULL)
```

## **Arguments**

colorList

character vector, list of colour names or R colour specification

#### Value

None

#### Author(s)

Henry Zhang

## **Examples**

setDefaultNaStrings

Setup Default NA Strings for caOmicsV Package

# Description

Set or change default NA strings used by caOmicsV Package.

# Usage

```
setDefaultNaStrings(nullStrings)
```

# Arguments

nullStrings character vector of null strings

#### Value

None

## Author(s)

Henry Zhang

## **Examples**

```
setDefaultNaStrings("null")
```

showBioMatrixPlotLayout

Display caOmicsV bioMatrix Layout

# Description

Display caOmicsV bioMatrix layout with default information for each plot area for purpose of optimizing the layout. A bioMatrix layout must be initialized first.

# Usage

```
showBioMatrixPlotLayout(geneNames, sampleNames, phenotypes, sampleColors=NULL,
    geneColors=NULL, phenoColors=NULL)
```

# **Arguments**

geneNames	character vector, gene names shown at left side of omics data plot area
sampleNames	character vector, sample names shown on the top of phenotype plot area
phenotypes	character vector, phenotype names listed on left side of phenotype plot area
sampleColors	character vector for colour name(s) of R colour specification for samples
geneColors	character vector of colour name or R colour specification for genes
phenoColors	character vector of colour name(s) or R colour specification for phenotypes

## Value

None

# Author(s)

Henry Zhang

```
initializeBioMatrixPlot(numOfGenes=1, numOfSamples=50)
showBioMatrixPlotLayout("Gene", paste("Sample", 1:50), "Diagnosis")
```

showBioNetNodesLayout Display Nodes Layout of caOmicsV bioNetCircos Plot

# Description

Display caOmicsV bioNetCircos layout with empty nodes and edges in order to optimize bioNetCircos layout. An igraph object must be initialized first.

# Usage

```
showBioNetNodesLayout(bgColor=grey(0.75, alpha=0.5))
```

## **Arguments**

bgColor

character vector for a color name or a R color specification

#### Value

None

#### Author(s)

Henry Zhang

# **Examples**

```
data(caOmicsV.bionet.eset)
expr <- caOmicsV.bionet.eset$heatmapData[[1]]
bioNet <- bc3net(expr)

initializeBioNetCircos(bioNet, totalSamples=60)
showBioNetNodesLayout(bgColor=grey(0.75, alpha=0.5))</pre>
```

 $\verb|showSupportedBioNetCircosPlotType| \\$ 

Display the Plot Types Supported by caOmicsV bioNetCircos Plot

# Description

Display plot types supported by current version of caOmicsV bioNetCircos plot. A bioNetCircos layout must be initialized first.

# Usage

```
show Supported Bio Net Circos Plot Type ()\\
```

# Value

None

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## Author(s)

Henry Zhang

# **Examples**

showSupportedBioNetCircosPlotType()

sortClinicalData

Sort Clinical Data by a Column

# Description

Sort sample information by a column defined by byItem (a column header). This function is for sorting sample information only.

## Usage

```
sortClinicalData(clinicalData, byItem)
```

## **Arguments**

clinicalData A data frame with rows for samples and columns for features. Sample names

must be in the first column.

by Item character vector of a feature (column header) by which the data will be sorted.

# Value

sampleData copy of the first argument with new row order.

# Author(s)

Henry Zhang

```
data(sampleDemoData)
sampleInfo <- sortClinicalData(sampleDemoData, colnames(sampleDemoData)[1])</pre>
```

sortOmicsDataByColumn Sort Omics Data by Column Header

## **Description**

Sort omics data by column header (sample names) based on the order of the second argument. The sample names in the first argument must have same items as the second argument except of order. After sorting, the sample names in omics data and the second argument will be in same order.

#### Usage

```
sortOmicsDataByColumn(omicsData, sampleNames)
```

#### **Arguments**

omicsData A data frame that holds genomic data such as gene expression, SNV, RNASeq

... The column headers must be the sample names that are same as the sample

names in clinical data.

sampleNames character vector, sample names in a given order (such as diagnosis). No redun-

dant entries allowed.

#### Value

omicsData copy of the first argument with columns in new order.

#### Author(s)

Henry Zhang

#### **Examples**

```
data(RNASeqDemoData)
sampleNames <- colnames(RNASeqDemoData)[-1]
sampleNames <- sampleNames[length(sampleNames):1]
expr <- sortOmicsDataByColumn(RNASeqDemoData, sampleNames)</pre>
```

sortOmicsDataByRow

Sort Omics Data by Row

# **Description**

Sort omics data by row (genes) in order to get omics data in a specific order, e.g., ordered by p values or by fold changes in expression data. After sorting, the order of row names in omics data will be same as the second argument.

## Usage

```
sortOmicsDataByRow(omicsData, geneNames)
```

## **Arguments**

omicsData a data frame that holds genomic data such as gene expression, SNV, RNASeq.

The first column must be gene names that are same as the second argument

except of order.

geneNames character vector, gene names in a given order (such as by p values). Redundant

gene names are allowed.

## Value

omicsData copy of the first argument with new row order.

## Author(s)

Henry Zhang

```
data(RNASeqDemoData)
geneNames <- as.character(RNASeqDemoData[,1])
geneNames <- geneNames[order(RNASeqDemoData[,2])]
expr <- sortOmicsDataByRow(RNASeqDemoData, geneNames)</pre>
```

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