

# ecoltk

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## R topics documented:

ecoli.m52.genome . . . . .	1
ecoligenomeBNUM2MULTIFUN . . . . .	2
ecoligenomeBNUM . . . . .	3
ecoligenome.operon . . . . .	3
gccontent . . . . .	4
linkedmultiget . . . . .	5
multiFun . . . . .	6
pointsCircle . . . . .	6
polar2xy . . . . .	8
polygonChrom . . . . .	9
wstringapply . . . . .	11

<b>Index</b>	<b>12</b>
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ecoli.m52.genome    *Escherichia coli* data

---

## Description

Meta-data related to *Escherichia coli*

## Usage

```
data(ecoli.m52.genome)
data(ecoligenomeCHRLOC)
data(ecoligenomeSYMBOL2AFFY)
data(ecoligenomeSYMBOL)
data(ecoligenomeSTRAND)
data(ecoligenome.operon)
ecoli.len
```

**Format**

The format for `ecoli.m52.genome` is character with genome sequence. The format for `ecoligenomeCHRLOC` is an environment (as a hash table). Each key is an Affymetrix probe set ID, and each value is vector of two integers (begining and end - see the details below) The format for `ecoligenomeSYMBOL2AFFFY` is an environment (as a hash table). Each key is a gene symbol name. The format for `ecoligenomeSYMBOL` is an environment (as a hash table). Each key is an Affymetrix probe set id `ecoli.len` is a variable containing the size of the genome in `ecoli.m52.genome`.

**Details**

The environments `ecoligenomeSYMBOL2AFFFY` and `ecoligenomeSYMBOL` are like the ones in the data packages built by `annBuilder`.

The environment `ecoligenomeCHRLOC` differs: two integers are associated with each key, one corresponds to the begining of the segment the other to the end.

The environment `ecoligenomeSTRAND` returns a logical. `TRUE` means that the orientation is '+', `FALSE` means that the orientation is '-' (and `NA` is used when irrelevant for the key).

**Source**

<http://www.genome.wisc.edu/sequencing/k12.htm> and [http://www.biostat.harvard.edu/complab/dchip/info\\_file.htm](http://www.biostat.harvard.edu/complab/dchip/info_file.htm)

**Examples**

```
data(ecoli.m52.genome)
```

---

```
ecoligenomeBNUM2MULTIFUN
      Environment
```

---

**Description**

An environment to store associations between 'bnum' identifiers (key) and 'MultiFun' identifiers (or strand information).

**Usage**

```
data(ecoligenomeBNUM2MULTIFUN)
```

**Format**

The format is: length 0 <environment> - attr(\*, "comments")= chr "GenProtEC: MultiFun assignments for E. coli modules September 17th, 2003"

**Details**

'MultiFun' is a classification scheme. The structure is 'approximately tree-like'. Several 'MultiFun' numbers can be assigned to one 'bnum'.

**Source**

"http://genprotec.mbl.edu/files/MultiFun.txt"

---

ecoligenomeBNUM *Environment for 'bnum' identifiers*

---

**Description**

Environments to associate Affymetrix probe set IDs with 'bnum' IDs

**Usage**

```
data(ecoligenomeBNUM)
data(ecoligenomeBNUM2SYMBOL)
data(ecoligenomeBNUM2ENZYME)
data(ecoligenomeBNUM2GENBANK)
data(ecoligenomeBNUM2GENEPRODUCT)
data(ecoligenomeSYMBOL2BNUM)
```

**Format**

These are environment objects.

**Details**

Escherichia coli genes are sometimes identified by 'bnum's. This identifier is typically a 'b' followed by digits.

**Source**

BNUM numbers were parsed out of the Affymetrix identifiers. BNUM2\* were obtained from the GenProtEC website.

---

ecoligenome.operon *Known operon in E.coli - data.frame*

---

**Description**

The known operon in the Escherichia coli genome.

**Usage**

```
data(ecoligenome.operon)
```

**Format**

A data frame with 932 observations (genes) on the following 4 variables.

**gene.name** a character vector

**gene.annotation** a character vector

**operon.name** a factor with levels the names of the operons

**operon.comments** a factor with levels the comments for the operons

## Details

For some operons, the source of information specifies the existence of regulating elements such as promoter, terminator, box, etc... In those cases, the `gene.name` is set to "Regulation", and the `gene.annotation` gives what kind of regulating element it is. If volunteers, it would be neat to map those on the genome... Besides that, not much to add. The data structure is fairly straightforward.

## Source

Built from the webpage: <http://www.cib.nig.ac.jp/dda/backup/taitoh/ecoli.operon.html>

## Examples

```
library(Biobase)
data(ecoligenome.operon)
data(ecoligenomeSYMBOL2AFFY)

## something that might be useful when working with Affymetrix data:
## get the Affymetrix identifiers for the probe sets bundled in operons
## (see the vignette for more details)
ecoligenome.operon$affyid <-
  unname(unlist(mget(ecoligenome.operon$gene.name,
                    ecoligenomeSYMBOL2AFFY, ifnotfound=NA)))
```

---

gccontent	<i>function to compute gccontent</i>
-----------	--------------------------------------

---

## Description

A simple R function to compute the GC content of a sequence

## Usage

```
gccontent(x)
```

## Arguments

`x` a vector of mode character

## Details

This a simple (and not particularly fast) function to compute the GC content of sequence. When speed is an issue, one should use the function in the package `matchprobes`. This function only exists to avoid dependency on this package.

## Value

The GC content (numeric)

---

linkedmultiget      *A function to look for values across linked environments*

---

## Description

A function to look for values across linked environments.

## Usage

```
linkedmultiget(x, envir.list = list(), unique = TRUE)
```

## Arguments

<code>x</code>	The keys in the first environment in the list.
<code>envir.list</code>	A list of environments.
<code>unique</code>	Simplify the list returned by ensuring that the values for each key are unique.

## Details

Environments can be considered as hashtables. The keys are obviously strings, but in some cases the associated values are also strings. This is the case for annotation environments (as built with the package `AnnBuilder`). This function helps to look for values across several environments: the keys have associated values in a first environment, these values are used as keys in the second environments, etc...

## Value

A list of length the length of `x`.

## Author(s)

Laurent Gautier

## See Also

[mget](#)

## Examples

```
data(ecoligenomeBNUM)
data(ecoligenomeBNUM2MULTIFUN)
data(multiFun)

## get 5 Affymetrix IDs
set.seed(456)
my.affyids <- sample(ls(ecoligenomeBNUM), 5)

## get the MULTIFUN annotations for them
r <- linkedmultiget(my.affyids, list(ecoligenomeBNUM,
                                     ecoligenomeBNUM2MULTIFUN, multiFun))

print(r)
```

---

 multiFun

*multiFun classification*


---

### Description

The MultiFun classification scheme

### Usage

```
data(multiFun)
data(ecoligenomeMULTIFUN2GO)
```

### Format

These are environments.

### Source

<http://genprotec.mbl.edu/files/MultiFun.txt>

### Examples

```
## To be done...
```

---

 pointsCircle

*Functions to plot circular related figures*


---

### Description

Functions to plot circular related figures

### Usage

```
linesCircle(radius, center.x = 0, center.y = 0, edges = 300, ...)
polygonDisk(radius, center.x = 0, center.y = 0, edges=300,
  ...)
arrowsArc(theta0, thetal, radius, center.x = 0, center.y = 0, edges = 10,
  length = 0.25, angle = 30, code = 2, ...)
pointsArc(theta0, thetal, radius, center.x = 0, center.y = 0, ...)
linesArc(theta0, thetal, radius, center.x = 0, center.y = 0, ...)
polygonArc(theta0, thetal, radius.in, radius.out,
  center.x = 0, center.y = 0,
  edges = 10,
  col = "black",
  border = NA,
  ...)
```

**Arguments**

`theta0, theta1` start and end angles for the arc  
`radius` radius of the circle  
`radius.in` inner radius  
`radius.out` outer radius  
`center.x, center.y` Coordinates for the center of the circle (default to (0, 0))  
`edges` number of edges the shape is made of  
`col` color  
`border` border (see [polygon](#))  
`length, angle, code` see the corresponding parameters for the function [arrows](#)  
`...` optional graphical paramaters

**Details**

Details to come... for now the best to run the examples and experiment by yourself...

**Value**

Function only used for their border effects.

**Author(s)**

laurent

**Examples**

```

par(mfrow=c(2,2))
n <- 10
thetas <- rev(seq(0, 2 * pi, length=n))

rhos <- rev(seq(1, n) / n)

xy <- polar2xy(rhos, thetas)
colo <- heat.colors(n)

plot(0, 0, xlim=c(-2, 2), ylim=c(-2, 2), type="n")
for (i in 1:n)
  linesCircle(rhos[i]/2, xy$x[i], xy$y[i])

plot(0, 0, xlim=c(-2, 2), ylim=c(-2, 2), type="n")
for (i in 1:n)
  polygonDisk(rhos[i]/2, xy$x[i], xy$y[i], col=colo[i])

plot(0, 0, xlim=c(-2, 2), ylim=c(-2, 2), type="n", xlab="", ylab="")
for (i in 1:n)
  polygonArc(0, thetas[i],
             rhos[i]/2, rhos[i],
             center.x = xy$x[i], center.y = xy$y[i], col=colo[i])

plot(0, 0, xlim=c(-2, 2), ylim=c(-2, 2), type="n", xlab="", ylab="")

```

```

for (i in (1:n)[-1]) {
  linesCircle(rhos[i-1], col="gray", lty=2)
  polygonArc(thetas[i-1], thetas[i],
             rhos[i-1], rhos[i], col=colo[i],
             edges=20)
  arrowsArc(thetas[i-1], thetas[i],
            rhos[i] + 1, col=colo[i],
            edges=20)
}

```

---

polar2xy

*Functions to perform polar coordinate related functions*


---

### Description

Functions to perform polar coordinate related functions

### Usage

```

polar2xy(rho, theta)
xy2polar(x, y)
rotate(x, y, alpha)

```

### Arguments

x	cartesian coordinate
y	cartesian coordinate
rho	polar radius rho
theta	polar angle theta
alpha	angle to perform rotation

### Details

`y` and `theta` can be respectively missing. In this case, `x` and `rho` are expected to be lists with entries `x`, `y`, `rho`, `theta` respectively.

### Examples

```

n <- 40
nn <- 2
thetas <- seq(0, nn * 2 * pi, length=n)

rhos <- seq(1, n) / n

plot(c(-1, 1), c(-1, 1), type="n")
abline(h=0, col="grey")
abline(v=0, col="grey")

xy <- polar2xy(rhos, thetas)

points(xy$x, xy$y, col=rainbow(n))

```



**Description**

Functions to plot circular chromosomes informations

**Usage**

```
cPlotCircle(radius=1, xlim=c(-2, 2), ylim=xlim, edges=300, main=NULL,
            main.inside, ...)
```

```
chromPos2angle(pos, len.chrom, rot=pi/2, clockwise=TRUE)
```

```
polygonChrom(begin, end, len.chrom, radius.in, radius.out,
              total.edges = 300,
              edges = max(round(abs(end - begin)/len.chrom *
                              total.edges), 2, na.rm = TRUE),
              rot = pi/2, clockwise = TRUE, ...)
```

```
linesChrom(begin, end, len.chrom, radius,
            total.edges = 300,
            edges = max(round(abs(end - begin)/len.chrom *
                            total.edges), 2, na.rm = TRUE),
            rot = pi/2, clockwise = TRUE, ...)
```

```
ecoli.len
```

**Arguments**

radius	radius
xlim, ylim	range for the plot. Can be used to zoom-in a particular region.
pos	position (nucleic base coordinate)
begin	beginning of the segment (nucleic base number).
end	end of the segment (nucleic base number).
len.chrom	length of the chromosome in base pairs
radius.in	inner radius
radius.out	outer radius
total.edges	total number of edges for the chromosome
edges	number of edges for the specific segment(s)
rot	rotation (default is $\pi / 2$ , bringing the angle zero at 12 o'clock)
clockwise	rotate clockwise. Default to TRUE.
main, main.inside	main titles for the plot
...	optional graphical parameters

**Details**

The function `chromPos2angle` is a convenience function. The variable `ecoli.len` contains the size of the Escheria coli genome considered (K12).

**Value**

Except `chromPos2angle`, the function are solely used for their border effects.

**Author(s)**

laurent <laurent@cbs.dtu.dk>

**Examples**

```
data(ecoligenomeSYMBOL2AFFY)
data(ecoligenomeCHRLOC)

## find the operon lactose ("lac*" genes)
lac.i <- grep("^lac", ls(ecoligenomeSYMBOL2AFFY))
lac.symbol <- ls(ecoligenomeSYMBOL2AFFY)[lac.i]
lac.affy <- unlist(lapply(lac.symbol, get, envir=ecoligenomeSYMBOL2AFFY))

beg.end <- lapply(lac.affy, get, envir=ecoligenomeCHRLOC)
beg.end <- matrix(unlist(beg.end), nc=2, byrow=TRUE)

lac.o <- order(beg.end[, 1])

lac.i <- lac.i[lac.o]
lac.symbol <- lac.symbol[lac.o]
lac.affy <- lac.affy[lac.o]
beg.end <- beg.end[lac.o, ]

lac.col <- rainbow(length(lac.affy))

par(mfrow=c(2,2))

## plot

cPlotCircle(main="lac genes")
polygonChrom(beg.end[, 1], beg.end[, 2], ecoli.len, 1, 1.2, col=lac.col)
rect(0, 0, 1.1, 1.1, border="red")

cPlotCircle(xlim=c(0, 1.2), ylim=c(0, 1.1))
polygonChrom(beg.end[, 1], beg.end[, 2], ecoli.len, 1, 1.1, col=lac.col)
rect(0.4, 0.8, 0.7, 1.1, border="red")

cPlotCircle(xlim=c(.45, .5), ylim=c(.85, 1.0))
polygonChrom(beg.end[, 1], beg.end[, 2], ecoli.len, 1, 1.03, col=lac.col)

mid.genes <- apply(beg.end, 1, mean)
mid.angles <- chromPos2angle(mid.genes, ecoli.len)
xy <- polar2xy(1.03, mid.angles)
xy.labels <- data.frame(x = seq(0.45, 0.5, length=4), y = seq(0.95, 1.0, length=4))
segments(xy$x, xy$y, xy.labels$x, xy.labels$y, col=lac.col)
text(xy.labels$x, xy.labels$y, lac.symbol, col=lac.col)
```

---

wstringapply      *Apply a function on a window sliding on a string*

---

**Description**

Apply a function on a window sliding on a string.

**Usage**

```
wstringapply(x, SIZE, SLIDE, FUN, ...)
```

**Arguments**

x	The string
SIZE	The size of the window (number of characters).
SLIDE	offset to move at each slide
FUN	The function to be applied
...	optional parameter for the function FUN

**Details**

Apply the function FUN to substrings of x of length SIZE.

**Value**

A list of size `nchar(x) - SIZE`.

**Author(s)**

L, Gautier

# Index

- \*Topic **aplot**
  - pointsCircle, 6
- \*Topic **datasets**
  - ecoli.m52.genome, 1
  - ecoligenome.operon, 3
  - ecoligenomeBNUM, 2
  - ecoligenomeBNUM2MULTIFUN, 2
  - multiFun, 5
- \*Topic **dplot**
  - polygonChrom, 8
- \*Topic **hplot**
  - polygonChrom, 8
- \*Topic **manip**
  - gccontent, 4
  - linkedmultiget, 4
  - polar2xy, 7
  - wstringapply, 10
  
- arrows, 6
- arrowsArc (pointsCircle), 6
  
- chromPos2angle (polygonChrom), 8
- cPlotCircle (polygonChrom), 8
  
- ecoli.len (ecoli.m52.genome), 1
- ecoli.m52.genome, 1
- ecoli.operon (ecoli.m52.genome), 1
- ecoligenome.operon, 3
- ecoligenomeBNUM, 2
- ecoligenomeBNUM2ENZYME
  - (ecoligenomeBNUM), 2
- ecoligenomeBNUM2GENBANK
  - (ecoligenomeBNUM), 2
- ecoligenomeBNUM2GENEPRODUCT
  - (ecoligenomeBNUM), 2
- ecoligenomeBNUM2MULTIFUN, 2
- ecoligenomeBNUM2STRAND
  - (ecoligenomeBNUM2MULTIFUN), 2
- ecoligenomeBNUM2SYMBOL
  - (ecoligenomeBNUM), 2
- ecoligenomeCHRLOC
  - (ecoli.m52.genome), 1
  
- ecoligenomeMULTIFUN2GO
  - (multiFun), 5
- ecoligenomeSTRAND
  - (ecoli.m52.genome), 1
- ecoligenomeSYMBOL
  - (ecoli.m52.genome), 1
- ecoligenomeSYMBOL2AFFY
  - (ecoli.m52.genome), 1
- ecoligenomeSYMBOL2BNUM
  - (ecoligenomeBNUM), 2
  
- gccontent, 4
  
- linesArc (pointsCircle), 6
- linesChrom (polygonChrom), 8
- linesCircle (pointsCircle), 6
- linkedmultiget, 4
  
- mget, 5
- multiFun, 5
  
- pointsArc (pointsCircle), 6
- pointsCircle, 6
- polar2xy, 7
- polygon, 6
- polygonArc (pointsCircle), 6
- polygonChrom, 8
- polygonDisk (pointsCircle), 6
  
- rotate (polar2xy), 7
  
- wstringapply, 10
  
- xy2polar (polar2xy), 7