# Package 'rtfbsdb'

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Title Parse TF mo	ifs from public databases, read into R, and scan using 'rtfbs'.
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License GPL vers	on 3 or newer
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LazyLoad yes	
R topics doc	imented:
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# **Description**

The motif library from CisBP web site.

Link: http://cisbp.ccbr.utoronto.ca/

## **Objects from the Class**

Objects can be created by calls of the form CisBP.extdata, CisBP.zipload, CisBP.download.

## **Slots**

```
species: String indicating the species name defined in the CisBP dataset.
zip.file: String indicating the filename of temporary data file.
zip.url: String indicating the download source.
```

file.tfinfo: String indicating the TF filename, defulat is TF\_Information.txt.

# **Extends**

```
Class "tfbs.db", directly.
```

## Methods

tfbs.createFromCisBP Build a tfbs object by querying the meta file of CisBP dataset and subsetting the results.

**CisBP.group** Get the statistical summary by grouping the field in the CisBP dataset.

# References

Weirauch, M. T., Yang, A., Albu, M., Cote, A. G., Montenegro-Montero, A., Drewe, P., ... & Hughes, T. R. (2014). Determination and inference of eukaryotic transcription factor sequence specificity. Cell, 158(6), 1431-1443.

# See Also

```
CisBP.group, tfbs.createFromCisBP
```

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#### **Examples**

```
showClass("CisBP.db")
```

CisBP.download

Download CisBP dataset.

## **Description**

Download TF data file from CisBP dataset and store it to temporary folder

## Usage

# **Arguments**

species	String, indicating the species name in the CisBP dataset
url	String, the URL of bulk dowbnloads from CisBP dataset,
	<pre>default is http://cisbp.ccbr.utoronto.ca/bulk_archive.php</pre>

#### **Details**

The dowload function has been confirmed in the web site of cisbp.ccbr.utoronto.ca o June, 2015.

#### Value

A CisBP object (class name: "CisBP.db") is returned with four items:

```
speciesString indicating the species namezip.fileString indicating the filename of temporary data file.zip.urlString indicating the download sourcefile.tfinfoString indicating the TF filename, default is TF_Information.txt.
```

# References

Weirauch, M. T., Yang, A., Albu, M., Cote, A. G., Montenegro-Montero, A., Drewe, P., ... & Hughes, T. R. (2014). Determination and inference of eukaryotic transcription factor sequence specificity. Cell, 158(6), 1431-1443.

# See Also

```
See Also as CisBP.zipload, CisBP.extdata.
```

```
#download human dataset
db1 <- CisBP.download("Homo_sapiens");
#download mouse dataset
db2 <- CisBP.download("Mus_musculus");</pre>
```

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CisBP.extdata Load internal CisBP dataset.

# Description

Build a CisBP object from the internal zip file stored in this package

# Usage

```
CisBP.extdata(species)
```

# **Arguments**

species

String, only valid for human and mouse species, i.e. Homo\_sapiens or Mus\_musculus

# **Details**

The CisBP data for Homo\_sapiens and Mus\_musculus are delivered by this package. When you use the newest dataset, you should download it from the website by CisBP.download.

# Value

A CisBP object (class name: "CisBP.db") is returned with four items:

species	String indicating the species name defined in the CisBP dataset.
zip.file	String indicating the filename of temporary data file.
zip.url	String indicating the download source
file.tfinfo	String indicating the TF filename, default is TF_Information.txt.

## See Also

```
See Also as CisBP.zipload, CisBP.download.
```

```
#reading data from inner file
db.human <- CisBP.extdata("Homo_sapiens")</pre>
```

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CisBP.group

Summarize the motif number.

## **Description**

Get the statistical summary by grouping the field in the CisBP dataset.

#### Usage

## **Arguments**

```
cisbp.db A CisBP object ("CisBP.db") including the TF_Information.txt.

group.by String, indicating which field will be used to group values. Available values are tf_name, tf_species, tf_status, family_name, motif_type and msource_id.

tf.information.type Number, indicating which TF meta file will be used. Available values are 1 for
```

Number, indicating which TF meta file will be used. Available values are 1 for TF\_Information.txt, 2 for TF\_Information\_all\_motifs.txt and 3 for F\_Information\_all\_motifs\_plus.txt.

## **Details**

Three TF information files in CisBP dataset.

- 1: TF\_Information.txt: (direct motifs) or (no direct but inferred motifs with 90%)
- 2: TF\_Information\_all\_motifs.txt: (direct motifs) and (inferred motifs above the threshold)
- 3: F\_Information\_all\_motifs\_plus.txt: All motifs

#### Value

A data frame returned includes two columns

```
group_by Values of grouping field number Counts of group value
```

## See Also

See Also as tfbs.createFromCisBP

```
# Load the internal CisBP dataset
db_human <- CisBP.extdata("Homo_sapiens");
# Group the motif count by the column of family_name in TF_Information.txt
gr1 <- CisBP.group(db_human, group.by="family_name", tf.information.type=1 );</pre>
```

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```
# Group the motif count by the column of tf_status in TF_Information.txt
gr2 <- CisBP.group(db_human, group.by="tf_status", tf.information.type=1);
# Group the motif count by the column of tf_status in TF_Information_all_motifs.txt
gr3 <- CisBP.group(db_human, group.by="tf_status", tf.information.type=2);
# Group the motif count by the column of tf_status in F_Information_all_motifs_plus.txt
gr4 <- CisBP.group(db_human, group.by="tf_status", tf.information.type=3);</pre>
```

CisBP.zipload

Load the zipped CisBP file.

## **Description**

Build a CisBP object from the zipped CisBP file.

#### Usage

```
CisBP.zipload(zip.file, species = "Homo_sapiens")
```

## **Arguments**

zip.file String, indicating the zipped file data
species String, indicating the species name in the CisBP database

#### **Details**

The zip data canbe downloaded from the web site, please check CisBP.download.

# Value

A CisBP object (class name: "CisBP.db") is returned with four items:

```
species String indicating the species name
zip.file String indicating the filename of temporary data file.
zip.url String indicating the download source
file.tfinfo String indicating the TF filename, default is TF_Information.txt.
```

## See Also

```
See Also as CisBP.extdata, CisBP.download.
```

```
# Download the dataset
db2 <- CisBP.download("Mus_musculus");

# Loading the zip file, the db2 and db3 have same TF data.
# Here is an example to show how to use CisBP.zipload.
# We dont nee to download it by CisBP.download and then load it by CisBP.zipload
db3 <- CisBP.zipload(db2@zip.file, species="Mus_musculus");</pre>
```

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```
print.tfbs.comparson
```

Print the brief comparson results

# Description

Print the brief comparson results.

# Usage

```
## S3 method for class 'tfbs.comparson'
print(x, ...)
```

# **Arguments**

- x The result obtained by tfbs.compareTFsite.
- ... Additional arguments affecting the print produced.

## **Details**

This command shows the calling parameters and significant motifs from the result object. The significant motifs are selected by the corrected p-value cutoff(0.05) and at most 20 significant motifs are listed. The adjust method of p-value is defined in the calling function.

# Value

No return values.

## See Also

```
See also as tfbs.compareTFsite.
```

# **Examples**

```
#See example in tfbs.compareTFsite
```

```
print.tfbs.finding Print scanning result of TF sites.
```

# **Description**

Print scanning result of TF sites.

# Usage

```
## S3 method for class 'tfbs.finding'
print(x, ...)
```

#### **Arguments**

x The result obtained by tfbs.scanTFsite.

... Additional arguments affecting the print produced.

# **Details**

This function shows a brief information including calling parameters and enriched motifs.

#### Value

No return values.

# See Also

```
See Also as tfbs.scanTFsite
```

## **Examples**

```
#See example in tfbs.scanTFsite
```

```
summary.tfbs.comparson
```

Summarize the comparson result

# **Description**

Return the significant motifs based on the adjust p-values using multiple comparisons.

# Usage

```
## S3 method for class 'tfbs.comparson'
summary(object, pv.cutoff = 0.05, pv.adj = NA, ...)
```

## **Arguments**

```
object The result obtained by tfbs.compareTFsite.
pv.cutoff The p-value cutoff for significant motifs.
pv.adj P-values adjust method for p.adjust function. The available values are "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr" or "none".
... Additional arguments affecting the summary produced.
```

# **Details**

A data frame with 6 colums is returned.

summary.tfbs.finding 9

## Value

The results is a data frame including 6 columns,

motif.id	Motif ID
tf.name	TF Name
Npos	Read count in positive loci.
Nneg	Read count in negative loci.

pv.adj p-value

es.ratio The ratio of read counts between positive loci and negative loci.

## See Also

See also as tfbs.compareTFsite.

```
summary.tfbs.finding
```

Summarize scanning results.

# **Description**

Return a data frame with summarized TF sites for every motif if the calling parameter is "matches".

# Usage

```
## S3 method for class 'tfbs.finding'
summary(object, ...)
```

# Arguments

object The result obtained by tfbs.scanTFsite.

... Additional arguments affecting the summary produced.

# **Details**

```
summary in class of tfbs.finding is returned.
```

## Value

This function will return a data frame with summarized TF sites for every motif if the calling parameter is "matches", otherwise, NULL will be returned.

## See Also

```
See Also as tfbs.scanTFsite
```

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tfbs

Create a tfbs object from the supplied PWM files.

# **Description**

Create a tfbs object from the supplied PWM files.

## Usage

```
tfbs(filenames,
    names,
    species="Homo_sapiens",
    extra_info = NULL, ...)
```

# **Arguments**

filenames	Vector of PWM files
names	Vector of unique gene symbols.
species	String indicating species name
extra_info	Data frame including meta information for all motifs., Default: NULL
• • •	Parameters, such as pseudocount, force_even, and the parameters used in read.table function.

## **Details**

Load the PWM files to build a "tfbs" object.

#### Value

A tfbs object (class: "tfbs") including all PWM matrics. The all attributes are as follows:

```
TFID
                 Vector of non-unique ID for TF.
                 String indicating the species name
species
                 Number of motifs in matrix.
ntfs
                 A list including PWM matics.
pwm
                 Vector of PWM filename.
filename
mgisymbols
                 Unique gene symbols for TF.
extra_info
                 Data frame, including extra information for PWMs, it maybe different with mo-
                 tif dataset, default:NULL.
distancematrix
                 Distance matrix between motifs returned by tfbs.getDistanceMatrix,
                 default:NULL.
expressionlevel
                 Data frame indicatig the result of expression level returned by tfbs.getExpression,
                 default:NULL.
```

The tfbs object can be created by the function of tfbs, tfbs.dirs, tfbs.createFromCisBP.

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#### See Also

```
tfbs,tfbs.dirs,tfbs.createFromCisBP
```

## **Examples**

tfbs-class

Class "tfbs"

#### **Description**

Tfbs object is a collection of motif PWM data. Some functions are provided based on the PWM and GENCODE data, such as clustering, search and compare.

# **Objects from the Class**

Objects can be created by calls of the function of tfbs.createFromCisBP, tfbs.dirs and tfbs.

## **Slots**

**TFID** Vector of non-unique ID for TF.

species String indicating the species name

ntfs Number of motifs in matrix.

pwm A list including PWM matics.

filename Vector of PWM filename.

mgisymbols Unique gene symbols for TF.

**extra\_info** Data frame, including extra information for PWMs, it maybe different with motif dataset, default:NULL.

**distancematrix** Distance matrix between motifs returned by tfbs.getDistanceMatrix, default:NULL.

**expressionlevel** Data frame indicatig the result of expression level returned by tfbs.getExpression, default:NULL.

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#### Methods

tfbs.getDistanceMatrix Calcuate a distance matrix with Pearson's R values

tfbs.getExpression Estimate gene expression of target TF.

tfbs.clusterMotifs Cluster the specified motifs and drawing the heatmap.

tfbs.scanTFsite Find TF sites from genome data within the BED ranges.

tfbs.compareTFsite Comparative TFBS search with the BED ranges

tfbs.selectByGeneExp Select the motifs with minimum p-value from each group of clustering.

tfbs.selectByRandom Select the motifs randomly from each group of clustering.

tfbs.drawLogosForClusters Draw the motif logos by one group per page.

tfbs.drawLogo Draw the logo for a single TF motif.

# See Also

The class definition of tfbs.

## **Examples**

```
showClass("tfbs")
```

tfbs.clusterMotifs *Clustering the specified motifs and drawing the heatmap*.

# **Description**

Clustering the specified motifs and drawing the heatmap.

## Usage

```
tfbs.clusterMotifs(tfbs,
    subset = NA,
    pdf.heatmap = NA,
    method = NA,
    group.k = NA)
```

## Arguments

 $\label{eq:continuous} A \ tfbs \ object \ ("tfbs") \ returned \ by \ tfbs. create From CisBP, tfbs. dirs$ 

or other functions.

subset Vector, the indexes of partial motifs if not all motifs are clustered.

pdf.heatmap String, a PDF filename for heatmap.

method String, availabe values are "agnes" and "cors".

group.k Integer, if the method of agnes is used to do clustering, the parameter of k is

optional to use as preset group number.

## **Details**

This result of clustering will be used in the

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#### Value

A matrix with 2 columns is returned, 1st column is the index of motifs and 2nd column is the group number of clustering.

#### See Also

See Also as tfbs.selectByGeneExp and tfbs.selectByRandom

# **Examples**

```
# Load the internal CisBP data set
db <- CisBP.extdata("Homo_sapiens");

# Create a tfbs object by querying the meta file of CisBP dataset.
tfs <- tfbs.createFromCisBP(db, motif_type="ChIP-seq", tf.information.type=1);

# Calculate the distance matrix
tfs <- tfbs.getDistanceMatrix( tfs, ncores=1);

# Cluster the motifs using the "cors" method
cluster1 <- tfbs.clusterMotifs(tfs, pdf.heatmap = "test-heatmap1.pdf", method="agnes");
show(cluster1);

# draw motif logos on one group per page.
tfbs.drawLogosForClusters(tfs, cluster1, "test-cluster1.pdf");</pre>
```

 ${\tt tfbs.compareTFsite}\ \textit{Comparative TS sites between positive and negative TRE loci}$ 

# **Description**

Comparative TS sites between positive and negative TRE loci for all motifs.

## Usage

```
tfbs.compareTFsite(tfbs,
    file.twoBit,
    positive.bed,
    negative.bed,
    file.prefix=NA,
    usemotifs=NA,
    ncores=3,
    negative.correction=FALSE,
    fdr=0.1,
    threshold=NA,
    gc.groups=4,
    background.order=2,
    background.length=100000,
    pv.adj = p.adjust.methods)
```

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#### **Arguments**

tfbs A tfbs object, see also "tfbs" String, the file name of genome data(e.g. hg19.2bit, mm10.2bit) file.twoBit positive.bed Data frame, bed-formatted TRE loci. negative.bed Data frame, bed-formatted background loci. file.prefix String, the prefix for outputted BED file, no bed files output if NA usemotifs Vector of index, the index of which motifs are used to compare. All motifs are used if NA. ncores Number, comupting nodes in parallel environment.(default=3) negative.correction Logical value, if the difference between positive and negative TREs is significant,the resampling will be applied to the correction for the negative TREs. (default=FALSE) Numeric value between 0 and 1, False Discovery Rate (FDR) of possible binding fdr sites in rtfbs package, only binding sites with FDR less than this value can be selected.(default=0.1) threshold Numeric value, only sites with scores above this threshold are returned, not be used if NA. (default = NA) Numeric value, indicating number of quantiles to group sequences into in rtfbs gc.groups package. (default = 4)background.order Number, order of Markov model to build background.(default=2). background.length Number, length of the sequence to simulate background.(default=100000). String, P-values correct method for p.adjust function. The available values pv.adj

are "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr" or "none".(default="bonferroni")

### **Details**

The difference of GC contents between positive.bed and negative.bed is checked before the comparson. The p-value of Wilcox test and a vioplot figure show this difference and help the user to determine whether the correction is necessary. If the difference is very significant, please set *background.correction* to do background correction by resampling the TREs from negative bed data based on the frequency of TREs in negative bed data.

#### Value

A object with the class name of "tfbs.comparson" will be resturned in this comparson function. It includes one list of parameters parm and one data frame of results result.

result is a data frame with the following columns:

motif.id Motif ID. tf.name TF name.

Npos TF site count found in positive ranges.

Nneg TF site count found in negative ranges.

es.ratio Enrichment score.

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```
pvalue p-value calculated by fisher test.
pv.adj p-value corrected by the multiple correction.
starch Binary filename of detected TF sites.
```

The result can be outputted to a report by the function tfbs.reportComparson.

#### See Also

```
print.tfbs.comparson, summary.tfbs.comparson, tfbs.reportComparson.
```

```
library(rtfbsdb);
file.twoBit
                <- "/local/storage/data/hg19/hg19.2bit"
db <- CisBP.extdata("Homo_sapiens");</pre>
tfs <- tfbs.createFromCisBP(db, family_name="AP-2");</pre>
#make two dummy BED data frame for positive loci and negative loci
pos.bed <- data.frame(chr="chr1",</pre>
start=round(runif(100,1000000, 2000000)),
stop=0,
name="",
score=0,
strand=".");
pos.bed$stop <- pos.bed$start + 3000;</pre>
neg.bed <- data.frame(chr="chr1",</pre>
start=round(runif(200, 800000, 1800000)),
stop=0,
name="",
score=0,
strand=".");
neg.bed$stop <- neg.bed$start + round(runif(200, 1000, 3000));</pre>
t1 <- tfbs.compareTFsite( tfs,</pre>
      file.twoBit,
      pos.bed,
      neg.bed,
      negative.correction=TRUE,
      ncores = 1); #ncores=3
#Show a brief result
#Show the comparson results of all motifs
show(t1$result);
#Output the result to one pdf report.
tfbs.reportComparson(tfs, t1, file.pdf="test-tfbs-comp.pdf", sig.only=FALSE);
```

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```
tfbs.createFromCisBP
```

Create TF object by querying the CisBP dataset.

# **Description**

Build a tfbs object by querying the meta file of CisBP dataset and subsetting the results.

## Usage

```
tfbs.createFromCisBP(cisbp.db,
      tf name = NULL
      tf status = NULL,
      family_name = NULL,
      motif_type = NULL,
      msource id = NULL,
      tf.information.type = 1,
      expressed.only=TRUE,
      include.DBID.Missing=TRUE,
      seq.datatype=NA,
      file.bigwig.plus=NA,
      file.bigwig.minus=NA,
      file.bam=NA,
      file.twoBit=NA,
      file.gencode.gtf=NA,
      ncores = 1)
```

# **Arguments**

```
A CisBP object("CisBP.db"), including the file of TF_Information.txt.
cisbp.db
                  String, indicating the TF_name field will be used to select motifs.
tf_name
tf_status
                  String, indicating the TF_Status field will be used to select motifs.
                  String, indicating the Family_Name field will be used to select motifs.
family_name
motif_type
                  String, indicating the Motif_Type field will be used to select motifs.
                  String, indicating the MSource_Identifier field will be used to select motifs.
msource_id
tf.information.type
                  Number, indicating which TF meta file will be used. Available values are 1
                  for TF_Information.txt, 2 for TF_Information_all_motifs.txt and 3 for
                  TF_Information_all_motifs_plus.txt.
expressed.only
                  Logical, indicating the only expressed TFs are selected to construct this object
                  based on the gene exprssion values.
include.DBID.Missing
                  Logical, indicating whether the TFs without association with GENCODE through
                  the DBID are selected.
seq.datatype String, indicating which kind of seq data is applied to this function, three values
```

are available: GRO-seq, PRO-seq and RNA-seq. Default: GRO-seq

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```
file.bigwig.plus
```

String, indicating bigwig file for strand plus(+) if seq.datatype is GRO-seq or PRO-seq.

file.bigwig.minus

String, indicating bigwig file for strand minus(-) if seq.datatype is GRO-seq or PRO-seq.

file.bam String, indicating BAM file for rna reads if seq.datatype is RNA-seq.

 $\verb|file.twoBit| String, indicating the binary data of sequence. (e.g. hg19.2bit, mm10.2bit)$ 

file.gencode.gtf

String, indicating Gencode GTF file downloaded from the Gencode web site.

ncores Number, comuputing nodes in parallel environment for gencode data converting.

#### **Details**

The function includes three steps to build a tfbs object:

- 1) Searching the TF information and PWM files in the CisBP dataset according to the criteria specified by the parameters of *tf\_name*, *tf\_status*, *family\_name*, *motif\_type* and *msource\_id*.
- 2) If seq.datatype is GRO-seq or PRO-seq and the bigwig files are provided, the gene expression values are calculated through querying the TREs region from the GENCODE database( for human, gencode.v21.annotation.gtf, for mouse: gencode.vM3.annotation.gtf) and querying the reads count in the plus and minus bigWig files.

If seq.datatype is RNA-seq and the BAM file is provided, read counts for each TRE regions will be queried from the BAM file.

3) If the expressed TFs only is used in the tfbs object, the TFs with p-values corrected by Bonfrroni less than 0.05 will be selected.

The following part explains how to calculate the gene expression.

For each motif, the occurance ranges can be queried by the gene ID After the searching, one range obtained from the merge of the multiple ranges will be used to detect the reads count in the specified bigwig files(including plus and minus). The probability of each motif can be calcuated by the reads count and lambda.

The lambda is determined by the following formulation:

```
r.lambda = 0.04 * sum(reads_in_all_chromosomes)/10751533/1000.
```

The dataset of GENECODE v21 (human) and vM3 (mouse) have been compiled into RDATA file and attached in this package.

The <code>gencode\_transcript\_ext</code> object can be accessed after the following command is executed successfully.

```
load( system.file("extdata", "gencode_v21_transcript_ext.rdata",
```

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#### Value

A tfbs object is returned with PWM matrices, see Also as "tfbs"

#### See Also

See Also as tfbs

#### **Examples**

tfbs.db-class

Class "tfbs.db"

# **Description**

Abstract class for motif dataset. The CisBP class is a son class of tfbs.db.

# **Objects from the Class**

Now code or function can be used to create this class.

#### **Slots**

```
species: Species name.
```

## Methods

No methods defined with class "tfbs.db" in the signature.

## See Also

```
"CisBP.db" inherits this class.
```

tfbs.dirs

## **Examples**

```
showClass("tfbs.db")
```

tfbs.dirs

Create a tfbs object from the folders.

## **Description**

Create a tfbs object from all the PWM files found in the supplied folders.

## Usage

```
tfbs.dirs(...,
    species = "Homo_sapiens",
    args.read.motif = NULL,
    pattern = glob2rx("*.pwm"),
    recursive = FALSE)
```

## **Arguments**

... Multiple strings, one or more folders can be used in this function.

species String, including the species name.

args.read.motif

List, including *pseudocount*, *force\_even* or other parameters used in read.table

function.

pattern String, a character vector specifying regular expression and wlidcards.

recursive Logical, indicating the loading recursively descends into subfolders or not, de-

fault: FALSE.

# **Details**

Two parameters in the list of args.read.motif can be used: pseudocount: log value for zero value in PWM matrix, default is -7. force\_even: whether the PWM matrix with odd size needs to be even.

## Value

A tfbs object collecting all the PWM files in the specified folders. For the details of tfbs object, please see tfbs

## See Also

The structure of tfbs object is described in "tfbs"

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	tfbs.drawLogo	Draw single motif logo.	
--	---------------	-------------------------	--

# **Description**

Draw the motif logos in two models, 1 logo within a page or 1 group within one page.

# Usage

# **Arguments**

tfbs	A tfbs object("tfbs")
file.pdf	String, the file name of PDF report.
index	Vector of number, indicating the motif index.
tf_id	Vector of string, indicating the TF_ID string, TF_ID is one motif attribute in TF_Information.txt. (Default=NULL).
motif_id	Vector of string, indicating the Motif_ID string, Motif_ID is one motif attribute in TF_Information.txt. (Default=NULL).
tf_name	Vector of string, indicating the TF_Name string, TF_Name is one motif attribute in TF_Information.txt. (Default=NULL).
family_name	Vector of string, indicating Family_Name string, Family_Name is one motif attribute in TF_Information.txt. (Default=NULL).
tf_status	String, indicating the TF_status value, TF_status is one motif attribute in TF_Information.txt. (Default=NULL).
groupby	String, indicating the group field is applied to print the motif, each group is printed in one page, the available values are NA, "Family_Name", "TF_Name", "TF_Status" or "Motif_Type". (Default=NA).

# **Details**

Multiple selection is provided for outputting logos. The selected motifs by each criteria will be combined into one set.

Draw the motif logos in two models:

(1) 1 logo within a page (2) 1 group within one page. The motif logos are splitted if motif count is greater than 10.

# Value

No return values.

# See Also

```
See Also as "tfbs"
```

#### **Examples**

```
db <- CisBP.extdata("Homo_sapiens");</pre>
tfs <- tfbs.createFromCisBP(db);</pre>
motif_id
         <- c( "M5604_1.01", "M5441_1.01", "M5162_1.01", "M5352_1.01");
          <- c( "T093250_1.01", "T093251_1.01", "T093252_1.01", "T093253_1.01");
family_name<- c( "p53", "Homeodomain", "Paired box", "Pipsqueak");</pre>
#Draw 10 motif logos from first one.
tfbs.drawLogo(tfs, file.pdf="tfbs.drawLogo1.pdf", index=c(1:10));
#Draw logos for specified Motif_ID, or TF_ID, or TF_Name, or Family_Name
tfbs.drawLogo(tfs, file.pdf="tfbs.drawLogo2.pdf",
motif_id=motif_id,
 tf_id=tf_id,
 tf_name="AP-2",
 family_name=family_name,
 groupby="TF_Status");
#Draw logos for specified TF_Status
tfbs.drawLogo(tfs, file.pdf="tfbs.drawLogo3.pdf", tf_status="D", groupby="TF_Status");
#unlink("tfbs.drawLogo1.pdf");
#unlink("tfbs.drawLogo2.pdf");
#unlink("tfbs.drawLogo3.pdf");
```

tfbs.drawLogosForClusters

Draw the motif logos by clustering.

## **Description**

Draw the motif logos by one cluster per page.

# Usage

```
tfbs.drawLogosForClusters(tfbs, cluster.mat, file.pdf)
```

## **Arguments**

```
tfbs A tfbs object("tfbs").

cluster.mat A matrix with 2 columns returned by tfbs.clusterMotifs, 1st column is the index of motifs and 2nd column is the group number of clustering.

file.pdf String indicating a PDF eilname.
```

# **Details**

It is different with tfbs.drawLogo which is capable of printing out motif logos in group. This group is calculated by the tfbs.clusterMotifs, not is classfied by any group filed.

#### Value

No return value.

#### See Also

```
See Also as tfbs.clusterMotifs
```

## **Examples**

```
# Load the internal CisBP data set
db <- CisBP.extdata("Homo_sapiens");

# Create a tfbs object by querying the meta file of CisBP dataset.
tfs <- tfbs.createFromCisBP(db, motif_type="ChIP-seq", tf.information.type=1 );

# Calculate the distance matrix
tfs <- tfbs.getDistanceMatrix( tfs, ncores=1);

# Cluster the motifs using the "cors" method
cluster1 <- tfbs.clusterMotifs(tfs, pdf.heatmap = "test-heatmap1.pdf", method="cors" );
show(cluster1);

# draw motif logos on one group per page.
tfbs.drawLogosForClusters(tfs, cluster1, "test-cluster1.pdf")</pre>
```

```
tfbs.getDistanceMatrix
```

Calcuate a distance matrix with Pearson's R values.

# **Description**

Compare any two motifs and return a matrix with Pearson's R values.

# Usage

```
tfbs.getDistanceMatrix(tfbs, ncores = 3, BG = log(c(0.25, 0.25, 0.25, 0.25)))
```

## **Arguments**

tfbs A tfbs object("tfbs").

ncores Number, the number of cores to use simultaneously.

BG The log value of probabilities for nucleotide A, C, G and T as Backgroud com-

puting.

# Details

Please do it parallel computation if you can use multi-cores because the calculation takes long time.

#### Value

A tfbs object with new distance matrix (@distancematrix).

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## See Also

```
"tfbs"
```

## **Examples**

```
db <- CisBP.extdata("Homo_sapiens");
tfs <- tfbs.createFromCisBP(db, family_name="AP-2");
tfs0 <- tfbs.getDistanceMatrix(tfs, ncores=1);</pre>
```

tfbs.getExpression Estimate gene expression of target TF.

# **Description**

Gets expression level of target TF.
USE extra\_info\$DBID to find gene information encoded by GENCODE V21

# Usage

```
tfbs.getExpression(tfbs,
    file.bigwig.plus, file.bigwig.minus,
    file.bam=NA,
    file.twoBit=NA,
    file.gencode.gtf=NA,
    seq.datatype=NA,
    ncores =3)
```

# Arguments

```
A tfbs object("tfbs").
tfbs
file.bigwig.plus
                 String, indicating bigwig file for strand plus(+) if seq.datatype is GRO-seq
                 or PRO-seq.
file.bigwig.minus
                 String, indicating bigwig file for strand minus(-) if seq.datatype is GRO-
                 seq or PRO-seq.
file.bam
                 String, indicating BAM file for rna reads if seq.datatype is RNA-seq.
                 String, indicating the binary data of sequence. (e.g. hg19.2bit, mm10.2bit)
file.twoBit
file.gencode.gtf
                 Gencode RDATA file encoded by ths package.
seq.datatype String, indicating which kind of seq data is applied to this function, three values
                 are available: GRO-seq, PRO-seq and RNA-seq. (Default=GRO-seq)
                 Number, comupting nodes in parallel environment.
ncores
```

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#### **Details**

For each motif, the occurance ranges can be queried by the gene ID in the GENCODE database( for human, gencode.v21.annotation.gtf, for mouse: gencode.vM3.annotation.gtf). After the searching, one range obtianed from the merge of the multiple ranges will be used to detect the reads count in the specified bigwig files(including plus and minus). The probability of each motif can be calcuated by the reads count and lambda.

The lambda is determined by the following formulation:

```
r.lambda = 0.04 * sum(reads_in_all_chromosomes)/10751533/1000.
```

The dataset of GENECODE v21 (human) and vM3 (mouse) have been compiled into RDATA file and attached in this package.

The gencode\_transcript\_ext object can be accessed after the following command is executed successfully.

```
load( system.file("extdata", "gencode_human21_transcript_ext.rdata",
```

packac

#### Value

A tbfs object with new expression data frame including the follwing columns:

Motif_ID	Motif_ID from CisBP dataset or other data source.
DBID	DBID from CisBP dataset or other data source.
chr	String, chromosome name.
start	Integer, start postion in which gene ID can be detected.
end	Integer, end postion in which gene ID can be detected.
strand	String, + or -, indicating the strand direction.
bed_length	Integer, the length of range which gene ID can be detected.

reads The reads number queried by BigWig function from the bigwig files( plus and

minus)

lambda The lambda parameter in poison distribution.

prob The probability calculated based on Poisson distribution.

# See Also

```
See Also as "tfbs"
```

```
# Load the internal CisBP data set
db.human <- CisBP.extdata("Homo_sapiens");

# Create a tfbs object by querying the meta file of CisBP dataset.
tfs <- tfbs.createFromCisBP(db.human, motif_type="ChIP-seq", tf.information.type=1 );</pre>
```

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tfbs.reportComparson

Output report for comparson results.

# **Description**

Output comparson results to a PDF report which includes motif names, counts of TF site, p-value, enrichment ratio and motif logos.

# Usage

```
tfbs.reportComparson(tfbs, r.comp,
    file.pdf = NA,
    report.size = "letter",
    report.title = "",
    sig.only = TRUE,
    pv.cutoff = NA,
    pv.adj = NA)
```

#### **Arguments**

tfbs	A tfbs object, see also "tfbs"
r.comp	A result object from the function of ${\tt tfbs.compareTFsite}$
file.pdf	String, the file name of PDF report.
report.size	String, the page size ( default="letter")
report.title	String, the report title.
sig.only	String, indicating whether only significant motifs are outputted or not.( default=TRUE).
pv.cutoff	Numeric value,indicating whether the different cutoff of p-value is applied to select the significant motifs.
pv.adj	String,indicating whether the different correction metod of p-value is applied to select the significant motifs.

# **Details**

The table with 7 columns is outputted into a PDF report within letter size.

Two color bars are used to display p-values and enrichment ratios. Motif logos are shown visually in each row.

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#### Value

No return values.

## See Also

```
tfbs.compareTFsite, summary.tfbs.comparson.
```

## **Examples**

```
# see examples in tfbs.compareTFsite
```

tfbs.reportFinding Make report for scanning results.

# **Description**

Output a PDF report includes motif names, counts of TF site and motif logos.

# Usage

```
tfbs.reportFinding(tfbs,
    r.scan,
    file.pdf = NA,
    report.size = "letter",
    report.title = "")
```

# **Arguments**

```
tfbs A tfbs object, see also "tfbs"

r.scan A result object from the function of tfbs.scanTFsite

file.pdf String, the file name of PDF report.

report.size String, the page size (default="letter")

report.title String, the report title.
```

# **Details**

The table with 4 columns is outputted into a PDF report within letter size. Motif logos are shown visually in each row.

## Value

No return values.

#### See Also

```
tfbs.scanTFsite, print.tfbs.finding
```

```
#See example in tfbs.scanTFsite
```

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tfbs.scanTFsite

Find TF sites from genome data within the BED loci

# Description

Find TF sites from genome data within the BED loci. Please notice that this package does not provided genome data such as hg19.2bit, mm10.2bit.

# Usage

```
tfbs.scanTFsite(tfbs,
    file.twoBit,
    tre.bed,
    return.type=c("matches", "posteriors", "maxposterior", "writedb"),
    file.prefix=NA,
    usemotifs = NA,
    ncores = 3,
    fdr = NA,
    threshold = 6,
    gc.groups = NA,
    background.order = 2,
    background.length = 100000)
```

# **Arguments**

A tfbs object ("tfbs") returned by tfbs.createFromCisBP, tfbs, tfbs.dirs		
String, the file name of genome data( e.g. hg19.2bit or mm10.2bit)		
Data frame, bed-formatted loci information with 6 columns		
String, four available values explained in th details(default = "matches")		
String, the prefix for outputted file, only used when the return.type is writedb		
Vector indicating indexes of motif to be used in scanning.		
Number, computing nodes in parallel environment (default = 3).		
Numeric value between 0 and 1, False Discovery Rate (FDR) of possible binding sites in rtfbs package, only binding sites with FDR less than this value can be selected. If fdr value is assigned, the threshold will be ignored.		
Numeric value, only sites with scores above this threshold are returned in $rtfbs$ package (default = 6).		
Numeric value, indicating number of quantiles to group sequences into in rtfbs package (default = 1).		
background.order		
Numeric value, indicating the order of Markov model to build in rtfbs package (default = 2).		
background.length		
Numeric value, indicating length of the sequence to simulate in rtfbs package (default = 100000)		

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#### **Details**

- (1) Four options are availabl for the function of tfbs.scanTFsite as follows.
  - matches: returns all matching TF sites for all motifs.
  - writedb: writes a bed file with matches sites. Assumes that sort-bed and starch tools are available in \$PATH
  - posteriors: returns the posteriors at each position in bed-formatted loci.
  - maxposterior: returns the max(posterior) in each position in bed-formatted loci.
- (2) In order to make the binary file with the parameter of writedb, make sure that starchcat and sort-bed command (in BEDOPS) can be accessed from R environment. If not, please put the folder in \$PATH.

#### Value

A list object will be returned with the class name of tfbs.finding. The object wraps four sublist as follows:

- 1) parm: Calling parameters (fdr, threshold), gc.groups...).
- 2) bed: Calling bed-formatted loci(tre.bed).
- 3) summary: A data frame including summrized information about matched TF sites for all motifs.
- 4) result: Scanning results which data type is depend on the parameter of return.type.

The option of *matches* returns a list including the result of every motif, which result is BED style data frame with the following columns.

chrom chromosome chromStart start position

chromEnd chromosome end position

name

score The score is given by the log likelihood ratio against the Marklov model(backgound).

strand strand

The option of writedb will return a binary BED filename in which store all bed ranges.

The option of *posteriors* will return a list for each motif returned by score.ms function. Scores represent the motif 'match score', or the product of the probability of observing each base under the motif or background models. Scores are returned under the motif model for all positions in the sequence, on both forward and reverse strands, and under the background model.

The option of *maxposterior* will return a probability matrix which the row indicates the target loci and the column indicates the motif.

#### See Also

```
print.tfbs.finding, summary.tfbs.finding, tfbs.reportFinding.
```

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```
library(rtfbsdb);
file.twoBit
                <- "/local/storage/data/hg19/hg19.2bit"
db <- CisBP.extdata("Homo_sapiens");</pre>
tfs <- tfbs.createFromCisBP(db, family_name="AP-2");</pre>
tre.bed <- data.frame(chr="chr1",</pre>
start=round(runif(10,1000000, 2000000)),
stop=0,
name="",
score=0,
strand=".");
tre.bed$stop <- tre.bed$start + 3000;</pre>
t1 <- tfbs.scanTFsite( tfs,</pre>
file.twoBit,
tre.bed,
file.prefix="test.db",
ncores = 1);
#show a brief information about the result
#show the summary information in the result
show(t1$summary);
#show the matched TF sites for first motif
show(t1$result[[1]]);
#Output a PDF report for all motifs.
tfbs.reportFinding(tfs, t1, file.pdf="Test Results");
t2 <- tfbs.scanTFsite( tfs,
file.twoBit,
tre.bed,
file.prefix="test.db",
return.type="writedb",
ncores = 1);
t2
t3 <- tfbs.scanTFsite( tfs,
file.twoBit,
tre.bed,
return.type="posteriors",
ncores = 1);
t3
t4 <- tfbs.scanTFsite( tfs,
file.twoBit,
tre.bed,
return.type="maxposterior",
ncores = 1);
```

```
t4;
t4$result;
```

```
tfbs.selectByGeneExp
```

Motif selection by gene expression level.

# **Description**

Select the motifs with minimum p-value from each group of clustering.

# Usage

```
tfbs.selectByGeneExp(tfbs, cluster.mat)
```

## **Arguments**

```
tfbs A tfbs object ("tfbs") with the data frame of gene expression level.

cluster.mat A matrix with 2 columns returned by tfbs.clusterMotifs, 1st column is the index of motifs and 2nd column is the group number of clustering.
```

#### **Details**

The function of tfbs.getExpression should be successfully called and the results of gene expression should be returned before this function is called. The indexes of selected motifs will be used in the function of tfbs.compareTFsite or tfbs.scanTFsite.

## Value

A vector of motif indices is returned.

# See Also

See Also as tfbs.selectByRandom, tfbs.getExpression

```
db <- CisBP.extdata("Homo_sapiens");

tfs <- tfbs.createFromCisBP(db, family_name="AP-2");

if(0)
{
    tfs <- tfbs.getExpression(tfs, file.bigwig.plus, file.bigwig.minus, file.hg19);

cluster1 <- tfbs.clusterMotifs(tfs, pdf.heatmap="test-AP2-heatmap.pdf" );

usemotif <- tfbs.selectByGeneExp(tf, cluster1);
}</pre>
```

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```
tfbs.selectByRandom
```

Random motif selection

# **Description**

Select the motifs randomly from each group of clustering.

# Usage

```
tfbs.selectByRandom(tfbs, cluster.mat)
```

# **Arguments**

```
tfbs A tfbs object("tfbs").

cluster.mat A matrix with 2 columns returned by tfbs.clusterMotifs, 1st column is the index of motifs and 2nd column is the group number of clustering.
```

## **Details**

The indexes of selected motifs can be used in the function of tfbs.compareTFsite or tfbs.scanTFsite.

#### Value

A vector of motif indices is returned.

# See Also

```
See Also as tfbs.selectByGeneExp, tfbs.getExpression
```

```
db <- CisBP.extdata("Homo_sapiens");

tfs <- tfbs.createFromCisBP(db, family_name="AP-2");

tfs <- tfbs.getDistanceMatrix(tfs, ncores=1);

cluster1 <- tfbs.clusterMotifs(tfs, pdf.heatmap="test-AP2-heatmap.pdf" );

usemotif <- tfbs.selectByRandom(tfs, cluster1);

show(usemotif);</pre>
```

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