

Package ‘REPTILE’

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Title Regulatory DNA Element Prediction

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Description Predicting regulatory DNA elements based on epigenomic signatures. This package is more of a set of building blocks than a direct solution. REPTILE regulatory prediction pipeline is built on this R package. See <<https://github.com/yupenghe/REPTILE>> for more information.

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URL <https://github.com/yupenghe/REPTILE>

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REPTILE-package	<i>Regulatory Element Prediction</i>
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Description

Predicting DNA regulatory elements based on epigenomic signatures. This package is more of a set of building blocks than a direct solution. REPTILE regulatory prediction pipeline is built on this R package. Please check the url below for details:

<https://github.com/yupenghe/REPTILE>

Details

Accurate enhancer identification is critical for understanding the spatiotemporal transcriptional regulation during development as well as the functional impact of disease-related non-coding genetic variants. REPTILE is a algorithm to identify the precise location of enhancers by integrating histone modification data and base-resolution DNA methylation profiles.

REPTILE was designed based on three observations: 1) regions that are differentially methylated (or differentially methylated regions, DMRs) across diverse cell and tissue types strongly overlap with enhancers. 2) With base-resolution DNA methylation data, the boundaries of DMRs can be accurately defined, circumventing the difficulty of determining enhancer boundaries. 3) DMR size is often smaller (~500bp) than known enhancers, known negative regions (regions with no observable enhancer activity) and genomic windows used in enhancer prediction (~2kb), all of which we termed as "query regions". Together with the association between transcription factor binding and DNA methylation level, DMRs may serve as high-resolution enhancer candidates and capture the local epigenomic patterns that would otherwise be averaged/washed out in analysis focusing on the query regions.

Running REPTILE involves four major steps. First, to identify DMRs, we compared the methylomes of target sample (where putative enhancers will be generated) and several other samples with different cell/tissue types (as reference). In the next step, input files for REPTILE are prepared, which store the information of query regions, DMRs and the epigenomic data. Taking these inputs, REPTILE represents each DMR or query region as a feature vector, where each element corresponds to either intensity or intensity deviation of one epigenetic mark. Intensity deviation is defined as the intensity in target sample subtracted by the mean intensity in reference samples (i.e. reference epigenome) and it captures the tissue-specificity of each epigenetic mark. In the third step, based on the feature vectors of known enhancers and negative regions as well as the feature vectors of the DMRs within them, we trained an enhancer model, containing two random forest classifiers, which respectively predict enhancer activities of query regions and DMRs. In the last step, REPTILE uses the enhancer model to calculate enhancer confidence scores for DMRs and query regions, based on which the final predictions are made.

The two key concepts on REPTILE are:

- Query regions - known enhancers, known negative regions and genomic windows used for enhancer prediction
- DMRs - differentially methylated regions

In REPTILE, DMRs are used as high-resolution candidates to capture the fine epigenomic signatures in query regions.

Author(s)

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References

He, Yupeng et al., *REPTILE: Regulatory Element Prediction based on Tissue-specific Local Epigenetic marks*, in preparation

Examples

```
library("REPTILE")
data("rsd")

## Training (needs a few minutes and ~1.8 Gb memory)
reptile.model <- reptile_train(rsd$training_data$region_epimark,
                             rsd$training_data$region_label,
                             rsd$training_data$DMR_epimark,
                             rsd$training_data$DMR_label,
                             ntree=50)

## Prediction
## - REPTILE
pred <- reptile_predict(reptile.model,
                      rsd$test_data$region_epimark,
                      rsd$test_data$DMR_epimark)

## - Random guessing
pred_guess = runif(length(pred$D))
names(pred_guess) = names(pred$D)

## Evaluation
res_reptile <- reptile_eval_prediction(pred$D,
                                     rsd$test_data$region_label)
res_guess <- reptile_eval_prediction(pred_guess,
                                   rsd$test_data$region_label)

## - Print AUROC and AUPR
cat(paste0("REPTILE\n",
          " AUROC = ",round(res_reptile$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_reptile$AUPR,digit=3))
    ,"\n")
cat(paste0("Random guessing\n",
          " AUROC = ",round(res_guess$AUROC,digit=3),
          "\n",
```

```

" AUPR = ",round(res_guess$AUPR,digit=3))
","\n")

```

```
calculate_epimark_deviation
```

Internal - calculating intensity deviation feature

Description

Internal function used to calculate the intensity deviation features. It is based on the epigenomic signatures of a given region in target sample, where prediction will be generated, and reference samples. Intensity deviation is defined as the intensity in target sample subtracted by the mean intensity in reference samples (i.e. reference epigenome) and it captures the tissue-specificity of each epigenetic mark.

Usage

```
calculate_epimark_deviation(data_info, x, query_sample,
                           ref_sample = NULL)
```

Arguments

data_info	data.frame instance generated by reading data information file specifying the samples and marks used in the analysis. The data.frame includes at least two columns named "sample" and "mark", corresponding to the samples and marks included.
x	data.frame instance generated by reading epimark file. The first four columns of the data.frame are "chr", "start", "end" and "id" of each region in the epimark file. The rest columns contain values of epigenetic marks in samples as specified in data_info and column names are under MARK_SAMPLE format, such as "H3K4me1_mESC".
query_sample	name of the target sample
ref_sample	a vector of names of the reference sample(s)

Value

data.frame instance containing intensity deviation values of each mark

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

See Also

[read_epigenomic_data](#)

get_option_parser_compute_score

Internal - parsing options for REPTILE_compute_score.R

Description

Internal function used to parsing options for "REPTILE_compute_score.R" script in the REPTILE enhancer prediction pipeline:

<https://github.com/yupenghe/REPTILE>

Usage

```
get_option_parser_compute_score()
```

Value

An instance of the OptionParser class.

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

get_option_parser_evaluation

Internal - parsing options for REPTILE_evaluate_prediction.R

Description

Internal function used to parsing options for "REPTILE_evaluate_prediction.R" script in the REPTILE enhancer prediction pipeline:

<https://github.com/yupenghe/REPTILE>

Usage

```
get_option_parser_evaluation()
```

Value

An instance of the OptionParser class.

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

`get_option_parser_training`*Internal - parsing options for REPTILE_train.R*

Description

Internal function used to parsing options for "REPTILE_train.R" script in the REPTILE enhancer prediction pipeline:

<https://github.com/yupenghe/REPTILE>

Usage

```
get_option_parser_training()
```

Value

An instance of the OptionParser class.

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

`read_epigenomic_data` *Reading epigenomic data from epimark file*

Description

Function to read epimark file from disk and generate data.frame instance. It is used to read epigenomic data from file on disk and generate the input data.frame instance to fuel the model training, prediction and other following steps. Epimark file is a tab-separated file with a header. The first four columns are "chr", "start", "end" and "id", specifying the chromosome, start, end and id of regions. Each of the remaining columns contain values of one epigenetic mark in one sample (condition, cell or tissue type, etc) and the column name follows "MARK_SAMPLE" format, such as "H3K4me1_mESC".

Usage

```
read_epigenomic_data(data_info, epimark_file, query_sample,  
                    ref_sample = NULL, incl_dev = T)
```

Arguments

data_info	data.frame generated by reading data information file specifying the samples and marks used in the analysis. The data.frame includes at least two columns named "sample" and "mark", corresponding to the samples and marks included.
epimark_file	name of epimark file
query_sample	name of the target sample
ref_sample	a vector of names of the reference sample(s)
incl_dev	logical value indicates whether to calculate the intensity deviation feature. Intensity deviation is defined as the intensity in target sample subtracted by the mean intensity in reference samples (i.e. reference epigenome) and it captures the tissue-specificity of each epigenetic mark.

Value

data.frame instance containing intensity and intensity deviation values of each mark for each region

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

See Also

[read_label](#)

read_label

Reading labels of regions from label file

Description

Function to read epimark file from disk and generate data.frame instance. It is used to read epigenomic data from file on disk and generate the input data.frame instance to fuel the model training, prediction and other following steps. Label file is a tab-separated file with a header. The first column contains the id of each region. The second or more columns specify whether a certain region is enhancer (1) or not (0) in a specific sample. Each of these columns corresponds to one sample and the name of the column is the sample name.

Usage

```
read_label(label_file, query_sample)
```

Arguments

label_file	name of label file on disk
query_sample	name(s) of sample(s), in which you would like to have label information

Value

an data.frame instance containing label of each region in query samples The possible values and their meanings of a label are:

NA - unknwon (will be ignored)

0 - not enhancer

1 - enhancer

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

See Also

[read_epigenomic_data](#)

reptile_eval_prediction

Evaluating the prediction results

Description

Function used to evaluate the predictions by comparing enhancer scores from reptile_predict or reptile_predict_genome_wide and the correct labels. Area under the Receiver Operating Characteristic (ROC) curve (AUROC) and Area under the Precision-Recall curve (AUPR) will be calculated.

Usage

```
reptile_eval_prediction(predictions, annotations)
```

Arguments

predictions	vector of enhancer scores for regions. The name of each value (score) corresponds to the id of the region.
annotations	vector of labels for regions with the same length as predictions. The name of each value (label) corresponds to the id of the region. Only two values are allowed in annotations: 0 (negative) and 1 (positive). No NA is allowed.

Value

A list containing two numbers

AUROC Area under the Receiver Operating Characteristic (ROC) curve

AUPR Area under the Precision-Recall curve

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

See Also

[reptile_predict](#), [reptile_predict_genome_wide](#)

Examples

```
library("REPTILE")
data("rsd")

## Training
rsd_model <- reptile_train(rsd$training_data$region_epimark,
                          rsd$training_data$region_label,
                          rsd$training_data$DMR_epimark,
                          rsd$training_data$DMR_label,
                          ntree=50)

## Prediction
## - REPTILE
pred <- reptile_predict(rsd_model,
                       rsd$test_data$region_epimark,
                       rsd$test_data$DMR_epimark)

## - Random guessing
pred_guess = runif(length(pred$D))
names(pred_guess) = names(pred$D)

## Evaluation
res_reptile <- reptile_eval_prediction(pred$D,
                                      rsd$test_data$region_label)
res_guess <- reptile_eval_prediction(pred_guess,
                                    rsd$test_data$region_label)

## - Print AUROC and AUPR
cat(paste0("REPTILE\n",
          " AUROC = ",round(res_reptile$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_reptile$AUPR,digit=3))
    ,"\n")
cat(paste0("Random guessing\n",
          " AUROC = ",round(res_guess$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_guess$AUPR,digit=3))
    ,"\n")
```

Description

Predicting enhancer activities of query regions based on the enhancer model from `reptile_train` in training step. This function calculates the combined enhancer score for each query region (given region) as the maximum among the score of whole query region and the scores of DMRs within it. This function is for generating genome-wide enhancer predictions.

Usage

```
reptile_predict(reptile_model,  
               epimark_region,  
               epimark_DMR = NULL,  
               family = "randomForest")
```

Arguments

<code>reptile_model</code>	Enhancer model from <code>reptile_train</code> . It is a list containing two objects of class <code>randomForest</code> or <code>glm</code> when <code>family</code> is set to be "Logistic"
<code>epimark_region</code>	data.frame instance from <code>read_epigenomic_data</code> , which containing intensity and intensity deviation values of each mark for each query region
<code>epimark_DMR</code>	data.frame instance from <code>read_epigenomic_data</code> , which containing intensity and intensity deviation values of each mark for each DMR
<code>family</code>	classifier family used in the enhancer model Default: <code>RandomForest</code> Classifiers available: - <code>RandomForest</code> : random forest - <code>Logistic</code> : logistic regression

Value

A list containing three vectors

D	Combined enhancer score of each query region
R	Enhancer score of each query region
DMR	Enhancer score of each DMR

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

See Also

[reptile_predict_genome_wide](#)
[reptile_train](#)
[read_epigenomic_data](#)
[read_label](#)

Examples

```

library("REPTILE")
data("rsd")

## Training
rsd_model <- reptile_train(rsd$training_data$region_epimark,
                          rsd$training_data$region_label,
                          rsd$training_data$DMR_epimark,
                          rsd$training_data$DMR_label,
                          ntree=50)

## Prediction
## - REPTILE
pred <- reptile_predict(rsd_model,
                       rsd$test_data$region_epimark,
                       rsd$test_data$DMR_epimark)

## - Random guessing
pred_guess = runif(length(pred$D))
names(pred_guess) = names(pred$D)

## Evaluation
res_reptile <- reptile_eval_prediction(pred$D,
                                      rsd$test_data$region_label)
res_guess <- reptile_eval_prediction(pred_guess,
                                    rsd$test_data$region_label)

## - Print AUROC and AUPR
cat(paste0("REPTILE\n",
          " AUROC = ",round(res_reptile$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_reptile$AUPR,digit=3))
    ,"\n")
cat(paste0("Random guessing\n",
          " AUROC = ",round(res_guess$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_guess$AUPR,digit=3))
    ,"\n")

```

reptile_predict_genome_wide

Predicting enhancer activity

Description

Predicting enhancer activities of query regions based on the enhancer model from `reptile_train` in training step. This function calculates the enhancer scores of DMRs and query regions. It does not try to generate combined enhancer scores.


```

                                rsd$training_data$DMR_label,
                                ntree=50)

## Prediction
## - REPTILE
pred <- reptile_predict(rsd_model,
                       rsd$test_data$region_epimark,
                       rsd$test_data$DMR_epimark)

## - Random guessing
pred_guess = runif(length(pred$D))
names(pred_guess) = names(pred$D)

## Evaluation
res_reptile <- reptile_eval_prediction(pred$D,
                                      rsd$test_data$region_label)
res_guess <- reptile_eval_prediction(pred_guess,
                                    rsd$test_data$region_label)

## - Print AUROC and AUPR
cat(paste0("REPTILE\n",
          " AUROC = ",round(res_reptile$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_reptile$AUPR,digit=3))
    ,"\n")
cat(paste0("Random guessing\n",
          " AUROC = ",round(res_guess$AUROC,digit=3),
          "\n",
          " AUPR = ",round(res_guess$AUPR,digit=3))
    ,"\n")

```

reptile_predict_one_mode

Internal - predicting enhancer activity of DMRs or query regions

Description

Internal function used to predict the enhancer activity of either DMRs or query regions.

Usage

```

reptile_predict_one_mode(reptile_classifier,
                        epimark,
                        family)

```

Arguments

reptile_classifier An object of class `randomForest` or `glm` when family is set to be "Logistic".

epimark `data.frame` instance from `read_epigenomic_data`, which containing intensity and intensity deviation values of each mark for each query region

family classifier family used in the enhancer model
 Default: RandomForest
 Classifiers available:
 - RandomForest: random forest
 - Logistic: logistic regression

Value

A vector of enhancer score of each query region or DMR

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

See Also

[reptile_predict](#), [reptile_predict_genome_wide](#)

 reptile_train

Learn a REPTILE enhancer model

Description

Learn a REPTILE enhancer model based on epigenomic signature of known enhancers.

Usage

```
reptile_train(epimark_region, label_region,
              epimark_DMR = NULL, label_DMR = NULL,
              family = "randomForest", ntree = 2000,
              nodesize = 1)
```

Arguments

epimark_region data.frame instance from read_epigenomic_data, which containing intensity and intensity deviation values of each mark for each query region.

label_region factor instance from read_label, containing the label of each query region. The possible values and their meanings of a label are: 0 (not enhancer), 1 (enhancer) and NA (unknown and it will be ignored).

epimark_DMR data.frame instance from read_epigenomic_data, which containing intensity and intensity deviation values of each mark for each DMR. If either this value or label_DMR is NULL, the output enhancer model will not include a classifier for predicting the enhancer activities of DMRs. Default: NULL


```

                                ntree=5)

print(rsd_model$D)
print(rsd_model$R)

```

reptile_train_one_mode

Internal - Learn single random forest classifier

Description

Internal function to learn a random forest classifier

Usage

```

reptile_train_one_mode(epimark, label,
                       family, ntree, nodesize)

```

Arguments

epimark	data.frame instance from read_epigenomic_data, which containing intensity and intensity deviation values of each mark for each DMR or query region.
label	factor instance from read_label, containing the label of each query region. The possible values and their meanings of a label are: 0 (not enhancer), 1 (enhancer) and NA (unknwon and will be ignored).
family	Classifier family used in the enhancer model Default: RandomForest Classifiers available: - RandomForest: random forest - Logistic: logistic regression
ntree	Number of tree to be constructed in the random forest model. See the function randomForest() in "randomForest" package for more information. Default: 2000
nodesize	Minimum size of terminal nodes. See the function randomForest() in "randomForest" package for more information. Default: 1

Value

An randomForest object or glm object when family is set to be "Logistic".

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

References

Breiman, L. (2001), *Random Forests*, Machine Learning 45(1), 5-32.
A. Liaw and M. Wiener (2002), *Classification and Regression by randomForest*, R News 2(3), 18–22.

See Also[reptile_train](#)

`rsd`*REPTILE sample data (rsd)*

Description

sample data for testing REPTILE training, prediction and evaluation.

Usage

```
data(rsd)
```

Format

A list containing two lists.

`training_data` is the data used for training REPTILE enhancer model. This list has four elements: `region_epimark`, `DMR_epimark`, `region_label` and `DMR_label`. The former two store the epigenomic signatures of query regions and DMRs. The latter two label which a certain query region or DMR is enhancer (1) or negative instance (0)

`test_data` is for training REPTILE enhancer model and it has four elements: `region_epimark`, `DMR_epimark` and `region_label`. The former two store the epigenomic signatures of query regions and DMRs. The `region_label` indicates whether a certain query region or DMR is enhancer (1) or negative instance (0)

Author(s)

Yupeng He <yupeng.he.bioinfo@gmail.com>

Source

`training_data` was based on the EP300 binding sites (positives), promoters (negatives) and randomly chosen genomic loci (negatives) in mouse embryonic stem cells.

The `test_data` data was constructed based on *in vivo* validated mouse sequences from VISTA enhancer browser (Oct 24th, 2015). The labels indicate the activity in mouse heart tissues from E11.5 embryo.

See the papers included in References for details.

References

He, Yupeng et al., *REPTILE: Regulatory Element Prediction based on Tissue-specific Local Epigenetic marks*, in preparation

Visel, Axel et al. (2007), *VISTA Enhancer Browser - a database of tissue-specific human enhancers* Nucleic acids research 35. suppl 1 <http://enhancer.lbl.gov/>

Examples

```

## Visualizing rsd data
library("REPTILE")
data(rsd)

## Epigenomic signature of query region grouped by labels
ind_pos = rsd$training_data$region_label == 1
pos_region = rsd$training_data$region_epimark[ind_pos,]
neg_region = rsd$training_data$region_epimark[!ind_pos,]

## Epigenomic signature of DMRs grouped by labels
ind_pos = rsd$training_data$DMR_label == 1
pos_DMR = rsd$training_data$DMR_epimark[ind_pos,]
neg_DMR = rsd$training_data$DMR_epimark[!ind_pos,]

## Prepare the data format required for plotting
n = ncol(rsd$training_data$DMR_epimark) ## Number of features
feature_data_DMR = list()
feature_data_region = list()
for(i in 1:n){
  feature_data_DMR <- append(feature_data_DMR,
                             list(neg_DMR[i],pos_DMR[i],
                                   NA,NA))
  feature_data_region <- append(feature_data_region,
                                list(neg_region[i],pos_region[i],
                                      NA,NA))
}

## Plot the feature distribution
par(mar=c(4,8,4,4))
## - query region
b <- boxplot(feature_data_region,
              xlab = "feature value",
              notch=TRUE,outline=FALSE,yaxt='n',
              xlim = c(1,n*4-2),ylim=c(-7,7),
              horizontal=TRUE,
              col=c(rgb(65,105,225,max=255),rgb(250,128,114,max=255)),
              main = "Feature value distribution in query regions"
            )
text(par("usr")[1]-0.2, seq(1.5,n*4-2,by=4),
     labels=gsub("_","-",colnames(rsd$training_data$region_epimark)),
     xpd = TRUE,adj=1)
legend(-8,4*n+4,c("negative","enhancer"),ncol=2,
      fill = c(rgb(250,128,114,max=255),rgb(65,105,225,max=255)),
      xpd=TRUE,bty='n')

## - DMR
b <- boxplot(feature_data_DMR,
              xlab = "feature value",
              notch=TRUE,outline=FALSE,yaxt='n',
              xlim = c(1,n*4-2),ylim=c(-7,7),
              horizontal=TRUE,

```

```
col=c(rgb(65,105,225,max=255),rgb(250,128,114,max=255)),
main = "Feature value distribution in DMRs"
)
text(par("usr")[1]-0.2, seq(1.5,n*4-2,by=4),
labels=gsub("_","-",colnames(rsd$training_data$DMR_epimark)),
xpd = TRUE,adj=1)
legend(-8,4*n+4,c("negative","enhancer"),ncol=2,
fill = c(rgb(250,128,114,max=255),rgb(65,105,225,max=255)),
xpd=TRUE,bty='n')
```

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