Introduction for FunTFPair package

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1 Introduction

Summary: Transcription factors (TFs) are fundamental regulators of gene expression and generally function in a complex and cooperative manner. Identifying context-dependent cooperative TFs is essential for understanding how cells respond to environmental change. The huge amount of various omics data currently available, providing genome-wide physical binding and functional effect information about transcription, holds a great opportunity to study TF cooperativity. Here we developed an R package, FunTFPair, which provides an easy and powerful way to identify condition-specific TF pairs by integrating transcription factor binding sites from ENCODE and gene expression profiles from GEO. Users only need to provide GEO ID that they are interested in. FunTFPair will automatically retrieve the expression profiles of the input GEO ID, get TF targets from ENCODE, and identify TF pairs whose common targets show statistically significant differences under changed experimental conditions or exhibit coordinated transcription in a specific condition. The functional TF pairs and their relative importance will be reported in a TF cooperation network. Two datasets from GEO are used as examples to demonstrate the usage and the reliability of the package.

Availability and Implementation: FunTFPair is implemented in R language. The source code is provided at https://github.com/slzhao/FunTFPair. A more detailed vignette is also released along the package.

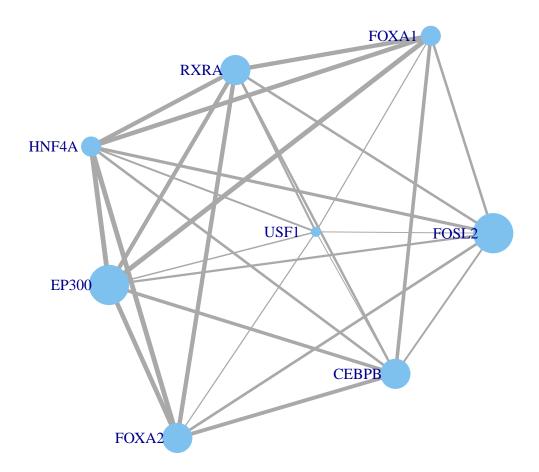
2 Functions in the package

- prepareGeoData: A function to download GDS or GSE data and convert ID if needed;
- differential Analysis: A function to perform differential analysis and then find the enrichment of target genes;

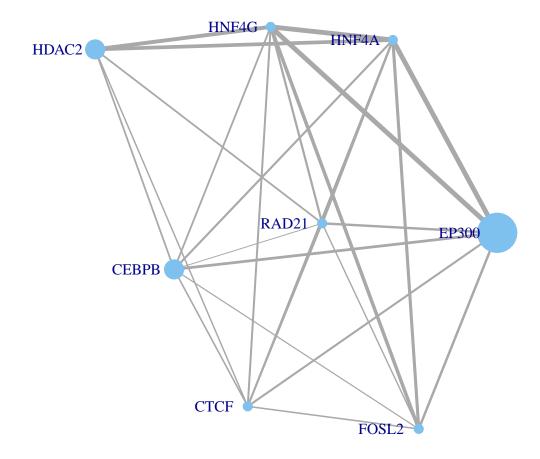
- correlationAnalysis: A function to perform differential analysis for expression data, to see if the shared targets of two tf were significantly changed between two conditions;
- network Vis: A function to perform functional TF network visualization;

3 Example for differential analysis based functional TF pairs identification

Here we used one dataset (GDS2213) profiles gene expression of HepG2 cells treated by 5-aza-2'-deoxycytidine (5-aza-dC), trichostatin A (TSA) or both, as an example of differential analysis-based functional TF pairs identification. The gene expression profiles under three kinds of treatment were compared with those in the control group, respectively. As a result, 24 TF pairs, involving 8 TFs, were identified to be functionally important under the treatment of 5-aza-dC, which inhibit DNA methylation. Consistently, most of TFs has been reported to be influenced by DNA methylation. More interestingly, HDAC2, an important histone deacetylase, joined the functional TF network when TSA, inhibitor of histone decetylation, was added.



Network visualization for azaAndTSA vs untreated
enrichTargetResultNetwork3 <- networkVis(enrichTargetResult[[3]])</pre>



Here the size of nodes indicated the importance of the TF in the cooprative network; while the width of the edges between pairs indicated the association between two TFs, which is the negative log 10 p value for the enrichement of shared targets in significantly changed genes.

4 Example for correlation analysis based functional TF pairs identification

You can also use *correlationAnalysis* function to perform correlation analysis based functional TF pairs identification. THe example codes can be viewed by following R codes.

example(correlationAnalysis, run.dontrun = TRUE)