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Abstract

R package PAGnet is an R package to provide the user with *Pseudomonas aeruginosa* genomic integrated regulatory network visualization, subnetwork filtering and master regulator analysis in his/her comoputer, consistent with the functionality which deployed on the online website (http://pagnetwork.org). The package could facilitate visualization and exploration of regulatory network, as well as master regulator analysis for identification of key transcription factors mediating a biological process or pathway.

Package

PAGnet 0.1.0

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1 Overview

Regulatory networks including virulence-related transcriptional factors (TFs) determine bacterial pathogenicity in response to different environmental cues. *Pseudomonas aeruginosa*, a Gram-negative opportunistic pathogen of humans, recruits numerous TFs in quorum sensing (QS) system, type III secretion system (T3SS) and Type VI secretion system (T6SS) to mediate the pathogenicity. Although many virulence-related TFs have been illustrated individually, very little is known about their crosstalks and regulatory network. Here, based on chromatin immunoprecipitation coupled with high-throughput sequencing (ChIP-seq) and transcriptome profiling (RNA-seq), we primarily focused on understanding the crosstalks of 20 virulence-related TFs, which led to construction of a virulence regulatory network named PAGnet (*Pseudomonas aeruginosa* genomic integrated regulatory network) including 82 crosstalk targets.

The PAGnet uncovered the intricate mechanism of virulence regulation and revealed master regulators in QS, T3SS and T6SS pathways. The package **PAGnet** is designed for network visualization, subnetwork filtering and master regulator analysis locally by running a local shniy GUI within R. In addition, the master regulator analysis can be performed in the R console without running the shiny GUI to provide more flexibility.

Before starting to use this package, you need to install the following packages:

```
library(shiny)
library(shinythemes)
library(scales)
```

Then load the package:

```
## loading package
library(PAGnet)
```

2 Quick Start

2.1 Master Regulator Analysis

The user can choose to use the default PAGnet or to upload their own regulatory network in a predefined format (Locus Tag).

A dataframe format of default **PAGnet** with two colmn transcription factors and targets is provided for MRA, user could also their own regulatory network in same format as input.

```
data(PAGnet)

##Use defaultPAGnet as regulatory network

head(pagnet)

## TranscriptionFactor Target

## 1 PA5261 PA0510

## 2 PA5261 PA1048
```

```
## 3 PA5261 PA1590

## 4 PA5261 PA3117

## 5 PA5261 PA3637

## 6 PA5261 PA3724
```

The users also need to provide a character vector of TFs (or only interested TFs).

```
## define transcription factors in regulatory network

head(tf)
## AlgR AmrZ BfmR CdpR ExsA FleQ
## "PA5261" "PA3385" "PA4101" "PA2588" "PA1713" "PA1097"
```

A character vector of locus tag should be provide as signature genes.

```
##Use QS related genes as signature genes

head(qs)
## rhlC rhlG rhlI rhlR rhlA rhlB
## "PA1130" "PA3387" "PA3476" "PA3477" "PA3479" "PA3478"
```

The function pagnet.mra is used to perform MRA over provided regulatory network. The MRA computes the overlap between the transcriptional regulatory unities (regulons) and the input signature genes using the hypergeometric distribution (with multiple hypothesis testing corrections). Having completed master regulator analysis, a table will be returned.

```
## Perform MRA and return results
 mra_results <- pagnet.mra(rnet=pagnet,tflist=tf,signature = qs,</pre>
                         pValueCutoff = 0.05)
 mra_results
   TF network.size regulon.size signature.size observed.signature.size
## 14 PA3477 351
                               5
## 9 PA1430
                   351
                               49
                                             13
## 15 PA1431
                  351
                               11
                                             13
                                                                    3
## 11 PA1003
                   351
                                17
                                                                    3
   Pvalue
## 14 4e-04
## 9 0.0044
## 15 0.0055
## 11 0.02
```

2.2 Local shiny interface

The function pagnet.mra.interface is used to call **local shiny GUI** (Figure 1).

```
## call local shony interfact
pagnet.mra.interface()
```



PAGnet: P. aeruginosa Genomic integrated regulatory network

(Verison 1.1)

Pseudomonas aeruginosa, a gram-negative opportunistic pathogen of humans, recruits more than twenty TFs in quorum sensing (QS) system, type III secretion system (T3SS) and type VI secretion system (T6SS) to mediate the virulence in P. aeruginosa. Here, based on chromatin immunoprecipitation coupled with high-throughput sequencing (ChIP-seq) and transcriptome profiling (RNA-seq) to construct a genomic integrated regulatory network named PAGnet. The current network includes regulatory relationships between 20 transcription factors and 348 their functional target genes, integrated based on our ChIP-seq and RNA-seq data analysis as well as literature curation.

Figure 1: A screenshot of local shiny interface of PAGnet

The shiny GUI provides visualization and exploration of PAGnet (Figure 2). It allows the user to filter the full network by selecting one or multiple transcription factor(s) to obtain a subnetwork of interest. A brief summary of the subnetwork is also provided with information about the transcription factors and their target genes.

The local shiny GUI also provides master regulator analysis for identification of key transcription factors mediating a biological process or pathway (Figure 3). First, the user can choose to use the default PAGnet or to upload their own regulatory network in a predefined format. Second, the user needs to specify a gene signature associated with a biological function or pathway of interest, either by selecting a gene set from public databases or uploading a user-customized gene list. In the current version, the platform provides gene sets in Gene Ontology (GO) and KEGG databases obtained from Pseudomonas Genome DB (Winsor et al. (2015)). Having completed master regulator analysis, a table will be returned with information about each transcription factor's corresponding gene ID, gene name, number of target genes, total number of hits (all signature genes in the network), observed hits (signature genes in the TF's regulon),

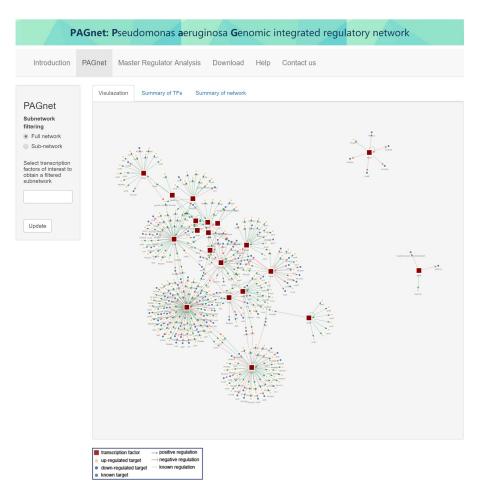


Figure 2: Visualization of PAGnet in local shiny GUI

and a p-value calculated based on a hypergeometric test. The table is sorted according to the statistical significance indicated by the p-values, and the top significant TFs can be prioritized as master regulators.

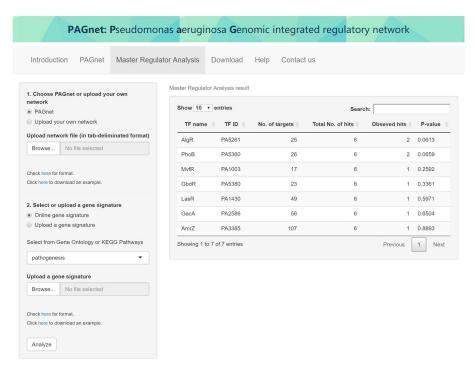


Figure 3: Master Regulator Analysis

3 Need help?

If you have any question/issue, please feel free to contact us.

```
Dr. Xin Deng (xindeng@cityu.edu.hk)

or

Dr. Xin Wang (xin.wang@cityu.edu.hk)
```

4 Session Information

```
## R version 3.5.3 (2019-03-11)
## Platform: x86_64-pc-linux-gnu (64-bit)
## Running under: Ubuntu 18.04.2 LTS
## Matrix products: default
## BLAS: /usr/lib/x86_64-linux-gnu/openblas/libblas.so.3
## LAPACK: /usr/lib/x86_64-linux-gnu/libopenblasp-r0.2.20.so
##
## locale:
## [1] LC_CTYPE=en_US.UTF-8
                                   LC_NUMERIC=C
## [3] LC_TIME=en_US.UTF-8
                                   LC_COLLATE=en_US.UTF-8
## [5] LC_MONETARY=en_US.UTF-8
                                   LC_MESSAGES=en_US.UTF-8
## [7] LC_PAPER=en_US.UTF-8
                                   LC_NAME=C
```

```
## [9] LC_ADDRESS=C
                                                 LC_TELEPHONE=C
## [11] LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C
## attached base packages:
## [1] stats
                      graphics grDevices utils
                                                                 datasets methods
                                                                                             base
## other attached packages:
## [1] PAGnet_0.1.0
                                   scales_1.0.0
                                                            shinythemes_1.1.2 shiny_1.2.0
## [5] BiocStyle_2.10.0
## loaded via a namespace (and not attached):
## [1] Rcpp_0.12.18 knitr_1.21 magrittr_1.5
## [4] munsell_0.5.0 colorspace_1.3-2 xtable_1.8-3
## [7] R6_2.3.0 stringr_1.3.1 tools_3.5.3
## [10] xfun_0.4 htmltools_0.3.6 yaml_2.2.0
## [13] digest_0.6.15 bookdown_0.8 BiocManager_1.30.4
## [16] later_0.7.3 promises_1.0.1 evaluate_0.12
## [19] mime_0.6 rmarkdown_1.11 stringi_1.2.3
## [22] compiler_3.5.3
                                     httpuv_1.4.4.1
```

References

Winsor, Geoffrey L, Emma J Griffiths, Raymond Lo, Bhavjinder K Dhillon, Julie A Shay, and Fiona SL Brinkman. 2015. "Enhanced Annotations and Features for Comparing Thousands of Pseudomonas Genomes in the Pseudomonas Genome Database." *Nucleic Acids Research* 44 (D1). Oxford University Press: D646–D653.