

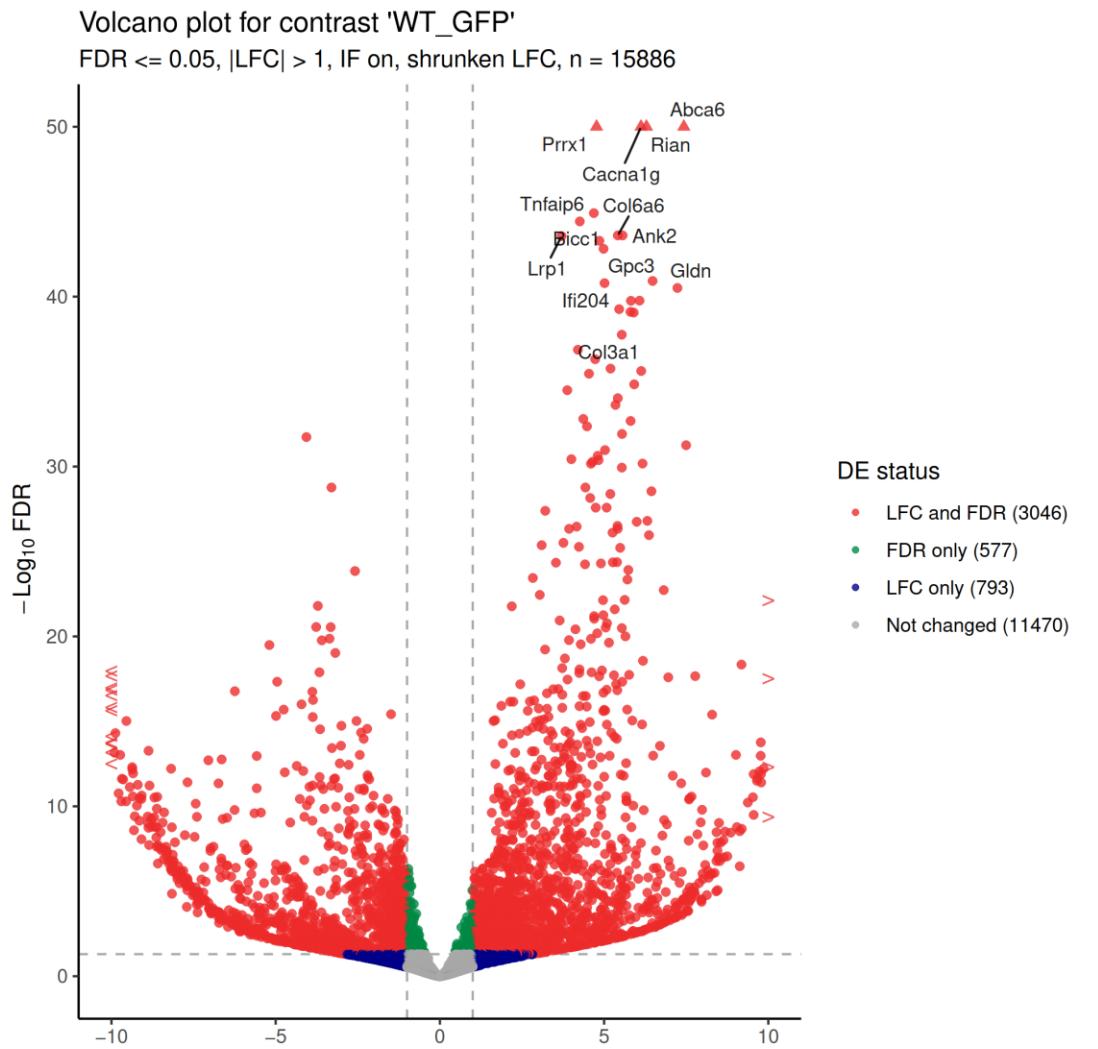


# Enrichment analysis

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with acknowledgements to Luciano Cascione and SIB



# Goal



Find biologically relevant **gene sets** among **deregulated genes** between **groups of cells**

# What is a gene set?

A gene set is an unordered collection of genes that are functionally related.

- Genes located in the same compartment in a cell (e.g. all proteins located in the cell nucleus)
- Proteins that are all regulated by a same transcription factor
- Custom gene list that comes from a publication and that are down-regulated in a mutant
- List of genes that contain SNPs associated with a disease
- ...etc!
- Several gene sets are grouped into Knowledge bases
- A pathway can be interpreted as a gene set by ignoring functional relationships among genes

# Gene set resources:

**Gene Ontology** -> <https://geneontology.org>

- Biological process
- Molecular function
- Cellular component

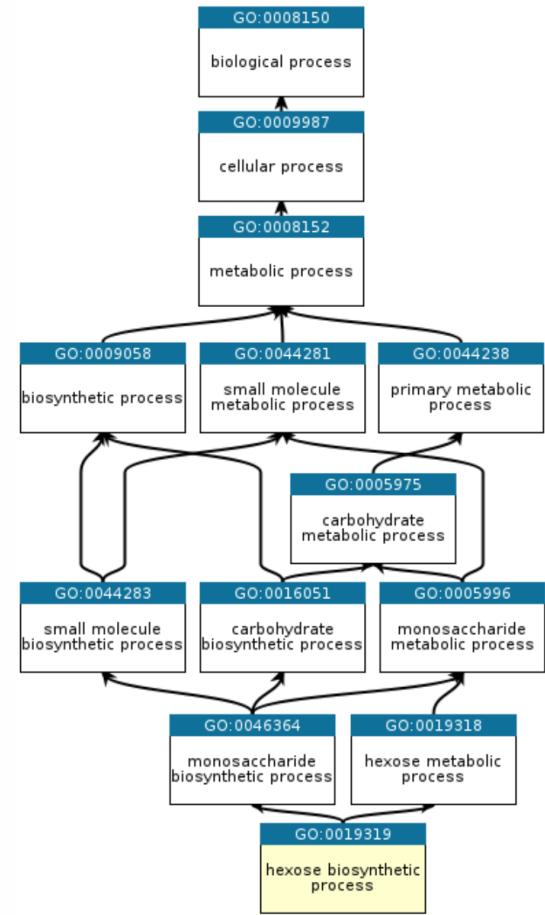
**KEGG** -> <https://www.kegg.jp/kegg/pathway.html>

**MSigDB** -> <https://www.gsea-msigdb.org/gsea/msigdb/index.jsp>

- Hallmark gene sets
- Positional gene sets
- Immunological sets etc.

**Reactome** -> <https://reactome.org/>

**WikiPathways** -> <https://www.wikipathways.org/index.php/WikiPathways>



# Methods

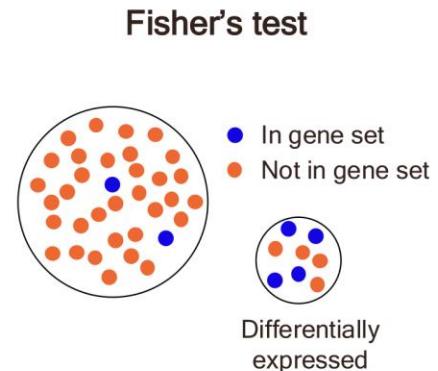
- Over-representation analysis (ORA)
  - Fisher's test of proportions of DEGs and gene set
- Gene set enrichment analysis (classical GSEA)
  - Enrichment score of gene set in ranked list of genes based on deregulation metric

# Over-representation analysis (ORA)

```
> cont.table<-matrix(c(2,3,5,12), ncol=2, byrow = T)  
> fisher.test(cont.table)
```

Fisher's Exact Test for Count Data

```
data: cont.table  
p-value = 1  
alternative hypothesis: true odds ratio is not equal to 1  
95 percent confidence interval:  
 0.1012333 18.7696686  
sample estimates:  
odds ratio  
 1.56456
```



$H_0$ : The proportion of genes in the gene set is the same for both groups

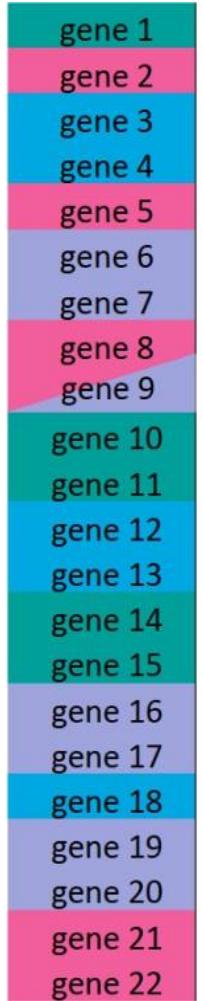
$H_a$ : The proportion of genes in the gene set is higher in the differentially expressed group

2x2 count table		Differentially expressed	Not Differentially expressed	total
blue	2	3	5	
Not blue	5	12	17	
total	7	15	22	

$$2/7 = 0.29$$

$$3/15 = 0.20$$

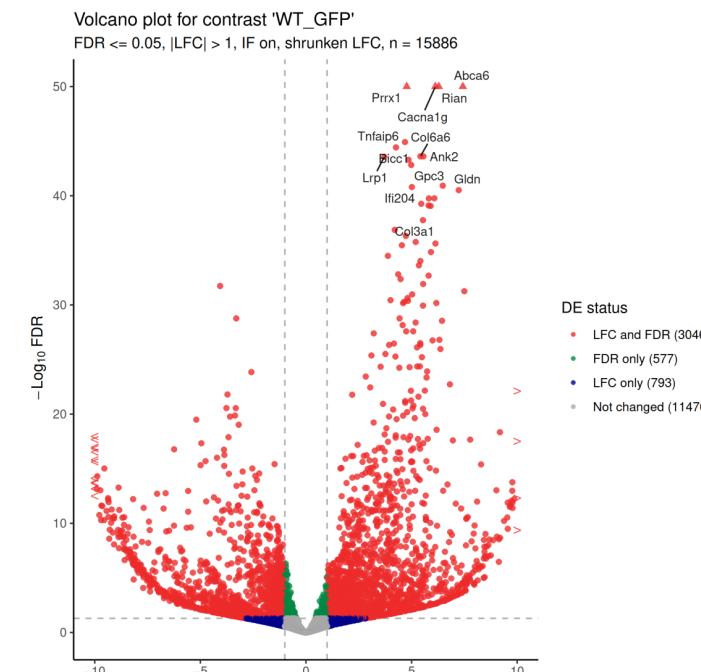
# Which gene-sets are enriched?



Run individual Fisher's exact tests for each gene set, **blue, pink, purple, green**

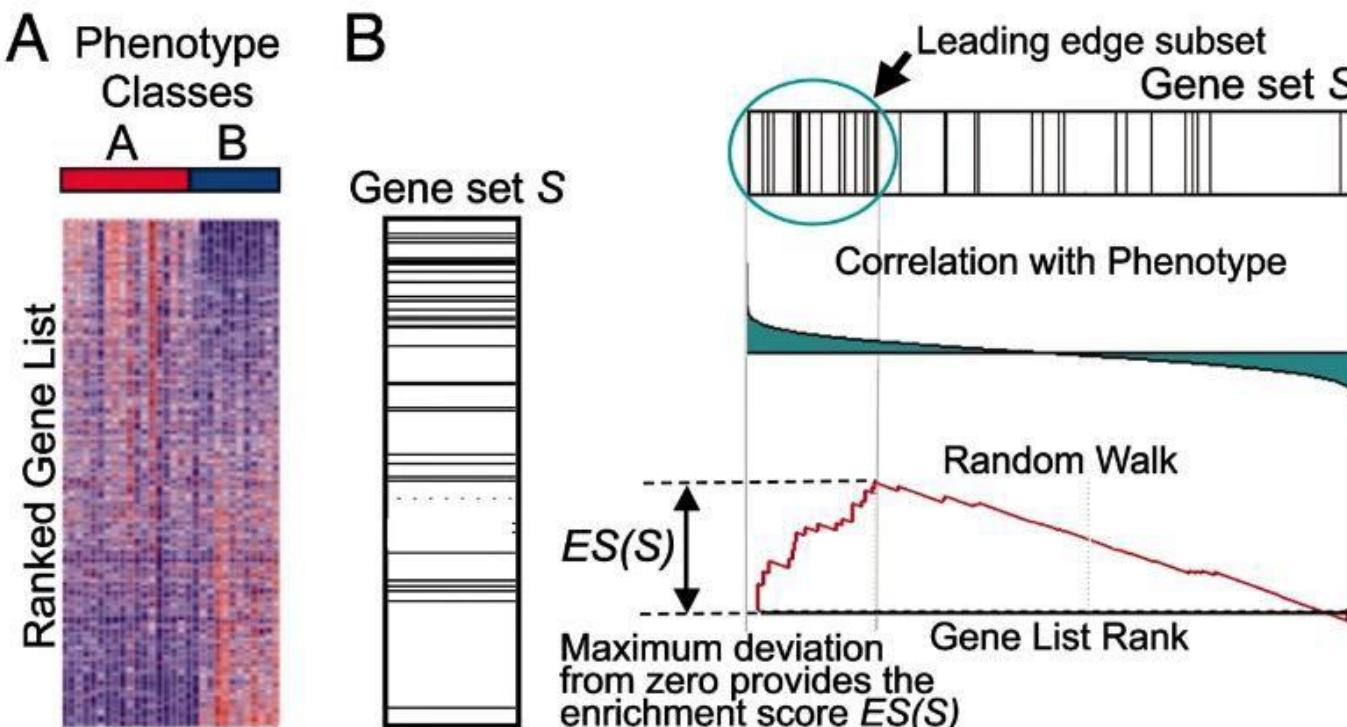
⇒ Multiple tests need **p-value adjustment.**

⇒ Fisher test is **threshold-based**



# Gene-Set Enrichment Analysis

ORA fails to detect situations where all genes in a predefined set change in a small but coordinated way

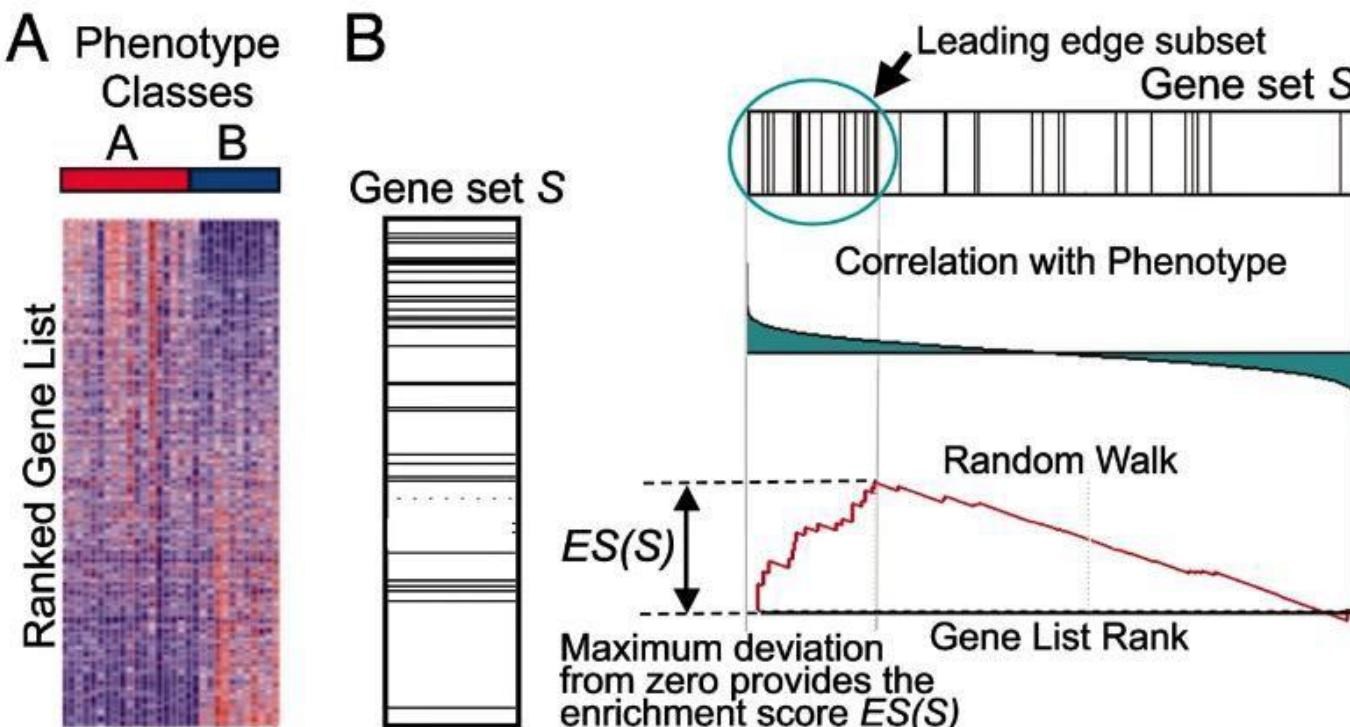


Genes are ranked based on their phenotypes.

Given apriori defined set of gene  $S$ , the goal of GSEA is to determine whether the members of  $S$  are randomly distributed throughout the ranked gene list ( $L$ ) or primarily found at the top or bottom.

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# How Can I Rank the Genes?

?



# How can I rank the genes?

Research Article | [Open access](#) | Published: 12 May 2017

## Ranking metrics in gene set enrichment analysis: do they matter?

[Joanna Zyla](#), [Michał Marczyk](#) , [January Weiner](#) & [Joanna Polanska](#)

*BMC Bioinformatics* **18**, Article number: 256 (2017) | [Cite this article](#)

**45k** Accesses | **43** Citations | **22** Altmetric | [Metrics](#)

### Abstract

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### Background

There exist many methods for describing the complex relation between changes of gene expression in molecular pathways or gene ontologies under different experimental conditions. Among them, Gene Set Enrichment Analysis seems to be one of the most commonly used (over 10,000 citations). An important parameter, which could affect the final result, is the choice of a metric for the ranking of genes. Applying a default ranking metric may lead to poor results.

# clusterProfiler

## A universal enrichment tool for interpreting omics data

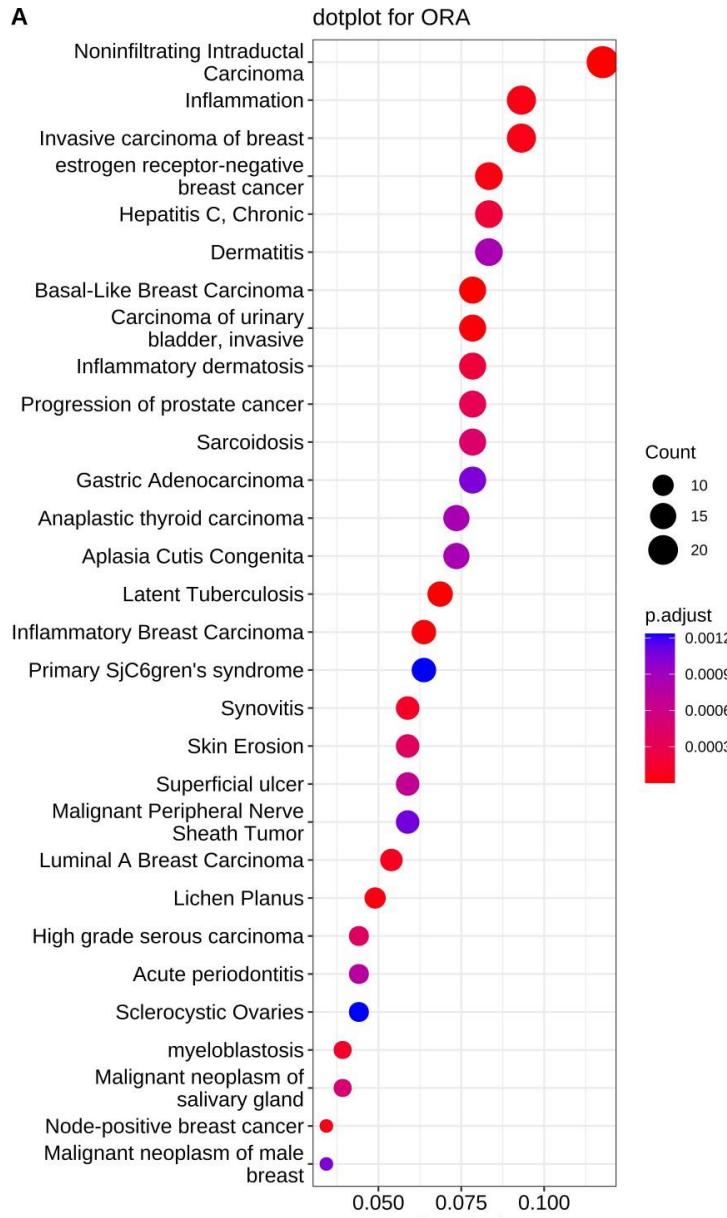
platforms all rank 36 / 2300 support 1 5 / 1 8 in Bioc 13.5 years build ok updated < 3 months dependencies 132

DOI: [10.18129/B9.bioc.clusterProfiler](https://doi.org/10.18129/B9.bioc.clusterProfiler)

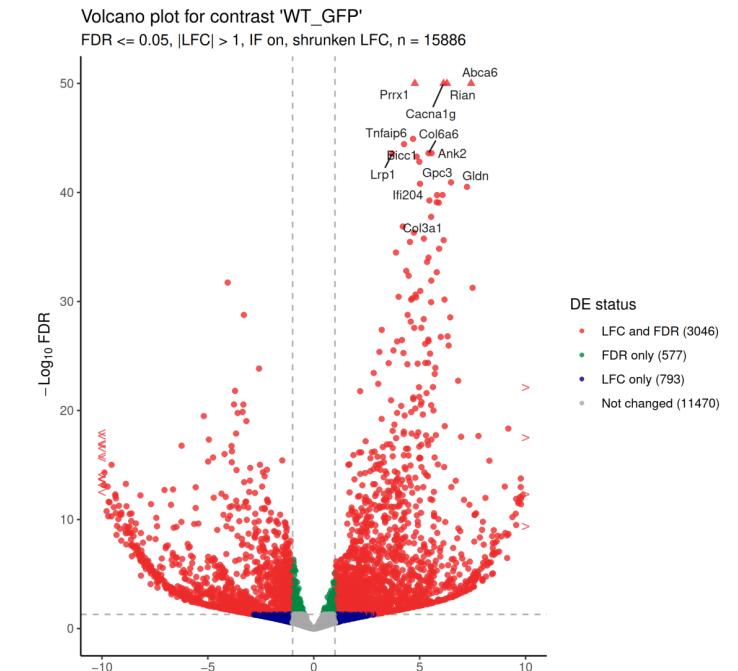
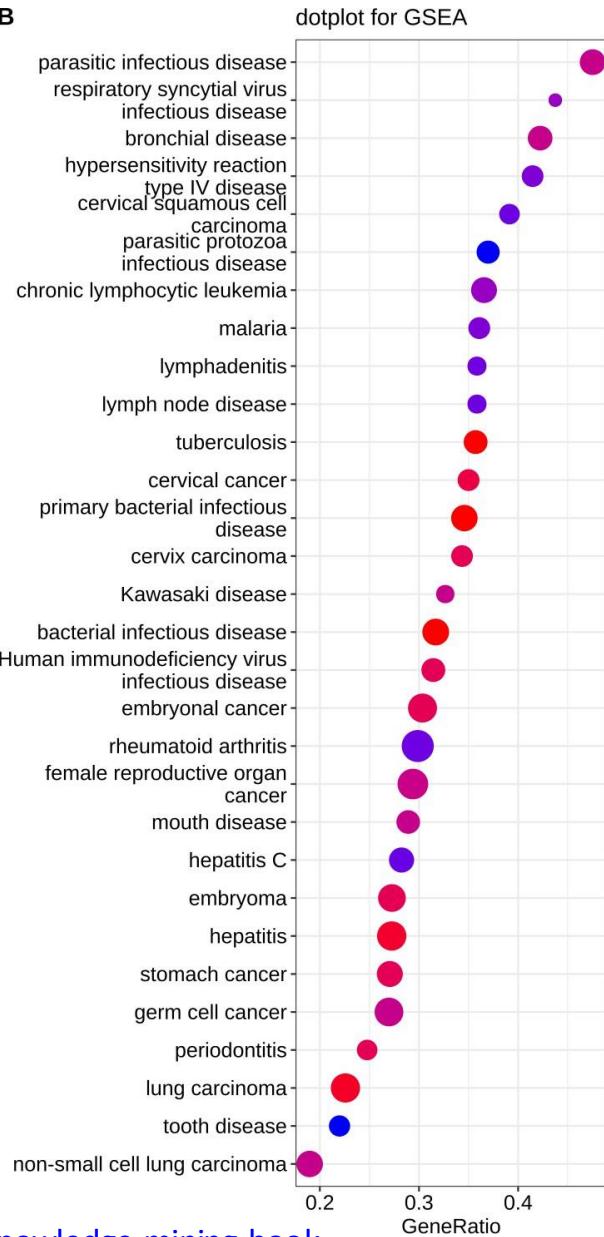
	KEGG	GO terms	custom
ORA	enrichKEGG()	enrichGO()	enricher()
GSEA	gseKEGG()	gseGO()	GSEA()

# Visualization

A

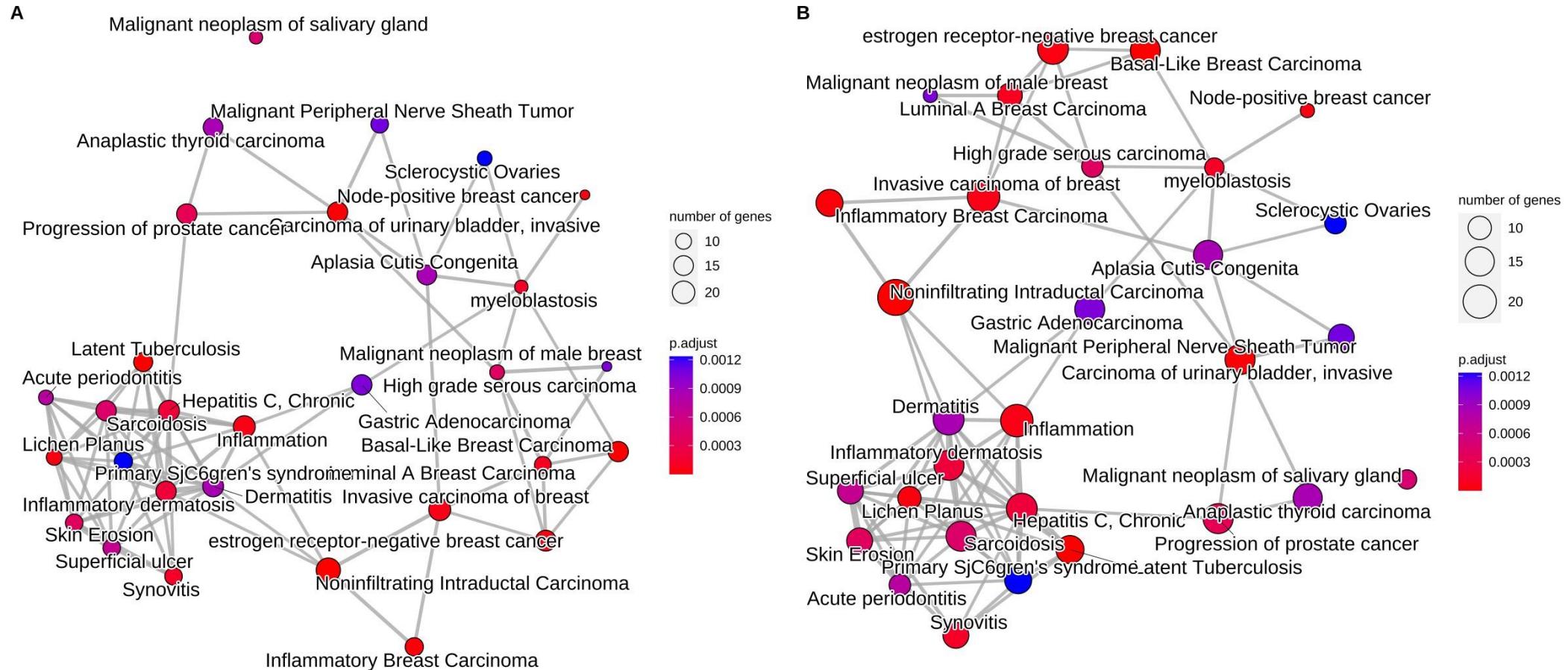


B



Adapted from: <https://yulab-smu.top/biomedical-knowledge-mining-book>

# Enrichment Map



Edges connect overlapping gene sets. In this way, overlapping gene sets tend to cluster together, making it easy to identify functional module.

*Thanks for your attention!*