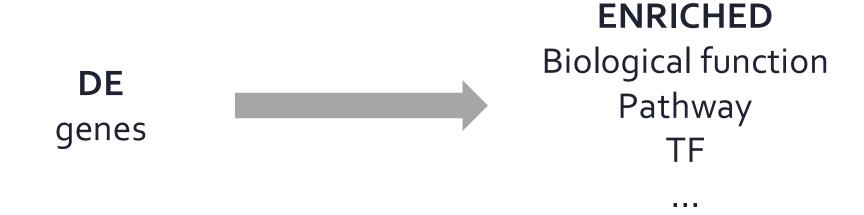
Introduction to RNA-Seq: Enrichment analysis

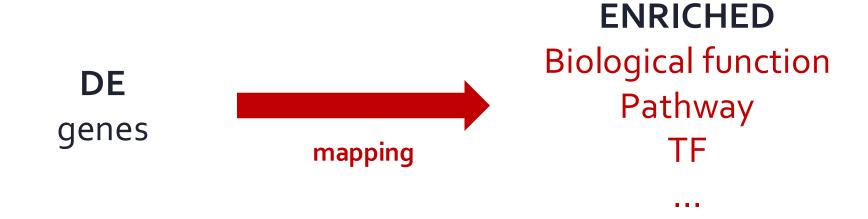
Wandrille Duchemin

What to do with your list of differentially expressed gene?

What to do with your list of differentially expressed gene?



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What to do with your list of differentially expressed gene?



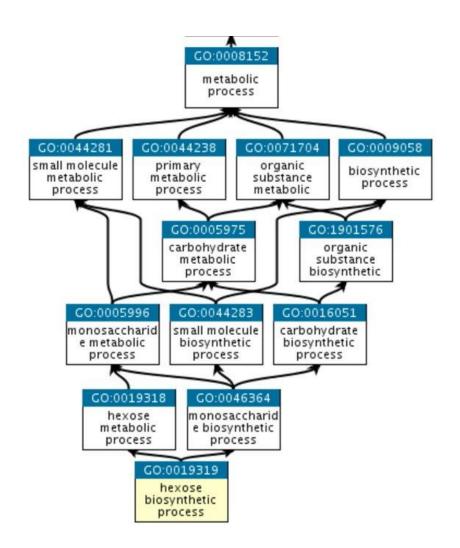
Regrouping genes together in meaningful sets

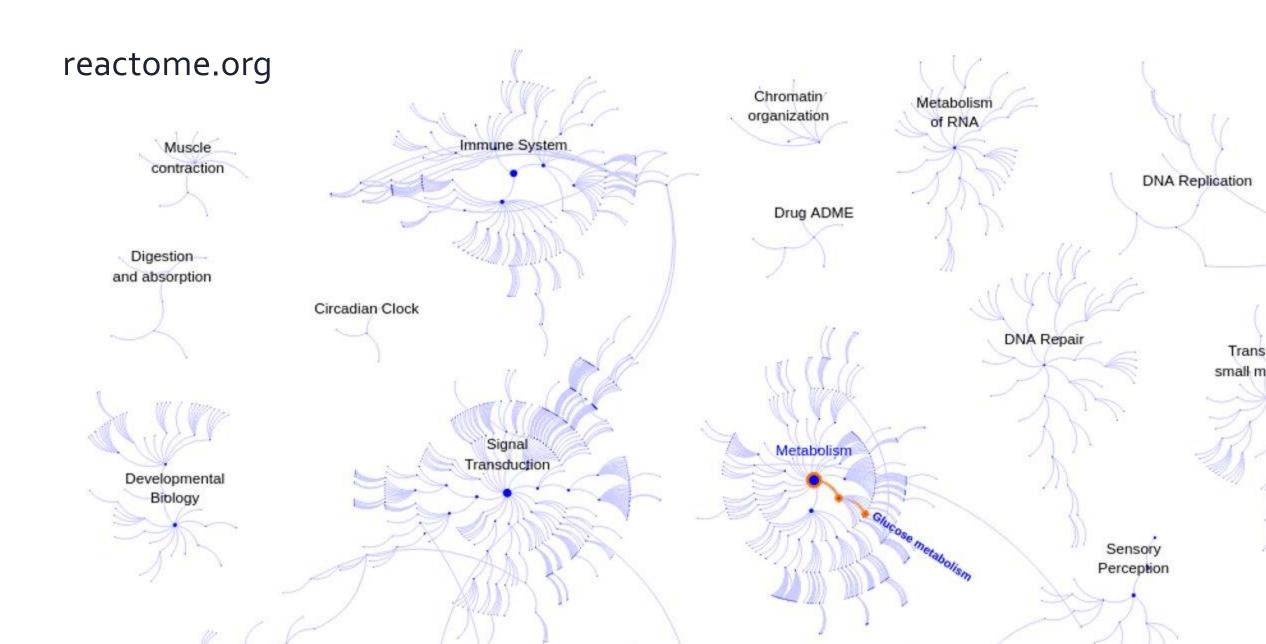
- Same pathway
- Same location in the cell
- Same molecular function
- •

- Gene Ontologies
- Reactome
- KEGG
- MSigDB
- Custom set
- ...

Gene Ontologies <u>geneontology.org</u> 3 domains of nested terms:

- Molecular Function
- Cellular Component
- Biological Process





MSigDB: http://www.gsea-msigdb.org/gsea/msigdb/index.jsp

Human, mouse, and rat only

Н

hallmark gene sets are coherently expressed signatures derived by aggregating many MSigDB gene sets to represent well-defined biological states or processes.

C1

positional gene sets for each human chromosome and cytogenetic band.

C2

curated gene sets from online pathway databases, publications in PubMed, and knowledge of domain experts.

C:

regulatory target gene sets based on gene target predictions for microRNA seed sequences and predicted transcription factor binding sites.

C4

computational gene sets defined by mining large collections of cancer-oriented microarray data.

C5

ontology gene sets consist of genes annotated by the same ontology term.

C6

oncogenic signature gene sets defined directly from microarray gene expression data from cancer gene perturbations.

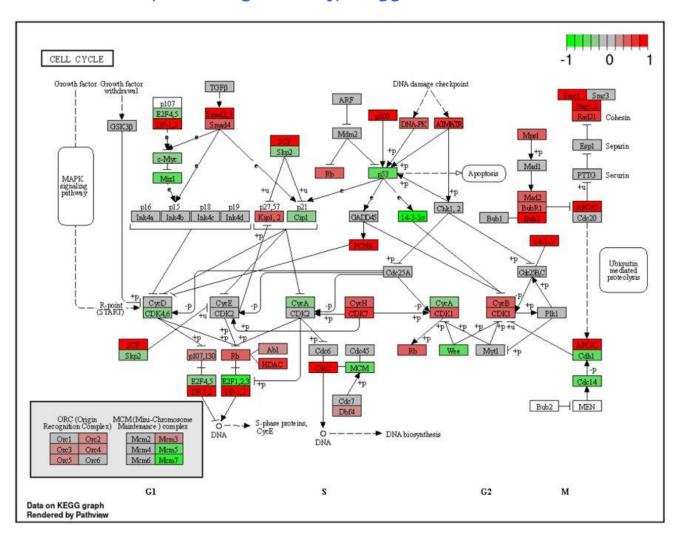
C7

immunologic signature gene sets represent cell states and perturbations within the immune system.

C8

cell type signature gene sets curated from cluster markers identified in single-cell sequencing studies of human tissue.

KEGG: http://www.genome.jp/keggKEGG PATHWAY



Projection of DE data onto a KEGG pathway map With R package pathview

https://bioconductor.org/packages/release/bioc/html/pathview.html

Custom gene sets

- Derived from specialized litterature
- Tentative annotation

Very important in non-model organisms

2 main approaches

- Over Representation Analysis (ORA)
- Gene Set Enrichment Analysis (GSEA)

Over Representation Analysis (ORA)

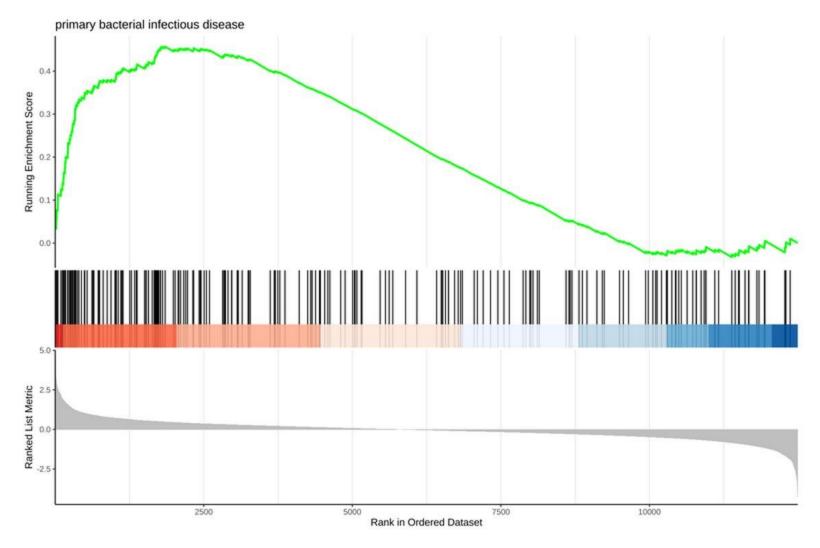
Basically a Fisher's exact test with p-value correction

	DE	Not DE
in gene set	А	В
not in gene set	С	D

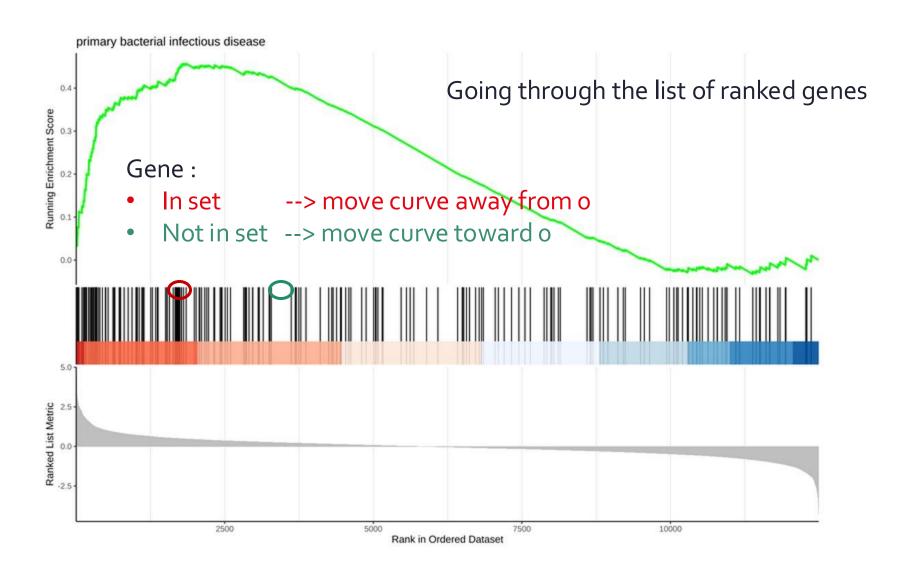
$$p=1-\sum_{i=0}^{k-1}rac{inom{M}{i}inom{N-M}{n-i}}{inom{N}{n}}$$

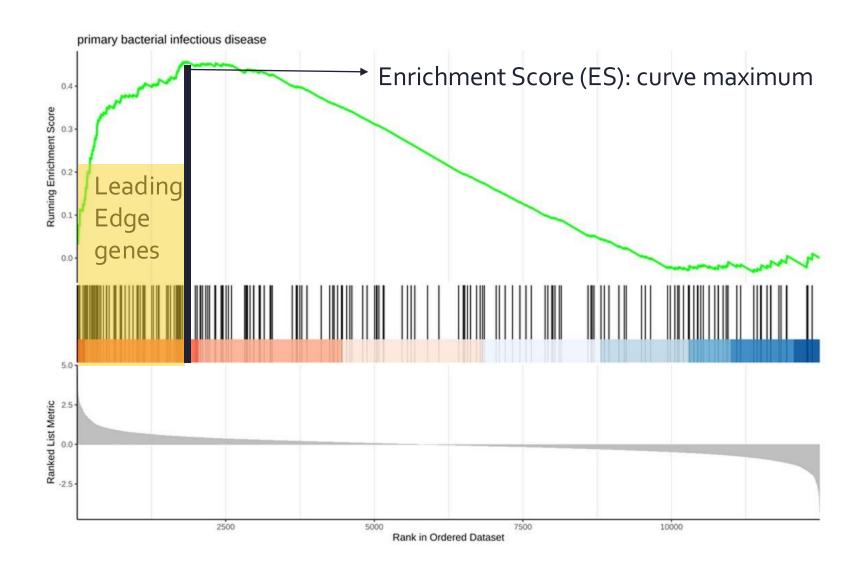
Gene Set Enrichment Analysis (GSEA)

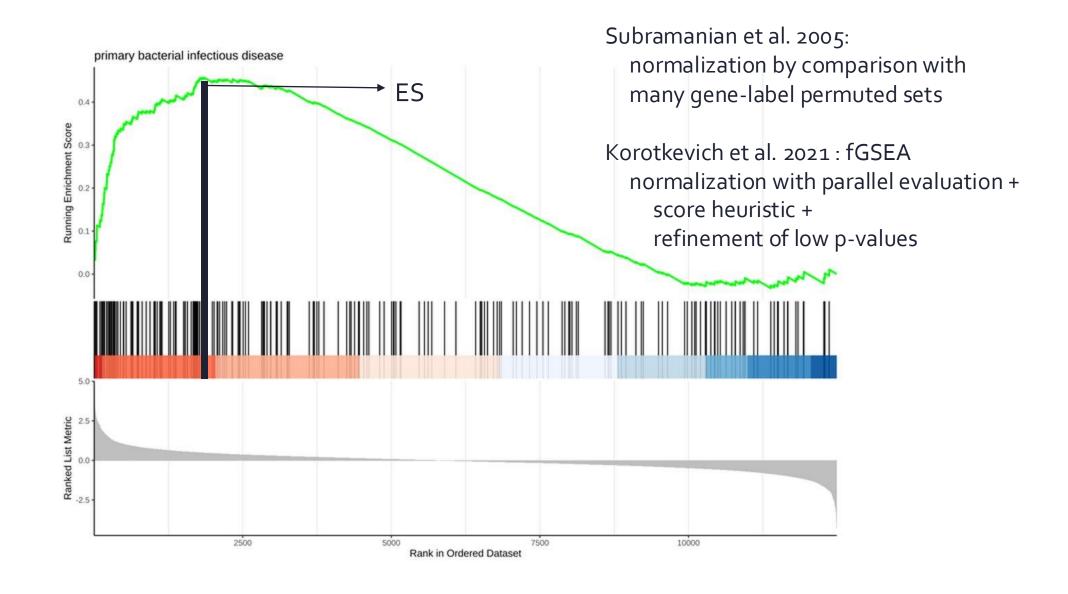
- Does not rely on o/1 DE
- Use ranking along a continuous measure (eg. log2FC)
- Compute Enrichment Score
- Estimate significance with a permutation scheme

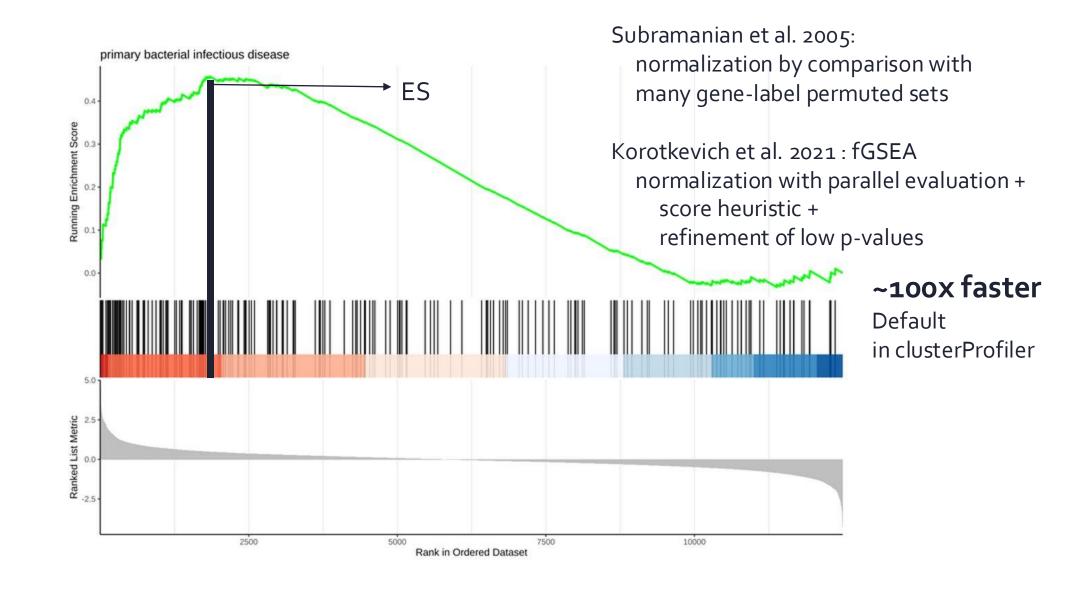


Enrichment Score visualized using functions from the R package enrichplot









Many more methods or implementations

- Signaling Pathway Impact Analysis
 https://bioconductor.org/packages/release/bioc/html/SPIA.html
- ISMARA (TF-based)
 ismara.unibas.ch/mara

Practical