## NGS2 course

Making Sense of Gene Lists

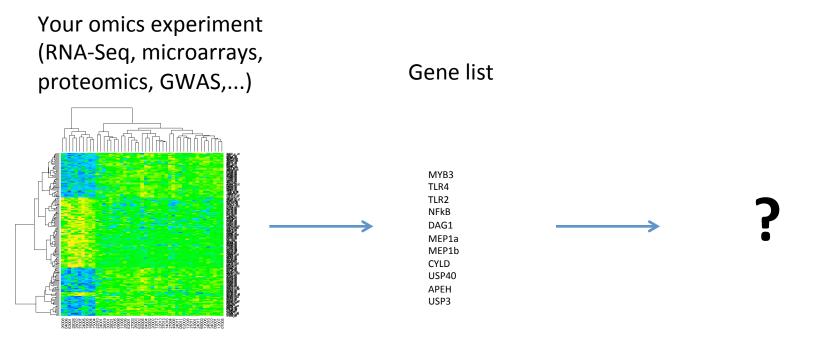
Stefan Wyder





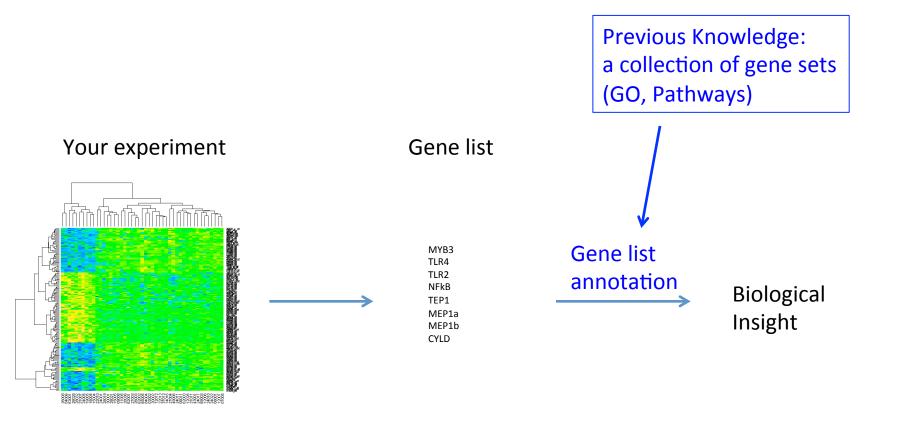
## **Gene List Annotation**

- You performed a genomic experiment and obtained a gene list
- Who wants to work through a list of hundreds of genes?
- What's next?



## Gene List Annotation

- We test whether the differentially expressed genes in our experiment are enriched in some predefined gene lists.
- Based on previous knowledge



# Obtaining Biological Insight

- to summarize gene lists
- to help and speed up the interpretation of an experiment
- to gain mechanical insight
- to find regulated processes/pathways
- to find involved regulatory elements (TF, miRNA)
- to identify new members of a pathway
- to find similar experiments
- •

Analysis based on gene lists is expected to be more robust and reproducible than single-gene analysis.

# **Enrichment Analysis**

#### **Over-Representation Analysis**

- hypergeometric aka Fisher's exact test
- input: 4 counts
- we need to set a cut-off a priori
- different results at different thresholds!

<b>Gene Set</b>	<b>Enrichment</b>	<b>Analysis</b>	(GSEA)
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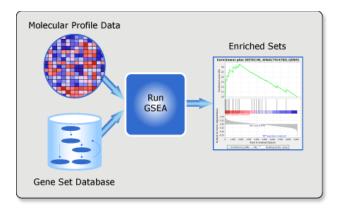
Subramanian et al. (2005) PNAS and many follow-up papers

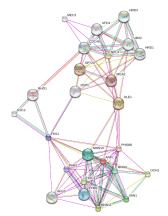
- bypasses the need for a cut-off
- input: list of all measured genes ranked by some statistics / effect size
- weak but consistent regulation of several members of a gene set can be detected

#### **Network Analysis**

- covers also the less well understood portion of gene interactions
- often inferred from co-expression data
- example: STRING (http://www.string-db.org/)
- combines info from co-expression, co-citation, PPI, ....

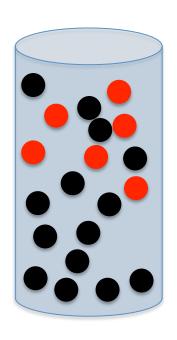
8	12
2	2412



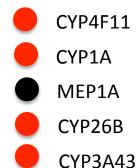


# Over-Representation Analysis

5000 black and 10 red balls in an urn each ball represents 1 gene
10 red balls ("Cytochromes")



Our list of differentially expr. genes: 4/5 balls are red

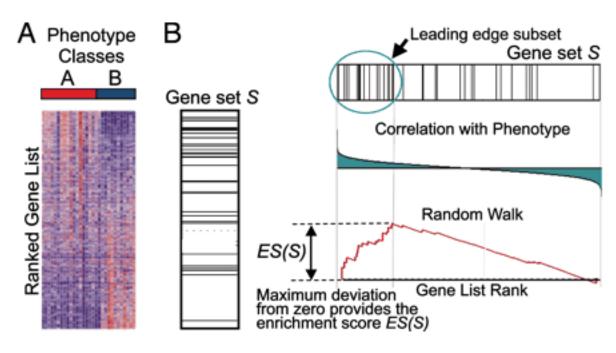


What is the probability? 2x2 contigency table

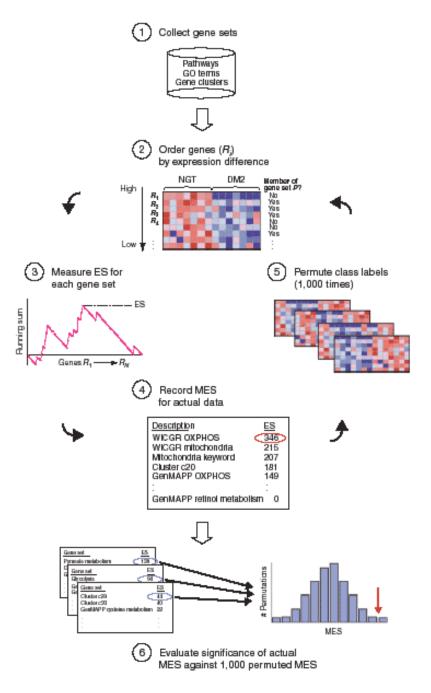
	Selected	Not
in category	4	6
not in category	1	4989

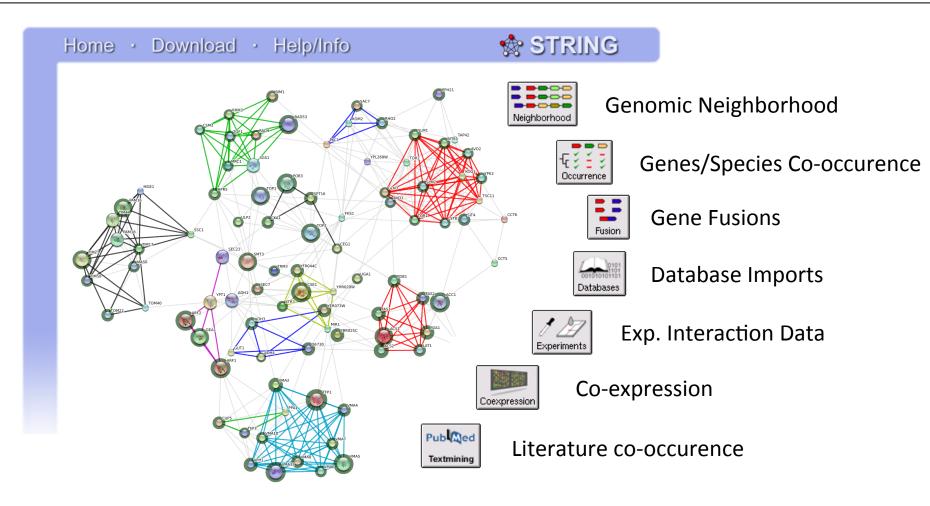
one-sided Fisher's exact test p-value = 4.03e-11

# Gene Set Enrichment Analysis



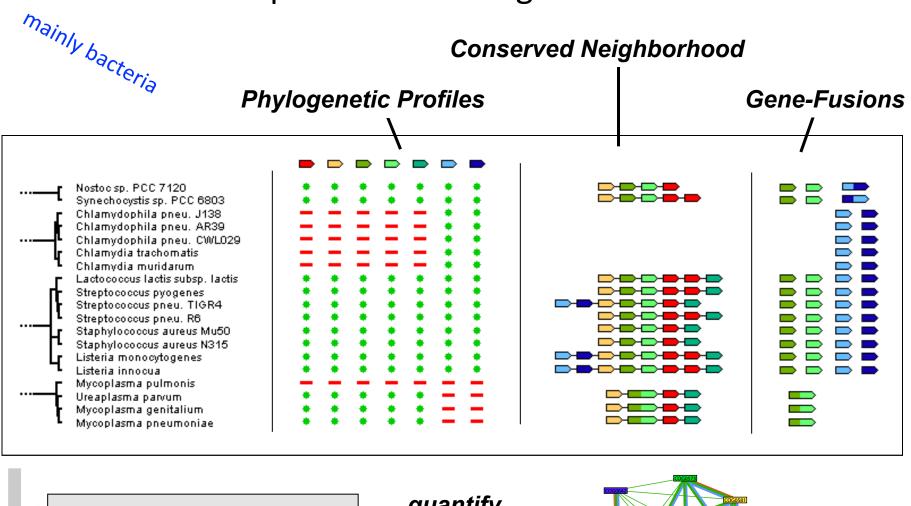
Subramanian et al. (2005) PNAS



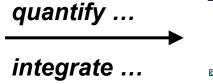


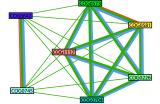
- functional association networks (physical or functional interactions)
- focus on useability and speed
- integrated scoring scheme (each interaction has confidence score)
- information transfer between species (>2000 species: Animals, Bact, Plants,...)

## Interaction prediction from genome information



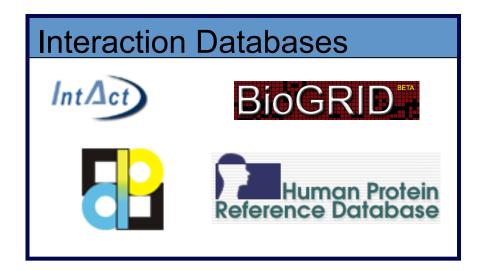
"genomic context"

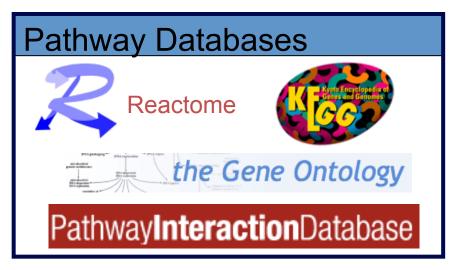


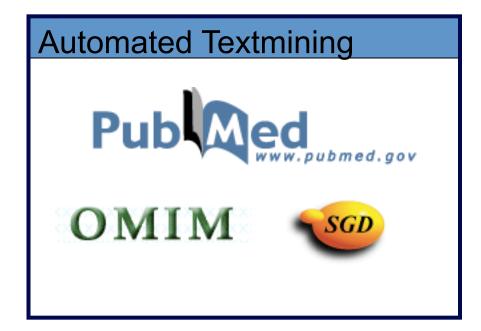


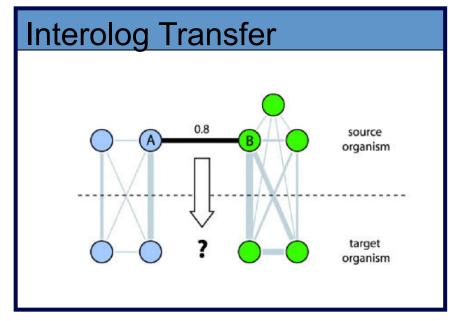
networks

### Other Interaction Sources

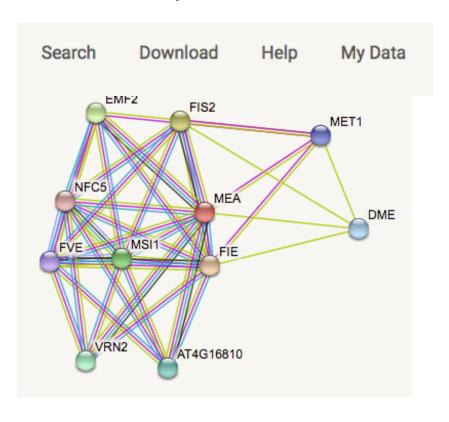


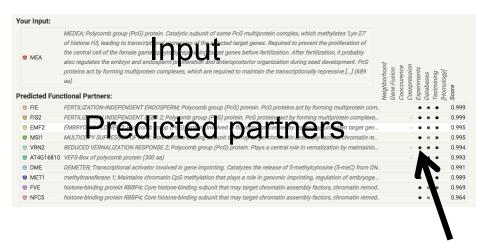






### Output







Textmining Experiments Databases Co-expression

Neighborhood Gene Fusion Co-occurrence

Switch
On/off
the input channel

Clickable

evidence

## **STRING**

- can do more than gene list annotation:
  - Predicting gene function
  - Identifying candidates for an unknown enzyme in a pathway
  - Identifying new member genes of a biological process
  - Finding relevant literature
- ID mapper engine understands a large number of gene formats
- STRING performs well compared with single-species databases
- R package to access STRING functionality from R
- available for download

## **Annotation Sources**

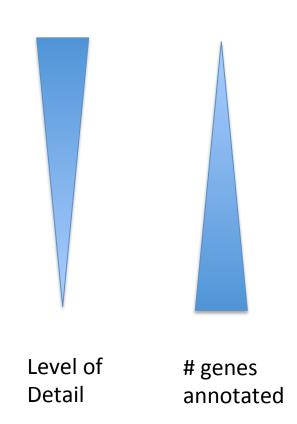
#### **Pathways**

KEGG, Reactome, BioCyc, ...

**Gene Ontology (GO)** 

**Gene/Protein Networks** 

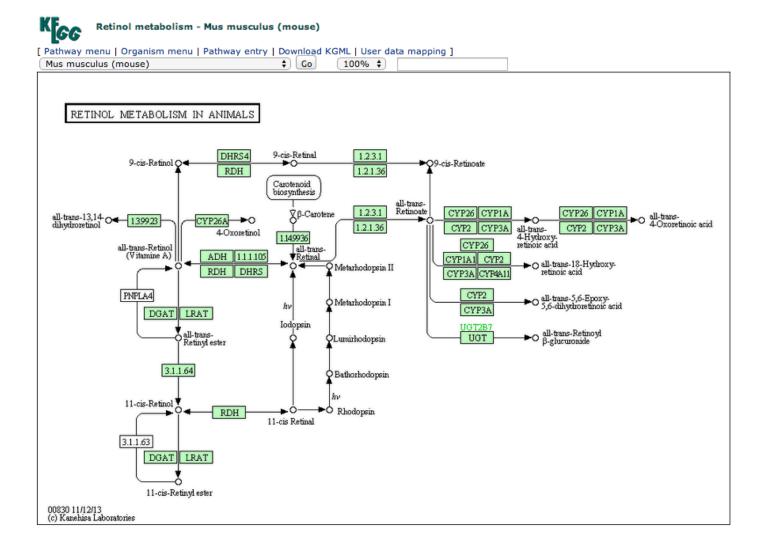
e.g. STRING



# **Pathways**

- pathway maps (aka reaction networks / wiring diagrams) represent experimental knowledge on metabolism and various other functions of the cell and the organism
- manually curated
- the main databases are KEGG and Reactome
- KEGG is free to use over the web but file download requires subscription
- KEGG covers >3'800 species (Archae, Bacteria, Plants, Animals) and Reactome covers 20 species (mostly mammals + fly + plants + E.coli) as of May 2015.

# **Example KEGG Pathway**



# **Gene Ontology**

#### **Gene Ontology (GO)**

http://www.geneontology.org/

- describes how gene products behave in a cellular context (BP, MF, C)
- controlled vocabulary of terms
- transparent (sources)
- manually curated lists for model species
- transfer to orthologs in other species (inferred annotation)

# Example

#### murine ADAM10

#### Molecular function

GO:0008237 metallopeptidase activity

GO:0042169 SH2 domain binding

. .

#### **Biological Process**

GO:0007220 Notch receptor processing

GO:0001701 in utero embryonic development

GO:0008284 positive regulation of cell proliferation

..

#### **Cellular Compartment**

GO:0005794 Golgi apparatus

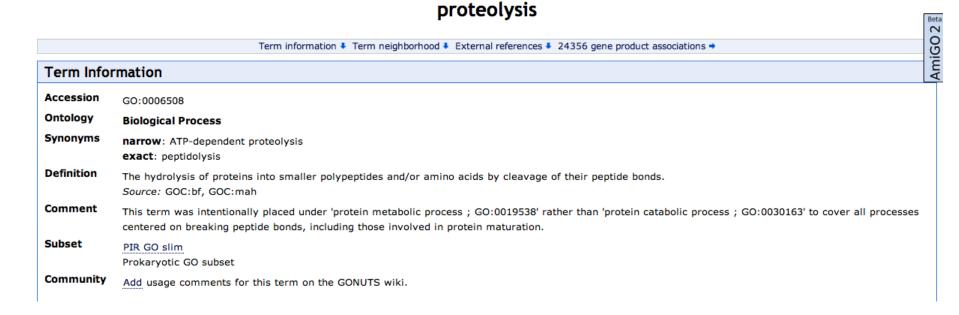
GO:0009986 cell surface

• •

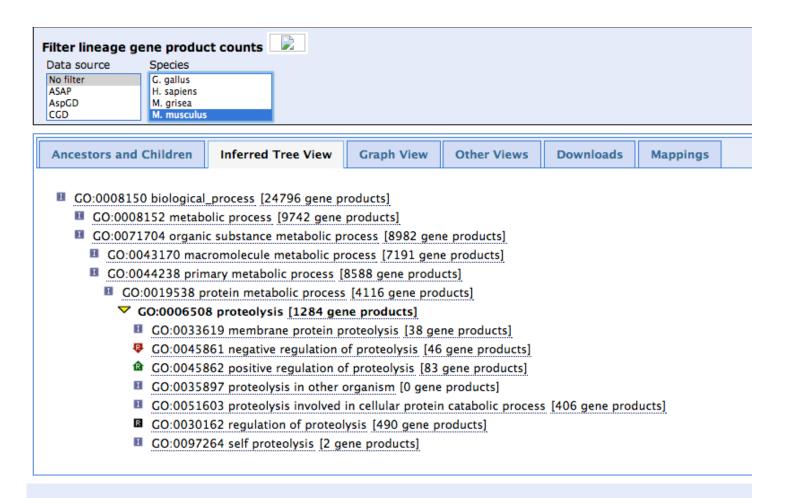
## Lookup of GO terms

# AmiGO http://amigo.geneontology.org

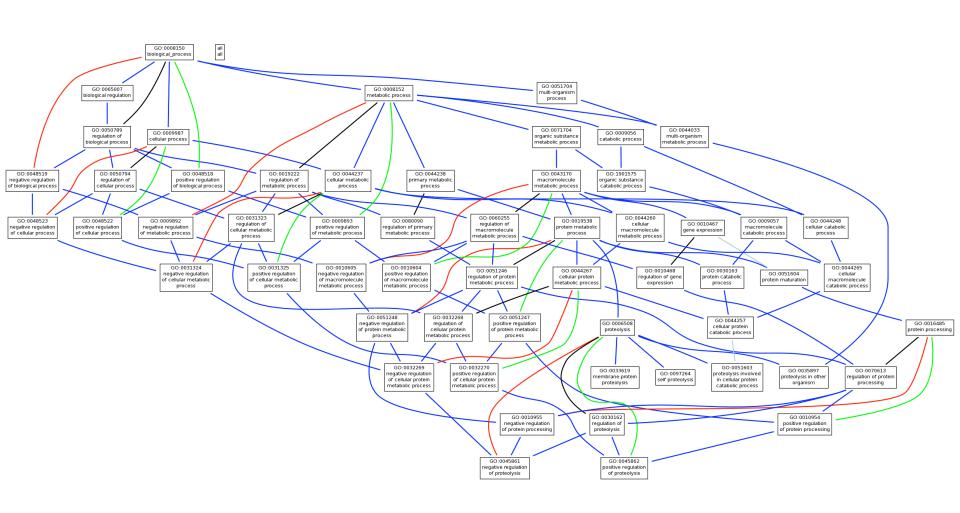




# GO Table View GO:0006508 Proteolysis



# Graphical View GO:0006508 Proteolysis



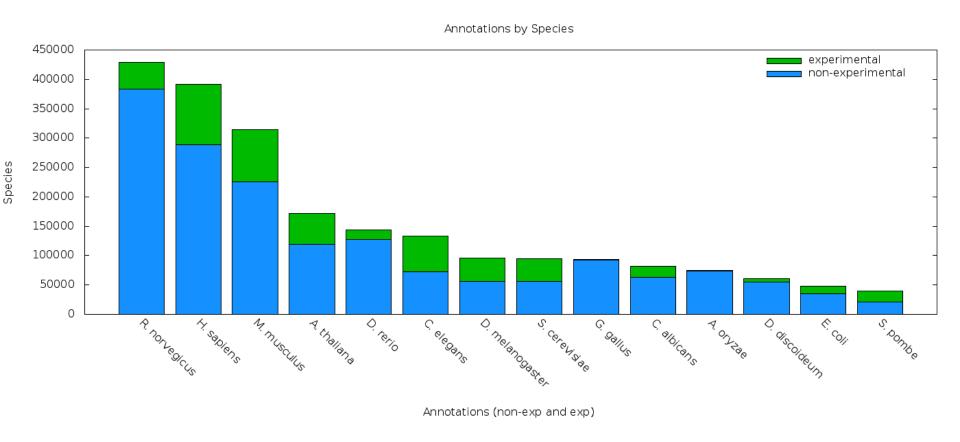
## **Ancestors and Children**

## AmiGO http://amigo.geneontology.org

Ancestors and Children	Inferred Tree View	Graph View	Other Views	Downloads	Mappings			
			Ancestors of pr	oteolysis (GO:0006	508)			
s	subject ¢		relation \$		o	bject ¢	ann	otations ¢
proteolysis		■ is	_a (inferred)	biological_p	ocess (GO:000	8150)	665024	
proteolysis		■ is	_a (inferred)	metabolic pr	ocess (GO:0008	8152)	368913	
proteolysis		■ is	_a (inferred)	organic subs	tance metaboli	c process (GO:0071704)	300256	
proteolysis		■ is	_a (inferred)	macromolec	ule metabolic p	rocess (GO:0043170)	202070	
proteolysis		■ is	_a (inferred)	primary met	abolic process (	(GO:0044238)	277534	
proteolysis		■ is	_a	protein meta	bolic process (	GO:0019538)	105597	
Children of proteolysis (GO:0006508)								
s	subject ø		relation ¢		o	bject ¢	ann	otations \$
membrane protein proteoly	ysis (GO:0033619)	Ⅱ is	_a	proteolysis			387	
negative regulation of prot	teolysis (GO:0045861)	<b>₽</b> ne	egatively_regulate:	s proteolysis			502	
positive regulation of prote	eolysis (GO:0045862)	<b>⊉</b> p	ositively_regulates	proteolysis			696	
proteolysis in other organi	sm (GO:0035897)	■ is	_a	proteolysis			83	
proteolysis involved in cell (GO:0051603)	ular protein catabolic pro	ocess I is	_a	proteolysis			8312	
regulation of proteolysis (C	GO:0030162)	<b>⊠</b> r€	egulates	proteolysis			4093	
self proteolysis (GO:00972	.64)	■ is	_a	proteolysis			38	

## **GO** statistics

Even in model organisms only a minority of genes has experimental GO annotation



http://beta.geneontology.org/user-story-tags/everybody?page=5

# False Discovery Rate (FDR)

Significance (alpha) level: probability of rejecting the null hypothesis given that it is true

Therefore at 5% significance level: for 100 tests where all null hypotheses are true, the expected number of incorrect rejections is 5

tests	incorrect rejections
100	5
10,000	500

#### **Multiple Testing Correction**

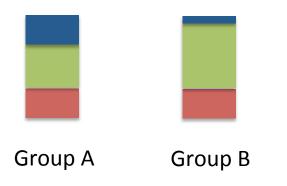
- Bonferroni
- False Discovery Rate (FDR): If we find 100 genes significantly differentially expressed at a 5% FDR, we expect at most 5 false discoveries in the list.

## Experimental design

Interpretability depends mostly on appropriate experimental design!

Randomize samples/treatments across lanes / flow cells

Multiple tissues/cell types/stages pooled in a sample -> complex and difficult to understand the ongoing processes (e.g. observed changes can simply be due to changes in relative abundance of different cell types independent of regulation)



Blood example during pregnancy

## Summary

- Gene list annotation with Pathways and Gene Ontology can help to obtain biological insight.
- 3 main methods: 1. Over-Representation Analysis, 2. Gene Set Enrichment Analysis (GSEA), 3. Network Analysis
- Biological interpretation requires broad knowledge of physiology & biochemistry and is often the most difficult and time-consuming step of an experiment.
- Even experts can usually not make sense of all the significantly enriched processes/pathways in well understood biological systems.
- Good experiments start with good experimental design! Think of possible confounders

## **URLs & Tips**

#### Main general Annotation Sources

- Gene Ontology (http://www.geneontology.org/)
  - AmiGO: http://amigo.geneontology.org
  - QuickGO: http://www.ebi.ac.uk/QuickGO/
  - Compilation of GO Tools: http://www.geneontology.org/GO.tools.shtml
- KEGG (http://www.genome.jp/kegg)
- Reactome (http://www.reactome.org)

- Most pathway databases offer also tools to colorize genes of interest on pathways
- Pathway analysis can also be done in R/bioconductor, see
   http://www.bioconductor.org/packages/release/BiocViews.html#\_\_\_Pathways