RNA-seq differential expression analysis

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What we will cover

We will cover differential expression analysis of RNA-seq data in R/Bioconductor.

We will start from a matrix of gene-level read counts.

We will cover the two most popular packages, DESeq2 and edgeR.

I will also show you how to deal with unwanted variation using the RUVSeq package.

What we will not cover

I will not talk about the preprocessing of RNA-seq data, i.e., what we do to obtain the gene-level read counts.

These steps are usually done with stand-alone software outside R.

I will not talk about isoform-level analysis and alternative splicing.

We will focus on gene-level differential expression.

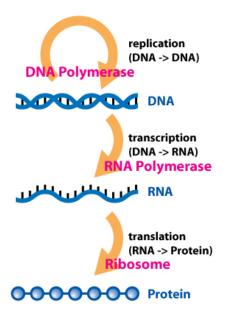
Where to find these slides

https://github.com/drisso/rnaseq_meetup

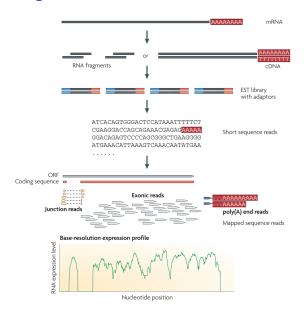
Where to find additional resources

- The edgeR user guide https://bioconductor.org/packages/edgeR
- ► The DESeq2 vignette https://bioconductor.org/packages/DESeq2
- ► The F1000 Research Bioconductor gateway https://f1000research.com/gateways/bioconductor
- https://support.bioconductor.org

From RNA to gene-level read counts



From RNA to gene-level read counts



From RNA to gene-level read counts

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                        CC3
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   ENSMUSG00000000049
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   ENSMUSG00000000056 3045 3106 3441 1940
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                        945 1031 1170
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The Poisson Model

When statisticians see counts, they immediately think about Simeon Poisson.



The Poisson Model

The Poisson distribution naturally arises from binomial calculations, with a large number of trials and a small probability.

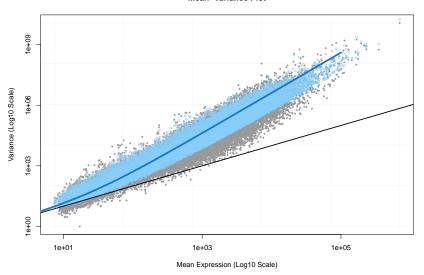
It has a rather stringent assumption: the variance is equal to the mean!

$$Var(Y_{ij}) = \mu_{ij}$$

In real datasets the variance is greater than the mean, a condition known as **overdispersion**.

A real example





The Negative Binomial Model

A generalization of the Poisson model is the negative binomial, that assumes that the variance is a quadratic function of the mean.

$$Var(Y_{ij}) = \mu_{ij} + \phi_j \mu_{ij}^2$$

where ϕ is called the **dispersion parameter**.

Both edgeR and DESeq2 assume that the data is distributed as a negative binomial.