

# Analysis of bulk RNA-Seq data

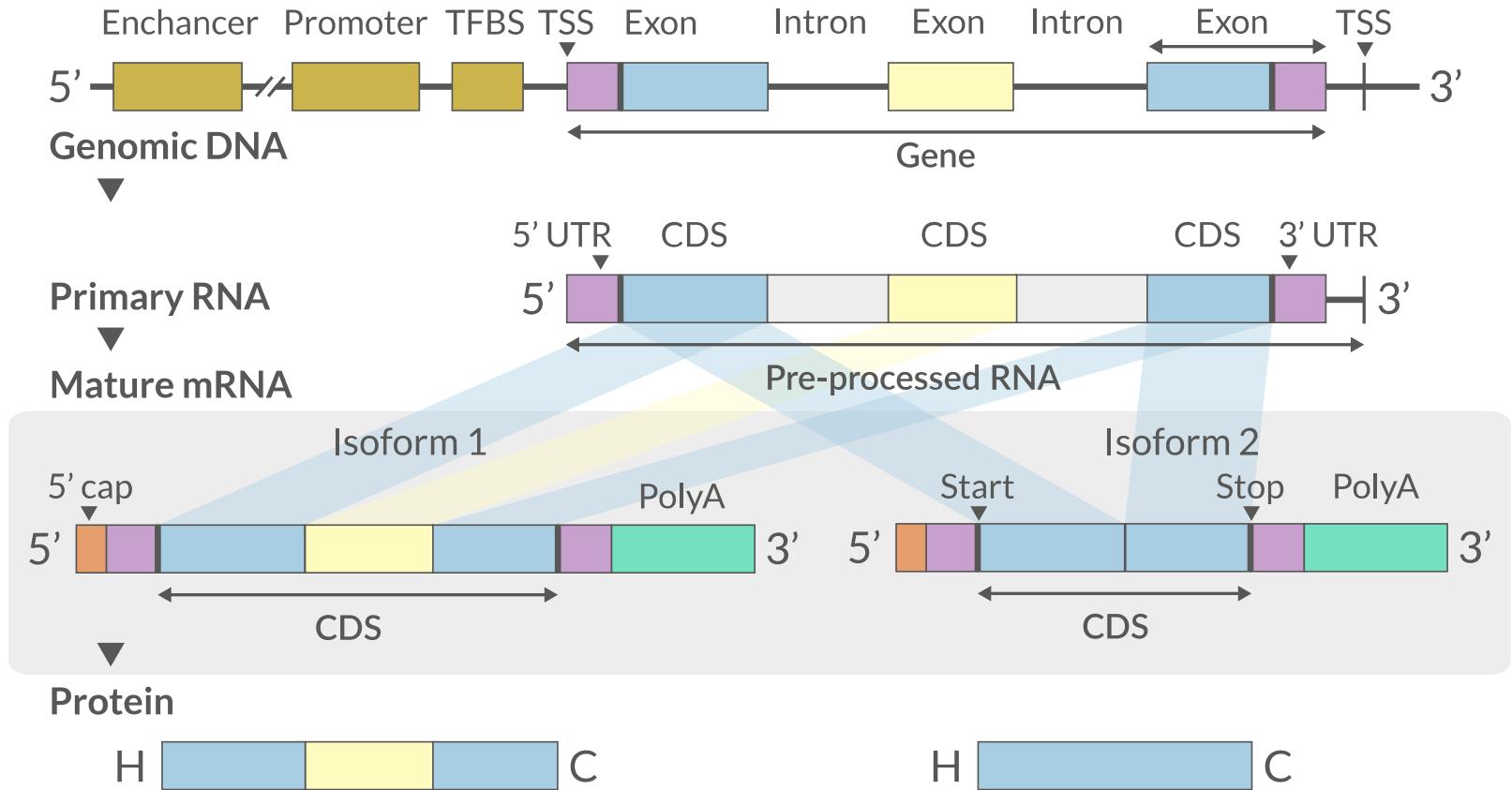
Introduction To Bioinformatics Using NGS Data

07-Oct-2022

# Contents

- RNA Sequencing
- Workflow
- DGE Workflow
- ReadQC
- Mapping
- Alignment QC
- Quantification
- Normalisation
- Exploratory
- DGE
- Functional analyses
- Summary
- Help

# What is RNA?



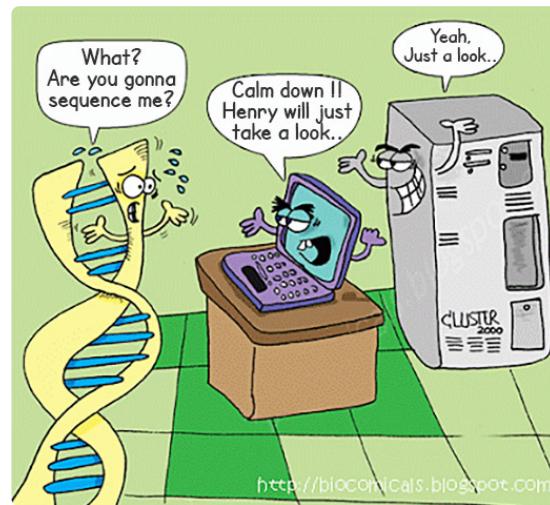
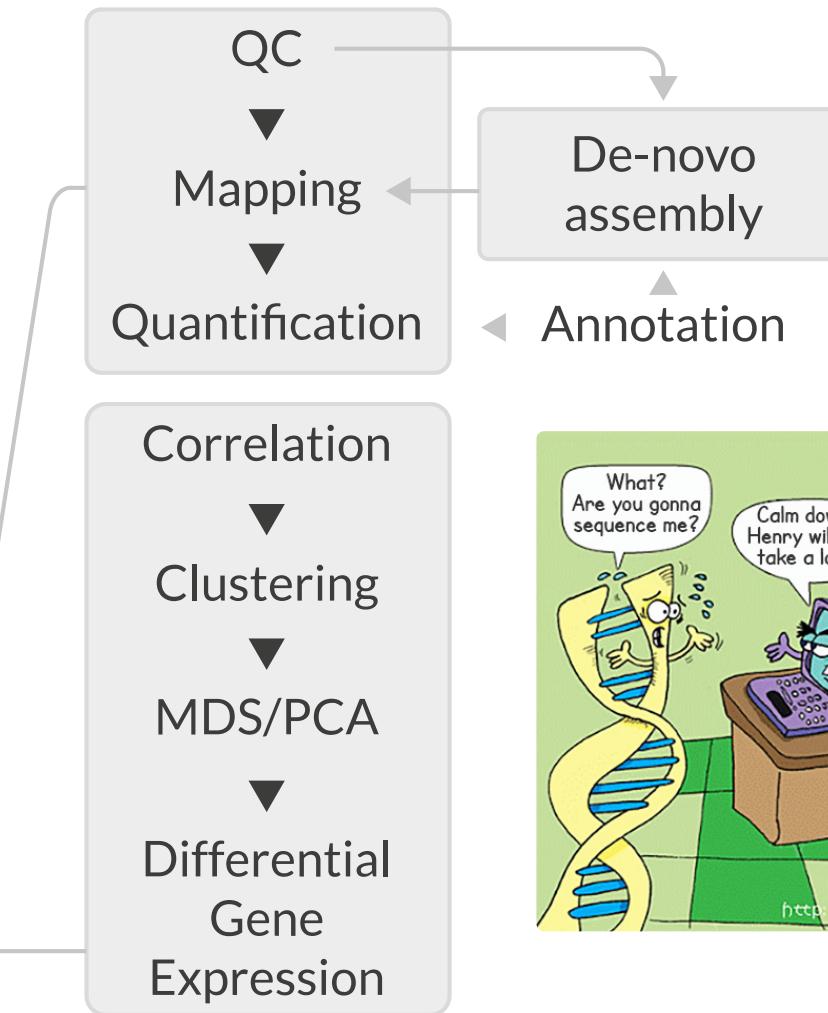
- The transcriptome is spatially and temporally dynamic
- Data comes from functional units (coding regions)
- Only a tiny fraction of the genome

# Applications

- Identify gene sequences in genomes (annotation)
- Learn about gene function
- Differential gene expression
- Explore isoform and allelic expression
- Understand co-expression, pathways and networks
- Gene fusion
- RNA editing

# Workflow

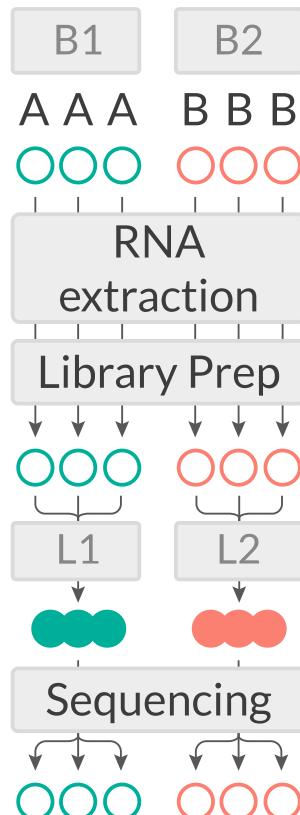
Experimental design  
▼  
RNA extraction  
▼  
Library preparation  
▼  
Sequencing  
▼  
Data processing  
▼  
Data analysis



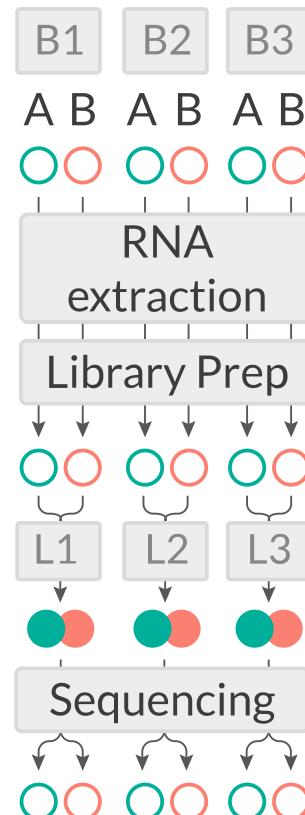
# Experimental design

- Biological replicates: 6 - 12 [SChurch et al, 2016](#)
  - Sample size estimation [Hart et al, 2013](#)
  - Power analysis [RNASeqPower](#) [RNASeqPower web app](#)
  - Balanced design to avoid batch effects
- [experDesign](#) [DeclareDesign](#)
- RIN values have strong effect [Romero et al, 2014](#)

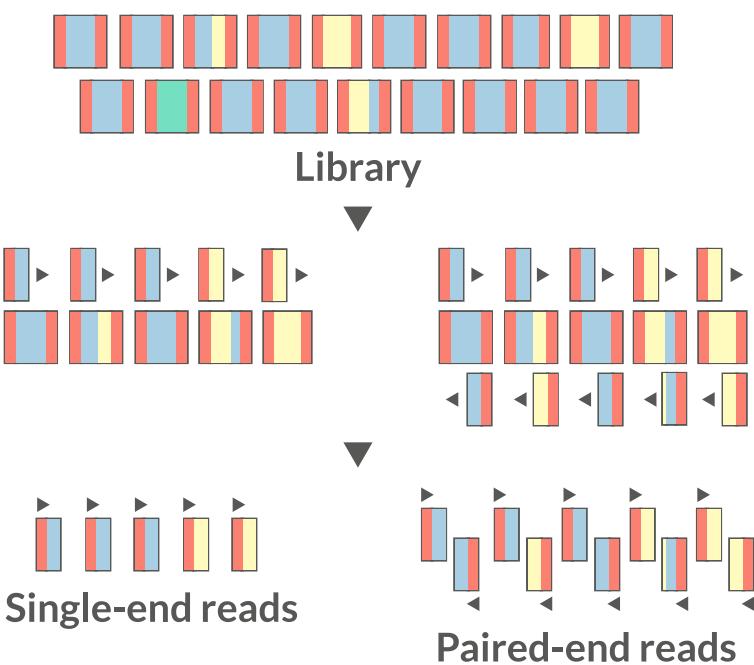
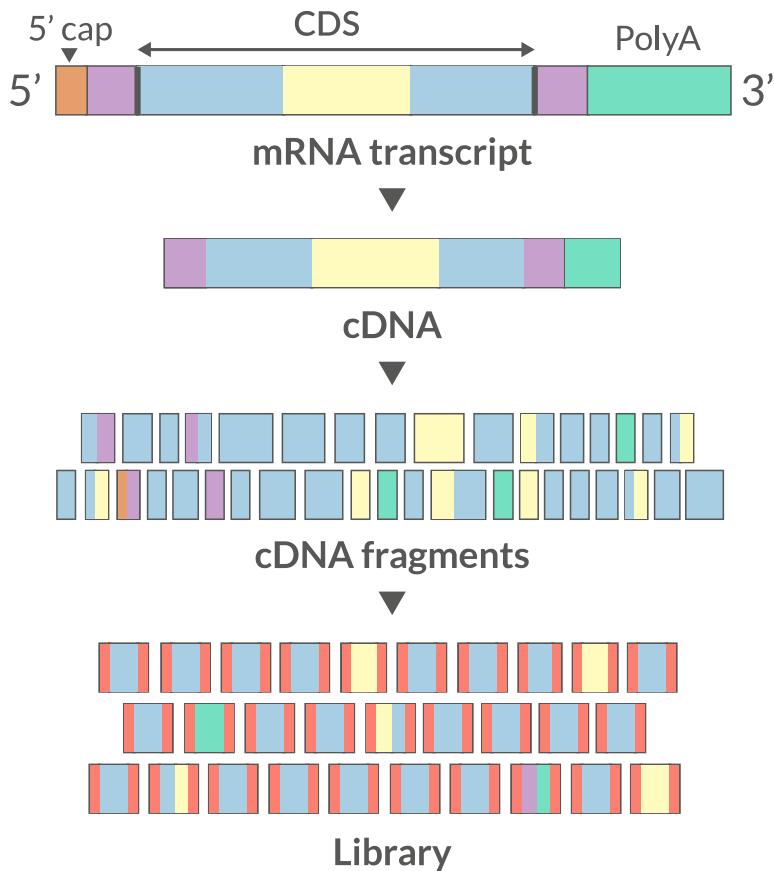
## Confounding



## Balanced

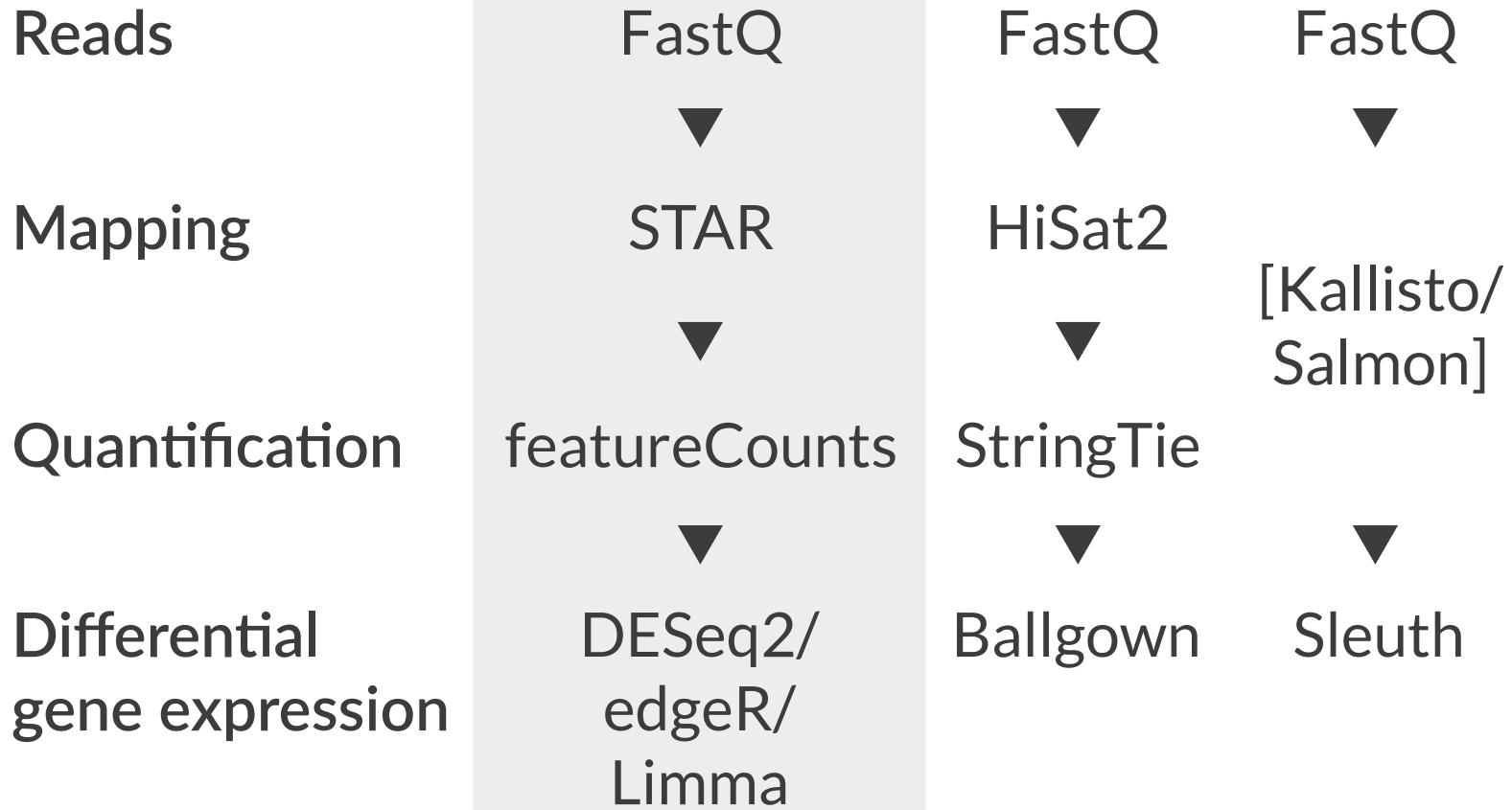


# Library & Sequencing



- polyA selection/Ribosomal RNA depletion
- single-end/Paired-end

# Workflow • DGE



# Read QC

- Number of reads
- Per base sequence quality
- Per sequence quality score
- Per base sequence content
- Per sequence GC content
- Per base N content
- Sequence length distribution
- Sequence duplication levels
- Overrepresented sequences
- Adapter content
- Kmer content

 FastQC  MultiQC

<https://sequencing.qcfail.com/>



 QCFAIL.com

Articles about common next-generation sequencing problems

# FastQC

## Good quality

**FastQC Report**

Thu 21 Dec 2017  
good\_sequence\_short.txt

**Summary**

- Basic Statistics
- Per base sequence quality
- Per tile sequence quality
- Per sequence quality scores
- Per base sequence content
- Per sequence GC content
- Per base N content
- Sequence Length Distribution
- Sequence Duplication Levels
- Overrepresented sequences
- Adapter Content

**Basic Statistics**

Measure	Value
Filename	good_sequence_short.txt
File type	Conventional base calls
Encoding	Illumina 1.5
Total Sequences	250000
Sequences flagged as poor quality	0
Sequence length	40
%GC	45

**Per base sequence quality**

Quality scores across all bases (illumina 1.5 encoding)

This figure is a Phred quality score plot. The x-axis represents the sequence length (40), and the y-axis represents the quality score (30 to 38). The plot shows a high proportion of high-quality bases (green) with a few low-quality bases (yellow).

## Poor quality

**FastQC Report**

Thu 21 Dec 2017  
bad\_sequence.txt

**Summary**

- Basic Statistics
- Per base sequence quality
- Per tile sequence quality
- Per sequence quality scores
- Per base sequence content
- Per sequence GC content
- Per base N content
- Sequence Length Distribution
- Sequence Duplication Levels
- Overrepresented sequences
- Adapter Content

**Basic Statistics**

Measure	Value
Filename	bad_sequence.txt
File type	Conventional base calls
Encoding	Illumina 1.5
Total Sequences	395288
Sequences flagged as poor quality	0
Sequence length	40
%GC	47

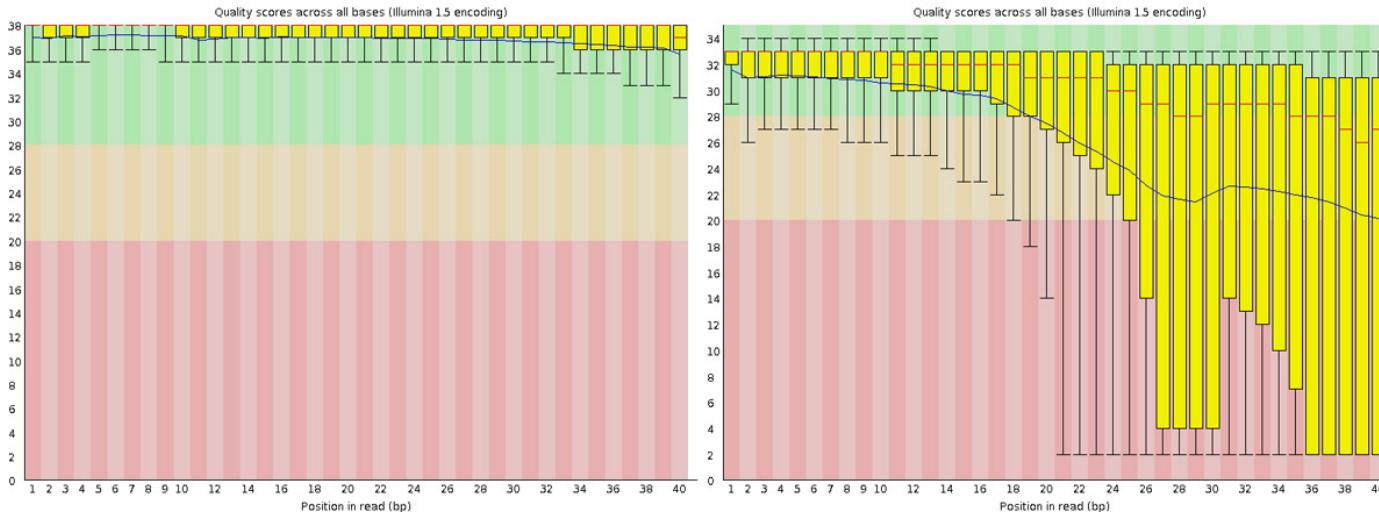
**Per base sequence quality**

Quality scores across all bases (illumina 1.5 encoding)

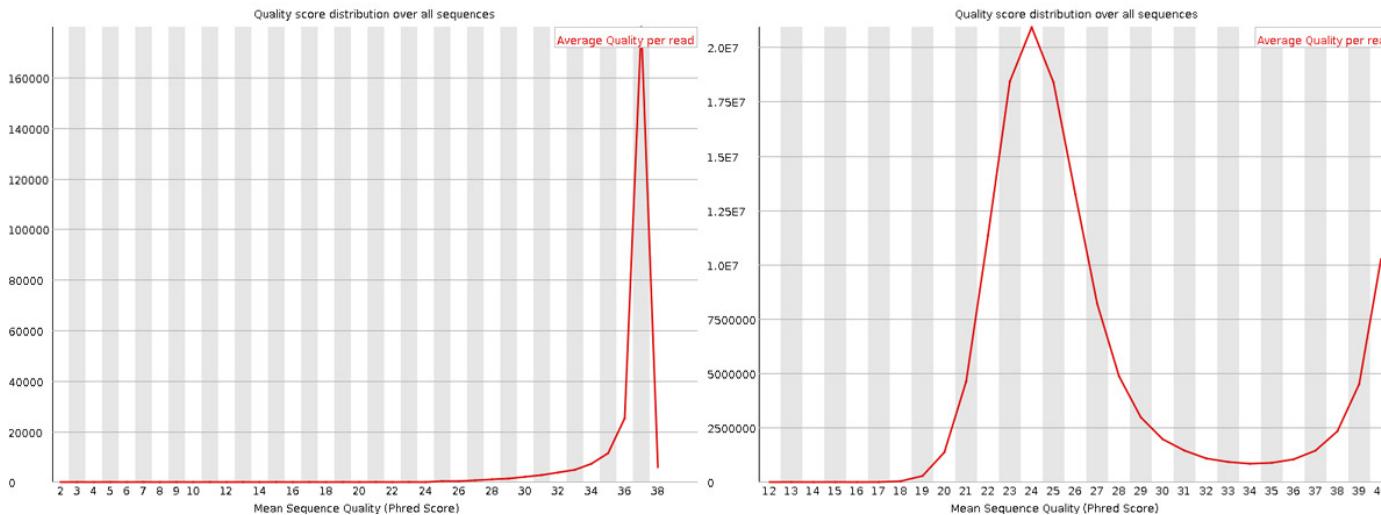
This figure is a Phred quality score plot. The x-axis represents the sequence length (40), and the y-axis represents the quality score (32 to 34). The plot shows a large number of low-quality bases (yellow) interspersed with some high-quality bases (green).

# Read QC • PBSQ, PSQS

## Per base sequence quality



## Per sequence quality scores



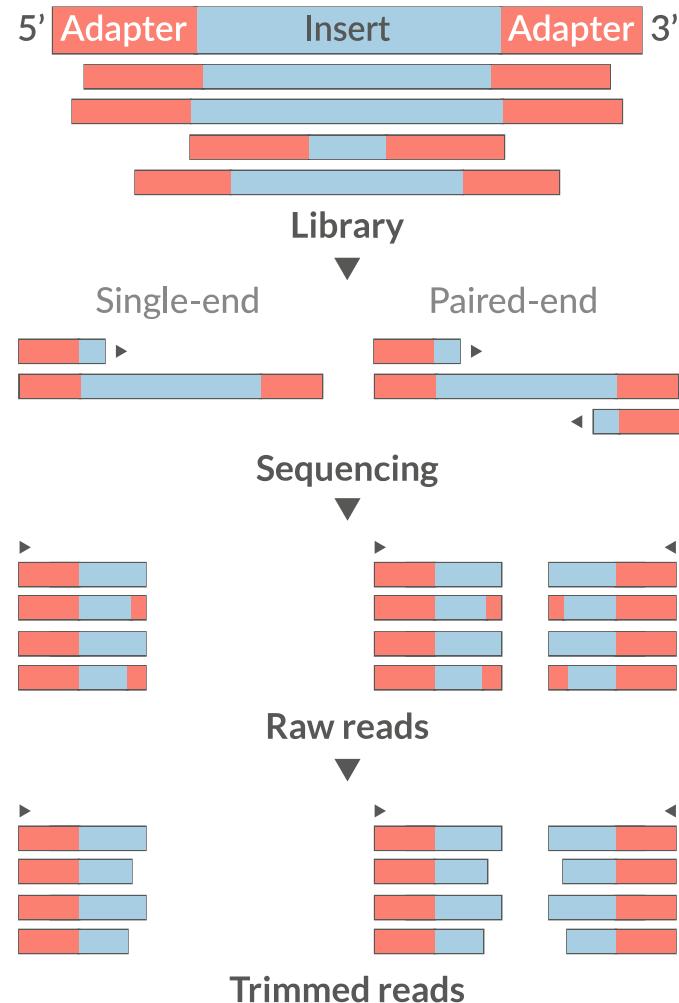
# Trimming

- Trimming reads to remove adapter/readthrough or low quality bases
- Related options are hard clipping, filtering reads
- Sliding window trimming
- Filter by min/max read length
  - Remove reads less than ~18nt
- Demultiplexing/Splitting

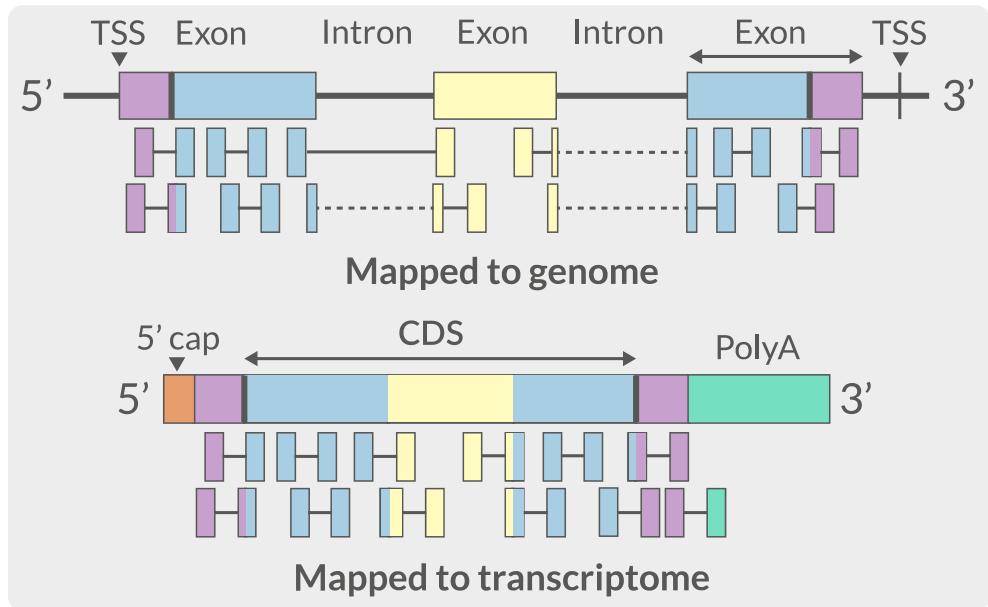
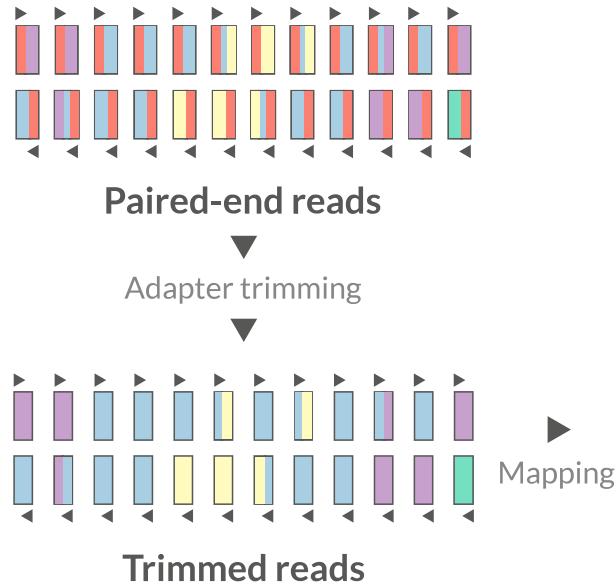
## When to avoid trimming?

- Read trimming may not always be necessary Liao et al, 2020
- Fixed read length may sometimes be more important
- Expected insert size distribution may be more important for assemblers

Cutadapt fastp Prinseq

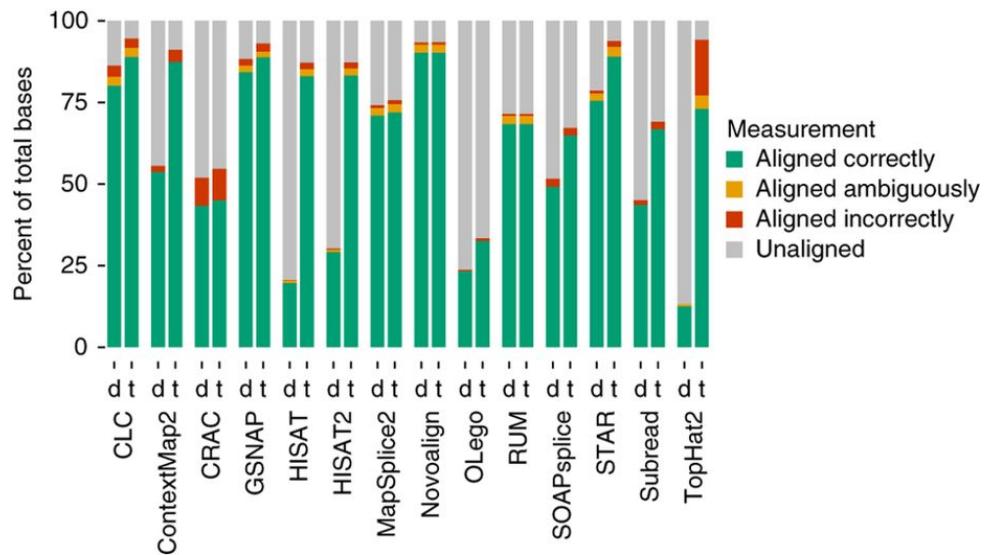
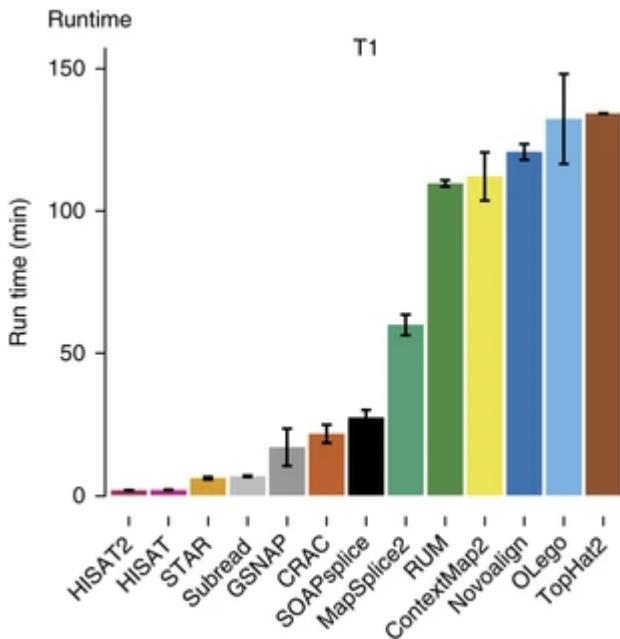


# Mapping



- Aligning reads back to a reference sequence
- Mapping to genome vs transcriptome
- Splice-aware alignment (genome) (STAR, HISAT2 etc)

# Aligners • Metrics



Increasing Accuracy  
↑

- Novel variants / RNA editing
- Allele-specific expression
- Genome annotation
- Gene and transcript discovery
- Differential expression

Baruzzo et al, 2017

# Mapping

- Reads (FASTQ)

```
@ST-E00274:179:HHYMLALXX:8:1101:1641:1309 1:N:0:NGATGT
NCATCGTGGTATTGCACATCTTTCTTATCAAATAAAAGTTAACCTACTCAGTTATGCCATACGTTTTGATGGCATTCCATAAA
+
#AAAFAYA<-AFFJJJAFA-FFJJJJFFF AJJJJ-<FFJJJ-A-F-7--FA7F7----FFFJFA<FFFFJ<AJ--FF-A<A-<JJ-7-7-
```

`@instrument:rnid:flowcellid:lane:tile:xpos:ypos read:isfiltered:controlnumber:sampleid`

- Reference Genome/Transcriptome (FASTA)

```
>1 dna:chromosome chromosome:GRCz10:1:1:58871917:1 REF
GATCTTAAACATTATTCCCCCTGCAAACATTTCATCATTGTCATTCCCCTC
CAAATTAAATTAGCCAGAGGCGCACACATACGACCTCTAAAAAGGTGCTGTAACATG
```

- Annotation (GTF/GFF)

```
#!genome-build GRCz10
#!genebuild-last-updated 2016-11
4      ensembl_havana  gene    6732     52059     .       -       .       gene_id "ENSDARG000
```

`seq source feature start end score strand frame attribute`

[Illumina FASTQ format](#) [GTF format](#)

# Alignment

- SAM/BAM (Sequence Alignment Map format)

```
ST-E00274:188:H3JWNCCXY:4:1102:32431:49900      163      1      1      60      8S139M4S
```

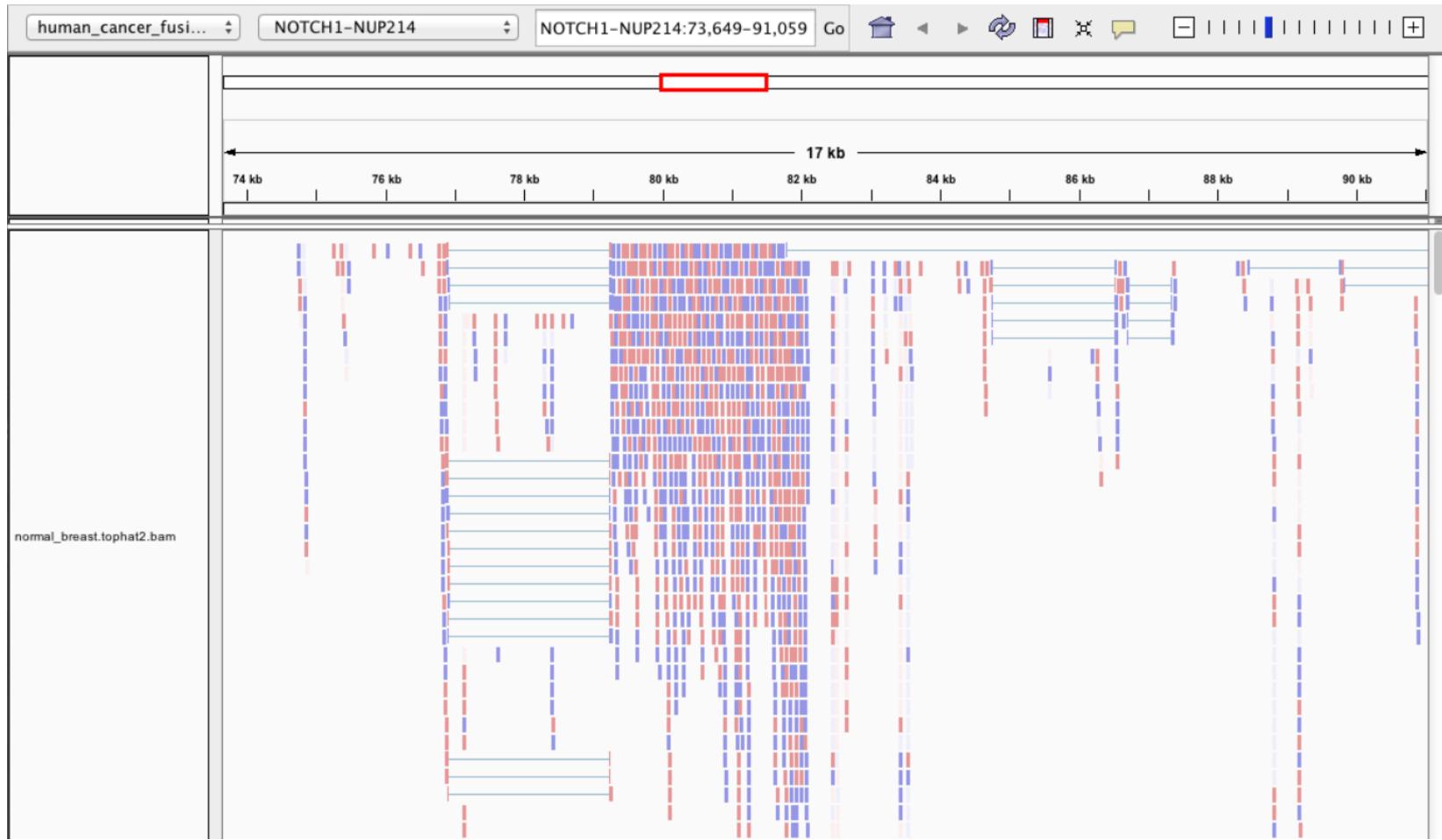
```
query flag ref pos mapq cigar mrnm mpos tlen seq qual opt
```

Never store alignment files in raw SAM format. Always compress it!

Format	Size_ GB
SAM	7.4
BAM	1.9
CRAM lossless Q	1.4
CRAM 8 bins Q	0.8
CRAM no Q	0.26

 SAM format

# Visualisation • IGV



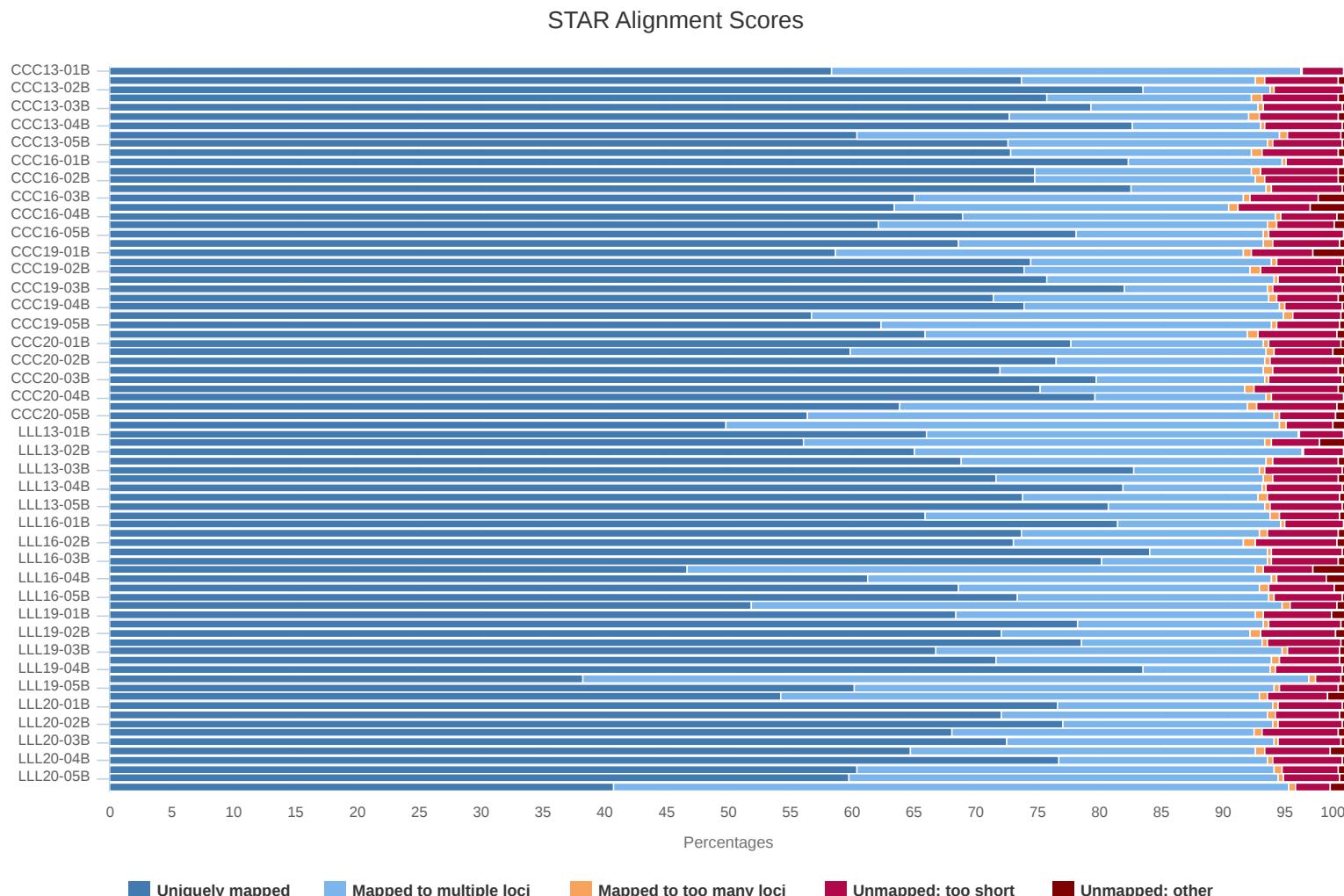
# Alignment QC

- Number of reads mapped/unmapped/paired etc
- Uniquely mapped
- Insert size distribution
- Coverage
- Gene body coverage
- Biotype counts / Chromosome counts
- Counts by region: gene/intron/non-genic
- Sequencing saturation
- Strand specificity

[!\[\]\(e2906a780c2bbcdc2a236d79598e58f1\_img.jpg\) STAR \(final log file\)](#) [!\[\]\(b46c7f04a8d398c60eb357f7415c967f\_img.jpg\) samtools stats](#) [!\[\]\(33ab5a78aa84cc698a8e599bbdc3ba81\_img.jpg\) bamtools stats](#) [!\[\]\(52b5558e6ed6332c21ac1989524fe688\_img.jpg\) QoRTs](#) [!\[\]\(c6ae25a62a813d4a80bdbcff336ea8c8\_img.jpg\) RSeQC](#) [!\[\]\(57d42b82c6f4e7c1587c95824bdeeac7\_img.jpg\) Qualimap](#)

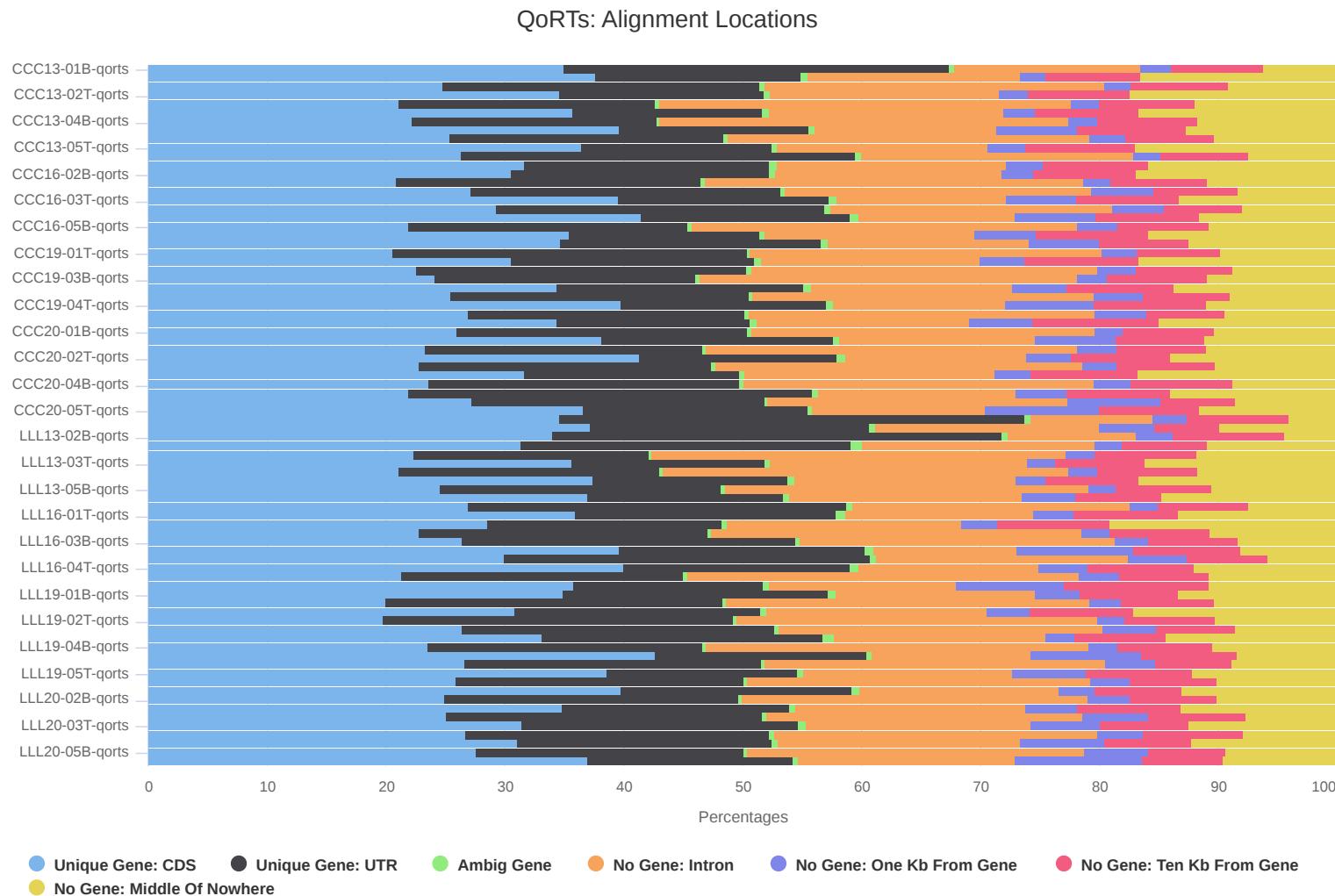
# Alignment QC • STAR Log

MultiQC can be used to summarise and plot STAR log files.

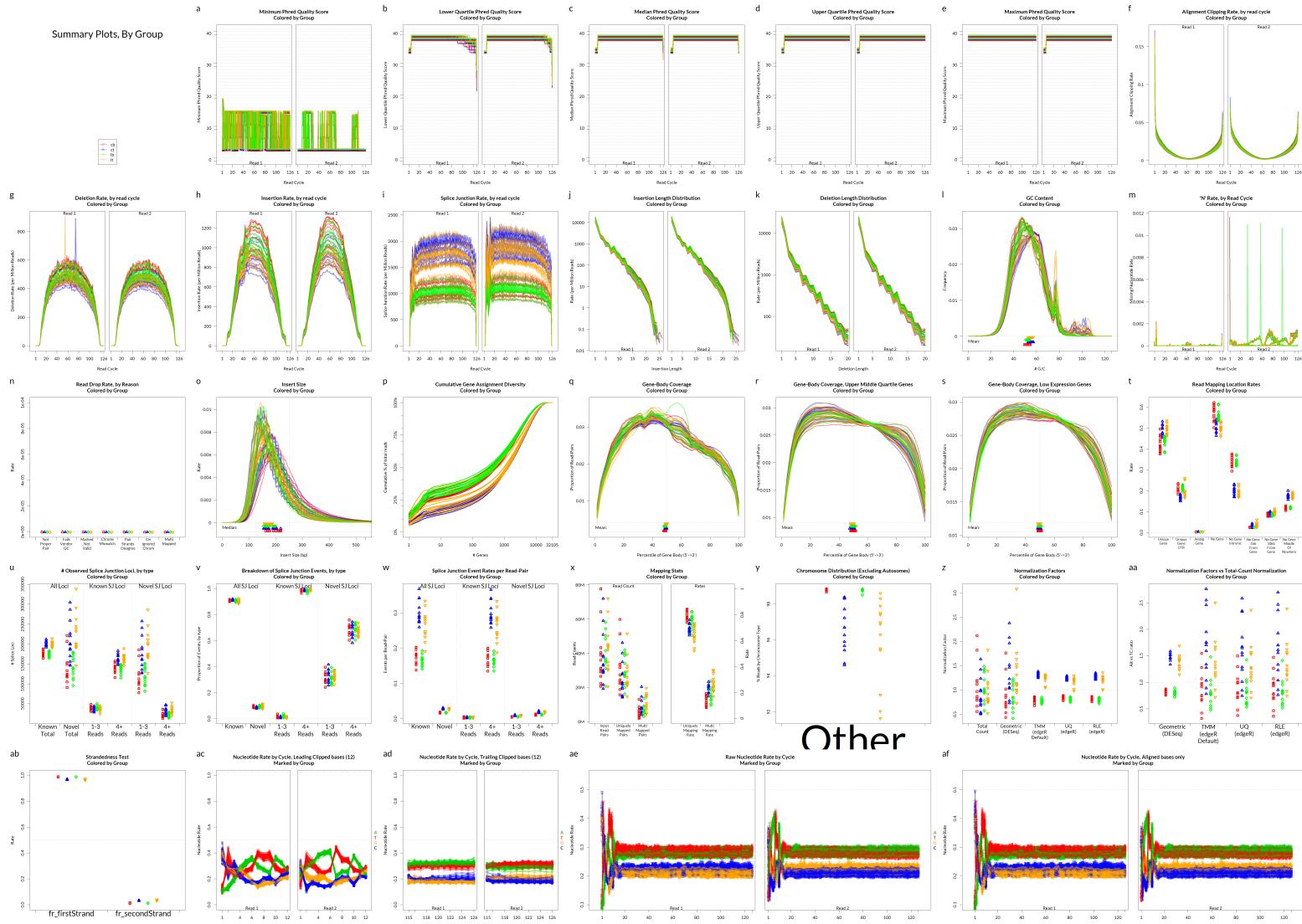


# Alignment QC • Features

QoRTs was run on all samples and summarised using MultiQC.



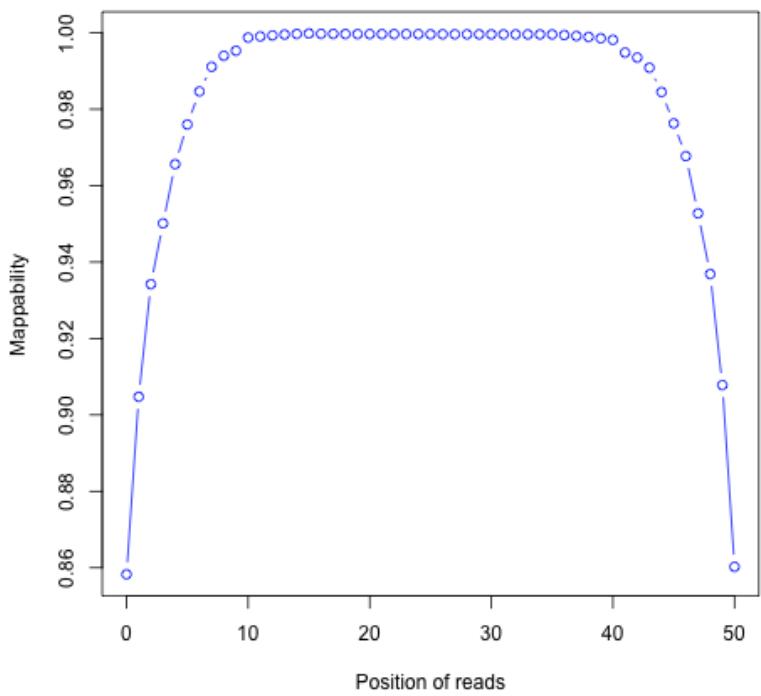
# Alignment QC • QoRTs



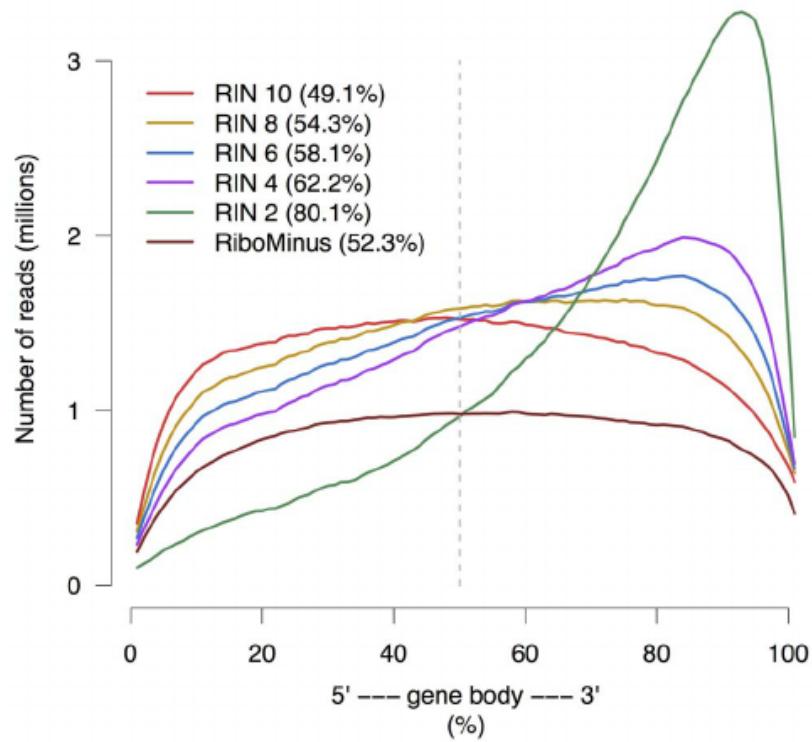
Other

# Alignment QC • Examples

Read mapping profile



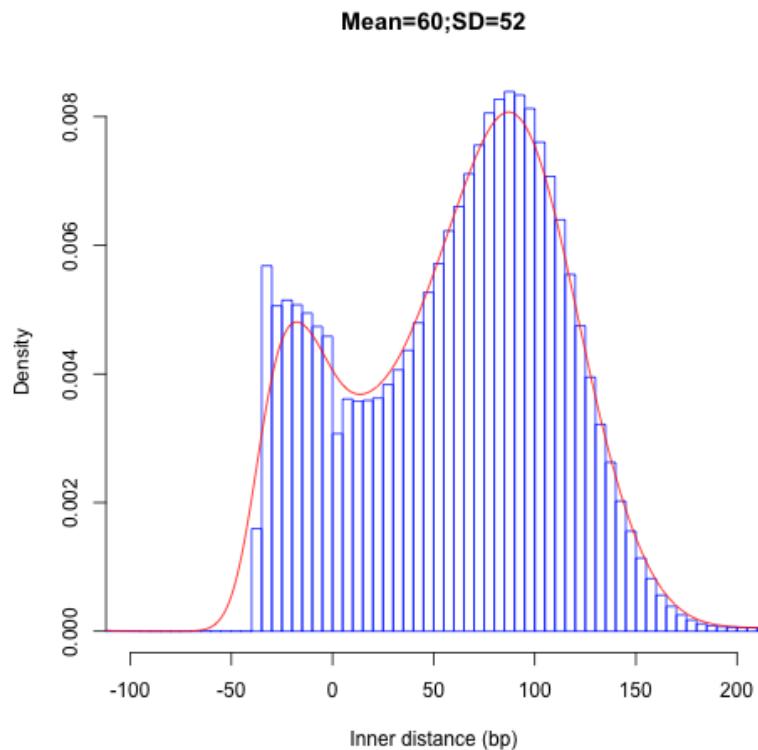
Gene body coverage



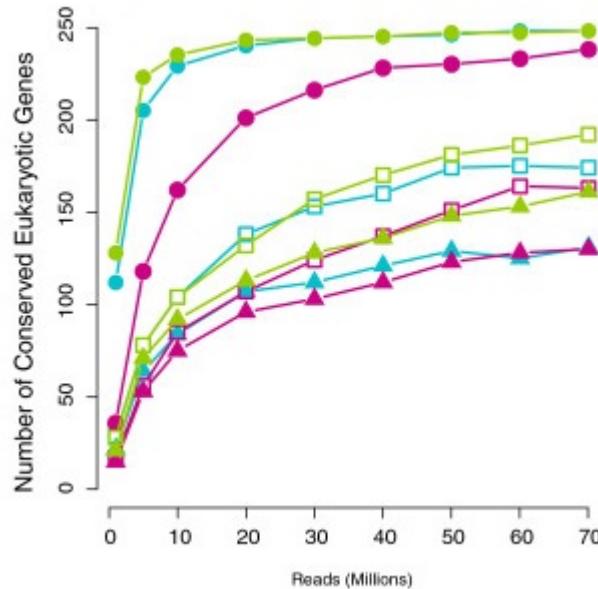
Sigurgeirsson et al, 2014

# Alignment QC • Examples

## Insert size



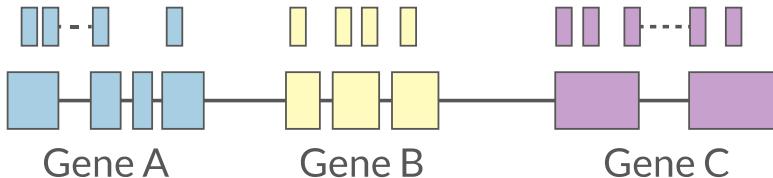
## Saturation curve



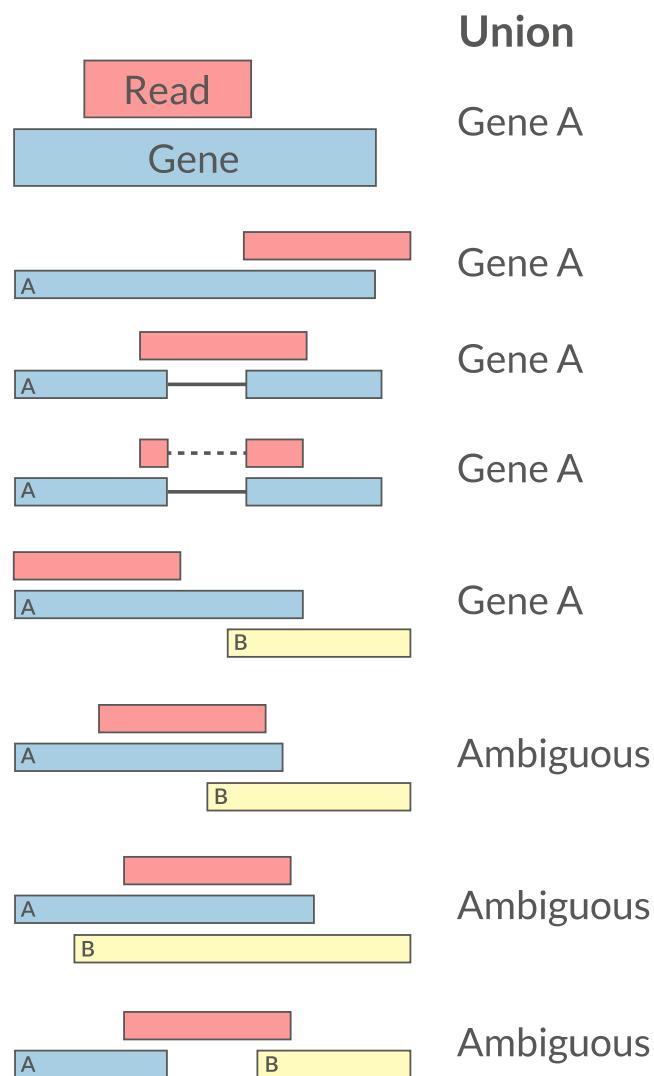
Francis et al, 2013

# Quantification • Counts

- Read counts = gene expression
- Reads can be quantified on any feature (gene, transcript, exon etc)
- Intersection on **gene models**
- Gene/Transcript level



featureCounts HT Seq



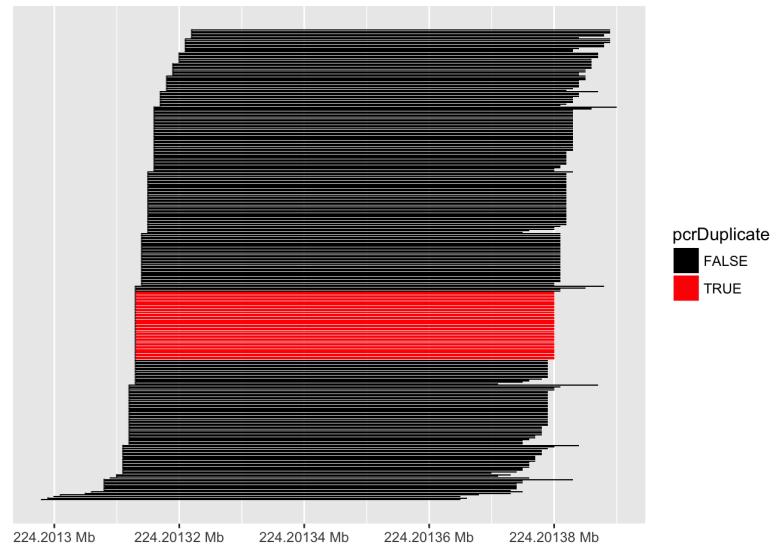
# Quantification

## PCR duplicates

- Computational deduplication not recommended [Klepikova et al, 2017](#) [Parekh et al, 2016](#)
- Use PCR-free library-prep kits
- Use UMIs during library-prep [Fu et al, 2018](#)

## Multi-mapping

- Added (BEDTools multicov)
- Discard (featureCounts, HTSeq)
- Distribute counts (Cufflinks, featureCounts)
- Rescue
  - Probabilistic assignment (Rcount, Cufflinks)
  - Prioritise features (Rcount)
  - Probabilistic assignment with EM (RSEM)



# Quantification • Abundance

- Count methods
  - Provide no inference on isoforms
  - Cannot accurately measure fold change
- Probabilistic assignment
  - Deconvolute ambiguous mappings
  - Transcript-level
  - cDNA reference

## Kallisto, Salmon

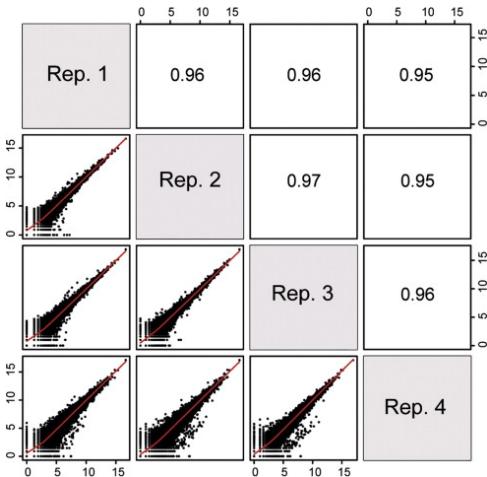
- Ultra-fast & alignment-free
- Bootstrapping & quantification confidence
- Transcript-level counts
- Transcript-level estimates improves gene-level estimates  [Soneson et al, 2015](#)  [tximport](#)
- Evaluation and comparison of isoform quantification tools  [Zhang et al, 2017](#)

 RSEM  Kallisto  Salmon

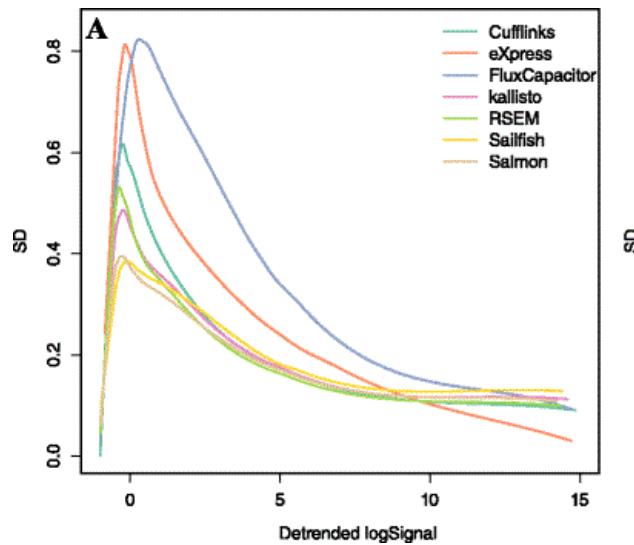
# Quantification QC

ENSG000000000003	140	242	188	143	287	344	438	280	253
ENSG000000000005	0	0	0	0	0	0	0	0	0
ENSG000000000419	69	98	77	55	52	94	116	79	69
ENSG000000000457	56	75	104	79	157	205	183	178	153
ENSG000000000460	33	27	23	19	27	42	69	44	40
ENSG000000000938	7	38	13	17	35	76	53	37	24
ENSG000000000971	545	878	694	636	647	216	492	798	323
ENSG000000001036	79	154	74	80	128	167	220	147	72

- Pairwise correlation between samples must be high (>0.9)



- Count QC using RNASeqComp



**MultiQC**  
v1.6

- General Stats
- featureCounts
- STAR
- Cutadapt
- FastQC
- Sequence Counts
- Sequence Quality Histograms
- Per Sequence Quality Scores
- Per Base Sequence Content
- Per Sequence GC Content
- Per Base N Content
- Sequence Length Distribution
- Sequence Duplication Levels
- Overrepresented sequences
- Adapter Content

# MultiQC

A modular tool to aggregate results from bioinformatics analyses across many samples into a single report.

Report generated on 2018-08-04, 01:51 based on data in: /Users/ewels/GitHub/MultiQC\_website/public\_html/examples/rna-seq

## General Statistics

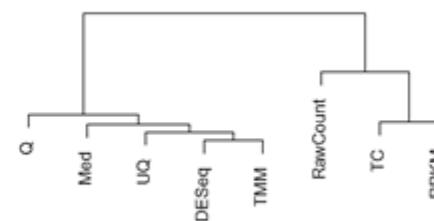
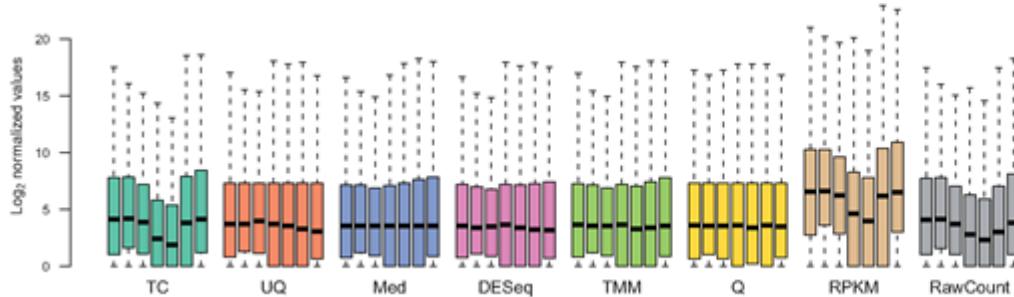
Copy table
Configure Columns
Plot
Showing 8/8 rows and 8/10 columns.

Sample Name	% Assigned	M Assigned	% Aligned	M Aligned	% Trimmed	% Dups	% GC	M Seqs
SRR3192396	67.5%	71.9	93.7%	97.8	4.0%	78.9%	51%	104.4
SRR3192397	66.6%	63.0	94.7%	87.1	3.5%	77.2%	49%	92.0
SRR3192398	50.9%	36.5	88.2%	58.7	5.0%	55.3%	47%	66.6
SRR3192399	52.3%	42.3	88.2%	65.6	5.0%	57.4%	47%	74.3
SRR3192400	70.3%	63.4	77.3%	73.4	7.2%	74.1%	45%	94.9
SRR3192401	71.2%	63.8	76.4%	72.8	6.3%	76.3%	45%	95.2
SRR3192657	73.1%	67.1	91.2%	85.0	3.1%	82.2%	51%	93.1
SRR3192658	71.2%	66.9	89.7%	87.1	3.4%	82.3%	52%	97.1

Toolbox
A
D
Download
H
?

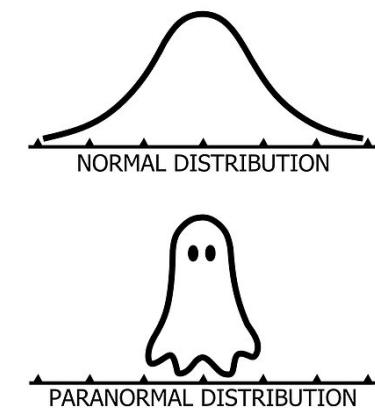
# Normalisation

- Control for Sequencing depth, compositional bias and more
- Median of Ratios (DESeq2) and TMM (edgeR) perform the best



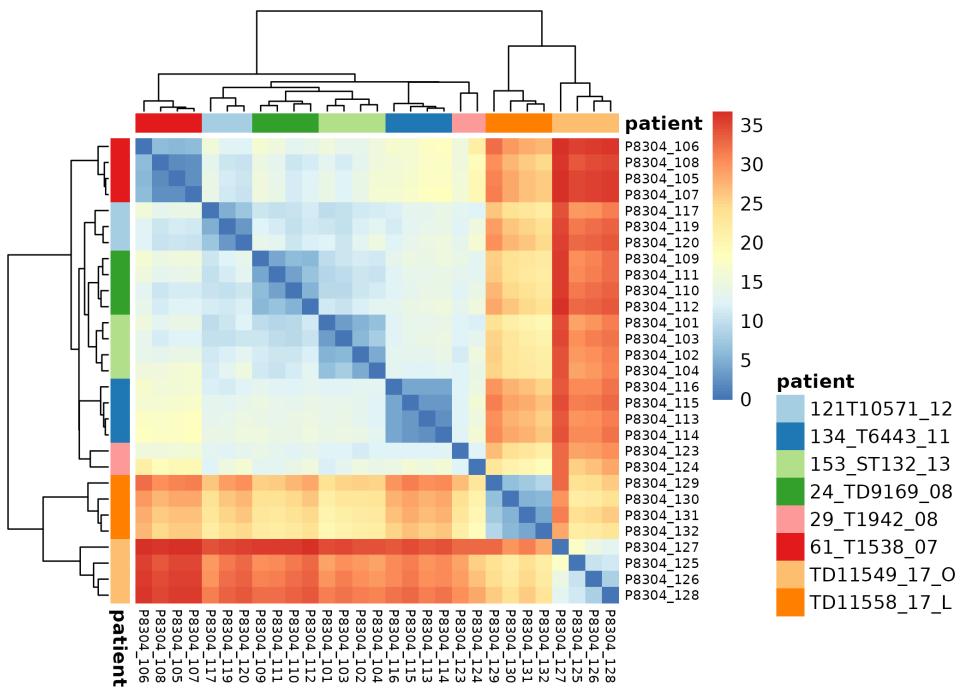
- For DGE using DGE packages, use raw counts
- For clustering, heatmaps etc use VST, VOOM or RLOG
- For own analysis, plots etc, use TPM
- Other solutions: spike-ins/house-keeping genes

Dillies et al, 2013 Evans et al, 2017 Wagner et al, 2012

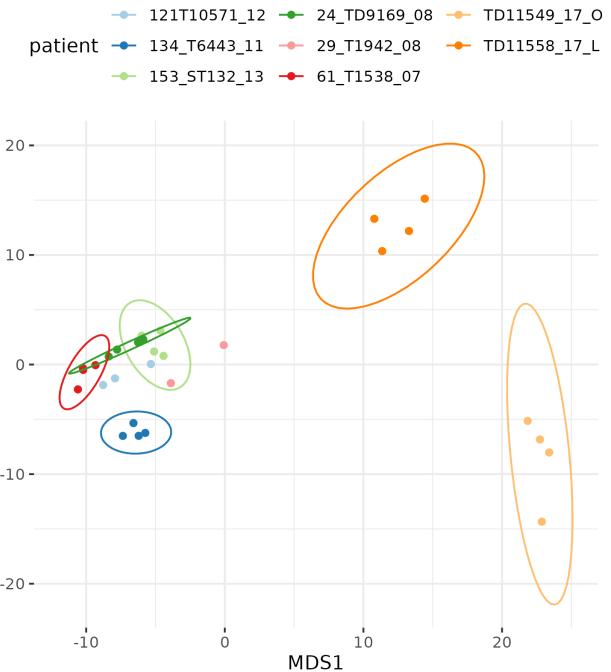


# Exploratory

- Remove lowly expressed genes
- Transform raw counts to VST, VOOM, RLOG, TPM etc
- Heatmaps, MDS, PCA etc.

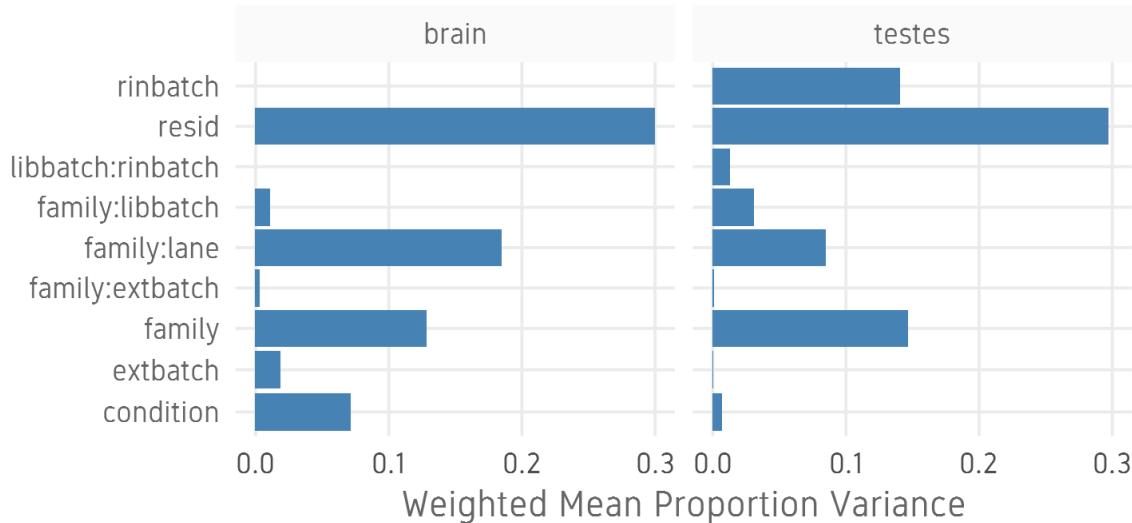


heatmaps



# Batch correction

- Estimate variation explained by variables (PVCA)  PVCA



- Find confounding effects as surrogate variables (SVA)  SVA
- Model known batches in the LM/GLM model
- Correct known batches (ComBat from SVA)(Only if you are desperate!  Zindler et al, 2020)
- Interactively evaluate batch effects and correction (BatchQC)  Manimaran et al, 2016  BatchQC

# Differential expression

		Gene A	Gene B	...	Gene N
Group 1	Sample 1	12	54	...	...
	Sample 2	8	47	...	...
	Sample 3	13	48	...	...
Group 2	Sample 1	22	50	...	...
	Sample 2	18	48	...	...
	Sample 3	25	41	...	...

- Univariate testing gene-by-gene
- More descriptive, less predictive

- Results `results()`

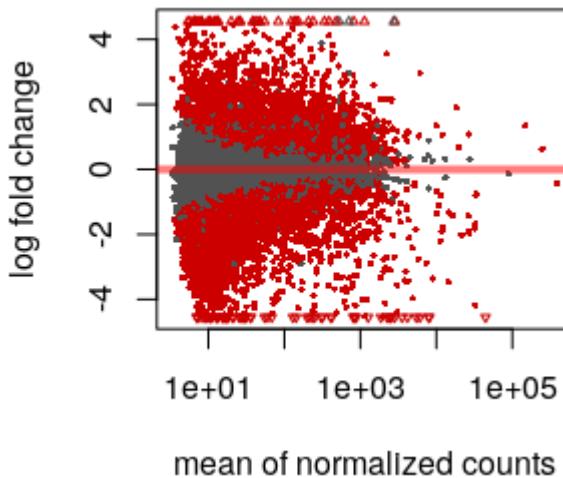
```
log2 fold change (MLE): type type2 vs control
Wald test p-value: type type2 vs control
DataFrame with 1 row and 6 columns
      baseMean    log2FoldChange        lfcSE
      <numeric>       <numeric>       <numeric>
ENSG000000000003 242.307796723287 -0.932926089608546 0.114285150312285
      stat          pvalue        padj
      <numeric>       <numeric>       <numeric>
ENSG000000000003 -8.16314356729037 3.26416150242775e-16 1.36240609998527e-14
```

- Summary `summary()`

```
out of 17889 with nonzero total read count
adjusted p-value < 0.1
LFC > 0 (up)      : 4526, 25%
LFC < 0 (down)     : 5062, 28%
outliers [1]       : 25, 0.14%
low counts [2]      : 0, 0%
(mean count < 3)
[1] see 'cooksCutoff' argument of ?results
[2] see 'independentFiltering' argument of ?results
```

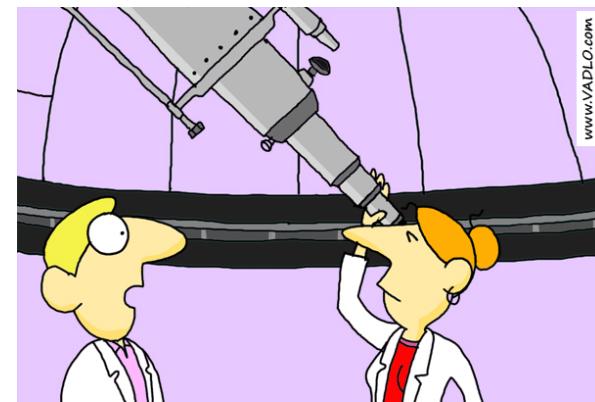
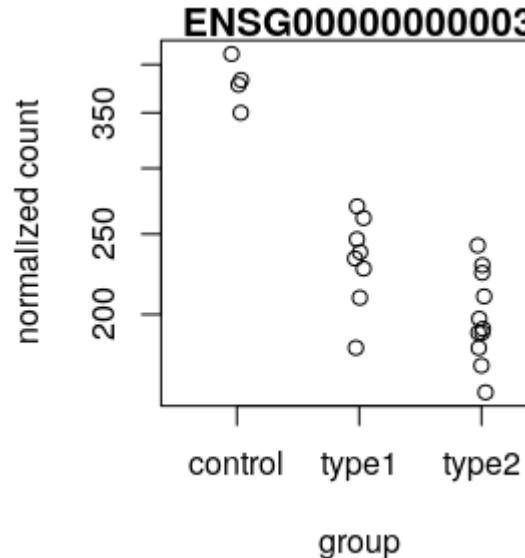
# DGE

- MA plot `plotMA()`



- Volcano plot

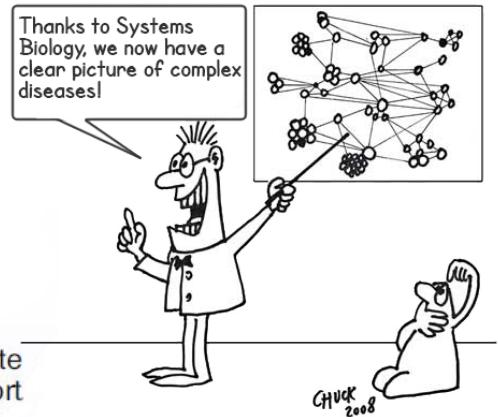
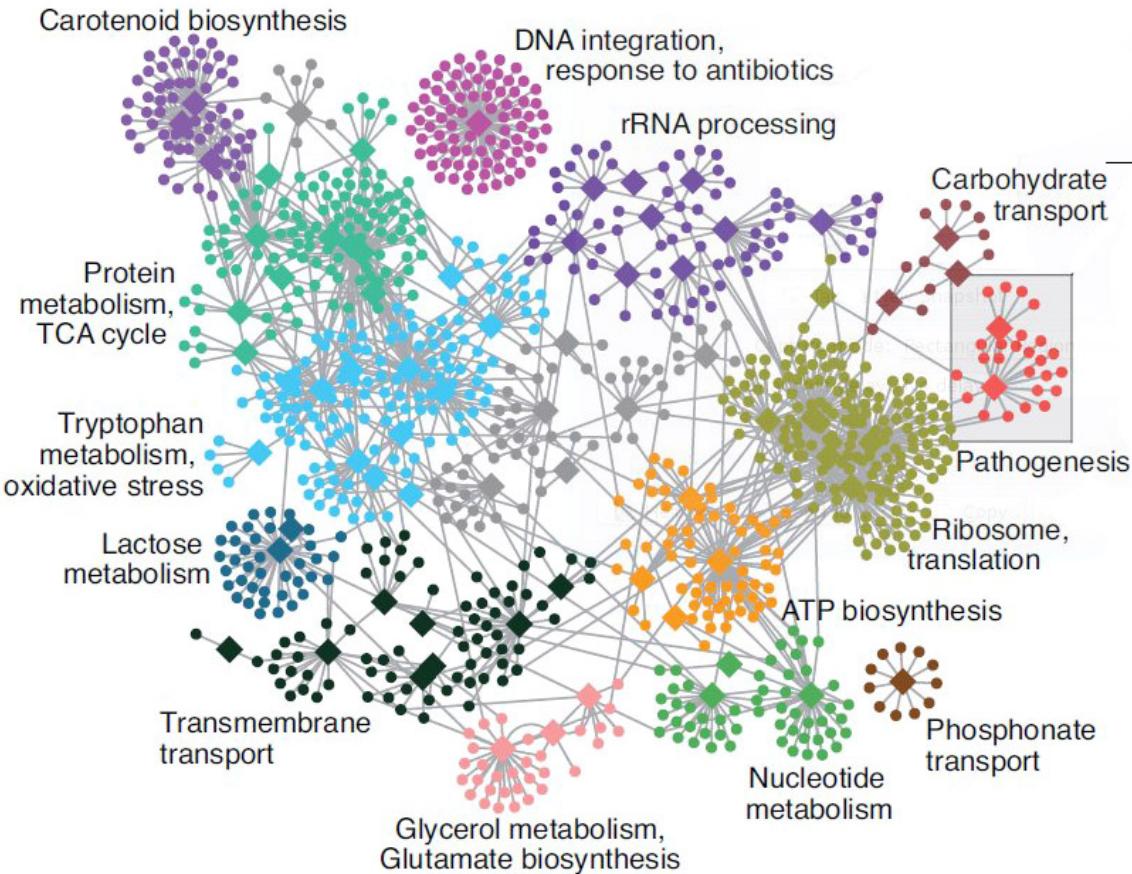
- Normalised counts `plotCounts()`



"Can you see the upper points of my scatter plot?"

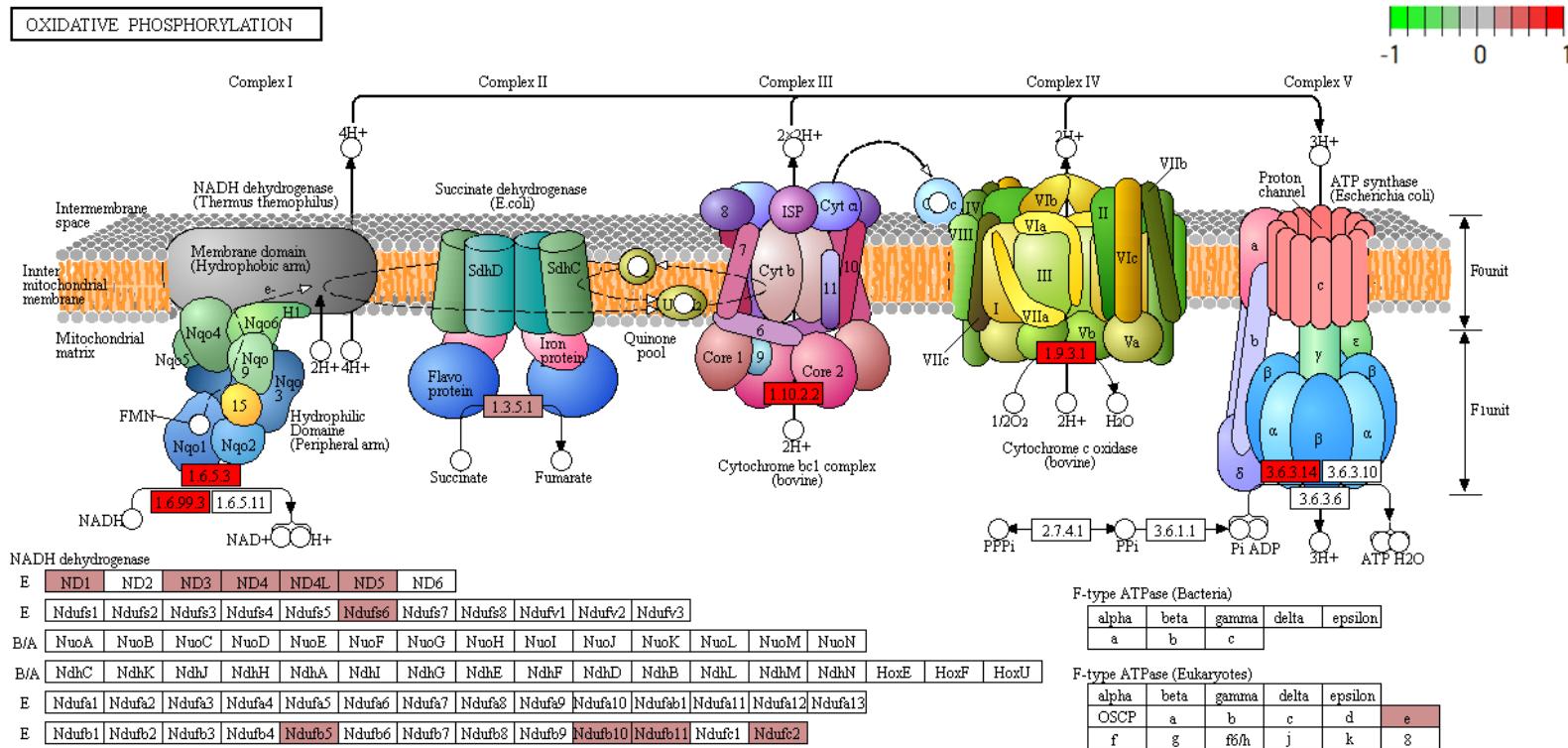
# Functional analysis • GO

- Gene set analysis (GSA)
- Gene set enrichment analysis (GSEA)
- Gene ontology / Reactome databases



# Functional analysis • Kegg

- Pathway analysis (Kegg)



# Summary

- Sound experimental design to avoid confounding
- Plan carefully about lib prep, sequencing etc based on experimental objective
- For DGE, biological replicates may be more important than other considerations (paired-end, sequencing depth, long reads etc)
- Discard low quality bases, reads, genes and samples
- Verify that tools and methods align with data assumptions
- Experiment with multiple pipelines and tools
- QC! QC everything at every step

Conesa, A., Madrigal, P., Tarazona, S., Gomez-Cabrero, D., Cervera, A., McPherson, A., ... & Mortazavi, A. (2016). A survey of best practices for RNA-seq data analysis. *Genome biology*, 17(1), 1-19.

Thank you. Questions?



# Hands-On tutorial

## Main exercise

- 01 Check the quality of the raw reads with **FastQC**
- 02 Map the reads to the reference genome using **HISAT2**
- 03 Assess the post-alignment quality using **QualiMap**
- 04 Count the reads overlapping with genes using **featureCounts**
- 05 Find differentially expressed genes using **DESeq2** in R

## Bonus exercises

- 01 Functional annotation of DE genes using **GO/Reactome/Kegg** databases
- 02 RNA-Seq figures and plots using **R**
- 03 Visualisation of RNA-seq BAM files using **IGV** genome browser

Data: </sw/courses/ngsintro/rnaseq/>

Work: </proj/snici2022-22-769/nobackup/user/rnaseq/>

# Hands-On tutorial

- Course data directory
- Your work directory

```
/sw/courses/ngsintro/rnaseq/
```

```
rnaseq/
+-- bonus/
|   +-- assembly/
|   +-- exon/
|   +-- funannot/
|   +-- plots/
+-- documents/
+-- main/
    +-- 1_raw/
    +-- 2_fastqc/
    +-- 3_mapping/
    +-- 4_qualimap/
    +-- 5_dge/
    +-- 6_multiqc/
    +-- reference/
        +-- mouse_chr19_hisat2/
+-- scripts/
```

```
/proj/snic2022-22-769/nobackup/user/rnaseq/
```

```
[user]/
rnaseq/
    +-- 1_raw/
    +-- 2_fastqc/
    +-- 3_mapping/
    +-- 4_qualimap/
    +-- 5_dge/
    +-- 6_multiqc/
    +-- reference/
        +-- mouse_chr19_hisat2/
    +-- scripts/
    +-- funannot/
    +-- plots/
```