

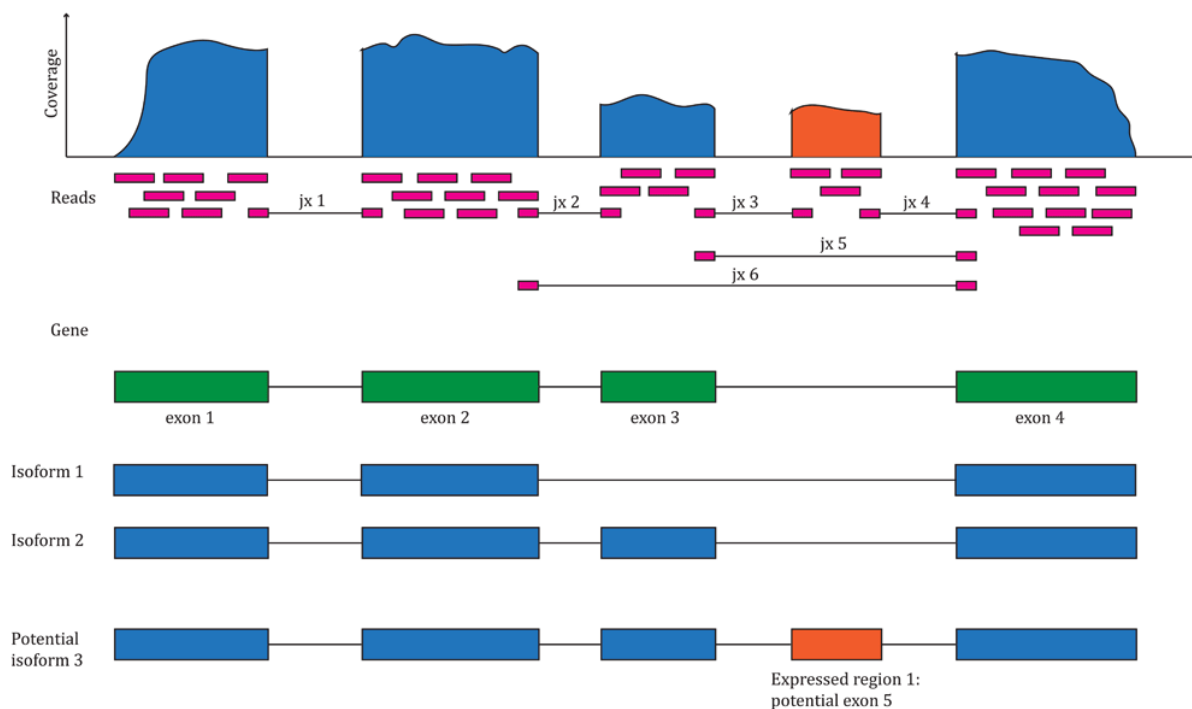
Introduction to RNAseq

Version 1

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Introduction to RNAseq



These are course notes for the “Introduction to RNAseq” course given by the Monash Bioinformatics Platform¹ for the Monash Data Fluency² initiative. Our teaching style is based on the style of The Carpentries³.

- PDF version for printing⁴

Source code

- GitHub page⁵

Authors and copyright

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¹<https://www.monash.edu/researchinfrastructure/bioinformatics>

²<https://monashdatafluency.github.io/>

³<https://carpentries.org/>

⁴<https://monashdatafluency.github.io/rnaseq-intro/rnaseq-intro.pdf>

⁵<https://github.com/MonashDataFluency/rnaseq-intro>



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Chapter 1

Genomics

Genome is a collection of all genetic material in the organism. To be more accurate each cell in our body contains a copy of entire genome. The study of genomics is all things to do with genome including gene expression, methylation profiles, transcriptional factor binding, chromatic arrangements and three dimensional structure of chromosomes and many more things.

In this course we are going to focus on one particular technique of genomics - RNA sequencing or RNAseq for short. This is a study of primarily but not limited to gene expression. Other common application of RNAseq are study of splicing and isoforms.

In genomics everything rotates around sequence information, let's look into biological sequence next.

1.1 Sequence

A sequence is a generic name describing the order of biological letters in molecules like DNA, RNA or ptiptide and protein. In the case of DNA and RNA letters mean nucleotide bases, **A**denine, **T**hymine, **C**ytosine **G**uanine or ATCG for short. Also know that mRNA contains **U**racil instead of Thymine. This would be one way to represent all five possible bases **A[T|U]CG** In the case of ptiptide/protein molecules each letter means amino acid type, I won't list discuss proteins any further in this book since it isn't a focus on this book.

Note that there is a greater nomenclature exists to cover all possible variances

##	Letter_code	Base_name
## 1	A	Adenine
## 2	C	Cytosine
## 3	G	Guanine
## 4	T U	Thymine Uracil
## 5	R	A G
## 6	Y	C T
## 7	S	G C
## 8	W	A T
## 9	K	G T
## 10	M	A C
## 11	B	C G T
## 12	D	A G T
## 13	H	A C T
## 14	V	A C G
## 15	N	any base

The reason we have sequence information of any kind is because someone somewhere made an afford to sequence some part of the molecule. In the case of whole genome sequencing an afford was made to sequence the entire genome, for human the entire genome is 3 billion bases long. However this is a sum of

all bases across all chromosomes, but chromosomes are not all equal, below is a sequence length of each chromosome.

##	Chrom	Length
## 1	1	248956422
## 2	2	242193529
## 3	3	198295559
## 4	4	190214555
## 5	5	181538259
## 6	6	170805979
## 7	7	159345973
## 8	8	145138636
## 9	9	138394717
## 10	10	133797422
## 11	11	135086622
## 12	12	133275309
## 13	13	114364328
## 14	14	107043718
## 15	15	101991189
## 16	16	90338345
## 17	17	83257441
## 18	18	80373285
## 19	19	58617616
## 20	20	64444167
## 21	21	46709983
## 22	22	50818468
## 23	X	156040895
## 24	Y	57227415
## 25	MT	16569

While sequence is a generic name and often times it needs some context around it, for example DNA sequence, RNA sequence or chromosome 3 sequence. There are other names like contig and a read, which are both sequences too, they mean more specific thing. In general both mean shorter sequence, where contig almost always should be longer than a read.

Let's talk about a read sequence next.

1.2 Read sequence

A read is product of a machine read out that takes some biological material i.e DNA or RNA sample and produces number of reads that resembles what you've put into the machine. There have been a few iteration of sequencing technologies, each iteration brings about higher yield (more reads) and longer reads. Current players in the genomics space are:

Short read technologies

- Illumina¹
 - MiSeq
 - HiSeq
 - NextSeq
 - NovaSeq
- BGI platforms²
 - BGISEQ-50
 - BGISEQ-500

¹<https://www.illumina.com/>

²<https://www.bgi.com/global/resources/sequencing-platforms/>

- Ion Torrent³

Medium to long read technologies

- PacBio⁴
 - Sequel System
- Oxford Nanopore⁵
 - MinION
 - GridION
 - PromethION

All of the above technologies will one way or another result in digital reads. As name implies short reads are typically 50-300 bases, whereas long reads can range from 1k to 10k or more. Usually the quantity and the length of the read are inversely proportional, however the throughput of a short reads technology at the moment is vastly greater than the long read and it doesn't suffer from the coverage issue. In terms of number of reads we are talking about millions to billions for short reads technology and in the range of 1 to 4 millions for long reads. This is a good plot showing nanopore output, which illustrates quantity vs length nicely.

Getting the reads is just a first step. You have to do something with your reads. The two path one can take are really assemble your reads into contigs and further into full genome or map your reads to an existing reference genome. In this course we'll be taking the later path - mapping our RNAseq reads to an existing reference genome. But let me just talk a little more about what a contig sequence is next.

1.3 Contig sequence

Contig has specific meaning in genome assembly. It is a larger fragment assembled from short (100-300 bp) reads that hasn't yet been merged into chromosome. However often times this term can be loosely used to mean any long stretch of sequence. I often (incorrectly) refer to chromosomes as different contigs in the FASTA file. With long read technologies advancing pretty rapidly it breaks conventional definitions of a read and contig, where a single read can be longer than a contig. However assembling two long reads would still result in a contig sequence. So once again it is important to give more context when using this term.

1.4 Summary

- **Genomics** is all things genome that rotates around genomic sequence
- **Sequence** is an ordered collection of nucleotide bases
- **Read** is an output from a sequencing machine that covered biological sample in digital sequence
- **RNAseq** focuses on gene expression and splicing isoforms

³<https://www.thermofisher.com/au/en/home/brands/ion-torrent.html>

⁴<https://www.pacb.com/>

⁵<https://nanoporetech.com/>

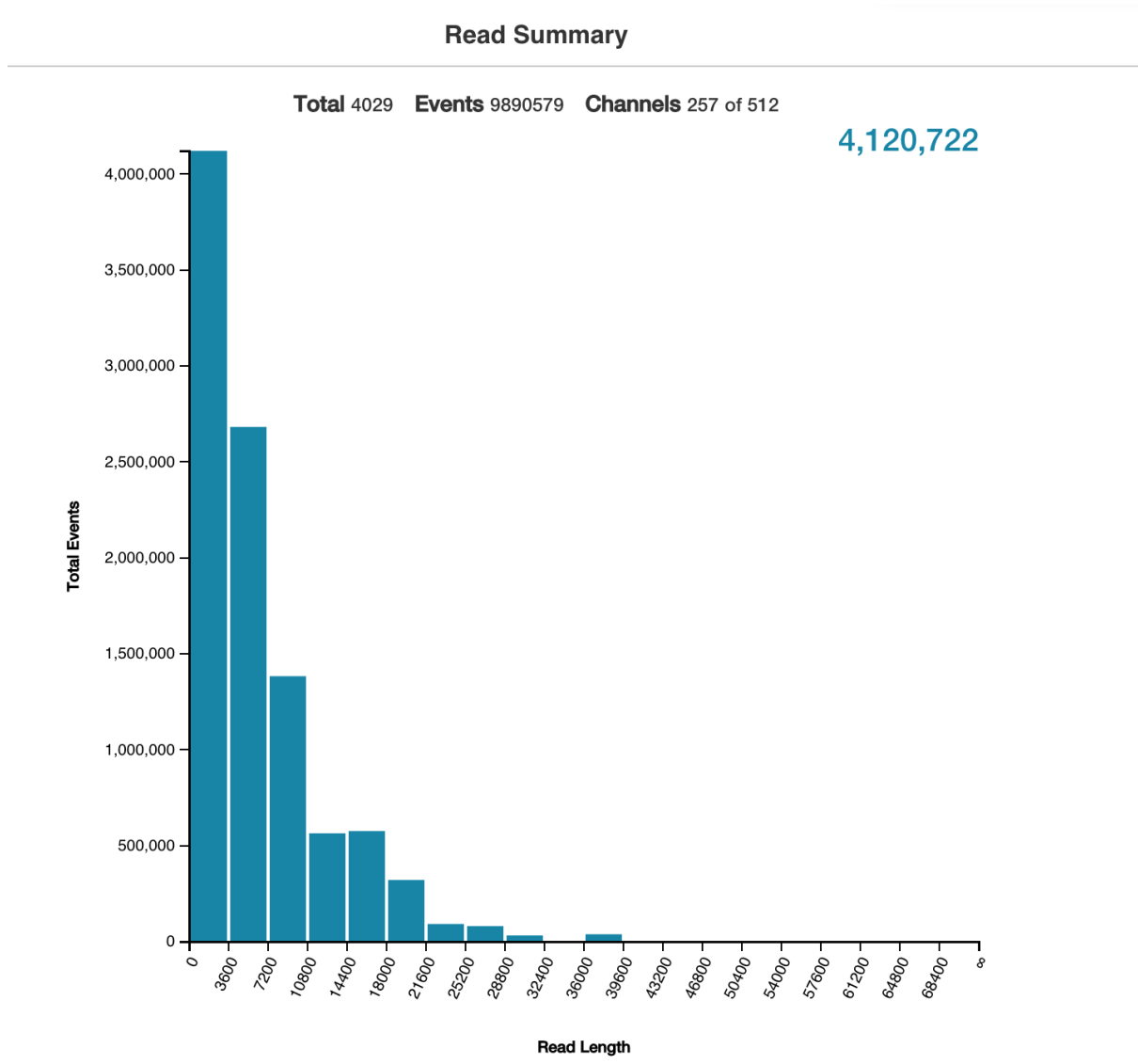


Figure 1.1: <https://bioinformatics.stackexchange.com/questions/3623/how-many-reads-has-my-sequencing-run-produced-on-minion>

Chapter 2

Reference sequence

In this course will be focusing on DNA/RNA molecules and hence our sequences will be nucleotides sequences. In the case of *Mus_musculus* (mouse) genome, we have 19 autosomal chromosomes and two sex chromosomes X and Y plus a mitochondrial chromosome (genome). This makes up 22 separate molecules and therefore 22 separate sequences. Typically that sequence is stored in **FASTA** file format

- FASTQ for raw short reads
- FAST5 for raw long reads
- FASTA for any other kind of sequence

The other possible bioinformatics files that we weren't focus on are, genbank and gff. Both of those files can hold annotation and sequence information in one file. It often works fine with small genomes, but large eukaryotic genomes are better split into two files reference sequence and annotation information.

2.0.1 FASTA

FASTA file typically can hold any type of sequence data. We are not going to talk about amino acid sequence in this course and so for your purpose we can think of FASTA as just holding nucleotides sequence. That sequence can be anything, primer or adapter sequence, individual transcripts (genes) sequence OR entire chromosome sequence. In the case when FASTA holds entire genome, that is all chromosomes for a given species, we tend to refer to that FASTA file as a reference genome file. If I'm asking biologist of a **reference genome**, I typically imply that I need a FASTA file with genomic sequence.

Be aware that there isn't such thing as "the genome" for a particular species. This is because any genome is a

There are a few different fasta files, but all of them will hold some sort of sequence. Sometime you'll see **.fna** to indicate nuclear sequence or **.faa** to indicate amino acid sequence, but more generally it'll be **.fa** or a longer version **.fasta**. Like any biological plain text file. This most certainly will be zipped so expect to see **.fa.gz** or **.fasta.gz**.

Note that our purpose we can think of FASTA file holding our reference genome sequence of interest e.g mouse genome.

This is previously assembled genome that was made publicly available. There are several different vendors that provide reference genomes. In general either one of these three will be a good place to start looking for your reference genome.

- Ensembl¹
- RefSeq²
- UCSC genomes³

¹<https://asia.ensembl.org/index.html>

²<https://www.ncbi.nlm.nih.gov/refseq/>

³<http://hgdownload.soe.ucsc.edu/downloads.html>

Those will have majority of species, however it is common to have a stand along community that host reference genomes from a different place. A good example is yeast genome database⁴. If you are working with yeast this is the place to look for reference genomes.

It is important to note that reference genome isn't a defined sequence. It is simply the best we have so far and it is a continuous effort to improve quality of the genomes. For widely used model organisms like mouse and yeast their genomes have been sequenced and assembled many times over the years and we have very good quality genomes. Species like axolotl or spiny mouse are less commonly used as model organism in comparison to mouse genomes and quality of those genomes is poorer. There are however other factors in place that influence genome quality e.g genome complexity.

2.1 Exercise

- go to Ensembl website⁵
- from the drop down on the left hand site select *Saccharomyces_cerevisiae*, which is a yeast species.
- click on download FASTA file. This will redirect you to ftp site
- find file **Saccharomyces_cerevisiae.R64-1-1.dna_sm.toplevel.fa.gz** and download it by clicking
- once downloaded double click on the file to open it and view the content.

depending on the operating system that you have you might have to unzip fasta file first before opening. It is best not to open with MS Office suite. Use text editor instead. On windows it'll be **notepad** on macOS it'll **textmate**.

At the very top you should see this

```
>I dna_sm:chromosome chromosome:R64-1-1:I:1:230218:1 REF
ccacaccacacccacacccacacaccacaccacacaccacacccacacacaca
caTCCTAACACTACCCTAACACAGCCCTAATCTAACCCCTGGCCAACCTGTCTCTCAACTT
ACCCCTCCATTACCCTGCCTCCACTCGTTACCCTGTCCATTCAACCATAACCACTCCGAAC
```

Note the very first line.

```
>I dna_sm:chromosome chromosome:R64-1-1:I:1:230218:1 REF
```

This is a FASTA header that indicates which sequence is follows. In this case it is I i.e sequence from chromosome one of yeast. If there are multiple chromosomes (usually the case for eukaryotic species) then you keep reading the sequence until you hit either the end of the file OR the next ">" sign

NOTE the actual sequence have been truncated

```
>I dna_sm:chromosome chromosome:R64-1-1:I:1:230218:1 REF
ccacaccacacccacacccacacaccacaccacacaccacacccacacacaca
caTCCTAACACTACCCTAACACAGCCCTAATCTAACCCCTGGCCAACCTGTCTCTCAACTT
ACCCCTCCATTACCCTGCCTCCACTCGTTACCCTGTCCATTCAACCATAACCACTCCGAAC
>II dna_sm:chromosome chromosome:R64-1-1:II:1:813184:1 REF
CCCACACACCACACCCACACCACACCCACACACCACACACCACACCCACACACCCACA
CCACACCACACCCACACCACACCCACACCCACACCCACACACCACACCCACACACACC
ACACCCACACACACCCACACCCACACACCACACCCACACACACACCACACCCACACACAC
CACACCACACCCACACCACACCCACACCCACACACCACACCCACACCCACACCCACACCC
```

The format of the FASTA header is rather loosely defined, but these are key points:

- must start with ">" sign
- the contig/chromosome name follows straight after ">" sign. Can be any string
- anything after a space is a comment/description

Below are all valid FASTA headers

```
>I
```

⁴<https://www.yeastgenome.org/>

⁵<https://asia.ensembl.org/index.html>

```
>chrI
>NC_000067.6
>I_named_it
```

Each header in a single FASTA file has to be unique in order to identify each sequence.

Are these two FASTA headers the same?

```
>I
>I_named_it
```

Are these two FASTA headers unique?

```
>I
>I_named_it
```

While this appears to be very small details, the reason it is important is because you want to know which chromosome (molecule) your reads came from.

2.2 Rule of thumb

Unless you are working with some exotic species and/or have a strong preference for genome vendor you can generally leave this part up to your bioinformatician to sort out. It is important to know that there are different vendors that have different versions (builds) of genomes.

. Often times bacterial RNAseq data come with some custom reference and annotation files. If that happens bioinformatician will need to convert everything to FASTA and GTF/GFF files.

Chapter 3

Gene annotation

As discussed in [reference genome] module. The reference genome is just a string of characters.

Chapter 4

File formats

4.1 Introduction

Bioinformatics file types don't differ from any other file type. Broadly any computer file is divided into either a plain text or binary files. The type doesn't matter; plain text files are "easy" to interact with because of their nature, whereas binary files require specialised tooling for interfacing with the file. Note that it is very common to zip plain text files, which simply means reducing the size of the file, which is better for storage purposes. Once zipped, the file is no longer plain text - it is binary now. It is a natural behaviour to unzip those files before looking inside. It is okay to do so, but be aware that this isn't the best practice, and you don't want to be storing your files in plain text, as this will eat up your disk space very quickly.

What is a file format?

It is a defined structure within the file. By far the most common structure of the plain text file is tabular, just like your conventional spreadsheet, where each column is separated by either a comma or a tab.

4.1.1 Comma Separated Variables (CSV)

Gene.ID	Chrom	Gene.Name	Biotype	K03_S3	K04_S4	WT1_S1	WT2_S2
ENSMUSG00000051951	1	Xkr4	protein_coding	0	0	0	0
ENSMUSG00000025900	1	Rp1	protein_coding	0	0	0	0
ENSMUSG000000109048	1	Rp1	protein_coding	0	0	0	0
ENSMUSG00000025902	1	Sox17	protein_coding	0	0	0	0
ENSMUSG00000033845	1	Mrpl15	protein_coding	934	1317	888	939
ENSMUSG00000025903	1	Lypla1	protein_coding	705	848	647	747
ENSMUSG000000104217	1	Gm37988	protein_coding	0	0	0	0
ENSMUSG00000033813	1	Tcea1	protein_coding	354	436	368	442
ENSMUSG00000002459	1	Rgs20	protein_coding	0	0	0	0

4.1.2 Tab Separated Variables (TSV)

Gene.ID	Chrom	Gene.Name	Biotype	K03_S3	K04_S4	WT1_S1	WT2_S2
ENSMUSG00000051951	1	Xkr4	protein_coding	0	0	0	0
ENSMUSG00000025900	1	Rp1	protein_coding	0	0	0	0
ENSMUSG000000109048	1	Rp1	protein_coding	0	0	0	0
ENSMUSG00000025902	1	Sox17	protein_coding	0	0	0	0
ENSMUSG00000033845	1	Mrpl15	protein_coding	934	1317	888	939
ENSMUSG00000025903	1	Lypla1	protein_coding	705	848	647	747
ENSMUSG000000104217	1	Gm37988	protein_coding	0	0	0	0
ENSMUSG00000033813	1	Tcea1	protein_coding	354	436	368	442

ENSMUSG00000002459,1,Rgs20,protein_coding,0,0,0,0

For the purpose of understanding RNAseq experiment the bare minimum you'll need to understand are these three file types:

- FASTQ
- FASTA
- GFF/GTF

4.1.2.1 FASTQ

This is a very important file(s) you don't mess directly with FASTQ files. Best not to even unzip those either. If think that you need to do something fastq files in terms of editing or some other out of ordinary manipulations. Make a copy and work on the copy. If you modify, corrupt or lose your FASTQ files, days, month or year of wet-lab work go down a toilet, let a lone large sum of money spend on RNA extraction, library preparation and sequencing.

4.1.2.2 FASTA

There are a few different fasta files, but all of them will hold some sort of sequence. Sometime you'll see `.fna` to indicate nuclear sequence or `.faa` to indicate amino acid sequence, but more generally it'll be `.fa` or a longer version `.fasta`. Like any biological plain text file. This most certainly will be zipped so expect to see `.fa.gz` or `.fasta.gz`.

For the purpose of RNAseq analysis know that FASTA file holds your reference sequence.

4.1.2.3 GTF/GFF

These are annotation files

There are many different biology / bioinformatics related file formats. You will naturally come across different files formats. It doesn't matter so much what they are called, i.e what they file extension is, rather what they hold.

Essentially all bioinformatic files are either plain text (a.k.a ASCII) or binary. Plain text files are "easy" to interface with. Unix/Linux command line provide many great tools that can work with plain text files. To list a few popular and powerful command line tools:

- `grep` search for a pattern
- `sed` search and replace
- `awk` search and replace and more
- `cat` look inside
- `cut` what the knife would do to bread

It is out of the scope of this course to go through any of the tools or command line in general, but if you going to do any kind of bioinformatics you'll eventually going to come across these tools.

The other file type is binary, this will requires designated tool for interfacing with the file for example SAM/BAM files have `samtools` that aid interaction with them.

4.2 Plain text files

Given that we know the structure of the file TSV or CSV we can use standard command line tools to get relevant/interesting bits e.g

```
~$ cut -f1,3,5-6 t.tsv
```

Gene.ID	Gene.Name	K03_S3	K04_S4
ENSMUSG000000051951	Xkr4	0	0

ENSMUSG00000025900	Rp1	0	0
ENSMUSG000000109048	Rp1	0	0
ENSMUSG00000025902	Sox17	0	0
ENSMUSG00000033845	Mrpl15	934	1317
ENSMUSG00000025903	Lypla1	705	848
ENSMUSG000000104217	Gm37988	0	0
ENSMUSG00000033813	Tcea1	354	436
ENSMUSG00000002459	Rgs20	0	0

4.3 Glossary

- interface: extract or insert relevant information

Chapter 5

All about RNAseq

5.1 General

- Functional annotation of the mouse (FANTOM)
- Encyclopedia of DNA elements (ENCODE)
- Single loci can lead to six different mRNA isoforms, on average, instead of single transcript

5.2 Technology

What is it for?

- what is expressed (exon locations /strand) ?
- how much is expressed (hits per transcripts) ?

old way to do DE - microarray

- do need as much starting material for RNAseq

Noise:

- biological
- poisson noise from sampling (have level of uncertainty at low level, low counts)
- technical
- Effect size (do the genes go up by a lot or not - high signal)
- amount of variation

get better depth using poly-A enriched, because rRNA depletion will leave lots of other RNA around therefore diluting mRNA by some percent

mean fragment size ~ 300 bp

measure of uncertainty of the base call

5.3 Differential expression (DE)

- library size is total number of mapped reads. Library size might vary for individual samples.
- Total number of reads per gene proportional to:
 - gene expression level (to get cpm (count per million) divide each read count by library size multiplied by million). This enables you to compare different library sized samples
 - transcript length

- sequencing depth of the library # Ideas
- have a url with multiqc report ready for browsing
- explain reference file (FASTA) and annotation files (GTF)

Chapter 6

Lesson plan

Perhaps it'll be good idea to introduce several different plain text files

- `txt`
- `csv/tsv`
- `gff/gff3/gtf`
- `fastq/fq`
- `fasta/fa/faa/fna`

DO NOT unzip your fastq file and store them as plain text ! DO NOT mess with your raw fastq files!

Chapter 7

Meeting notes

This is overall them of the day

How to instruct them to be good RNAseq user?

have a url with multiqc page ready?

This is

explain reference file (FASTA) and annotation files (GTF)

This if you can set up ftp link so that users can pont and click through directory structure to interact with the sikRun output.

Mentione salmon

Good to mention in parsing - salmon if you are using salmon you not going to see DNA contamination. so something to be aware.

Also mentione that this is for model organism that are well annotation/assembled.

Use Chiara's data to study chromosome 21. From multiqc report chromosome 21 has a lot more reads. It looks like ribosomal contamination even though library was polyA pooled.

Take all reads from chr21 and do feature counts and look where reads go. Suspecting that reads go to ribosomal genes. If so this would be great case study for the workshop.

<https://www.ncbi.nlm.nih.gov/pubmed/29788454>

genotify

Reproducible research using the same reference files

TCGA for example is using a single reference for the whole cohort

If you gonna do second study e.g chipSeq you gotta have it on the same reference as your RNAseq

there is no such thing as "the reference genome" there are different builds, it is work in progress