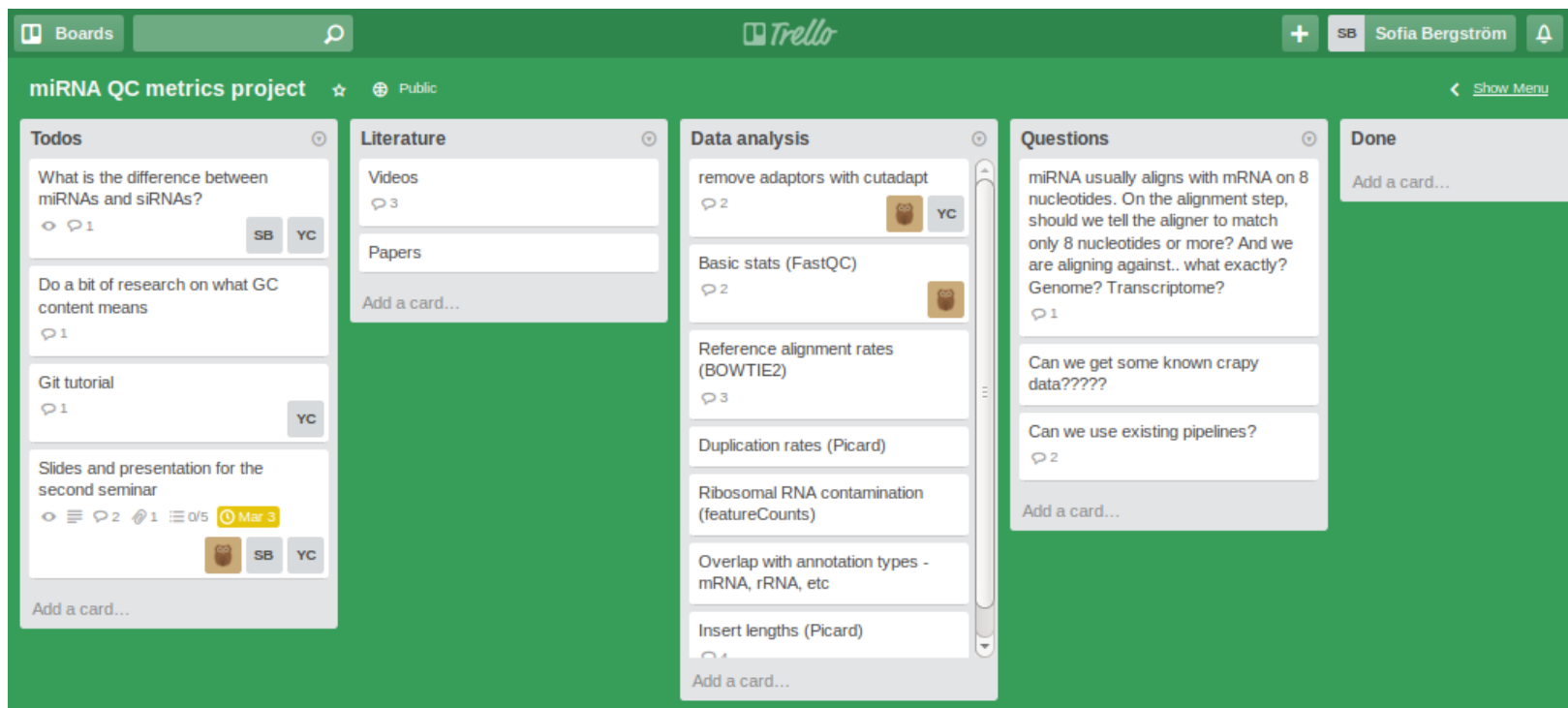

miRNA QC metrics

Guillermo Carrasco
Sofia Bergström
Yim Wing Chow

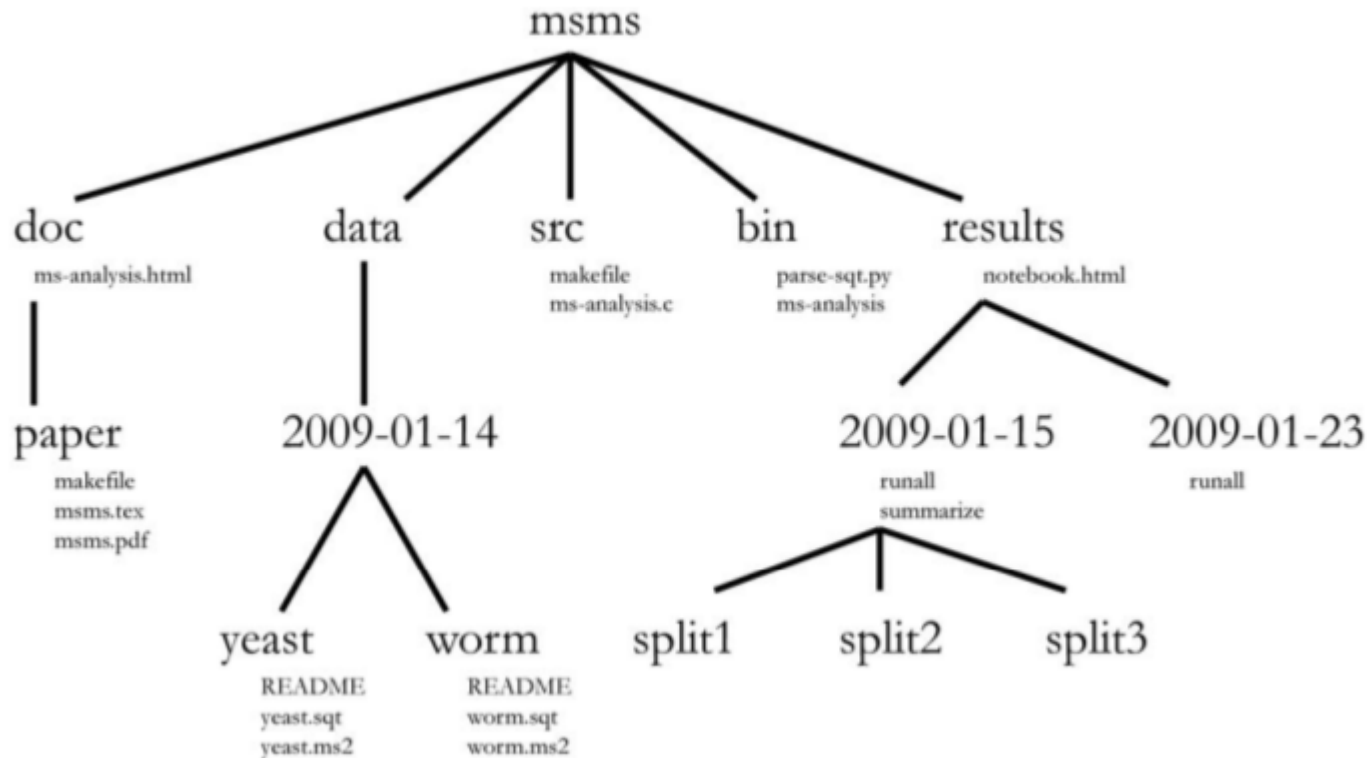
Project structure

Trello



Project structure

Github



Pipeline

https://github.com/SciLifeLab/miRNA_processing

- Remove adapters (**Cutadapt**)
 - Basic stats (**FastQC**)
 - Read alignment (**Bowtie**)
 - Duplication rates (**Picard**)
 - rRNA contamination (**featureCounts**)
 - Overlap with annotation types; mRNA, rRNA
 - Insert and alignment length (**Picard**)
-













Pipeline

To be done:

- Automated search against miRBase
 - Make it more easy to test different parameters
 - Options in configuration file?
-

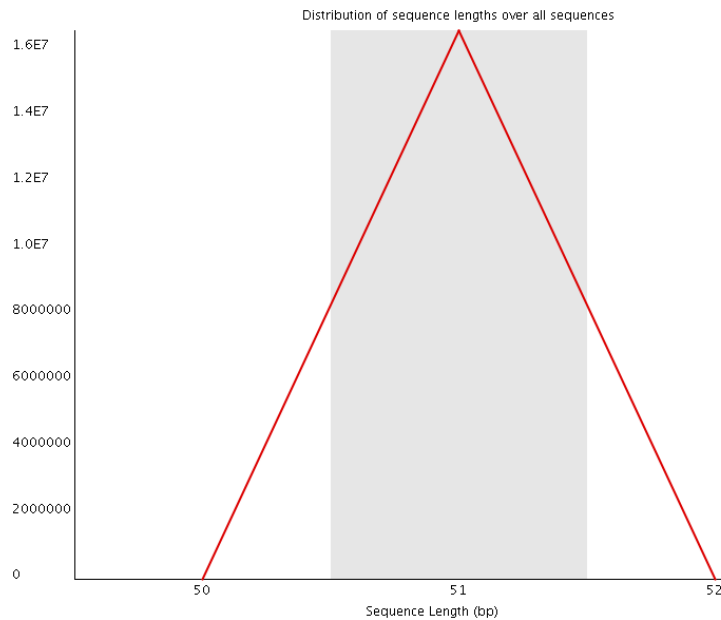
FastQC

- FastQC is a quality control tool for NGS data

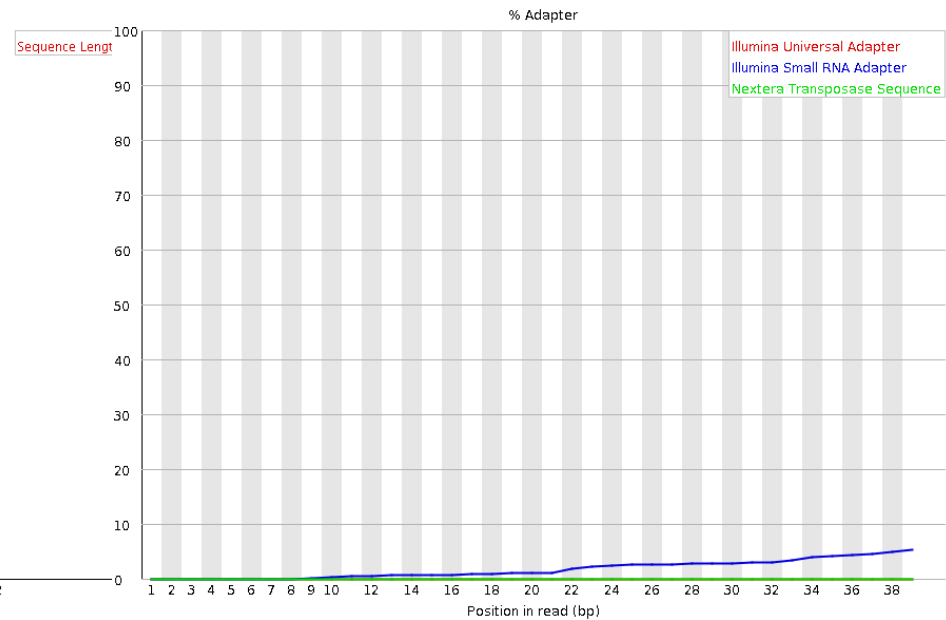
-  [Basic Statistics](#)
-  [Per base sequence quality](#)
-  [Per tile sequence quality](#)
-  [Per sequence quality scores](#)
-  [Per base sequence content](#)
-  [Per sequence GC content](#)
-  [Per base N content](#)
-  [Sequence Length Distribution](#)
-  [Sequence Duplication Levels](#)
-  [Overrepresented sequences](#)
-  [Adapter Content](#)
-  [Kmer Content](#)

FastQC

First run on one sample. No adapter trimming



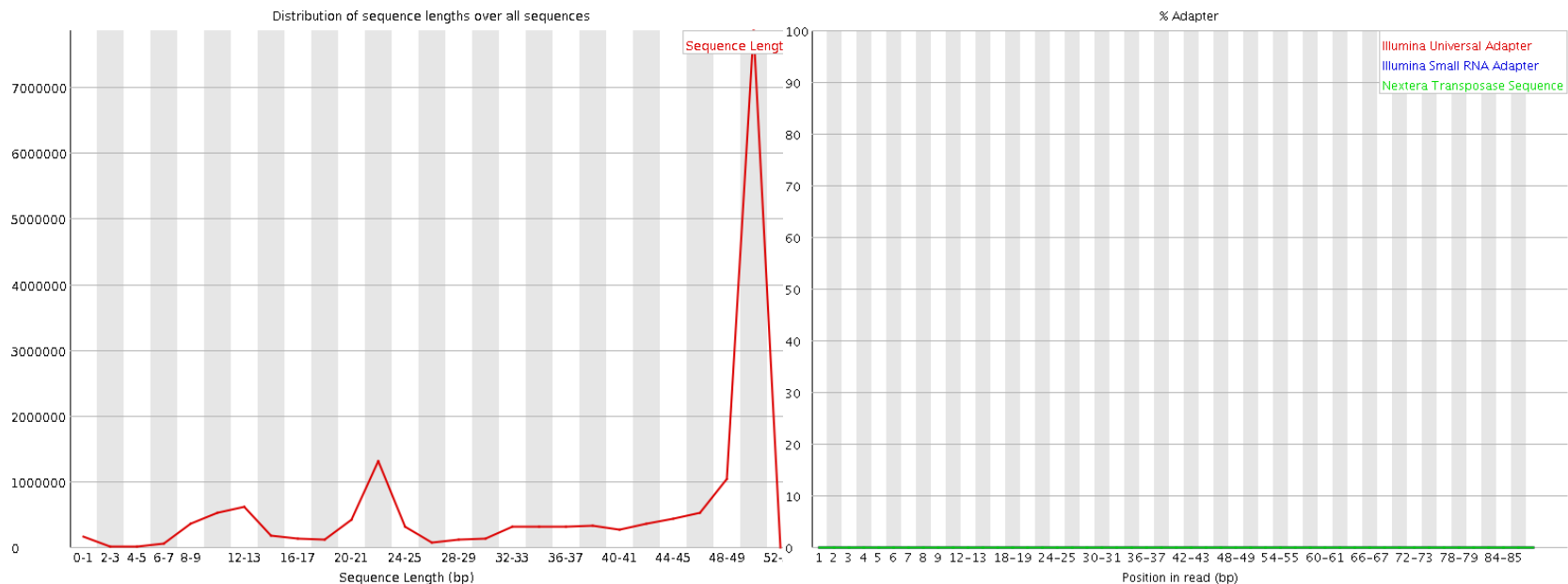
Length distribution



Adapter content

FastQC

First run on one sample. With adapter trimming



Length distribution

Adapter content

Project plan

- Week 10:
 - Small pipeline fixes and improvements
 - Continue testing/understanding tools
 - Try to run the analysis in all the samples
 - Week 11:
 - Study and compare results
 - Try to find reasonable thresholds for different QC metrics
 - Poster preparation
 - Week 12:
 - Conclusions
 - Finish the poster and presentation
-

Thank you!

Questions?

Guillermo's diary

Yim's diary

Sofia's diary
