



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

General Statistics

Copy table

Configure columns

Scatter plot

Violin plot

Showing 18/18 rows and 6/9 columns.

Export as CSV

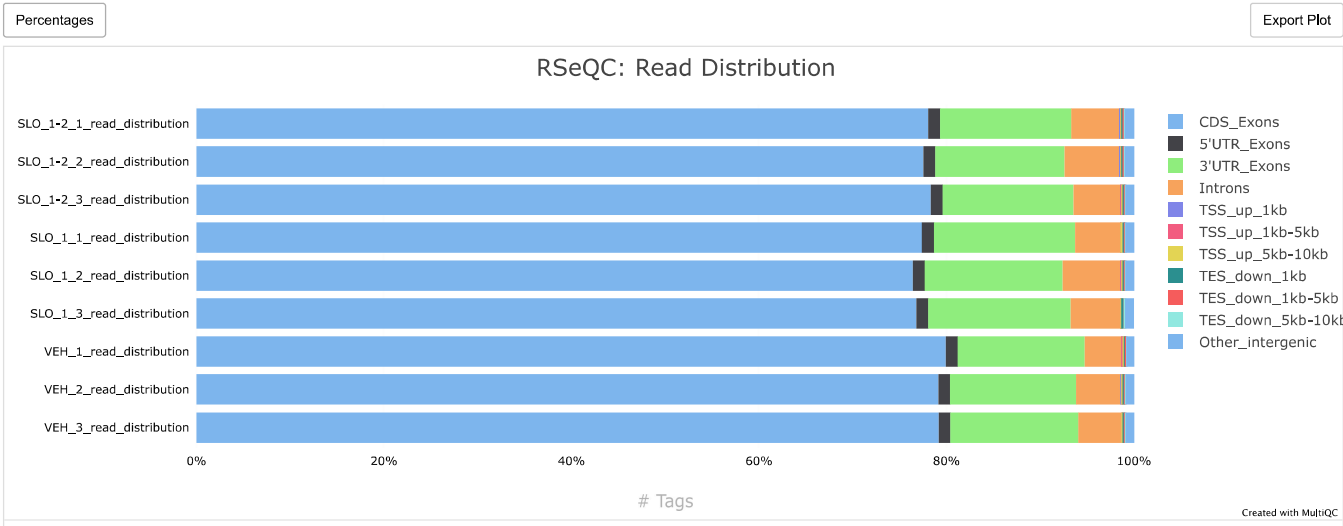
Sample Name	% Aligned	M Aligned	% Dropped	% Dups	% GC	M Seqs
SLO_1-2_1	91.1 %	19.8 M				
SLO_1-2_1_S35_R1_001			0.0 %	52.1 %	46 %	21.8 M
SLO_1-2_2	91.1 %	19.8 M				
SLO_1-2_2_S36_R1_001			0.0 %	51.7 %	46 %	21.8 M
SLO_1-2_3	91.2 %	18.6 M				
SLO_1-2_3_S37_R1_001			0.0 %	51.9 %	46 %	20.4 M
SLO_1_1	91.7 %	18.2 M				
SLO_1_1_S38_R1_001			0.0 %	50.3 %	46 %	19.8 M
SLO_1_2	91.9 %	18.8 M				
SLO_1_2_S39_R1_001			0.0 %	49.3 %	46 %	20.4 M
SLO_1_3	91.9 %	16.7 M				
SLO_1_3_S40_R1_001			0.0 %	48.9 %	46 %	18.2 M
VEH_1	90.7 %	18.0 M				
VEH_1_S32_R1_001			0.0 %	53.0 %	46 %	19.8 M
VEH_2	90.7 %	17.7 M				
VEH_2_S33_R1_001			0.0 %	51.9 %	46 %	19.5 M
VEH_3	90.5 %	22.0 M				
VEH_3_S34_R1_001			0.0 %	55.1 %	46 %	24.3 M

RSeQC

RSeQC package provides a number of useful modules that can comprehensively evaluate high throughput RNA-seq data. DOI: 10.1093/bioinformatics/bts356.

Read Distribution

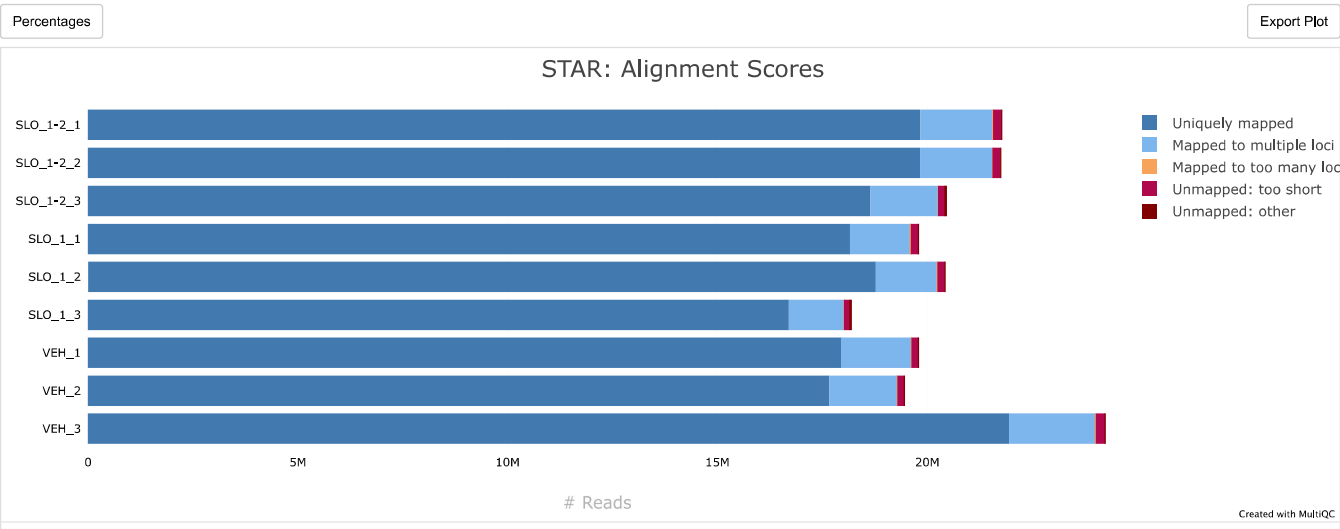
Read Distribution calculates how mapped reads are distributed over genome features.



# STAR

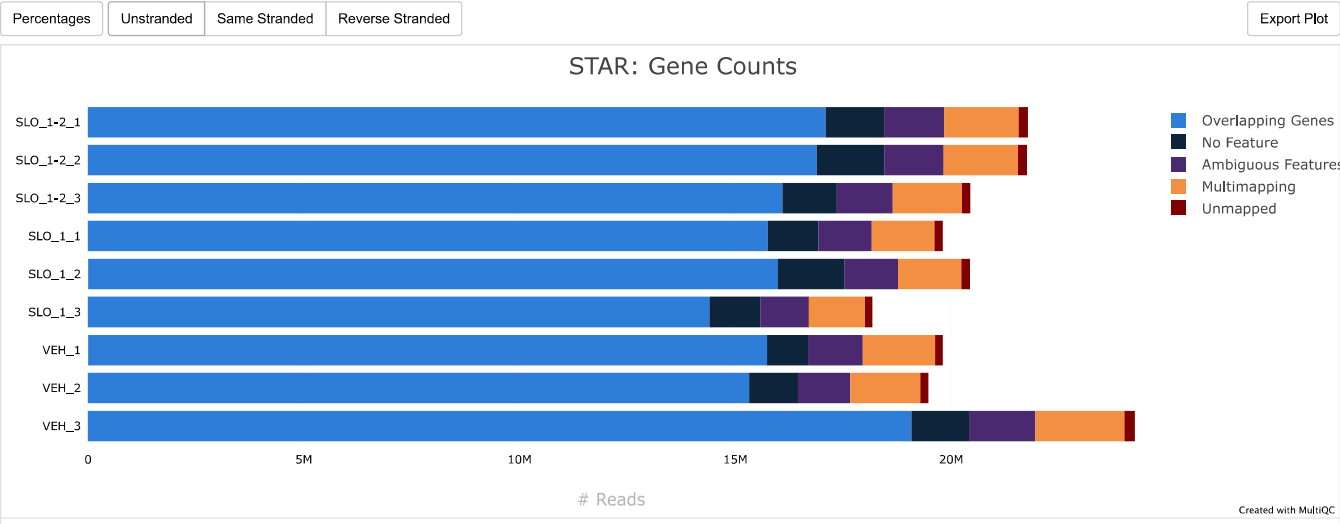
STAR is an ultrafast universal RNA-seq aligner. DOI: 10.1093/bioinformatics/bts635.

## Alignment Scores



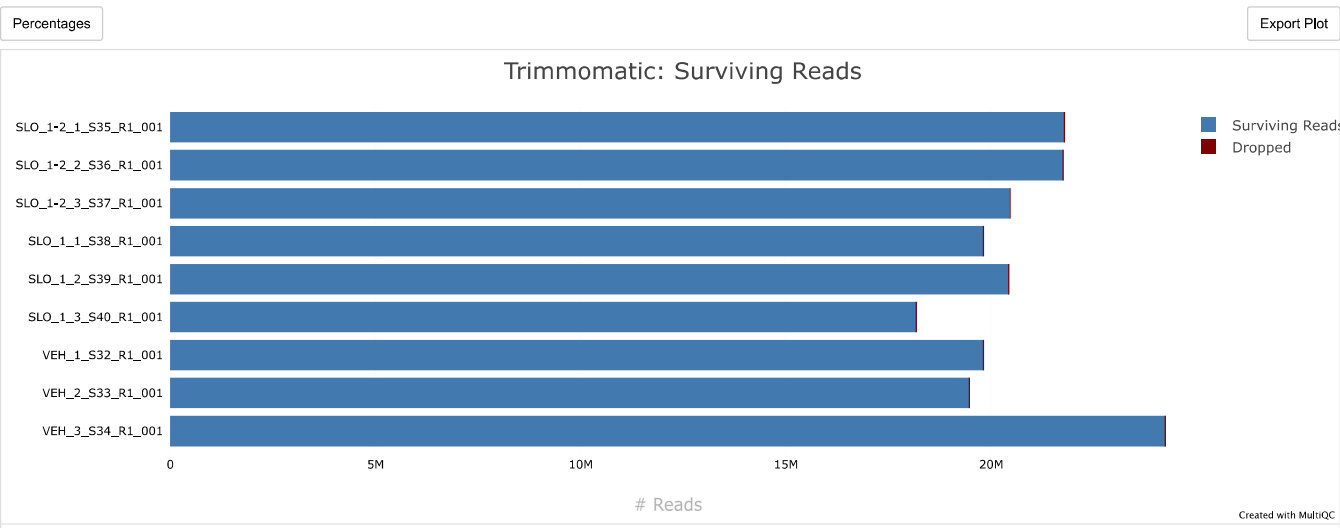
## Gene Counts

Statistics from results generated using --quantMode GeneCounts . The three tabs show counts for unstranded RNA-seq, counts for the 1st read strand aligned with RNA and counts for the 2nd read strand aligned with RNA.



# Trimmomatic

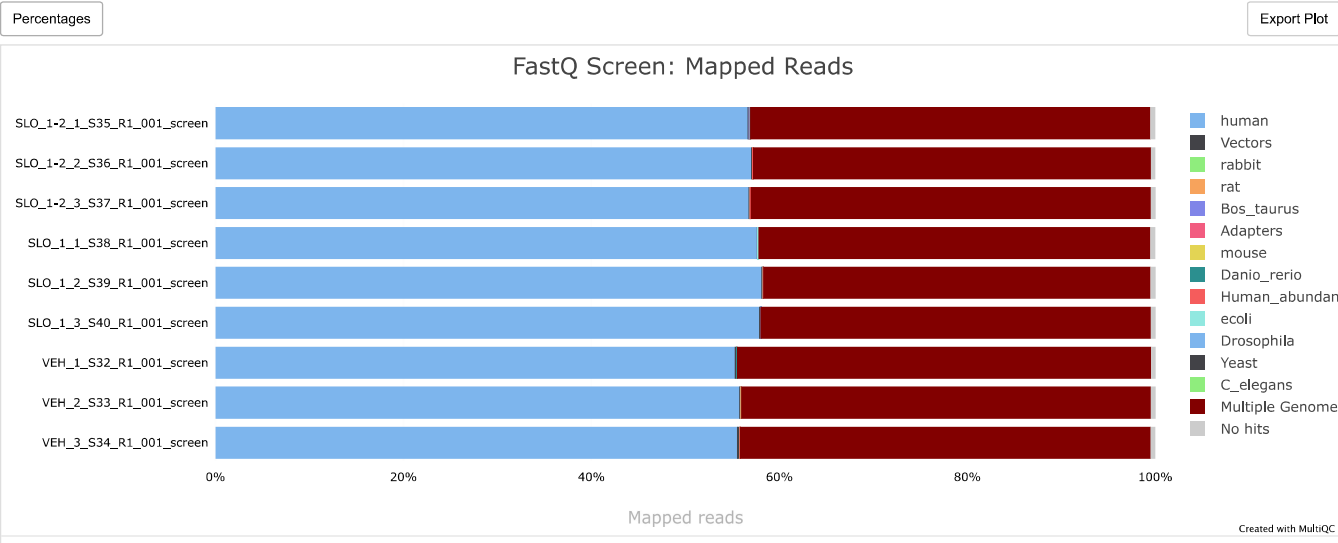
Trimmomatic is a flexible read trimming tool for Illumina NGS data. DOI: 10.1093/bioinformatics/btu170.



# FastQ Screen Version: 0.14.0

FastQ Screen allows you to screen a library of sequences in FastQ format against a set of sequence databases so you can see if the composition of the library matches with what you expect. DOI: 10.12688/f1000research.15931.2.

## Mapped Reads

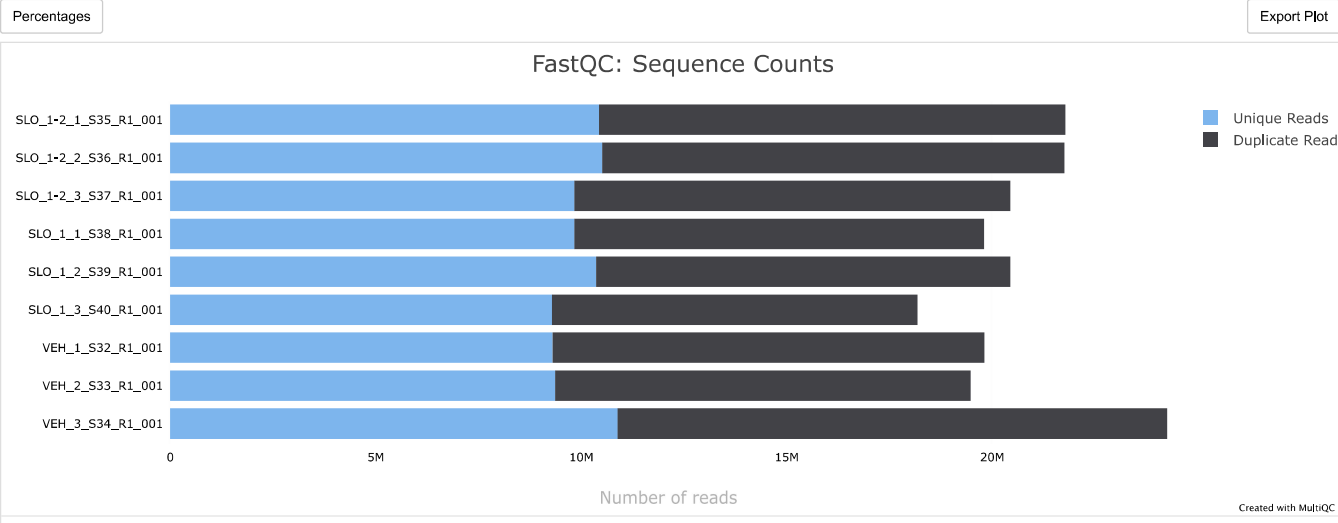


# FastQC Version: 0.11.9

FastQC is a quality control tool for high throughput sequence data, written by Simon Andrews at the Babraham Institute in Cambridge.

## Sequence Counts

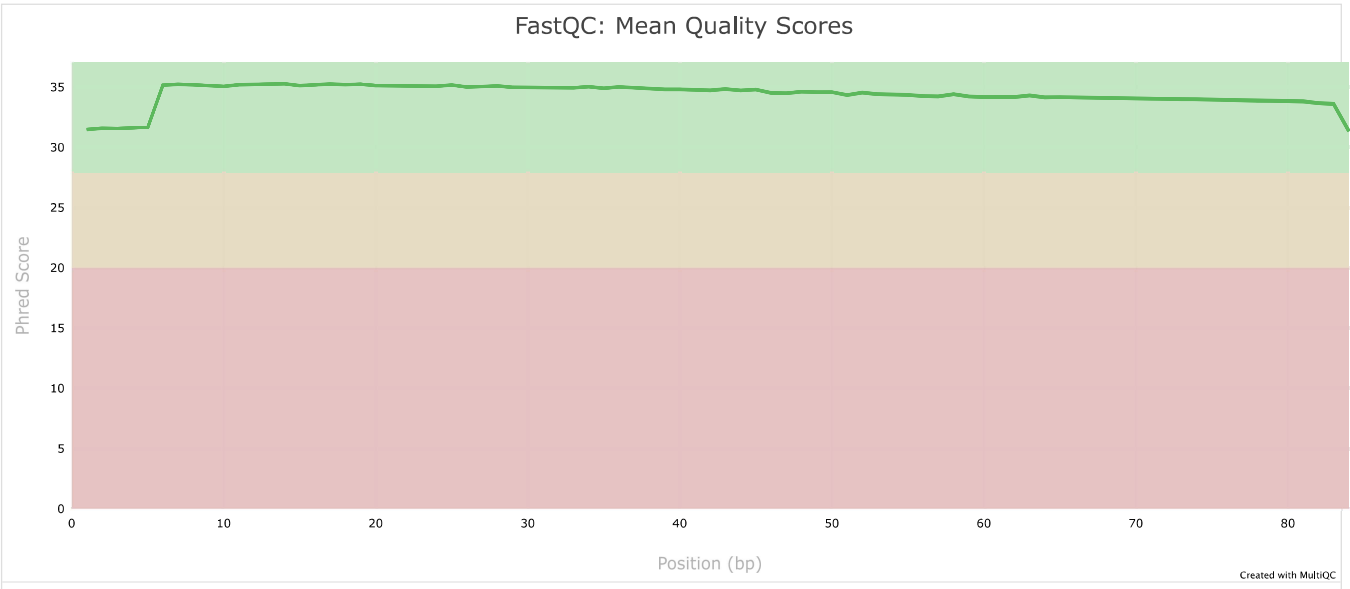
Sequence counts for each sample. Duplicate read counts are an estimate only.



Sequence Quality Histograms 9

The mean quality value across each base position in the read.

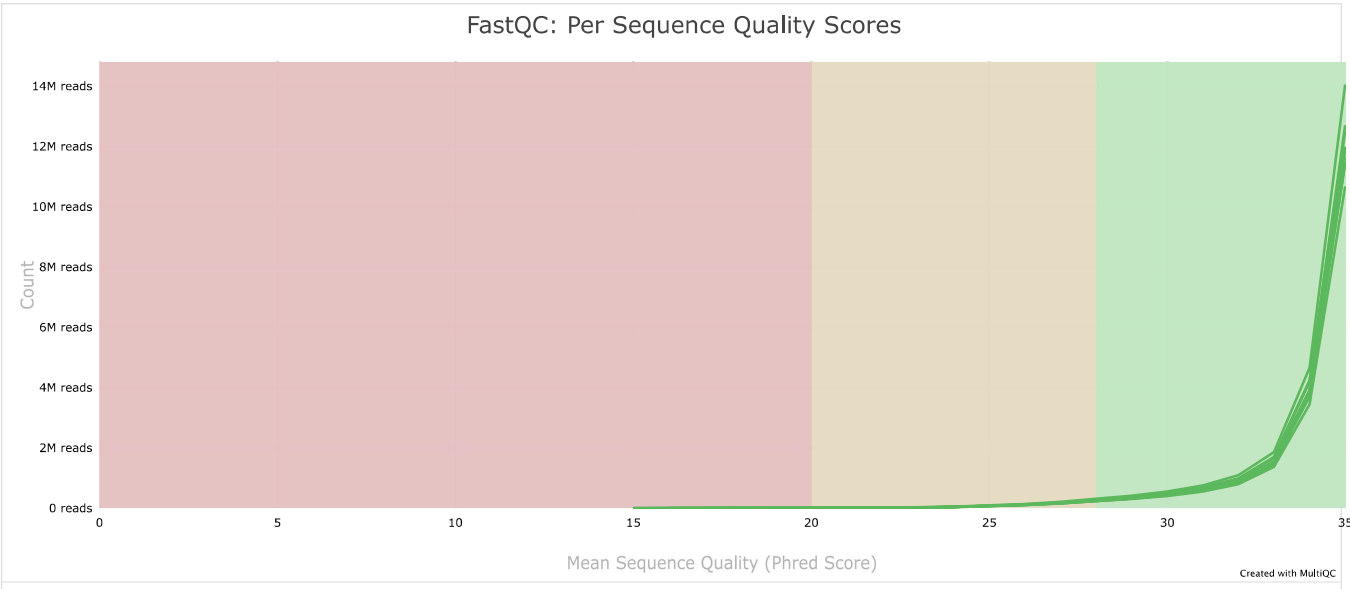
Export Plot



Per Sequence Quality Scores 9

The number of reads with average quality scores. Shows if a subset of reads has poor quality.

Export Plot



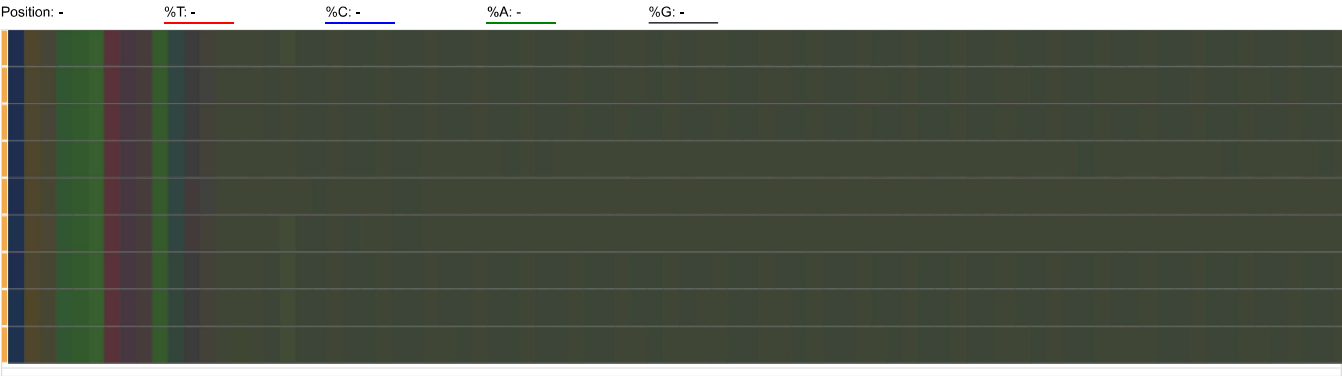
Per Base Sequence Content

9

The proportion of each base position for which each of the four normal DNA bases has been called.

Click a sample row to see a line plot for that dataset.

Rollover for sample name



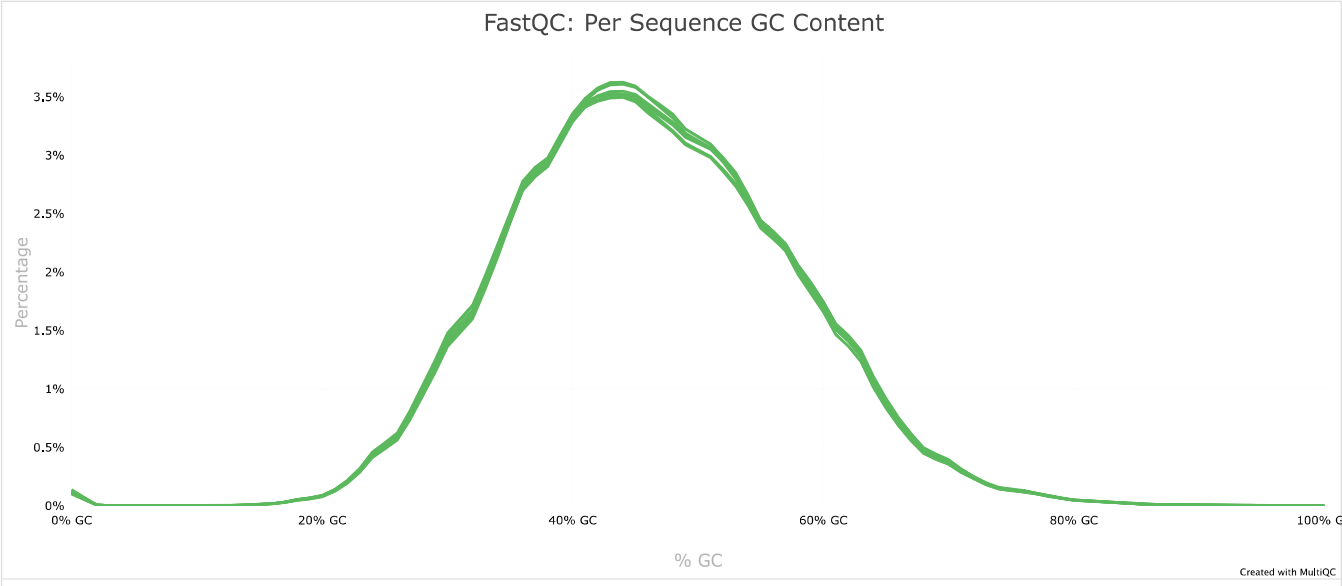
Per Sequence GC Content

9

The average GC content of reads. Normal random library typically have a roughly normal distribution of GC content.

Percentages   Counts

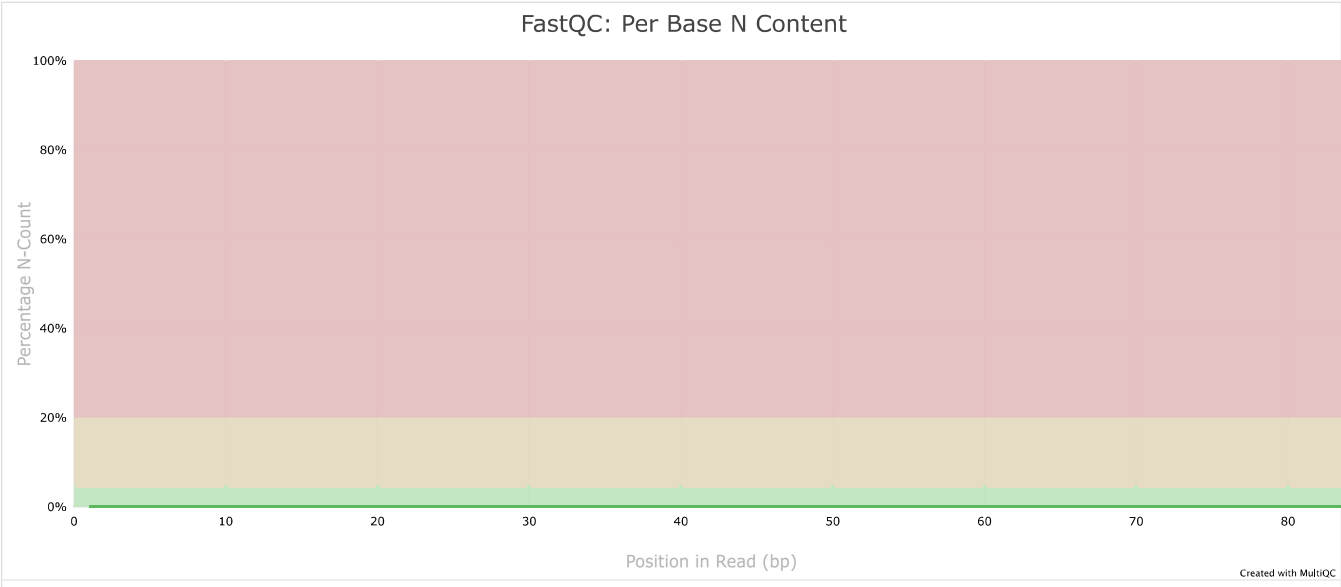
Export Plot



Per Base N Content 9

The percentage of base calls at each position for which an N was called.

Export Plot



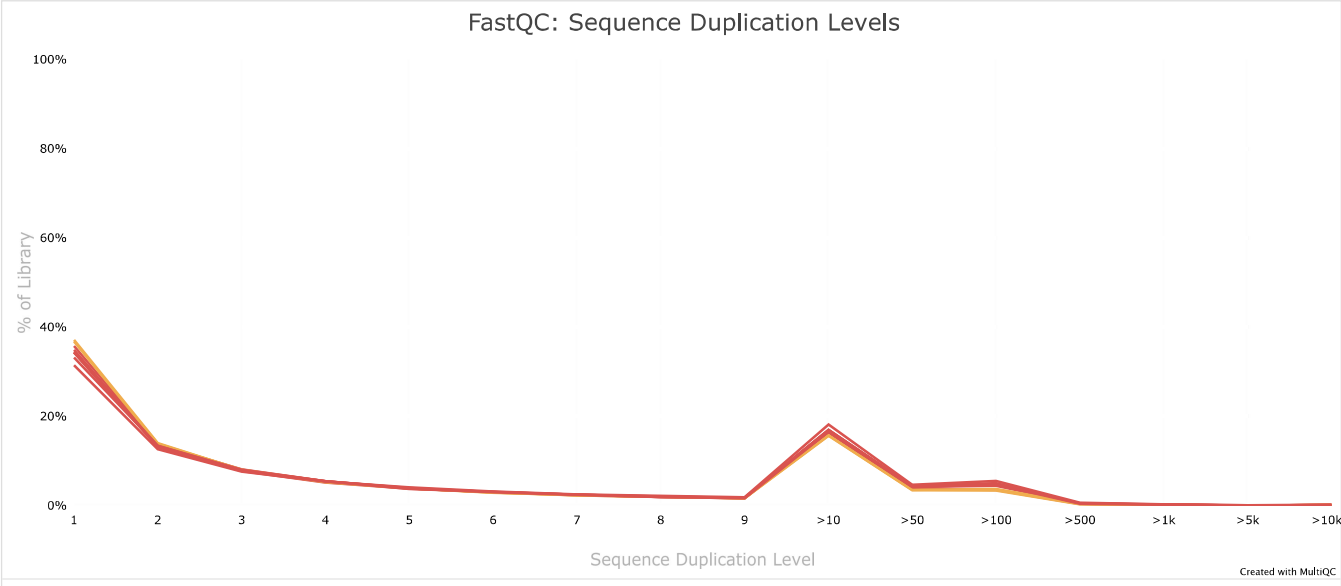
Sequence Length Distribution 9

All samples have sequences of a single length (84bp).

Sequence Duplication Levels 2

The relative level of duplication found for every sequence.

Export Plot



Overrepresented sequences by sample 9

The total amount of overrepresented sequences found in each library.

9 samples had less than 1% of reads made up of overrepresented sequences

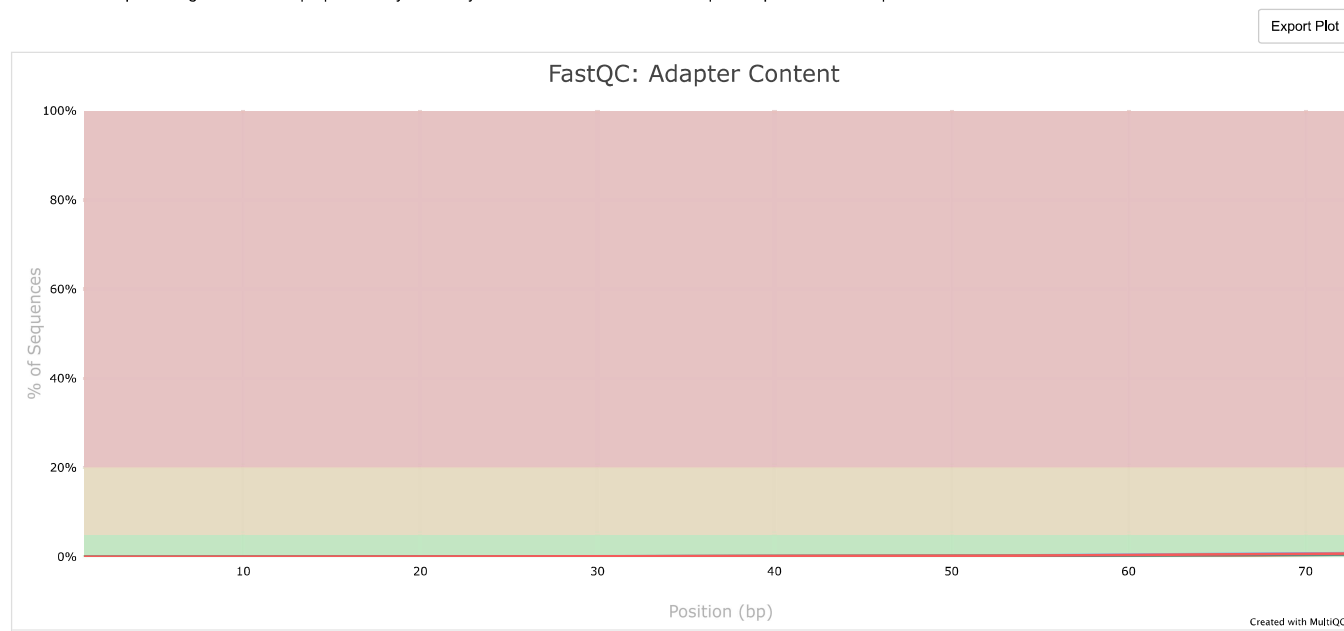
Top overrepresented sequences across all samples. The table shows 20 most overrepresented sequences across all samples, ranked by the number of samples they occur in.

Top overrepresented sequences across all samples. The table shows 20 most overrepresented sequences across all samples, ranked by the number of samples they occur in.

Overrepresented sequence	Samples	Occurrences	% of all reads
TTTTTTTTTTTTTTTTTTTTTTTTTTT	9	226 467	0.1218 %

## 9

The cumulative percentage count of the proportion of your library which has seen each of the adapter sequences at each position.



Status for each FastQC section showing whether results seem entirely normal (green), slightly abnormal (orange) or very unusual (red).

	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
SLO_1-2_1_S35_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
SLO_1-2_2_S36_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
SLO_1-2_3_S37_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
SLO_1_1_S38_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
SLO_1_2_S39_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
SLO_1_3_S40_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
VEH_1_S32_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
VEH_2_S33_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass
VEH_3_S34_R1_001	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Pass	Warning	Warning	Pass

# Software Versions

Software Versions lists versions of software tools extracted from file contents.

 Copy table

Software	Version
FastQ Screen	0.14.0
FastQC	0.11.9

**MultiQC v1.21** - Written by [Phil Ewels](#), available on [GitHub](#).  
This report uses [HighCharts](#), [jQuery](#), [jQuery UI](#), [Bootstrap](#), [FileSaver.js](#) and [clipboard.js](#).

