

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

Report generated on 2019-10-23, 13:24 based on data in:
/data/storage/SAYRES/NASONIA/Heini/Processing/Sayres/RNA/raw_FastQC

Welcome! Not sure where to start?

Watch a tutorial video

don't show again

(6:06)

General Statistics

Copy table

Configure Columns

Plot

Showing 24/24 rows and 3/5 columns.

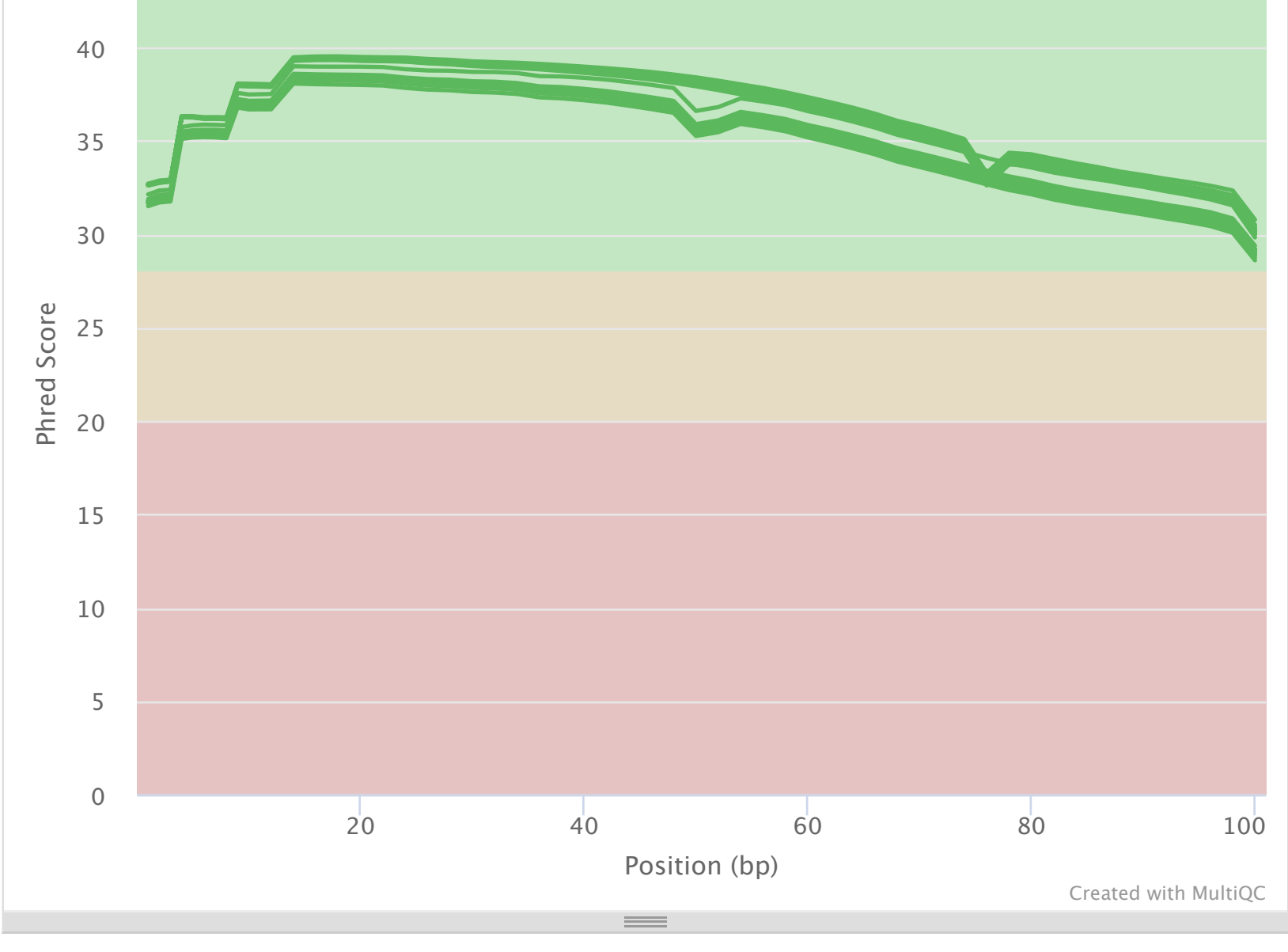
Sample Name	% Dups	% GC	M Seqs
014444_A2_ATCACG_run384_L002_R1_001	66.6%	40%	16.6
014444_A2_ATCACG_run384_L002_R2_001	58.9%	41%	16.6
014445_A4_CGATGT_run384_L002_R1_001	66.6%	40%	16.1
014445_A4_CGATGT_run384_L002_R2_001	59.0%	41%	16.1
014446_A5_TTAGGC_run384_L002_R1_001	61.9%	41%	12.7
014446_A5_TTAGGC_run384_L002_R2_001	55.0%	42%	12.7
014447_B2_TGACCA_run384_L002_R1_001	65.2%	40%	17.9
014447_B2_TGACCA_run384_L002_R2_001	59.2%	41%	17.9
014448_B4_ACAGTG_run384_L002_R1_001	66.2%	40%	15.6
014448_B4_ACAGTG_run384_L002_R2_001	59.8%	41%	15.6
014449_B6_GCCAAT_run384_L002_R1_001	69.4%	40%	21.5
014449_B6_GCCAAT_run384_L002_R2_001	63.4%	41%	21.5

FastQC

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

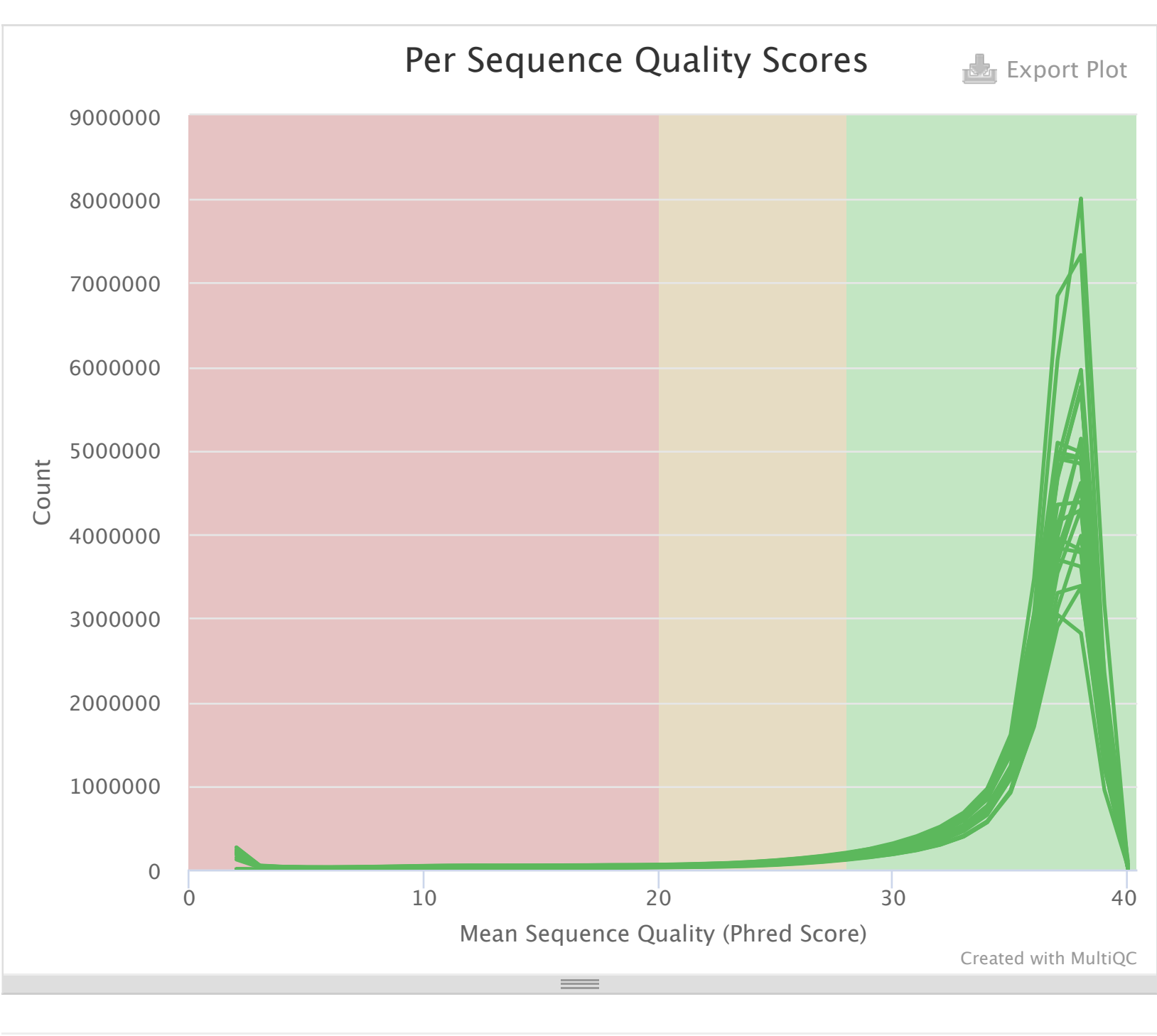
Sequence Quality Histograms

The mean quality value across each base position in the read. See the [FastQC help](#)-Limits: ☒ on



Per Sequence Quality Scores

The number of reads with average quality scores. Shows if a subset of reads has poor quality. See the [FastQC help](#). Y-Limits: ☒ on



Per Base Sequence Content

The proportion of each base position for which each of the four normal DNA bases has been called. See the [FastQC help](#).

