Rfastp Report

Summary

0.21.0 (https://github.com/OpenGene/fastp)
single end (50 cycles)
50bp
49bp
32.619322% (may be overestimated since this is SE data)
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGGAATCTAATGTC

Before filtering 23.379546 M total reads: total bases: 1.168977 G 1.150622 G (98.429834%) Q20 bases: 1.081053 G (92.478555%) Q30 bases: 46.123001% GC content:

After filtering

23.325007 M total reads: total bases: 1.166046 G Q20 bases: 1.147887 G (98.442700%) Q30 bases: 1.078491 G (92.491327%)

46.155371% GC content: Filtering result 23.325007 M (99.766723%) reads passed filters: reads with low quality: 6.241000 K (0.026694%)

0 (0.000000%)

reads with too many N: reads too short:

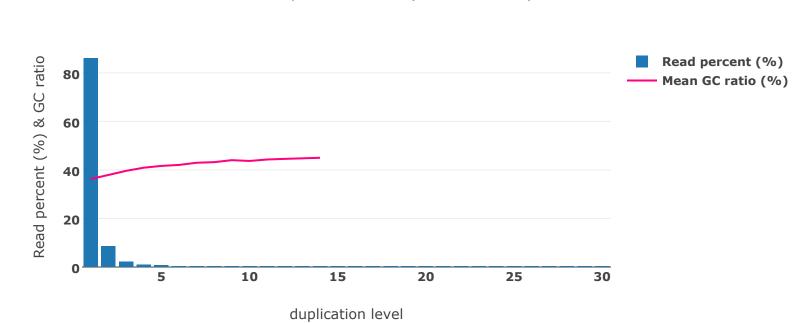
48.298000 K (0.206582%) Adapters

Adapter or bad ligation of read1

The input has little adapter percentage (~0.230334%), probably it's trimmed before.		
CGAAA	14851	
CGAAAG	4240	
AGAAAGCA	1403	
GGAAAGCA	1316	
CGAAAGCAT	802	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAG	6738	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGG	10086	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGGA	14176	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGGAATCTAA	5543	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGGAATCTAAT	4722	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGGAATCTAATG	1606	
CGAAAGCAGGTCAATTATATTCAATATGGAAAGAATAAAAGAACTAAGGAATCTAATGT	2085	
other adapter sequences	11906	

Duplication

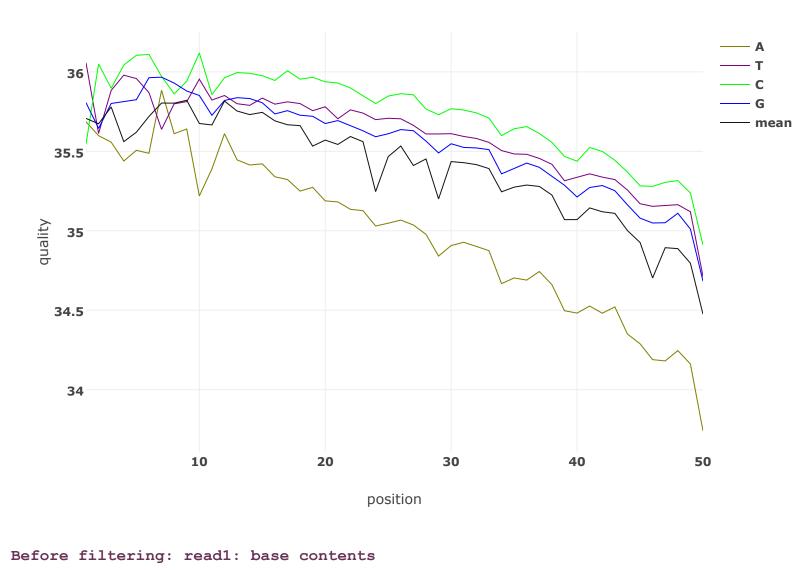
duplication rate (32.619322%)



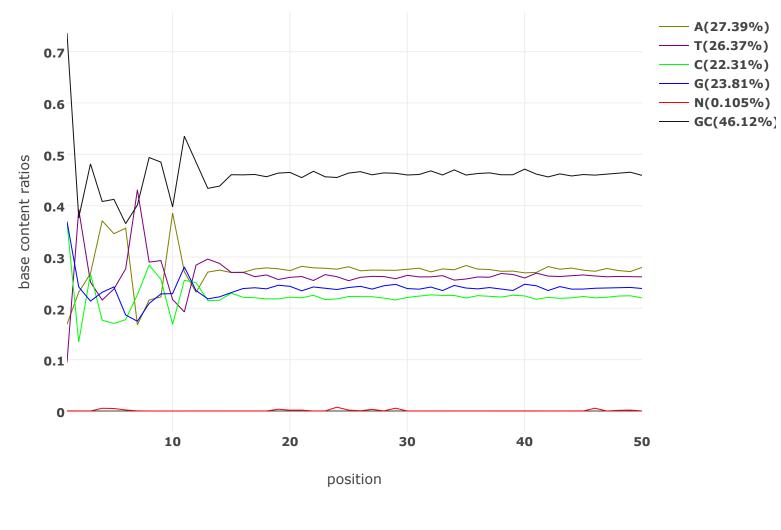
Before filtering

Before filtering: read1: quality

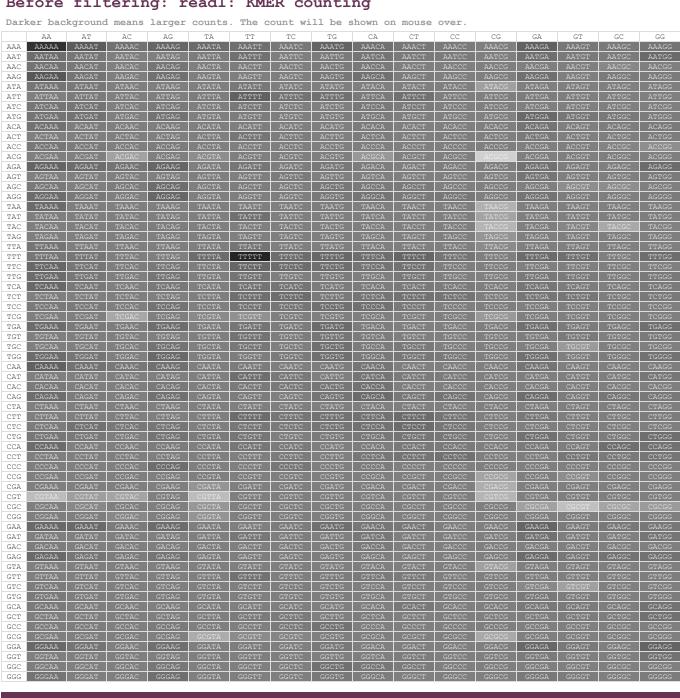
Value of each position will be shown on mouse over.



Value of each position will be shown on mouse over

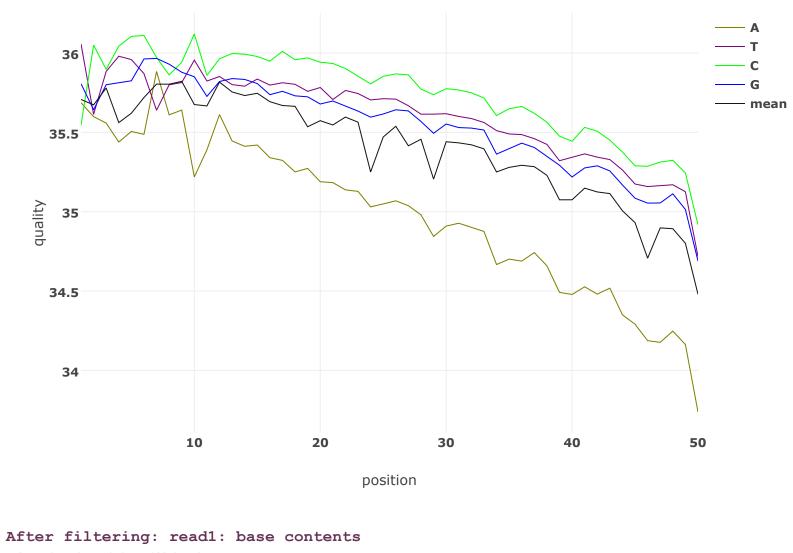


Before filtering: read1: KMER counting



After filtering

After filtering: read1: quality Value of each position will be shown on mouse over



Value of each position will be shown on mouse over.

