

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2022/04/19 00:40:04

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR053629.sorted.bam -c -nw 400 -hm 3
```

1.2. Alignment

Command line:	/home/luna/Desktop/Software/bwa/bwa mem -t 16 -k 30 -R @RG\tID:Singlecell2022\tLB:Library\tPL:ILLUMINA\tSM:sample_SRR053629 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR053629.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Apr 19 00:40:04 CST 2022
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR053629.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	11,997,232
Mapped reads	9,003,753 / 75.05%
Unmapped reads	2,993,479 / 24.95%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	259 / 0%
Read min/max/mean length	30 / 48 / 48
Duplicated reads (estimated)	2,548,307 / 21.24%
Duplication rate	19.18%
Clipped reads	769,752 / 6.42%

2.2. ACGT Content

Number/percentage of A's	125,967,407 / 29.62%
Number/percentage of C's	81,672,135 / 19.21%
Number/percentage of T's	127,636,734 / 30.02%
Number/percentage of G's	89,734,038 / 21.1%
Number/percentage of N's	198,619 / 0.05%
GC Percentage	40.31%

2.3. Coverage

Mean	0.1374

Standard Deviation	1.1197
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2.4. Mapping Quality

Mean Mapping Quality	45.57
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2.5. Mismatches and indels

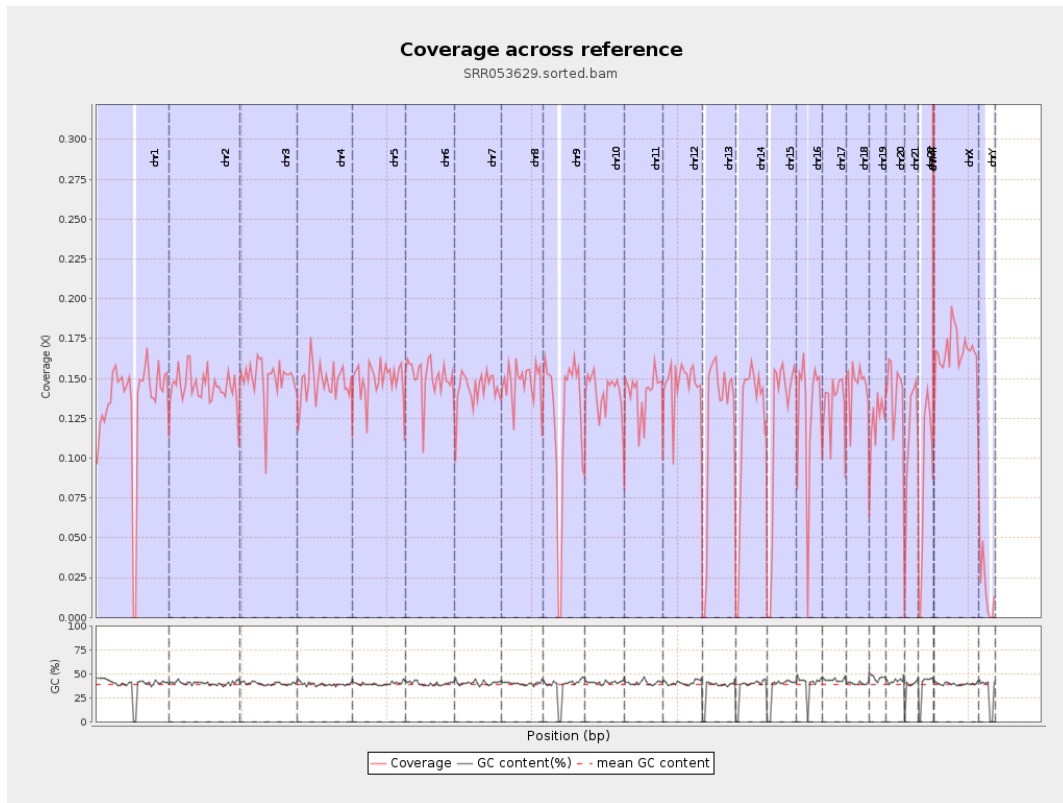
General error rate	0.6%
Mismatches	2,535,712
Insertions	18,075
Mapped reads with at least one insertion	0.2%
Deletions	54,918
Mapped reads with at least one deletion	0.61%
Homopolymer indels	47.31%

2.6. Chromosome stats

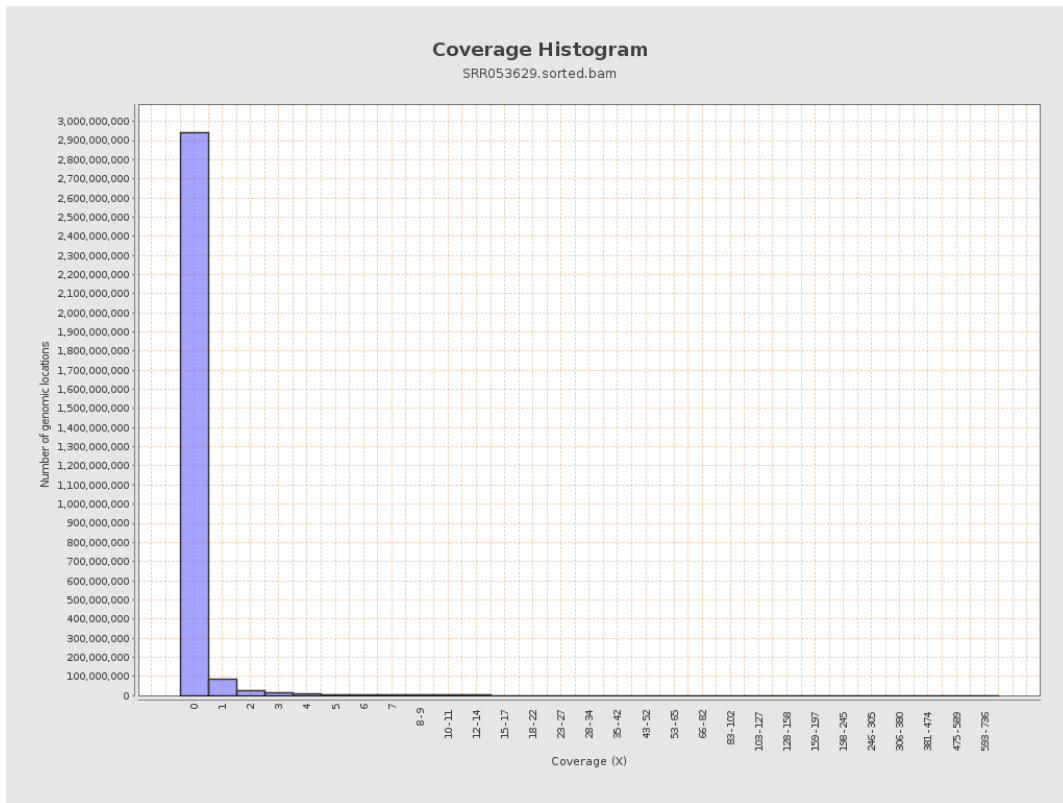
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	33296567	0.1336	1.1515
chr2	243199373	35327877	0.1453	1.2825
chr3	198022430	29642997	0.1497	1.1189
chr4	191154276	28096489	0.147	1.1516
chr5	180915260	26991141	0.1492	1.1001
chr6	171115067	25712921	0.1503	1.1848
chr7	159138663	22930192	0.1441	1.1571

chr8	146364022	21590941	0.1475	1.1628
chr9	141213431	17712089	0.1254	1.0292
chr10	135534747	19186585	0.1416	1.1612
chr11	135006516	18923766	0.1402	1.1493
chr12	133851895	19612734	0.1465	1.0855
chr13	115169878	14122005	0.1226	0.9992
chr14	107349540	12951322	0.1206	1.0485
chr15	102531392	12341897	0.1204	0.9745
chr16	90354753	11583756	0.1282	1.0361
chr17	81195210	10876095	0.1339	1.0206
chr18	78077248	11433578	0.1464	1.1975
chr19	59128983	7329434	0.124	1.1489
chr20	63025520	8860479	0.1406	1.0758
chr21	48129895	5385673	0.1119	1.023
chr22	51304566	4469114	0.0871	0.8004
chrMT	16571	52235	3.1522	6.9106
chrX	155270560	25789624	0.1661	1.2467
chrY	59373566	1069836	0.018	0.3844

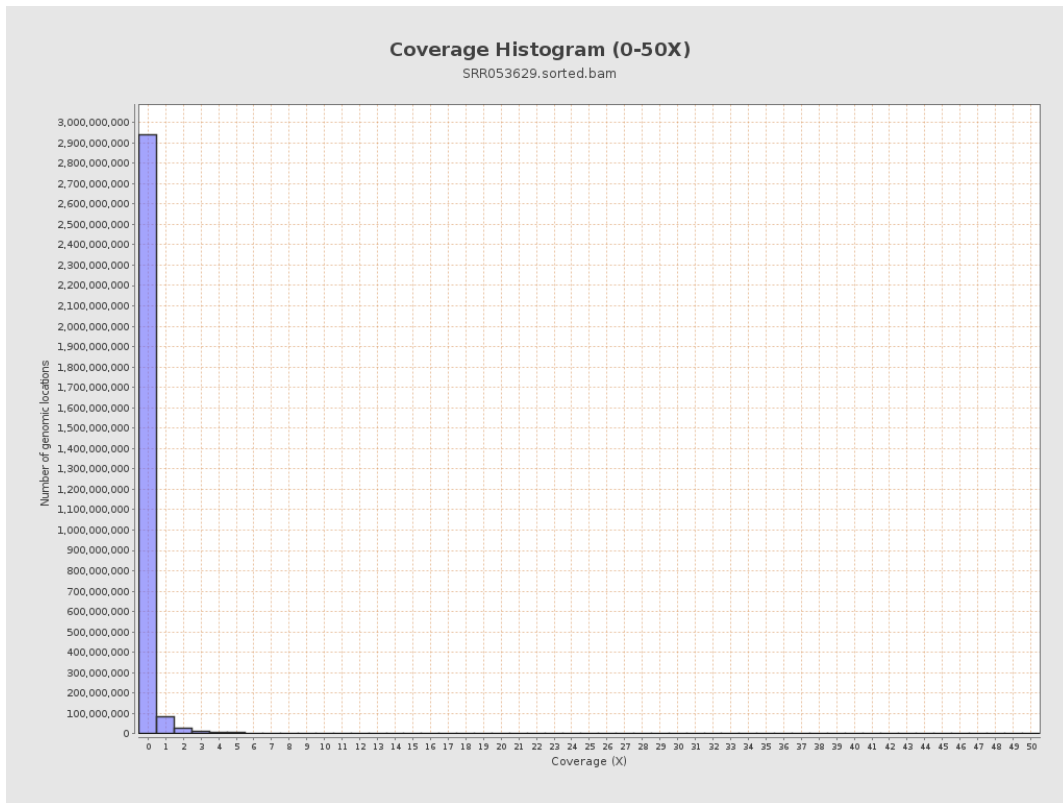
3. Results : Coverage across reference



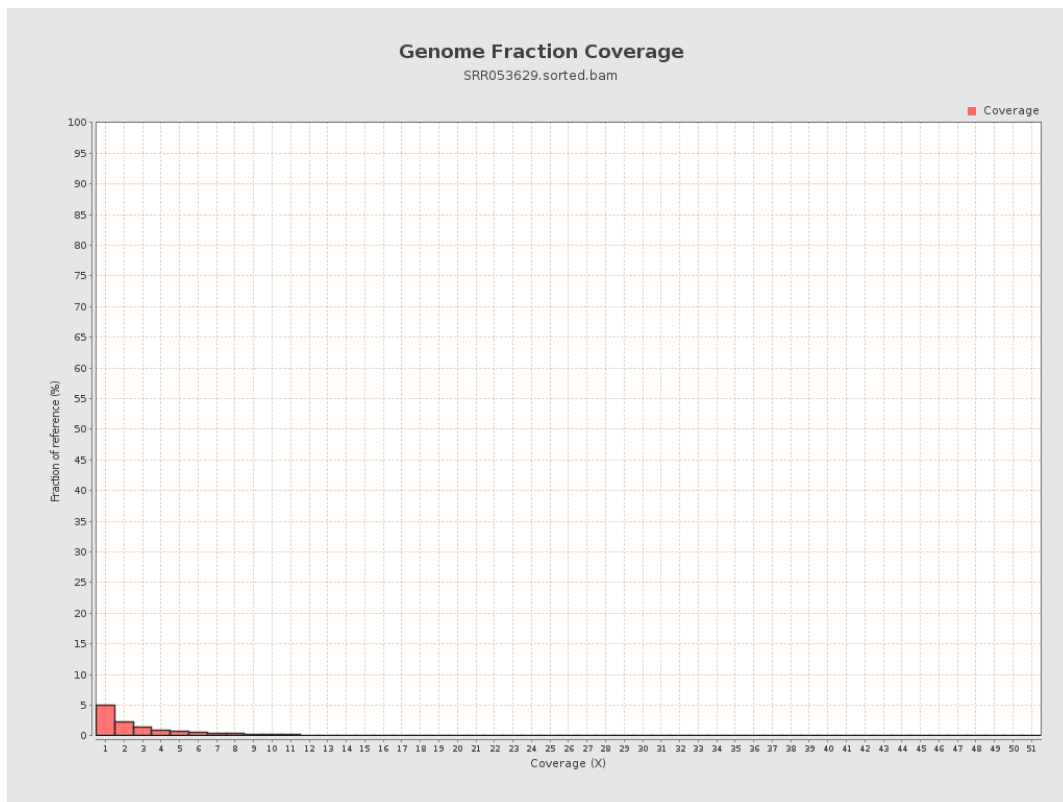
4. Results : Coverage Histogram



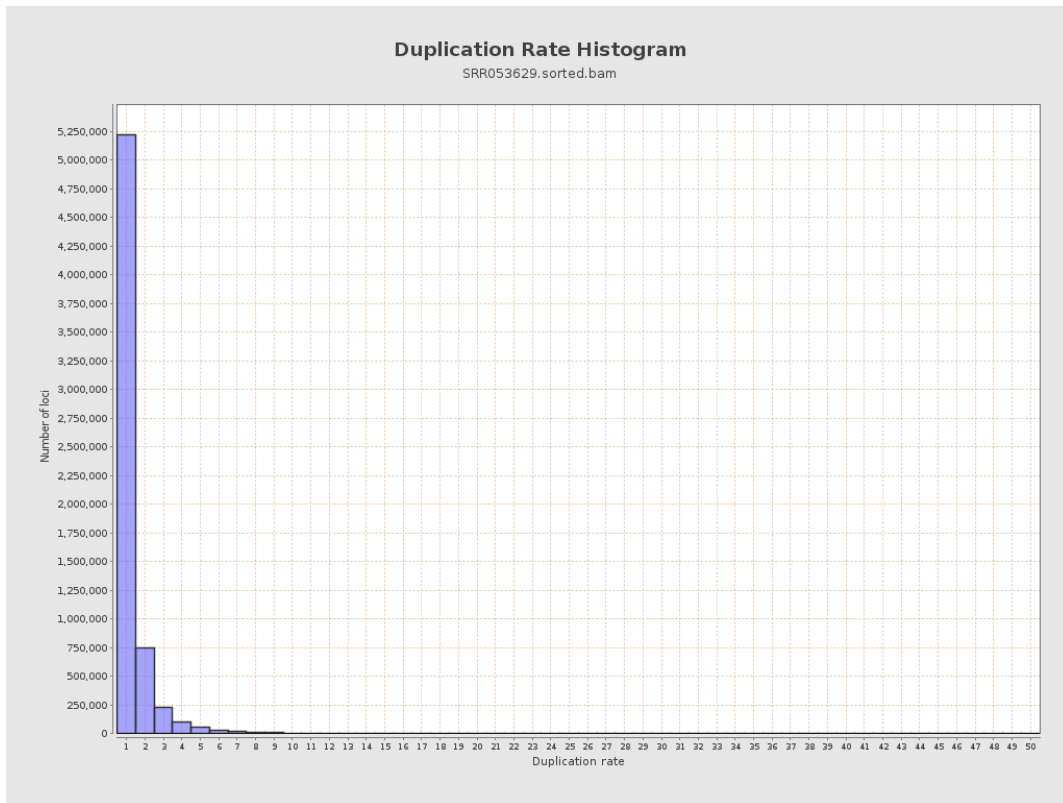
5. Results : Coverage Histogram (0-50X)



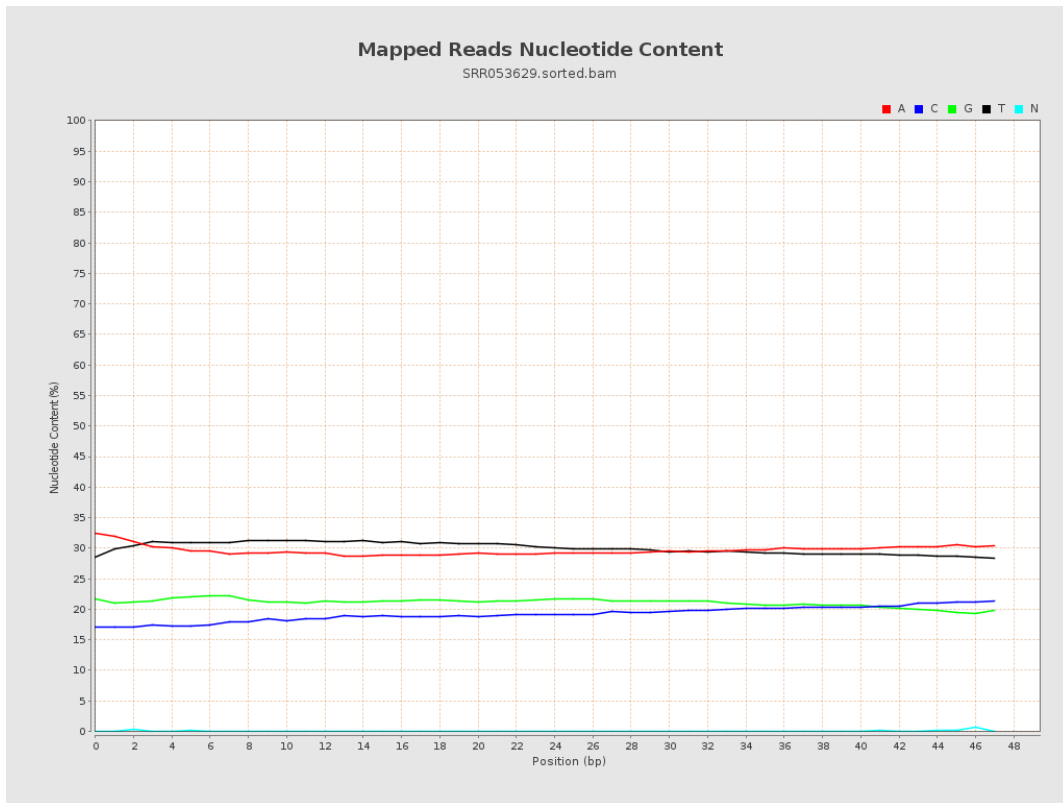
6. Results : Genome Fraction Coverage



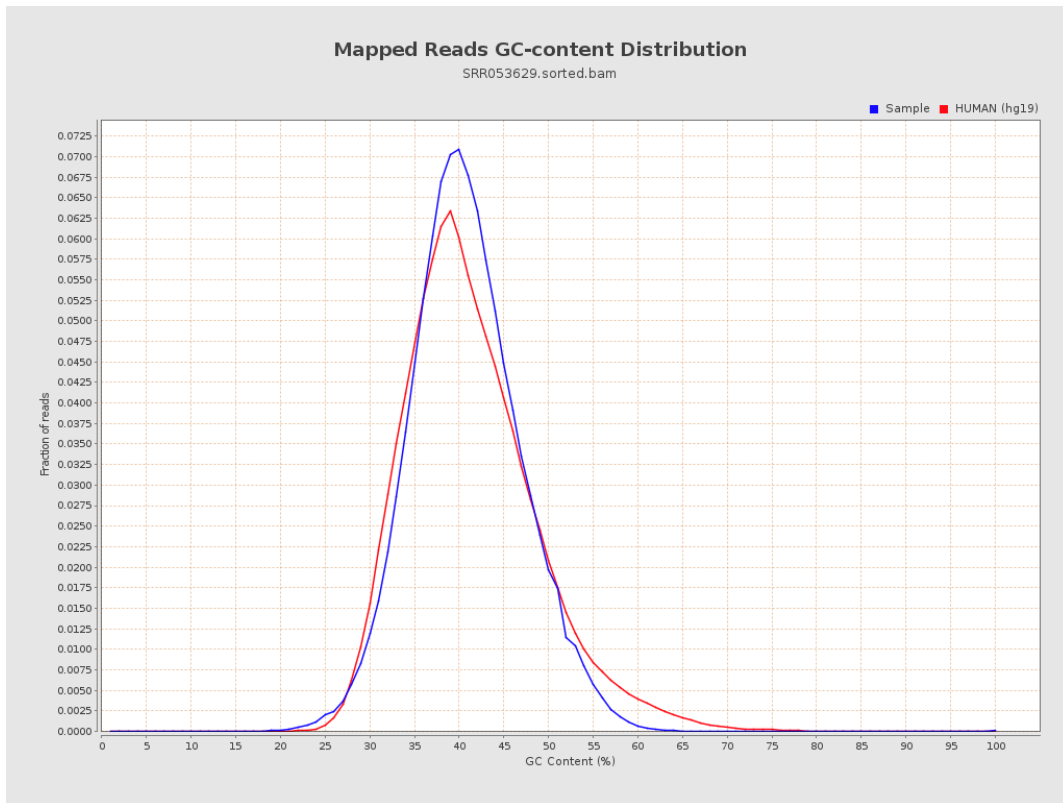
7. Results : Duplication Rate Histogram



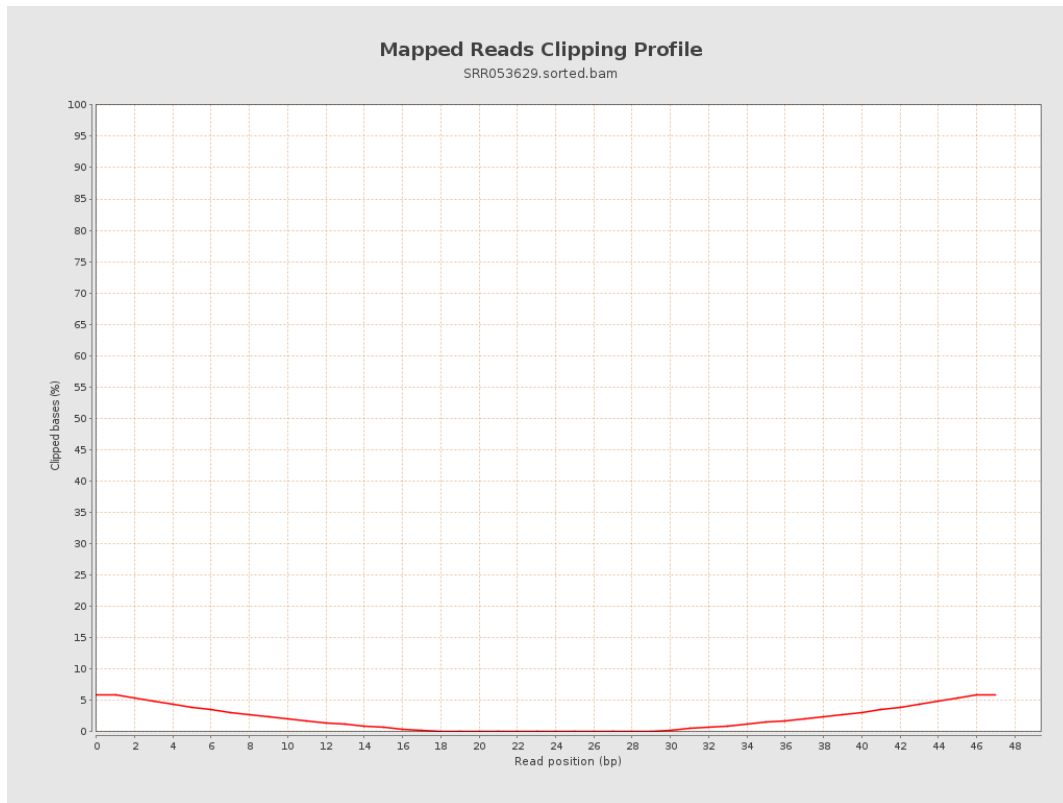
8. Results : Mapped Reads Nucleotide Content



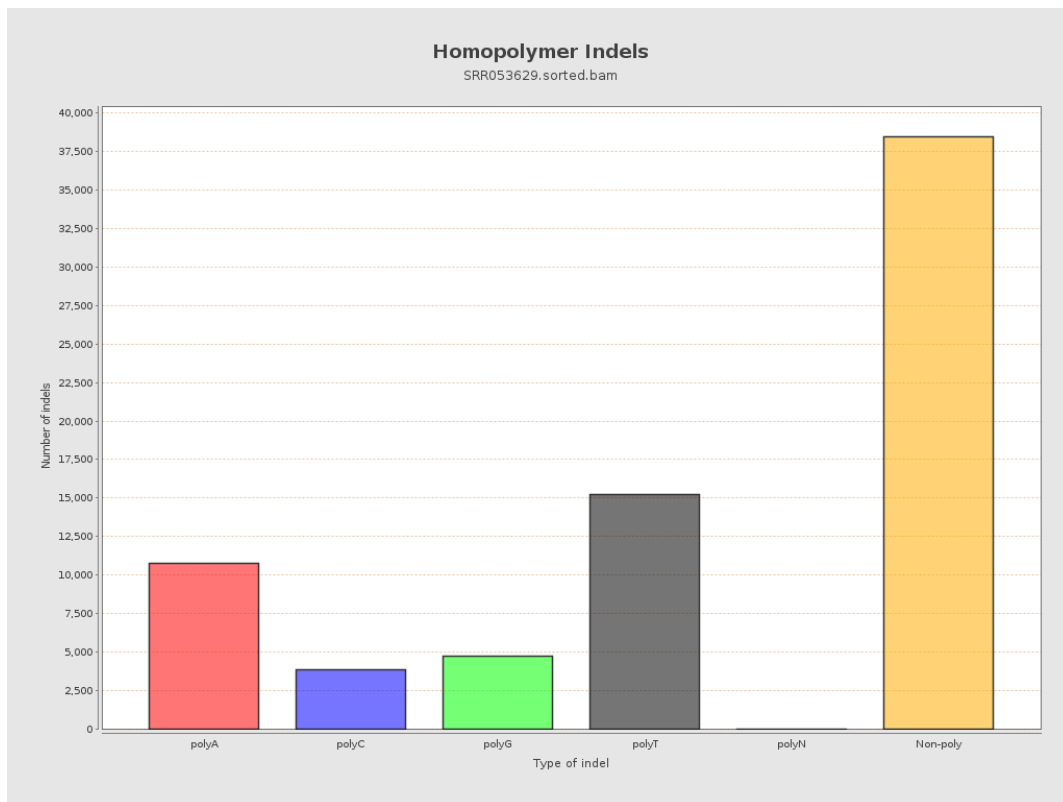
9. Results : Mapped Reads GC-content Distribution



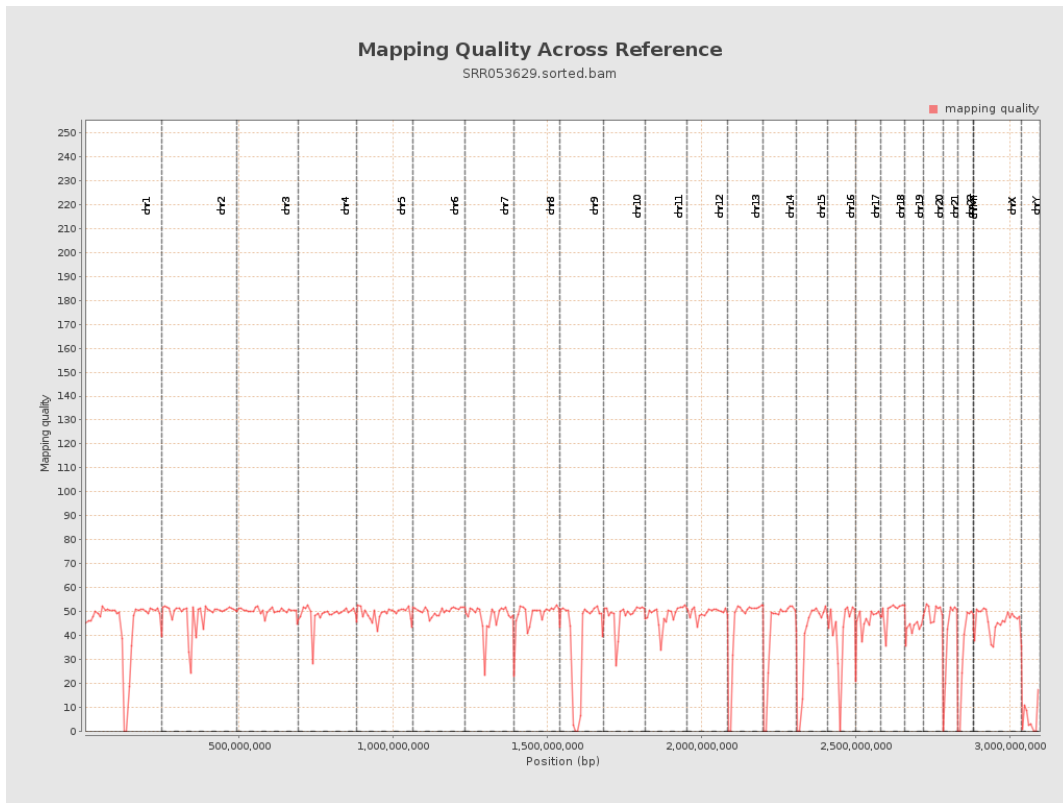
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

