

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/17 23:10:58

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6238627.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_SRR6238627 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6238627.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Sep 17 23:10:58 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6238627.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	4,774,684
Mapped reads	4,210,385 / 88.18%
Unmapped reads	564,299 / 11.82%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	43,413 / 0.91%
Read min/max/mean length	30 / 76 / 76.32
Duplicated reads (estimated)	2,061,319 / 43.17%
Duplication rate	22.86%
Clipped reads	2,842,737 / 59.54%

2.2. ACGT Content

Number/percentage of A's	64,116,036 / 25.03%
Number/percentage of C's	44,206,603 / 17.26%
Number/percentage of T's	87,131,319 / 34.01%
Number/percentage of G's	60,702,530 / 23.7%
Number/percentage of N's	5,735 / 0%
GC Percentage	40.95%

2.3. Coverage

Mean	0.0828

Standard Deviation	1.422
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2.4. Mapping Quality

Mean Mapping Quality	40.28
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2.5. Mismatches and indels

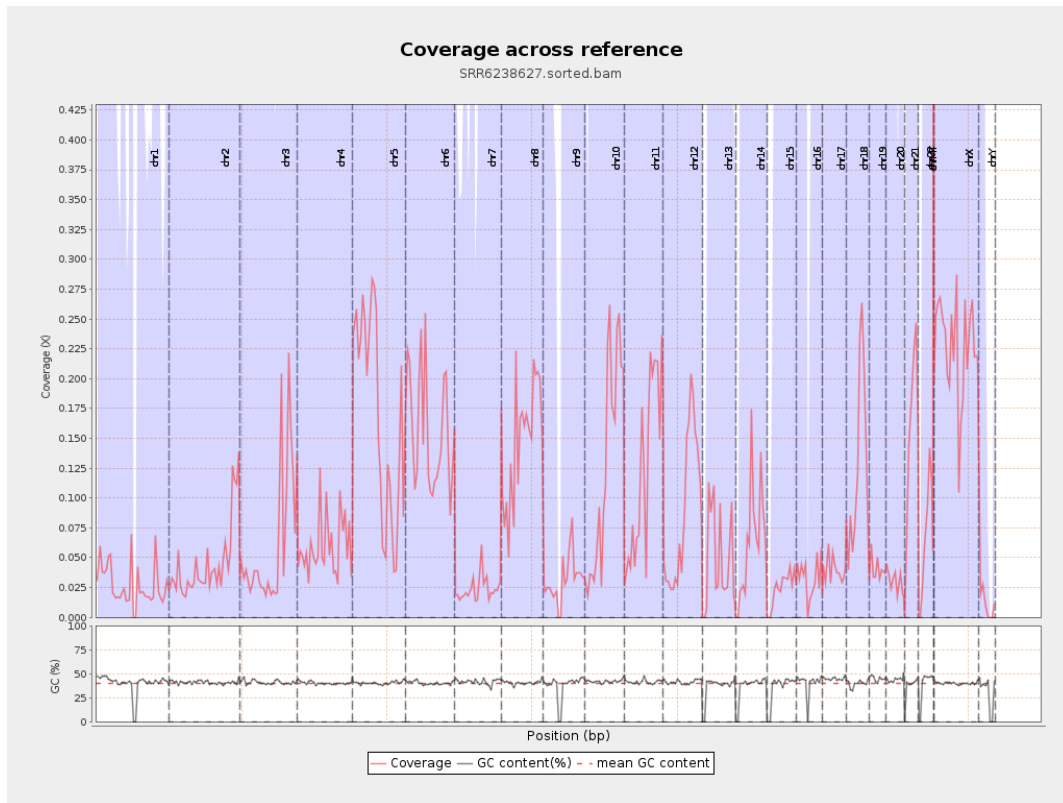
General error rate	0.57%
Mismatches	1,427,086
Insertions	16,356
Mapped reads with at least one insertion	0.38%
Deletions	66,861
Mapped reads with at least one deletion	1.57%
Homopolymer indels	42.01%

2.6. Chromosome stats

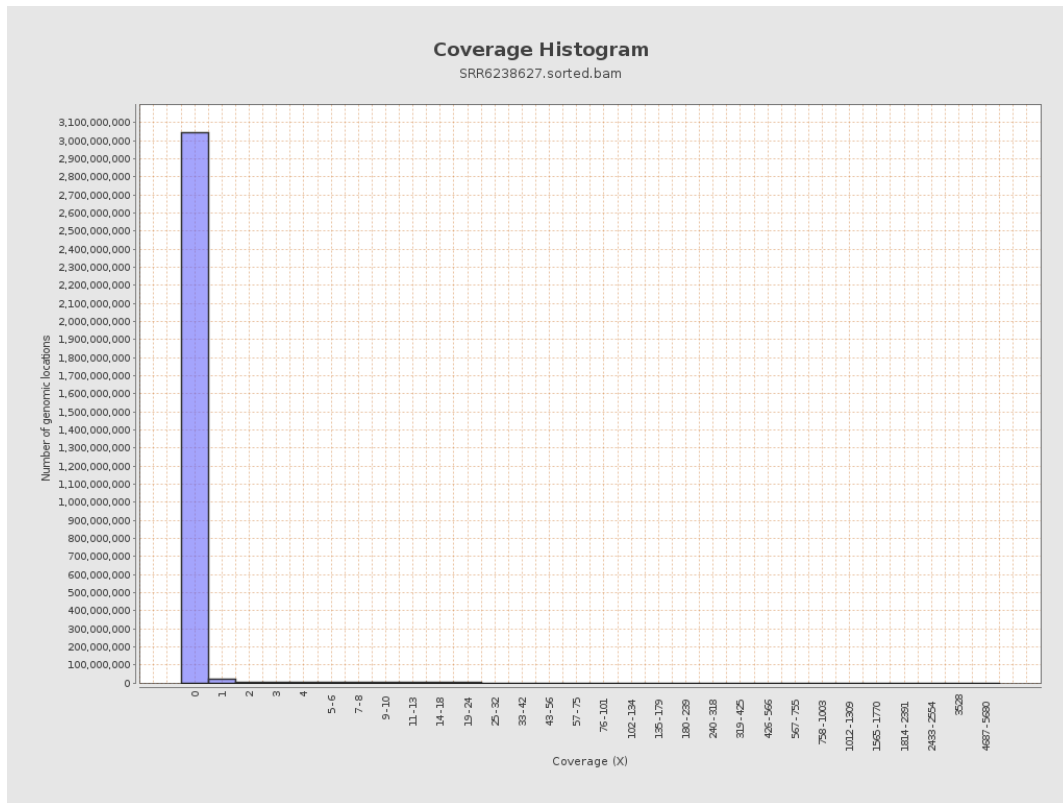
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	6715699	0.0269	1.0823
chr2	243199373	10962880	0.0451	2.6135
chr3	198022430	12023454	0.0607	0.9131
chr4	191154276	11305355	0.0591	0.8927
chr5	180915260	30653986	0.1694	1.5576
chr6	171115067	26632884	0.1556	1.7631
chr7	159138663	3808860	0.0239	0.6778

chr8	146364022	21771379	0.1487	1.629
chr9	141213431	4404300	0.0312	0.7034
chr10	135534747	16910004	0.1248	1.3062
chr11	135006516	16538640	0.1225	1.3522
chr12	133851895	12007221	0.0897	1.1238
chr13	115169878	5552272	0.0482	1.0538
chr14	107349540	6901691	0.0643	1.0331
chr15	102531392	2525438	0.0246	0.7942
chr16	90354753	3009063	0.0333	0.7067
chr17	81195210	3269685	0.0403	0.7214
chr18	78077248	10180857	0.1304	2.9135
chr19	59128983	2410929	0.0408	1.0223
chr20	63025520	1901245	0.0302	0.6509
chr21	48129895	7374999	0.1532	1.4668
chr22	51304566	3457064	0.0674	0.9483
chrMT	16571	11455	0.6913	3.078
chrX	155270560	34999249	0.2254	1.824
chrY	59373566	946658	0.0159	0.5294

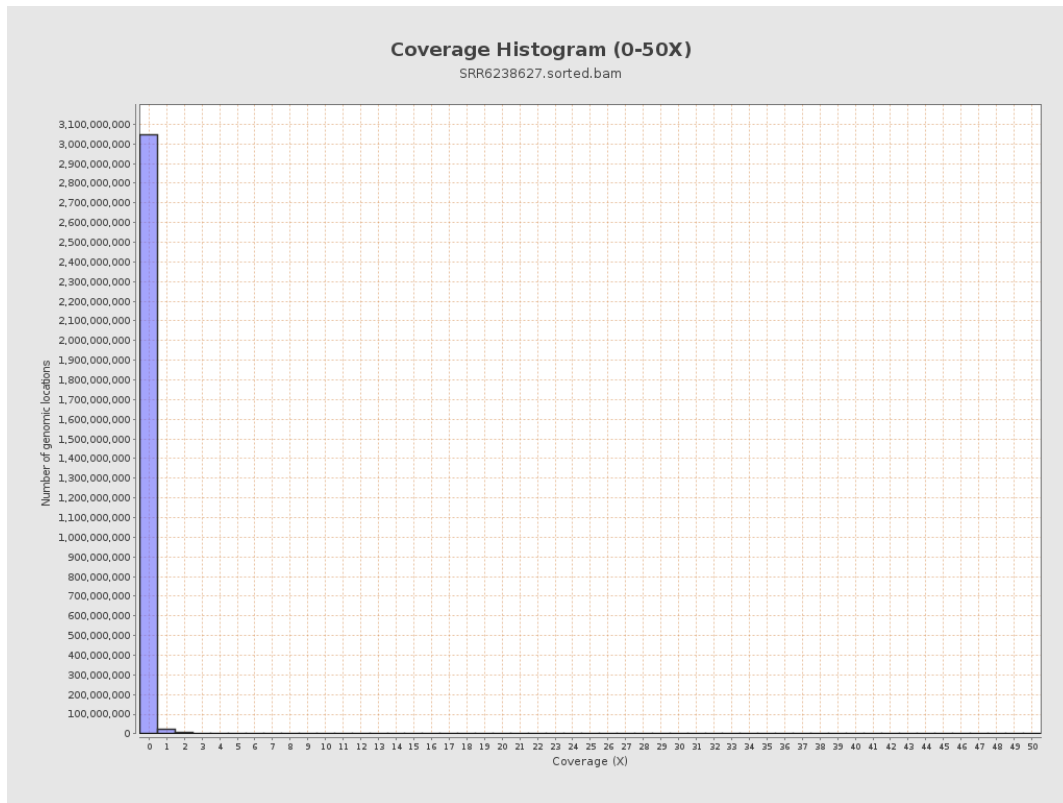
3. Results : Coverage across reference



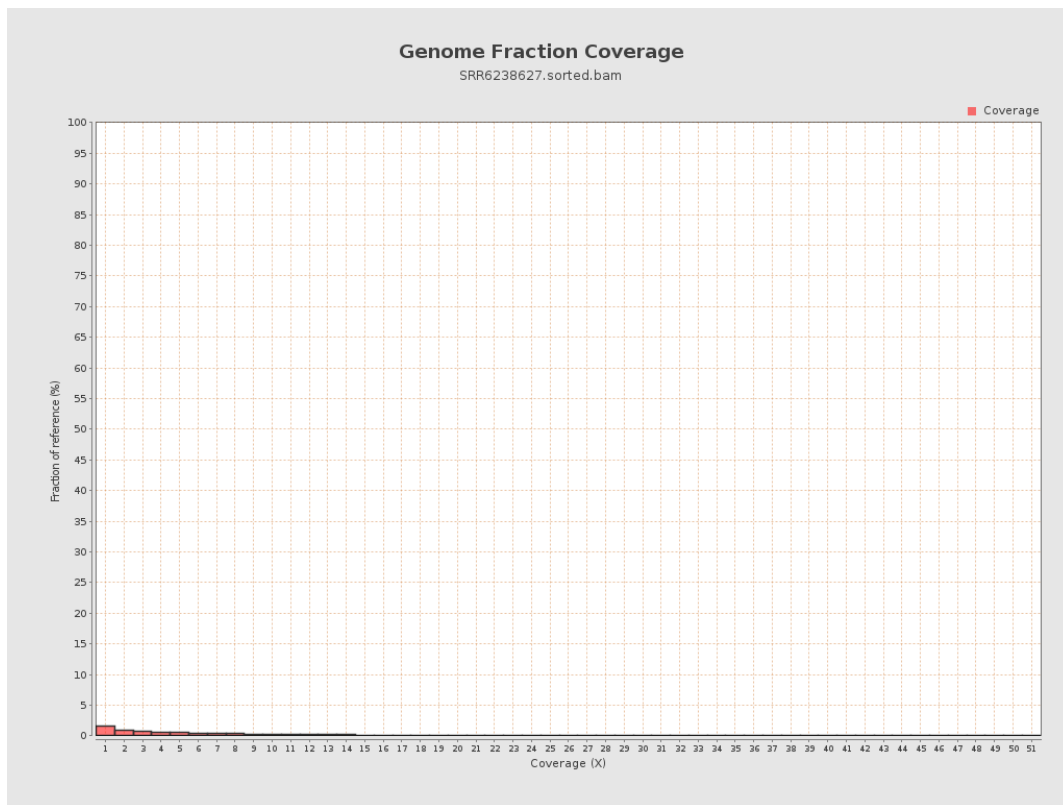
4. Results : Coverage Histogram



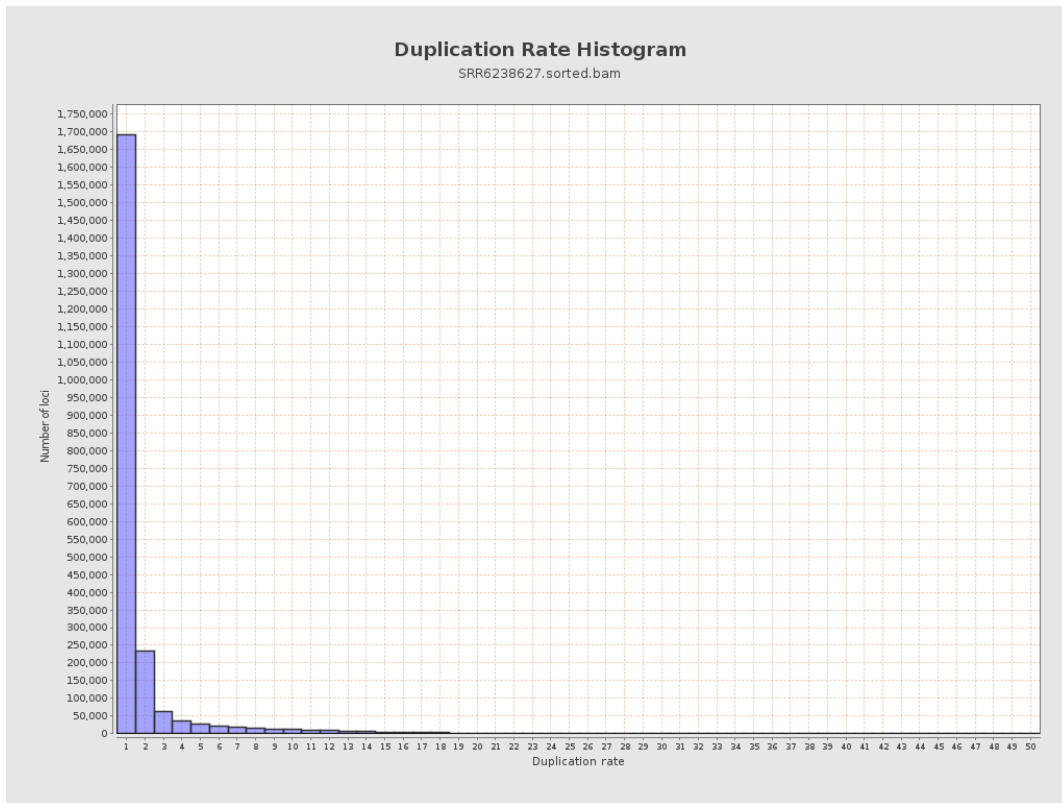
5. Results : Coverage Histogram (0-50X)



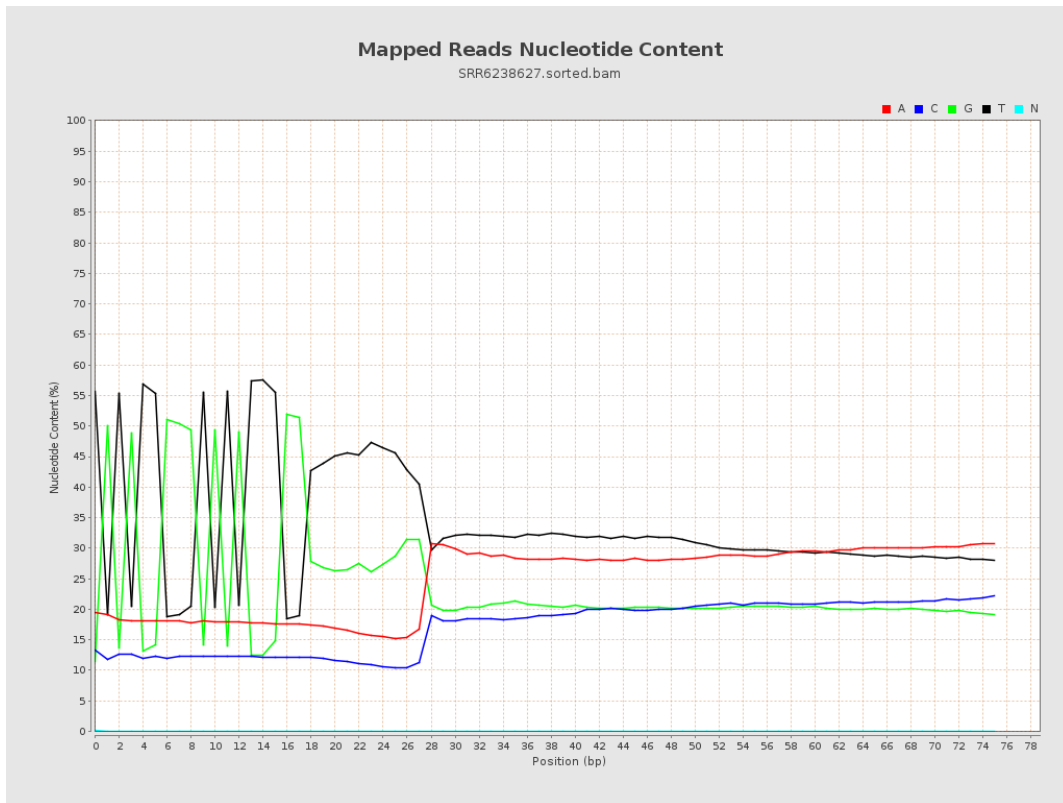
6. Results : Genome Fraction Coverage



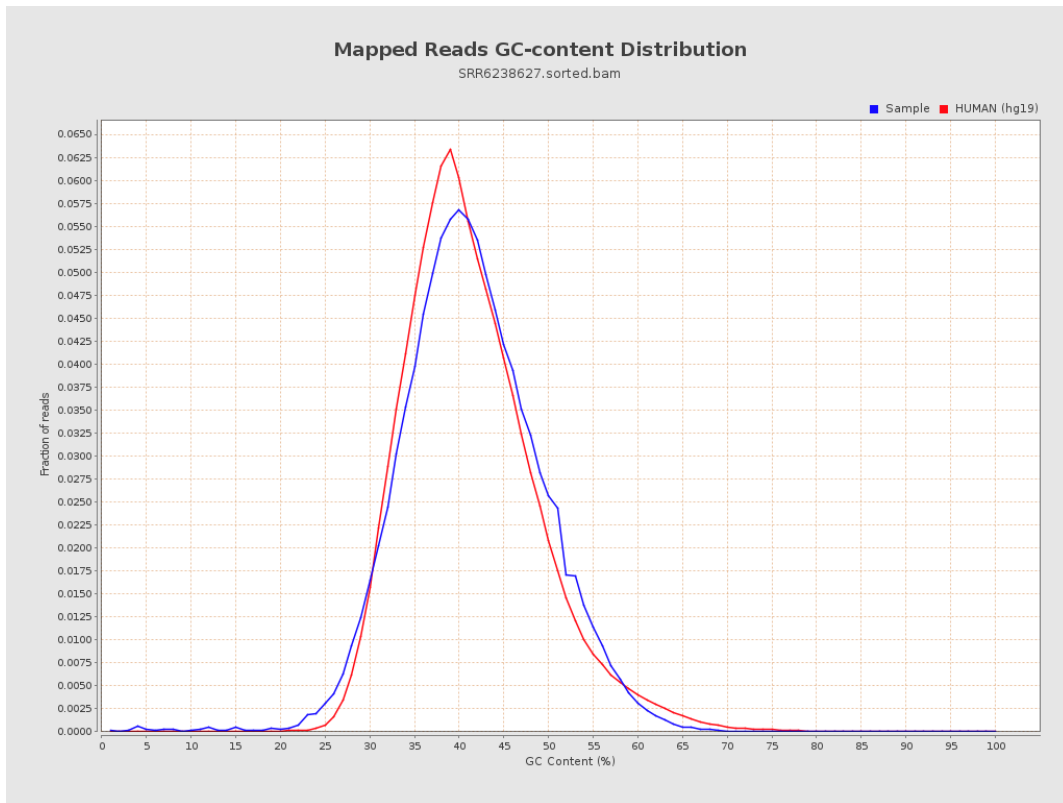
7. Results : Duplication Rate Histogram



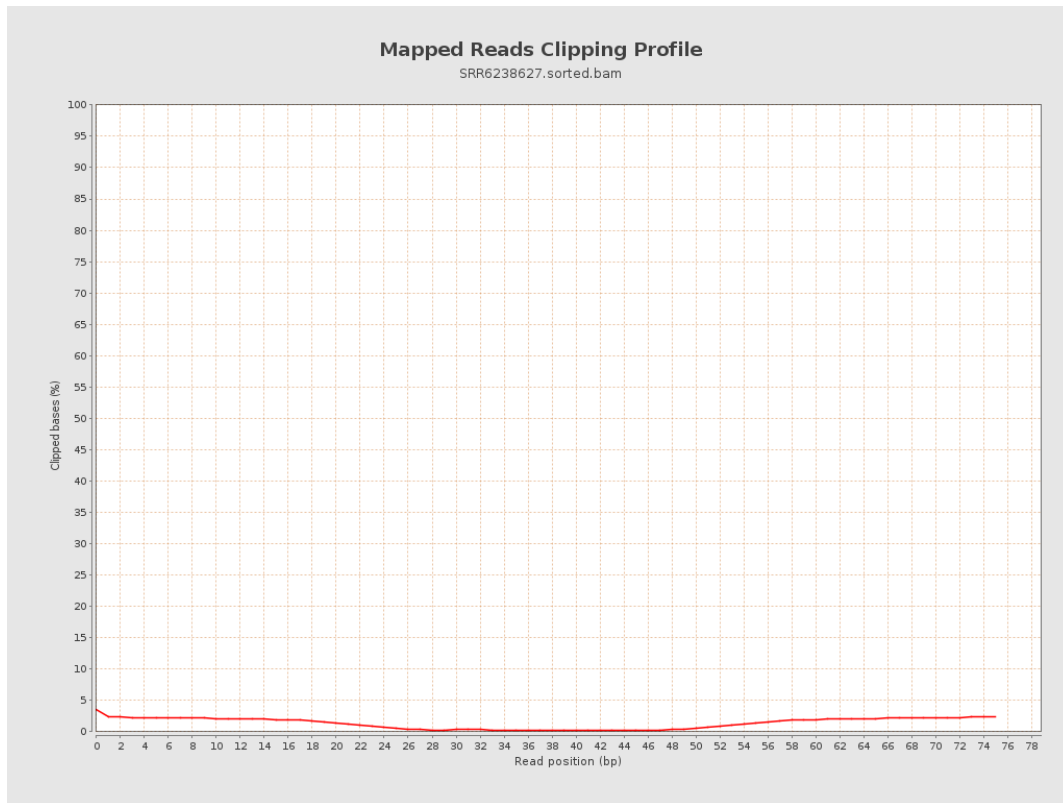
8. Results : Mapped Reads Nucleotide Content



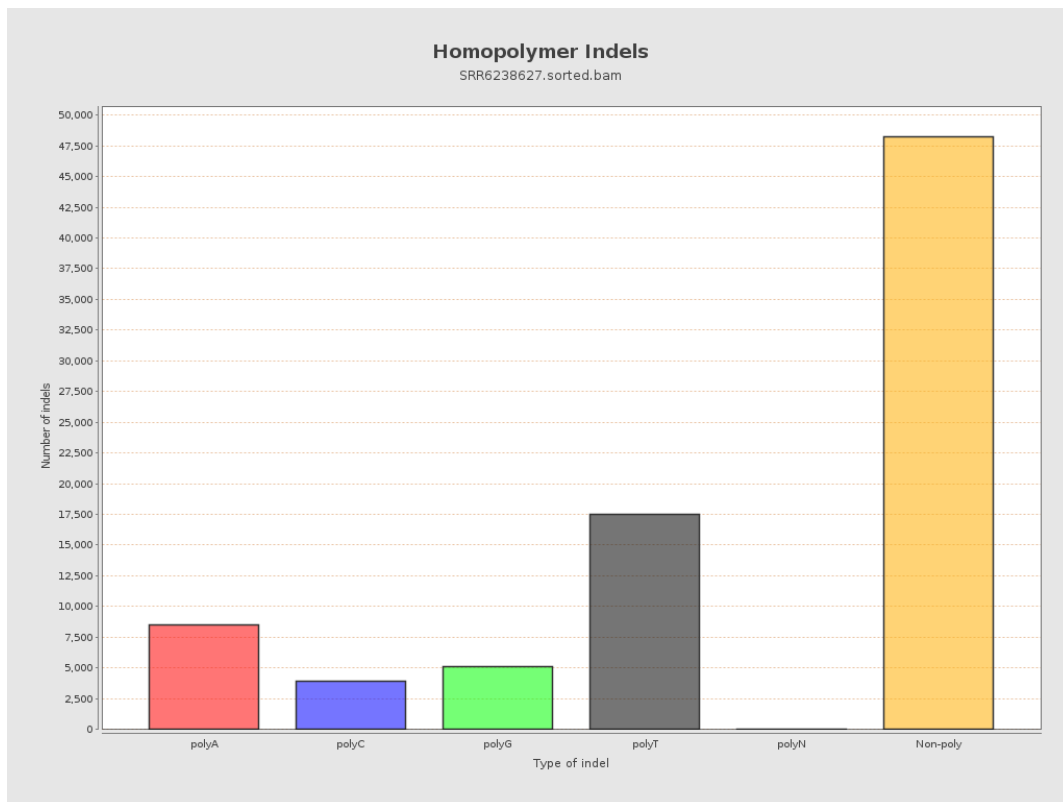
9. Results : Mapped Reads GC-content Distribution



10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

