

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/15 01:11:50

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6063538.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_SRR6063538 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6063538.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sun Sep 15 01:11:49 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6063538.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,797,237
Mapped reads	3,578,553 / 94.24%
Unmapped reads	218,684 / 5.76%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	38,454 / 1.01%
Read min/max/mean length	30 / 76 / 76.35
Duplicated reads (estimated)	908,312 / 23.92%
Duplication rate	17.04%
Clipped reads	1,766,984 / 46.53%

2.2. ACGT Content

Number/percentage of A's	62,178,547 / 26.58%
Number/percentage of C's	41,882,084 / 17.9%
Number/percentage of T's	76,248,413 / 32.6%
Number/percentage of G's	53,583,562 / 22.91%
Number/percentage of N's	26,791 / 0.01%
GC Percentage	40.81%

2.3. Coverage

Mean	0.0756

Standard Deviation	0.8558
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2.4. Mapping Quality

Mean Mapping Quality	45.94
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2.5. Mismatches and indels

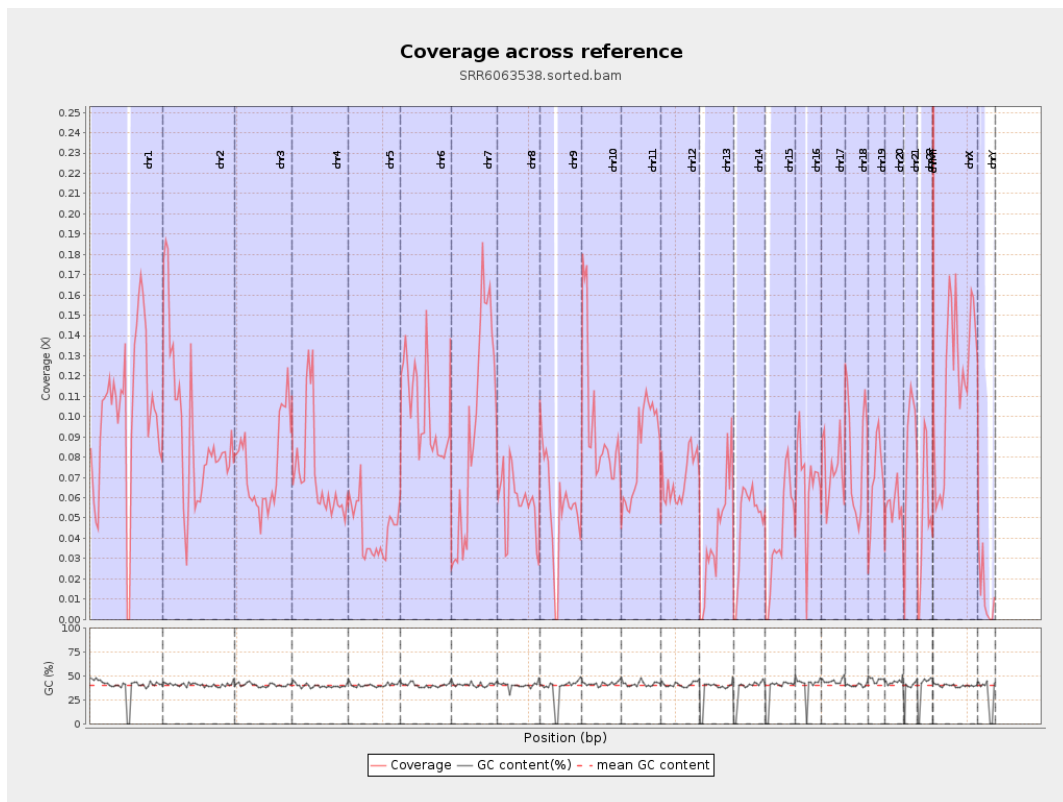
General error rate	0.52%
Mismatches	1,186,441
Insertions	16,301
Mapped reads with at least one insertion	0.45%
Deletions	47,259
Mapped reads with at least one deletion	1.31%
Homopolymer indels	46.32%

2.6. Chromosome stats

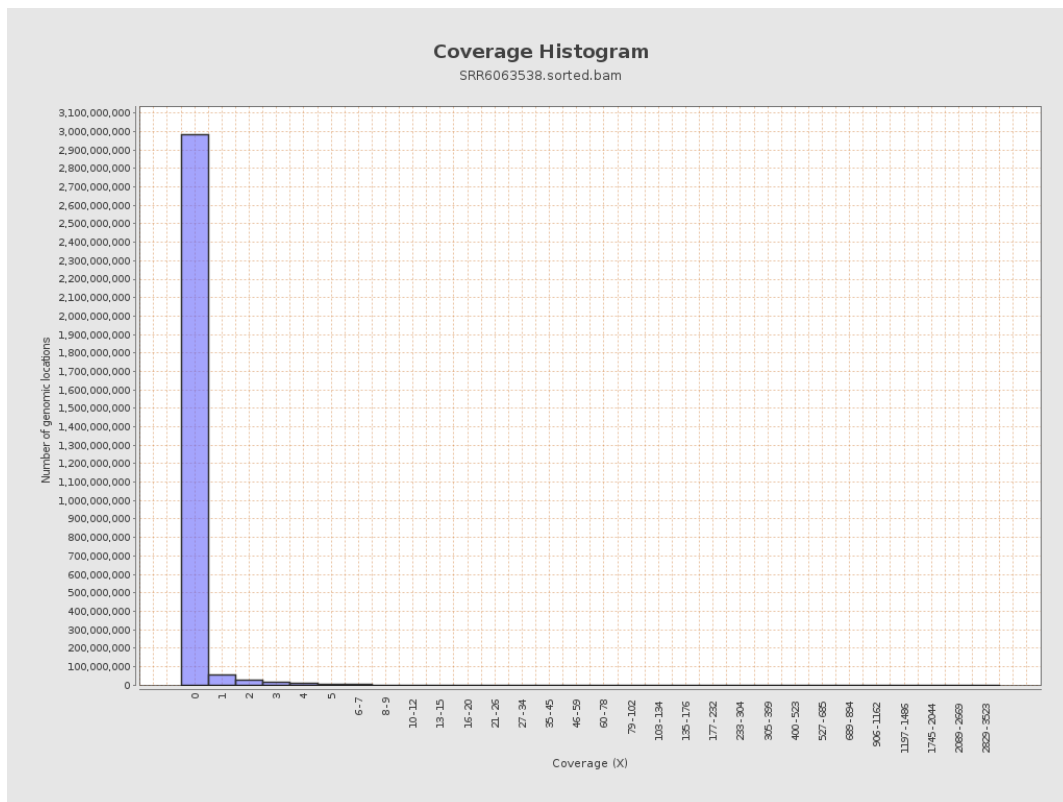
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	25165864	0.101	1.1614
chr2	243199373	22507173	0.0925	1.2646
chr3	198022430	14972036	0.0756	0.5034
chr4	191154276	13794727	0.0722	0.5296
chr5	180915260	8022988	0.0443	0.3806
chr6	171115067	17694077	0.1034	0.6655
chr7	159138663	15039347	0.0945	0.8041

chr8	146364022	8357438	0.0571	2.0436
chr9	141213431	7768558	0.055	0.5861
chr10	135534747	13110585	0.0967	0.6875
chr11	135006516	11275575	0.0835	0.5812
chr12	133851895	9323145	0.0697	0.4866
chr13	115169878	4964127	0.0431	0.4098
chr14	107349540	5190482	0.0484	0.42
chr15	102531392	4196707	0.0409	0.3699
chr16	90354753	6067456	0.0672	0.471
chr17	81195210	6062483	0.0747	0.4975
chr18	78077248	6310565	0.0808	0.9848
chr19	59128983	4091683	0.0692	0.8076
chr20	63025520	3551336	0.0563	0.4475
chr21	48129895	4076950	0.0847	0.5481
chr22	51304566	2516379	0.049	0.3901
chrMT	16571	1088283	65.6739	49.8536
chrX	155270560	18115068	0.1167	0.6748
chrY	59373566	739283	0.0125	0.3136

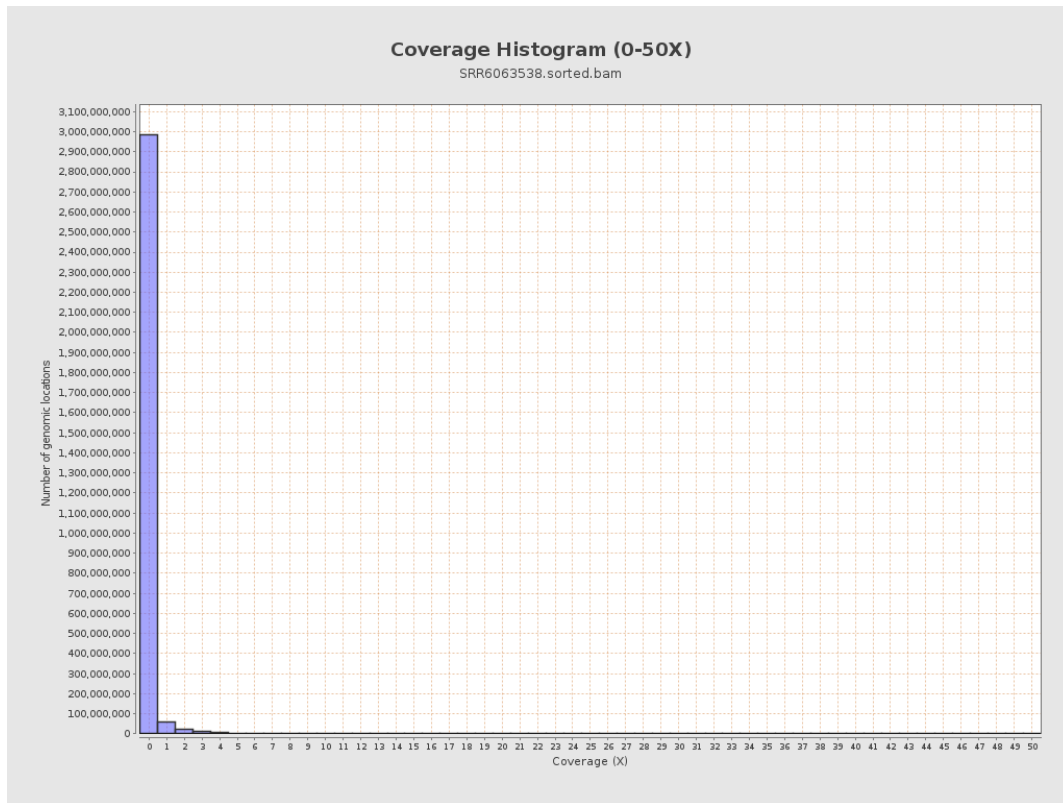
3. Results : Coverage across reference



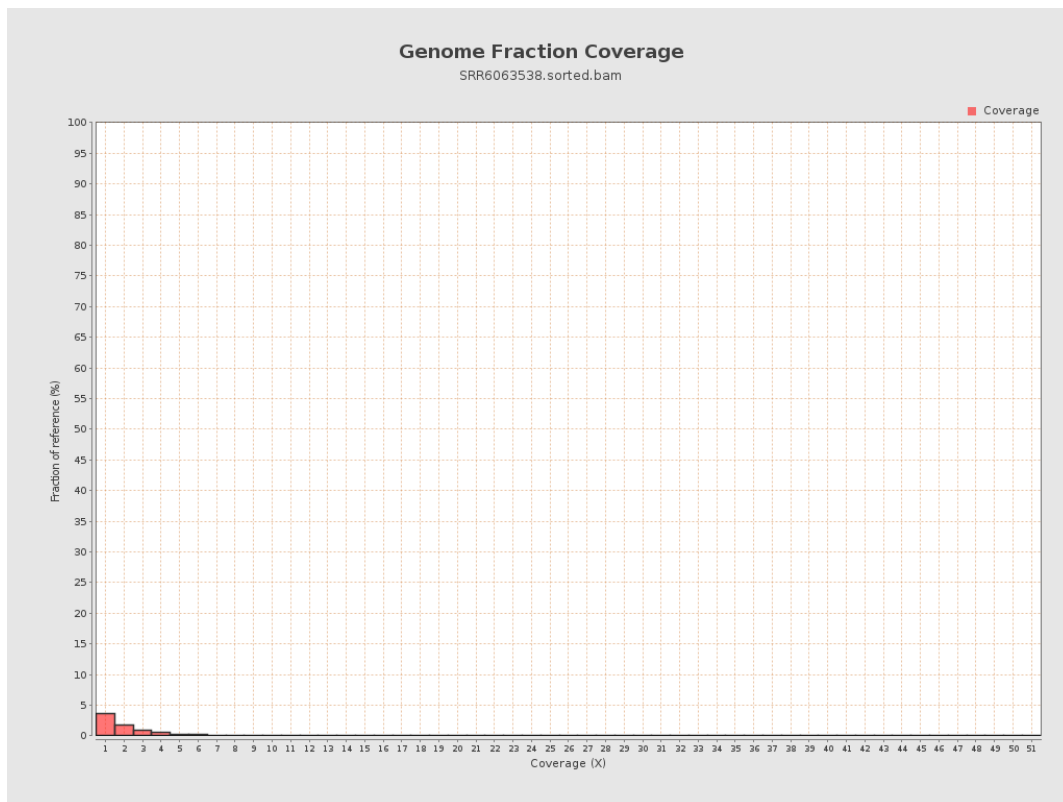
4. Results : Coverage Histogram



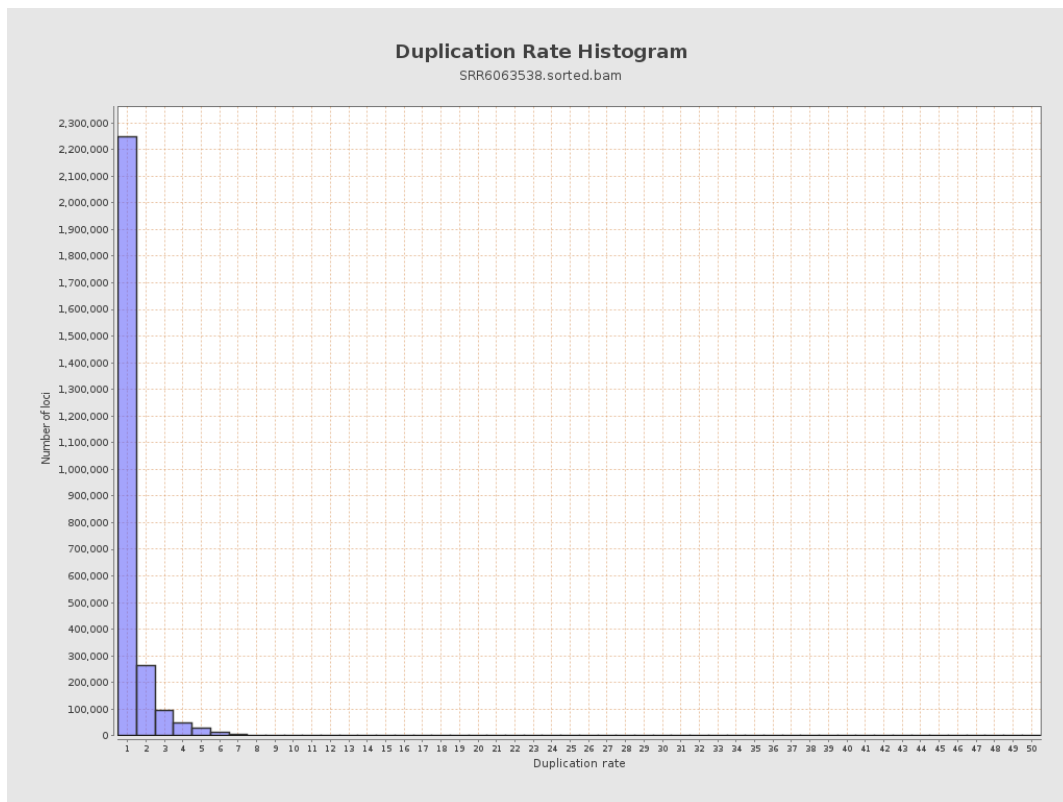
5. Results : Coverage Histogram (0-50X)



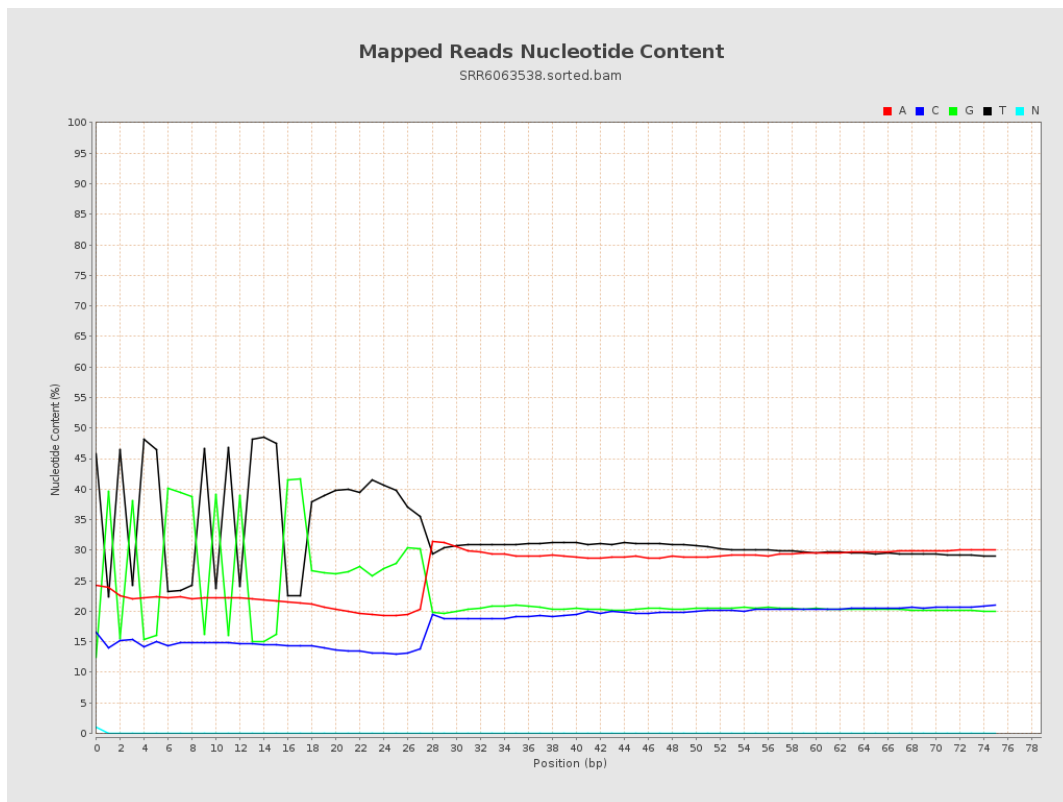
6. Results : Genome Fraction Coverage



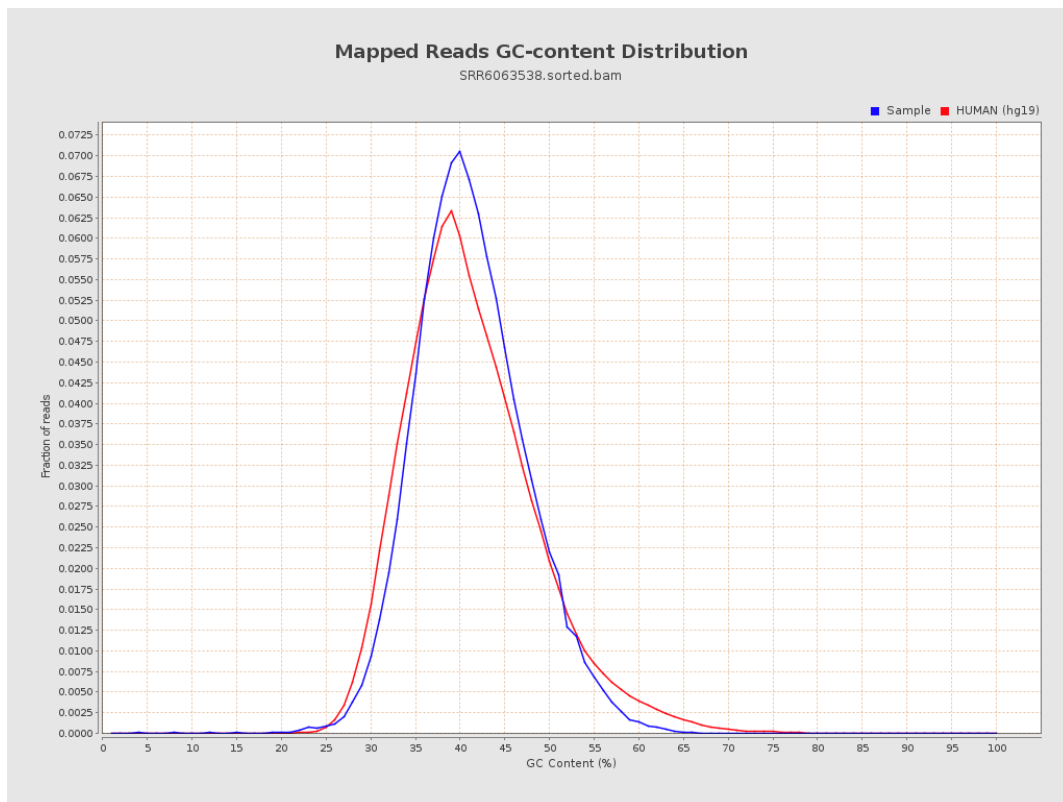
7. Results : Duplication Rate Histogram



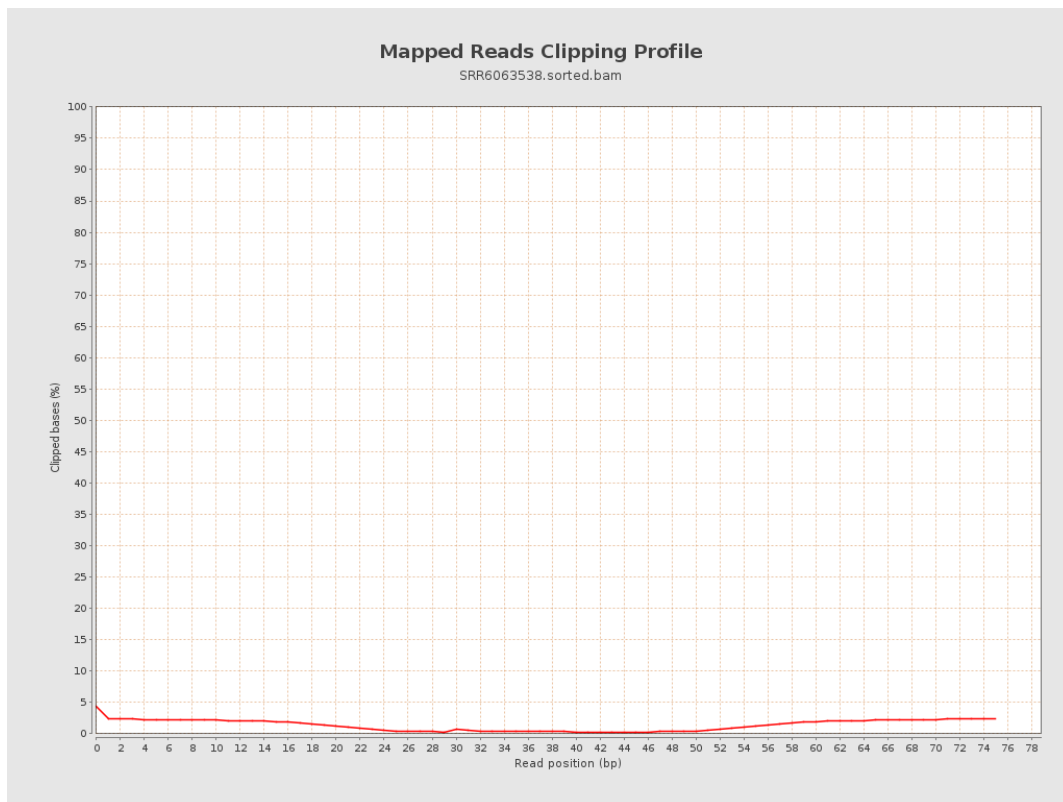
8. Results : Mapped Reads Nucleotide Content



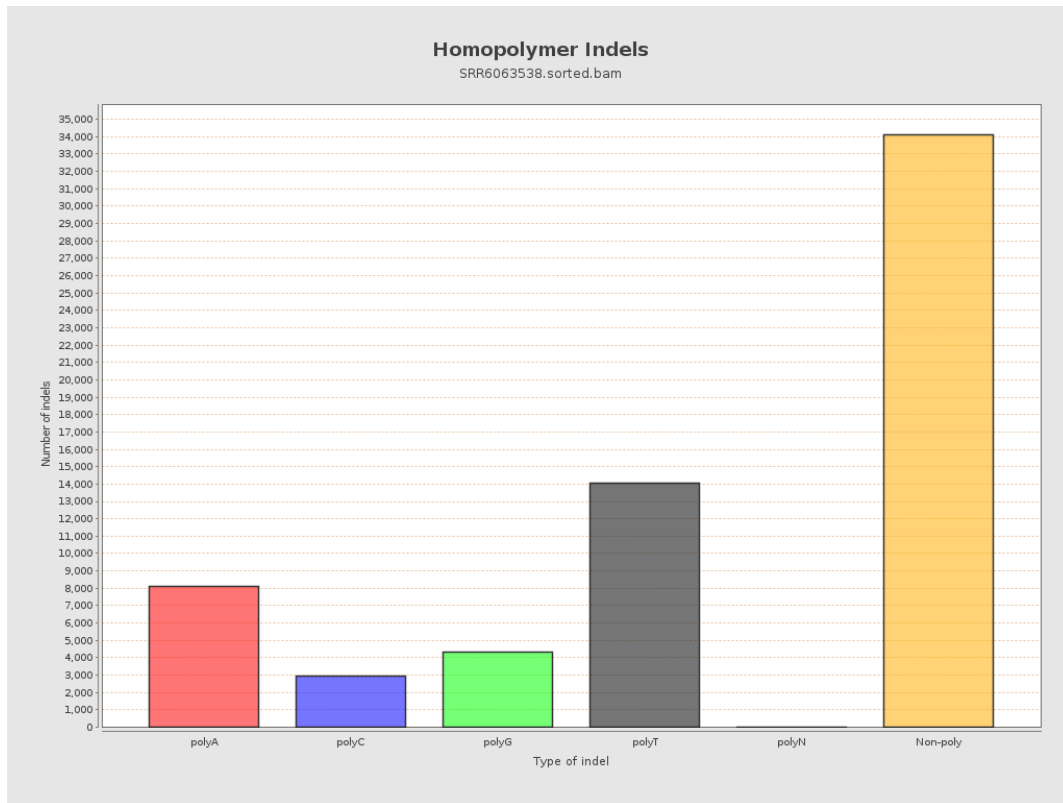
9. Results : Mapped Reads GC-content Distribution



10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

