

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/25 11:33:17

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR3084386.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_SRR3084386 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR3084386.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sun Aug 25 11:33:15 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR3084386.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	1,665,810
Mapped reads	1,533,815 / 92.08%
Unmapped reads	131,995 / 7.92%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	7,818 / 0.47%
Read min/max/mean length	30 / 76 / 76.16
Duplicated reads (estimated)	47,035 / 2.82%
Duplication rate	2.58%
Clipped reads	623,713 / 37.44%

2.2. ACGT Content

Number/percentage of A's	29,399,723 / 28.32%
Number/percentage of C's	19,441,586 / 18.73%
Number/percentage of T's	32,589,948 / 31.39%
Number/percentage of G's	22,249,601 / 21.43%
Number/percentage of N's	138,587 / 0.13%
GC Percentage	40.16%

2.3. Coverage

Mean	0.0335

Standard Deviation	0.2598
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2.4. Mapping Quality

Mean Mapping Quality	46.74
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2.5. Mismatches and indels

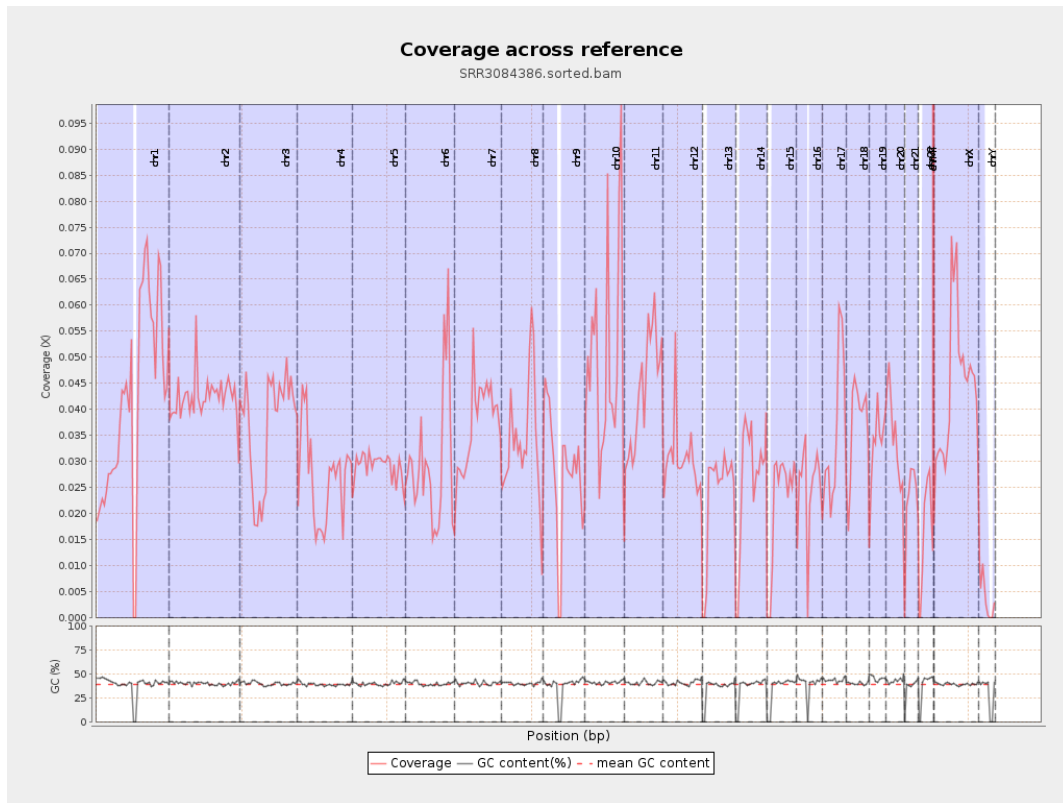
General error rate	0.99%
Mismatches	1,014,556
Insertions	7,295
Mapped reads with at least one insertion	0.47%
Deletions	20,491
Mapped reads with at least one deletion	1.32%
Homopolymer indels	47.71%

2.6. Chromosome stats

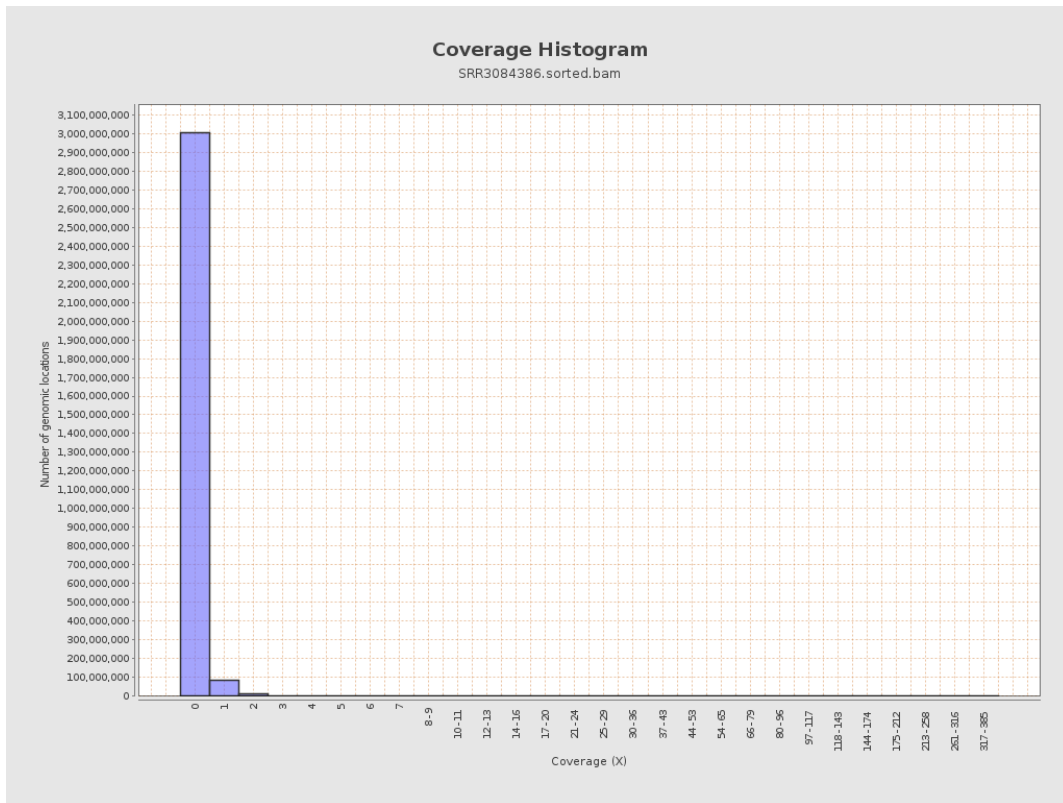
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	10297791	0.0413	0.4054
chr2	243199373	10290383	0.0423	0.2925
chr3	198022430	7270493	0.0367	0.2145
chr4	191154276	5150529	0.0269	0.1894
chr5	180915260	5259231	0.0291	0.1886
chr6	171115067	5048073	0.0295	0.2104
chr7	159138663	5971290	0.0375	0.3466

chr8	146364022	4928936	0.0337	0.2448
chr9	141213431	3954335	0.028	0.2328
chr10	135534747	6772146	0.05	0.3744
chr11	135006516	5851139	0.0433	0.2701
chr12	133851895	4076824	0.0305	0.1941
chr13	115169878	2683619	0.0233	0.1687
chr14	107349540	2855977	0.0266	0.1854
chr15	102531392	2291980	0.0224	0.1661
chr16	90354753	2255371	0.025	0.187
chr17	81195210	2857462	0.0352	0.2189
chr18	78077248	2878398	0.0369	0.3555
chr19	59128983	2013752	0.0341	0.301
chr20	63025520	2155909	0.0342	0.2059
chr21	48129895	1079545	0.0224	0.1738
chr22	51304566	830460	0.0162	0.1394
chrMT	16571	20011	1.2076	1.3868
chrX	155270560	6841592	0.0441	0.2515
chrY	59373566	219322	0.0037	0.0865

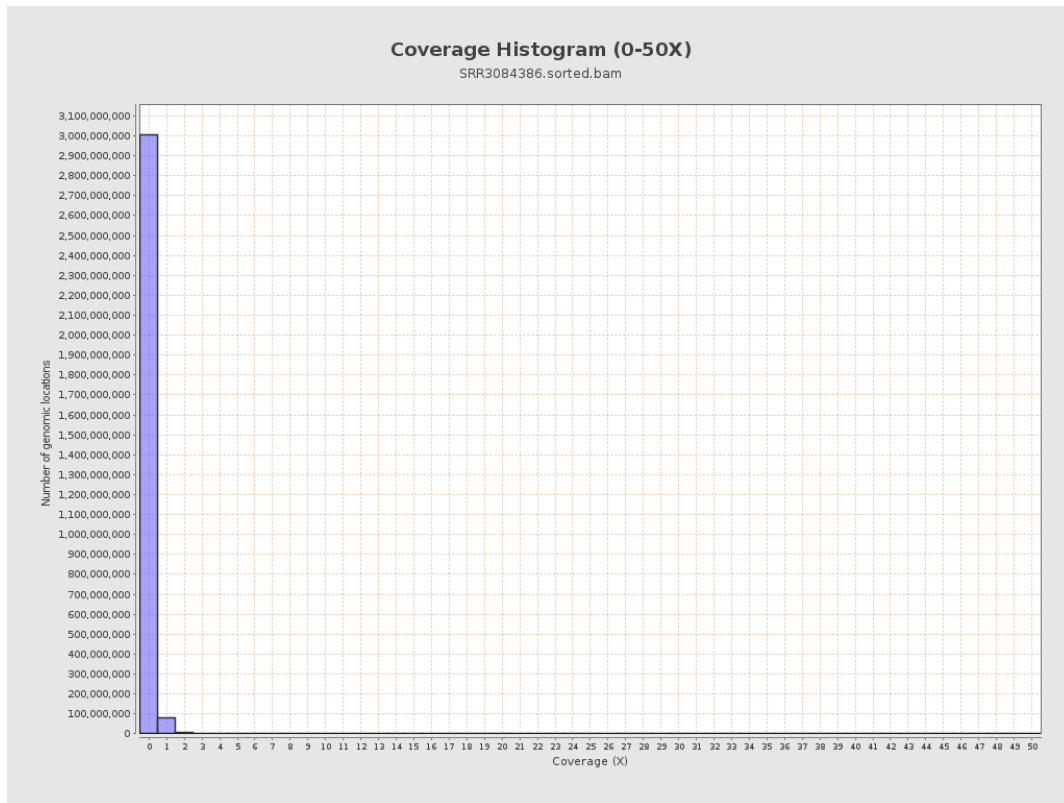
3. Results : Coverage across reference



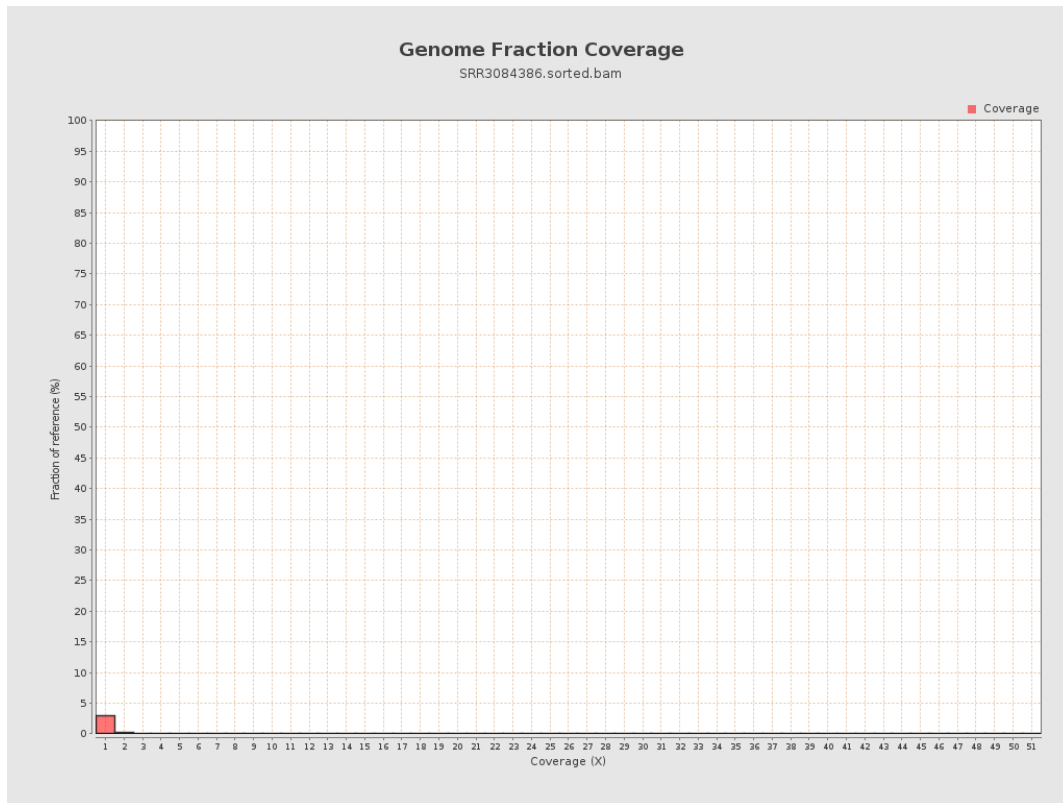
4. Results : Coverage Histogram



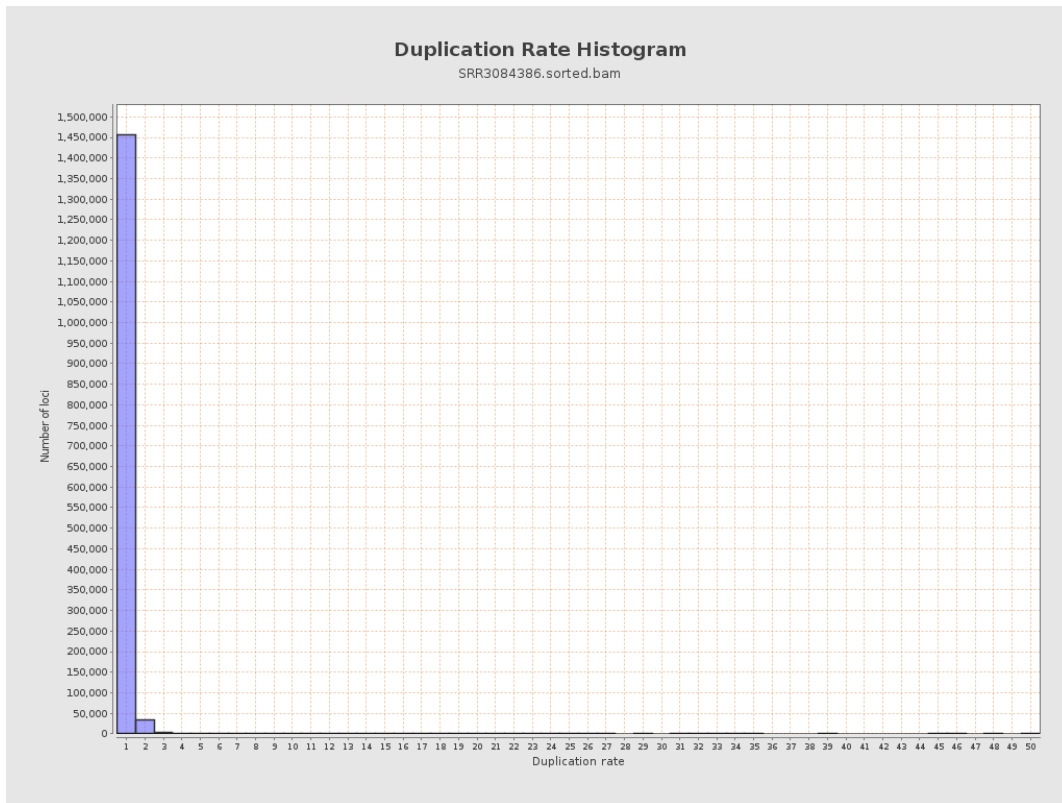
5. Results : Coverage Histogram (0-50X)



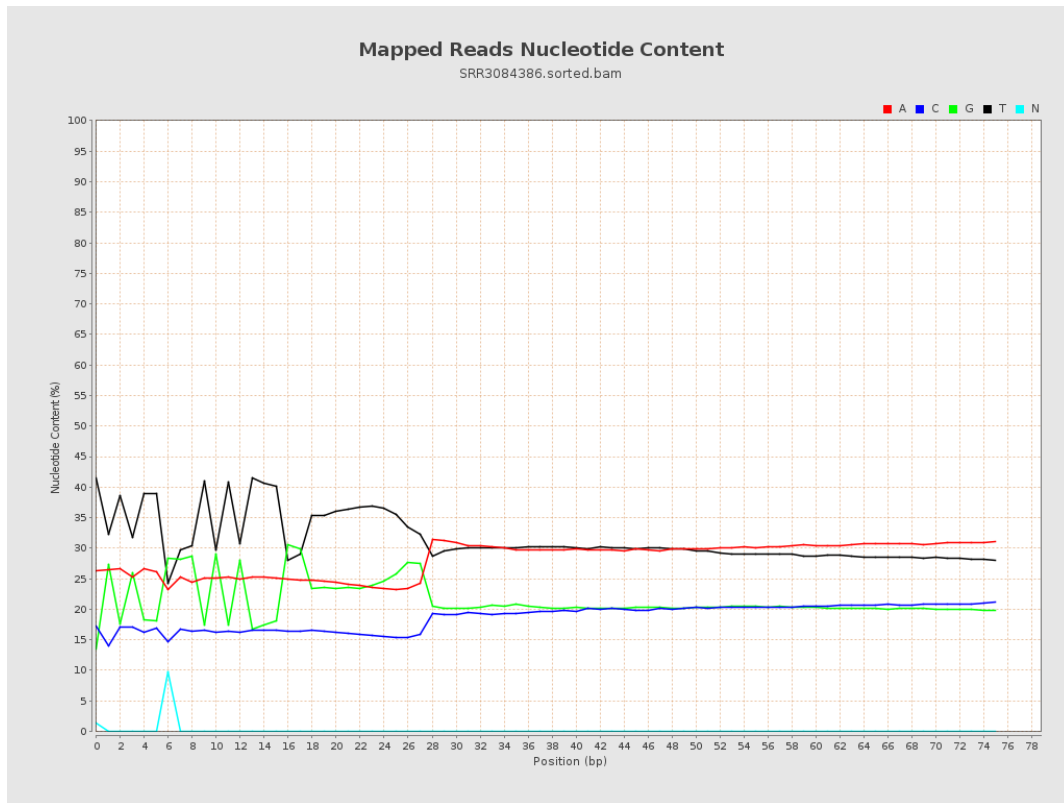
6. Results : Genome Fraction Coverage



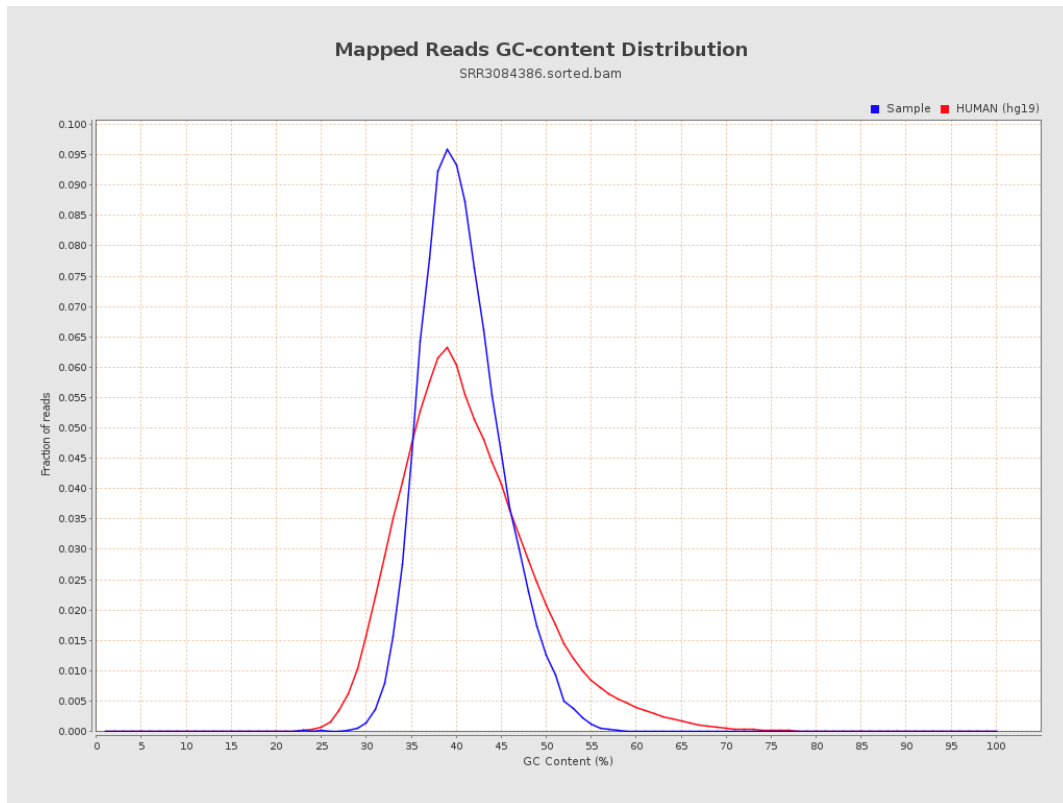
7. Results : Duplication Rate Histogram



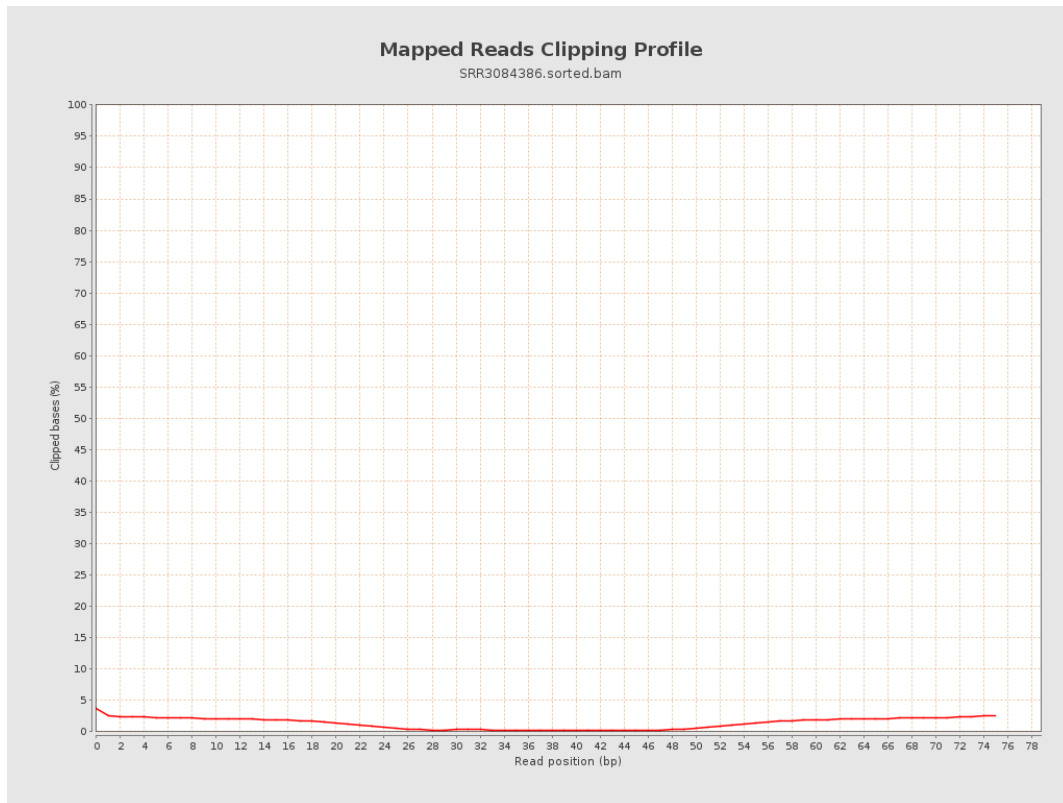
8. Results : Mapped Reads Nucleotide Content



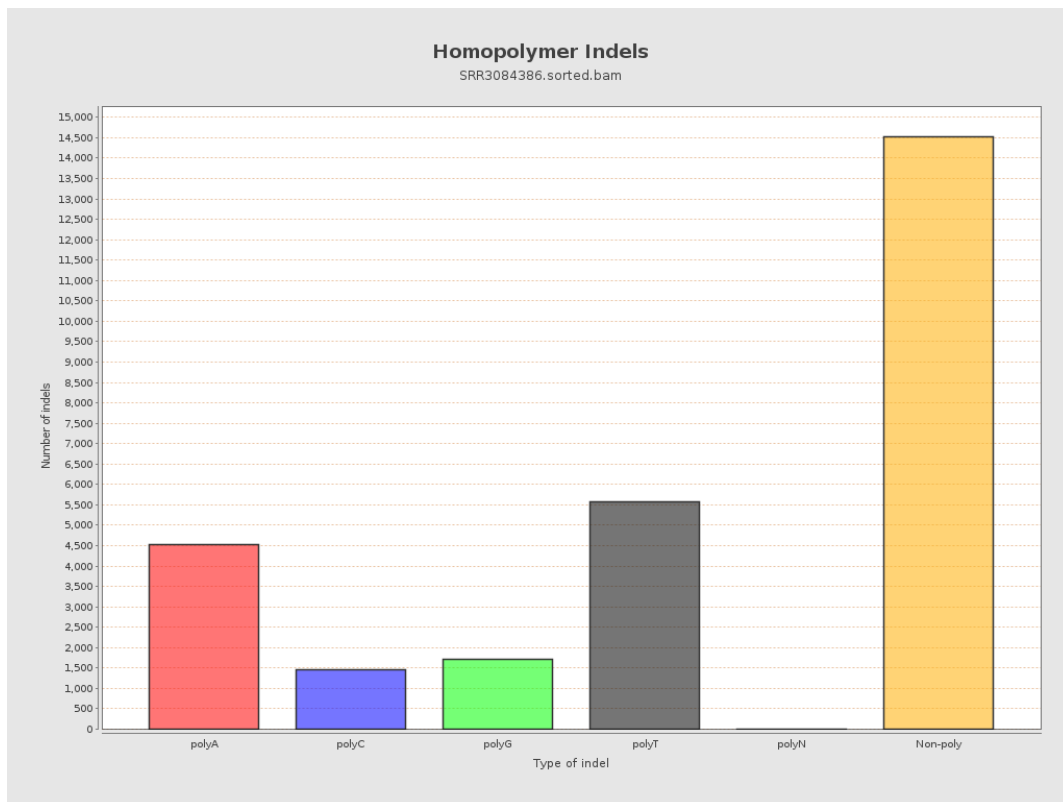
9. Results : Mapped Reads GC-content Distribution



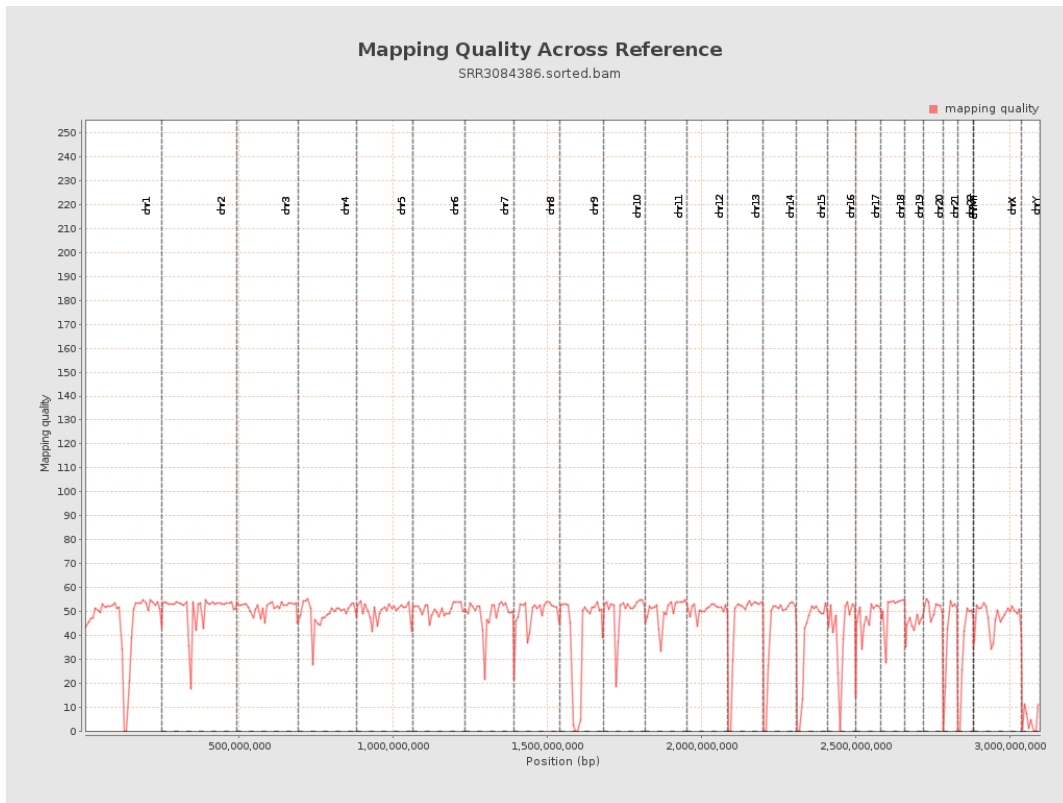
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

