

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/24 22:43:35

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR3082979.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_SRR3082979 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR3082979.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sat Aug 24 22:43:33 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR3082979.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,555,530
Mapped reads	2,849,468 / 80.14%
Unmapped reads	706,062 / 19.86%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	18,726 / 0.53%
Read min/max/mean length	30 / 76 / 76.18
Duplicated reads (estimated)	105,198 / 2.96%
Duplication rate	2.96%
Clipped reads	1,071,097 / 30.12%

2.2. ACGT Content

Number/percentage of A's	56,739,207 / 28.95%
Number/percentage of C's	36,202,348 / 18.47%
Number/percentage of T's	62,700,871 / 32%
Number/percentage of G's	40,301,298 / 20.57%
Number/percentage of N's	19,813 / 0.01%
GC Percentage	39.04%

2.3. Coverage

Mean	0.0633

Standard Deviation	0.4394
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2.4. Mapping Quality

Mean Mapping Quality	47.14
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2.5. Mismatches and indels

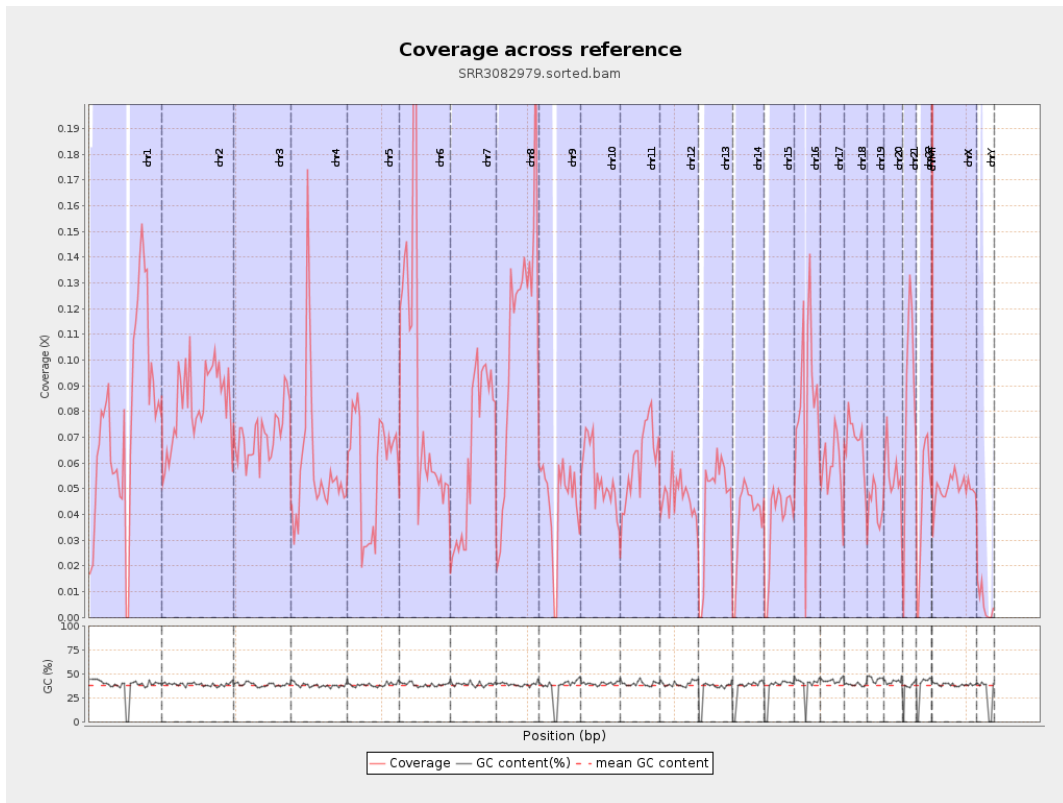
General error rate	0.94%
Mismatches	1,818,118
Insertions	16,588
Mapped reads with at least one insertion	0.58%
Deletions	43,530
Mapped reads with at least one deletion	1.51%
Homopolymer indels	48.07%

2.6. Chromosome stats

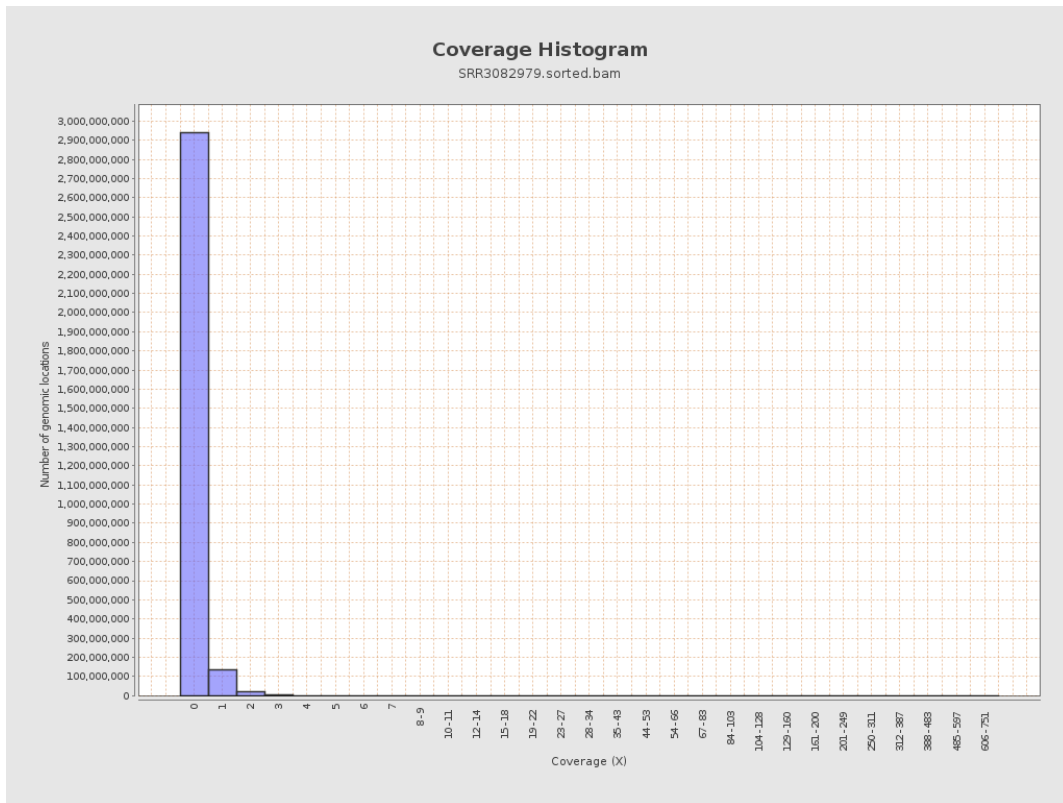
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	18976049	0.0761	0.6962
chr2	243199373	20237003	0.0832	0.5359
chr3	198022430	14033825	0.0709	0.3139
chr4	191154276	11277007	0.059	0.2979
chr5	180915260	10433453	0.0577	0.2851
chr6	171115067	15130519	0.0884	0.493
chr7	159138663	10199141	0.0641	0.3872

chr8	146364022	15989735	0.1092	0.6208
chr9	141213431	6394270	0.0453	0.3904
chr10	135534747	7386915	0.0545	0.3659
chr11	135006516	8249928	0.0611	0.3793
chr12	133851895	6255283	0.0467	0.2617
chr13	115169878	5214375	0.0453	0.2508
chr14	107349540	4167526	0.0388	0.2879
chr15	102531392	3725690	0.0363	0.2233
chr16	90354753	7504722	0.0831	0.3943
chr17	81195210	4815382	0.0593	0.3349
chr18	78077248	5368148	0.0688	0.9669
chr19	59128983	2601471	0.044	0.5204
chr20	63025520	3523077	0.0559	0.3038
chr21	48129895	4024662	0.0836	0.3543
chr22	51304566	2309184	0.045	0.2489
chrMT	16571	134352	8.1077	6.1588
chrX	155270560	7774704	0.0501	0.2982
chrY	59373566	311701	0.0052	0.1098

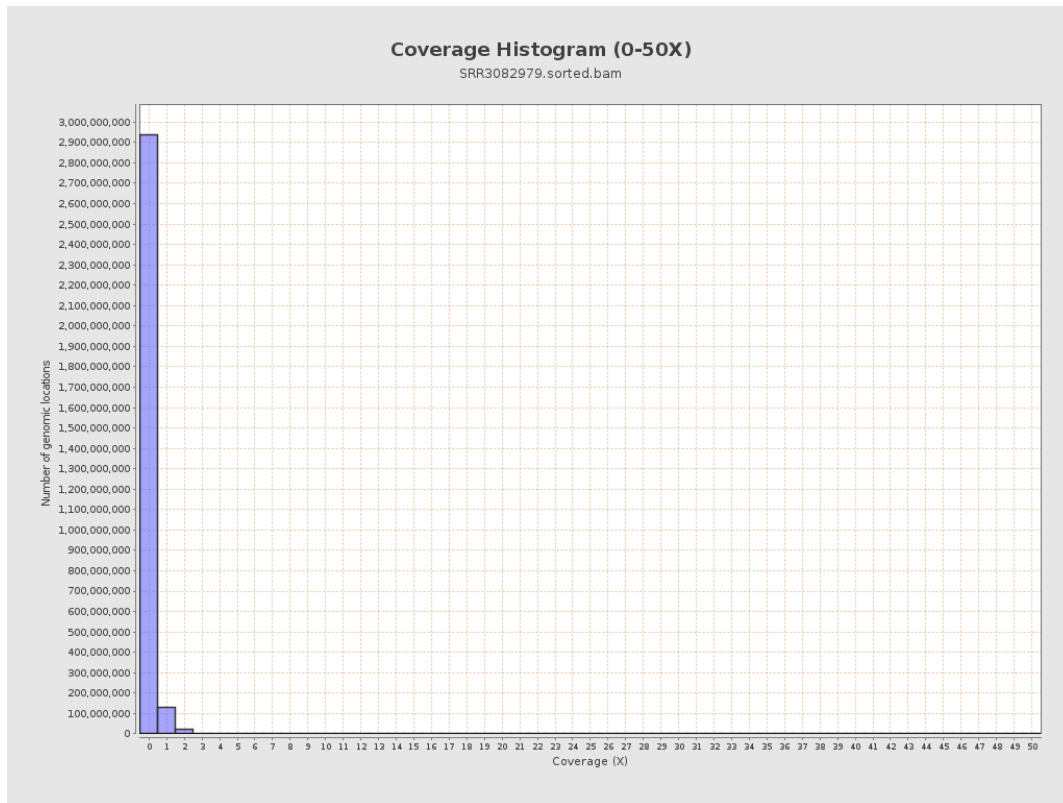
3. Results : Coverage across reference



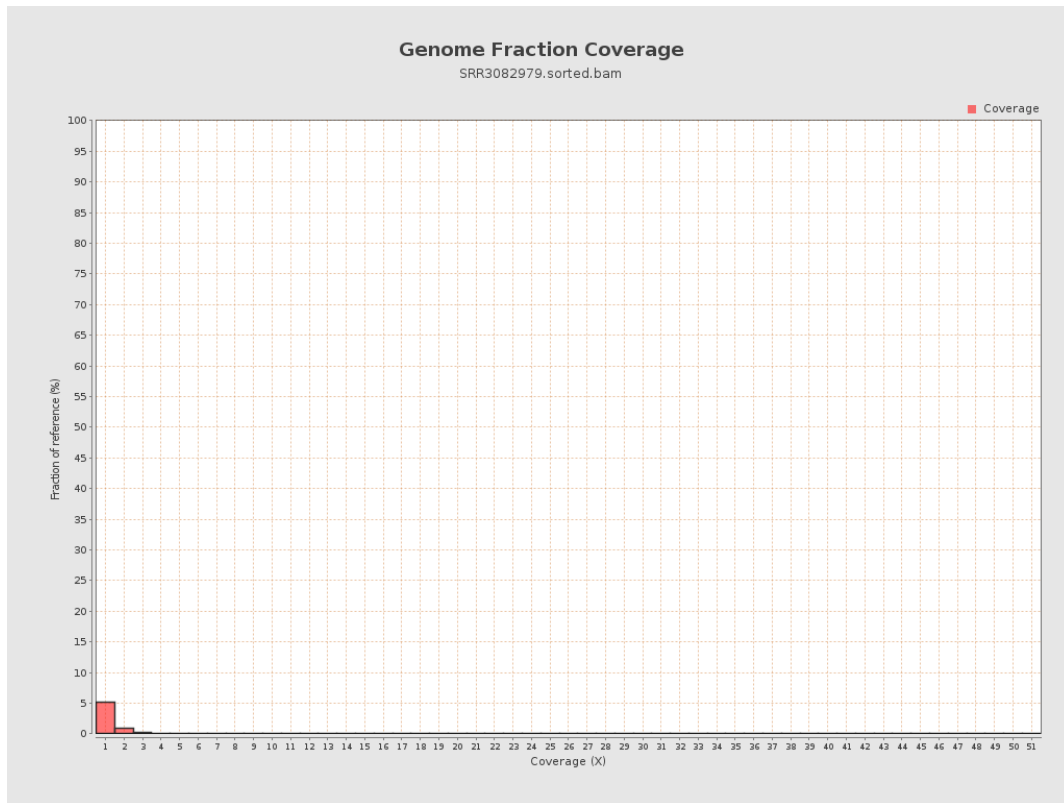
4. Results : Coverage Histogram



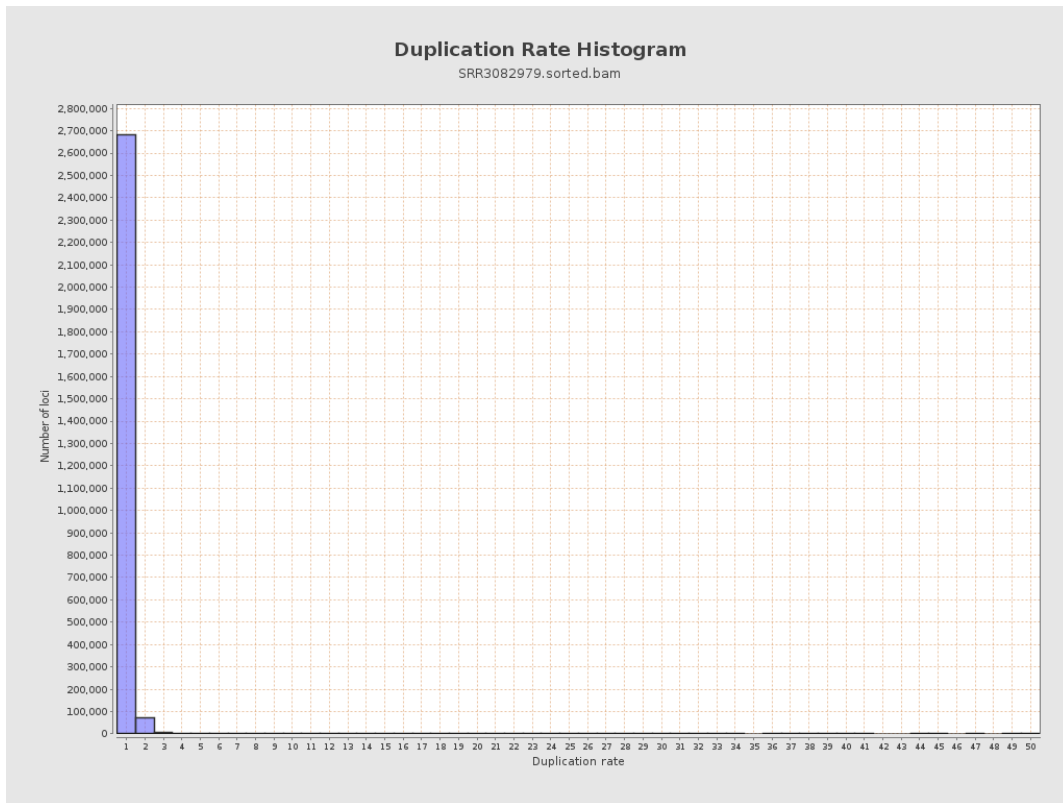
5. Results : Coverage Histogram (0-50X)



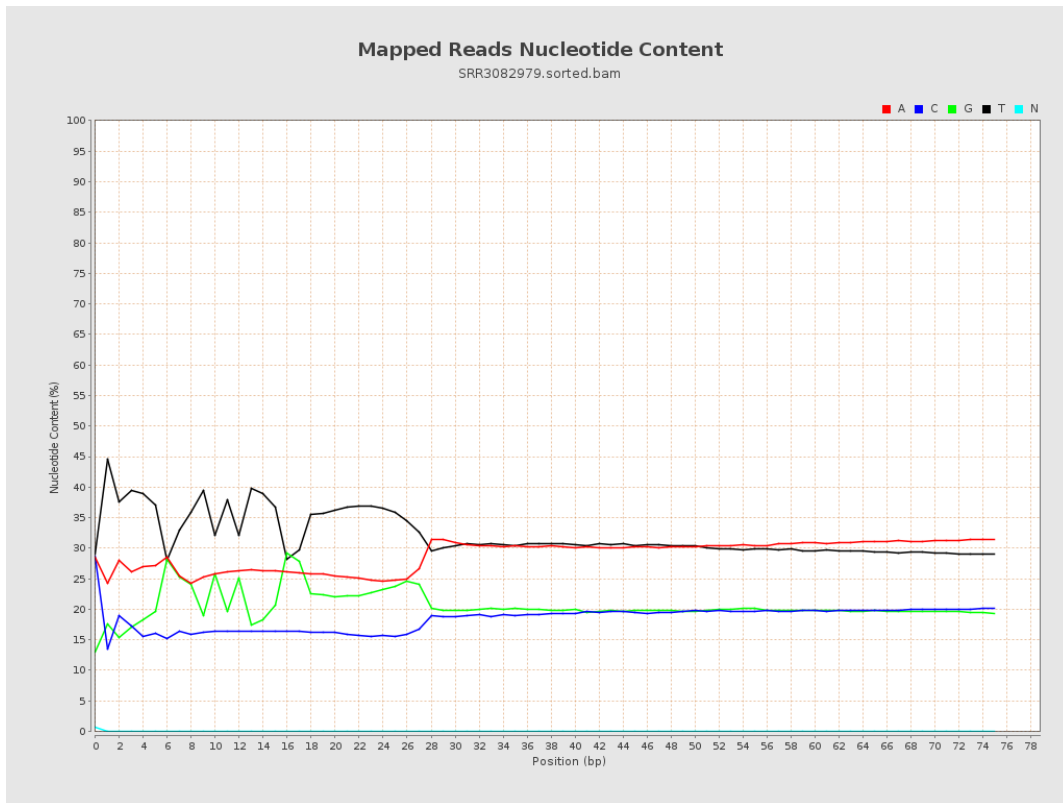
6. Results : Genome Fraction Coverage



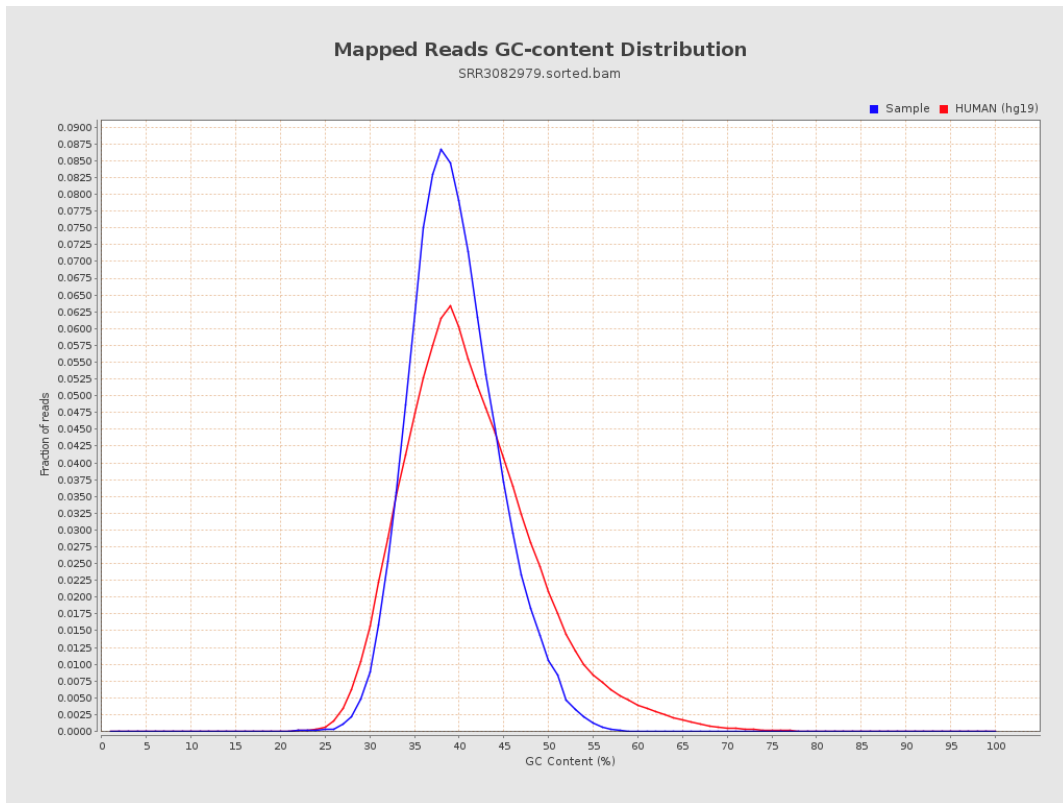
7. Results : Duplication Rate Histogram



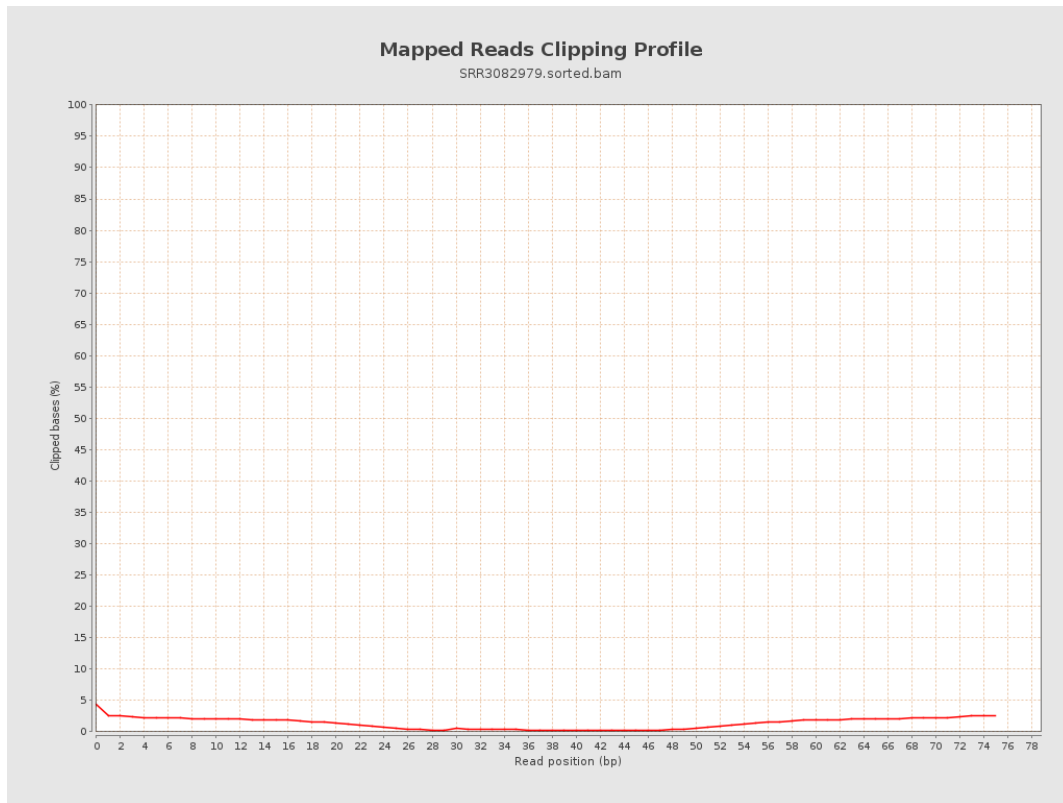
8. Results : Mapped Reads Nucleotide Content



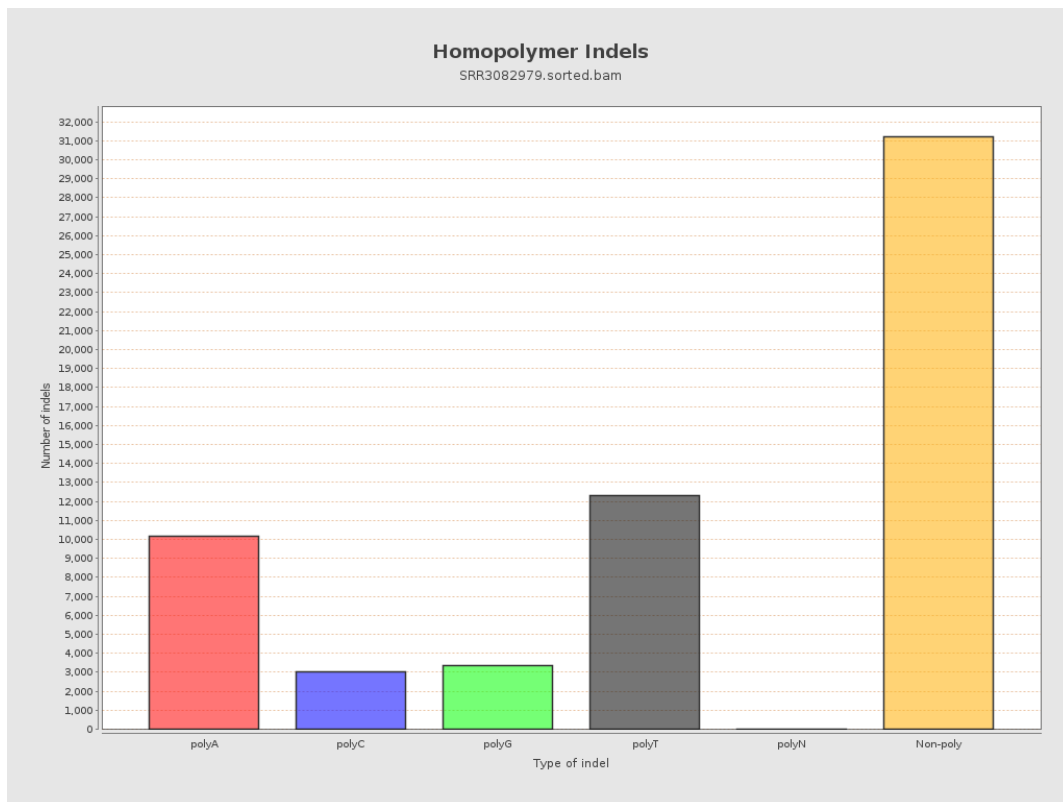
9. Results : Mapped Reads GC-content Distribution



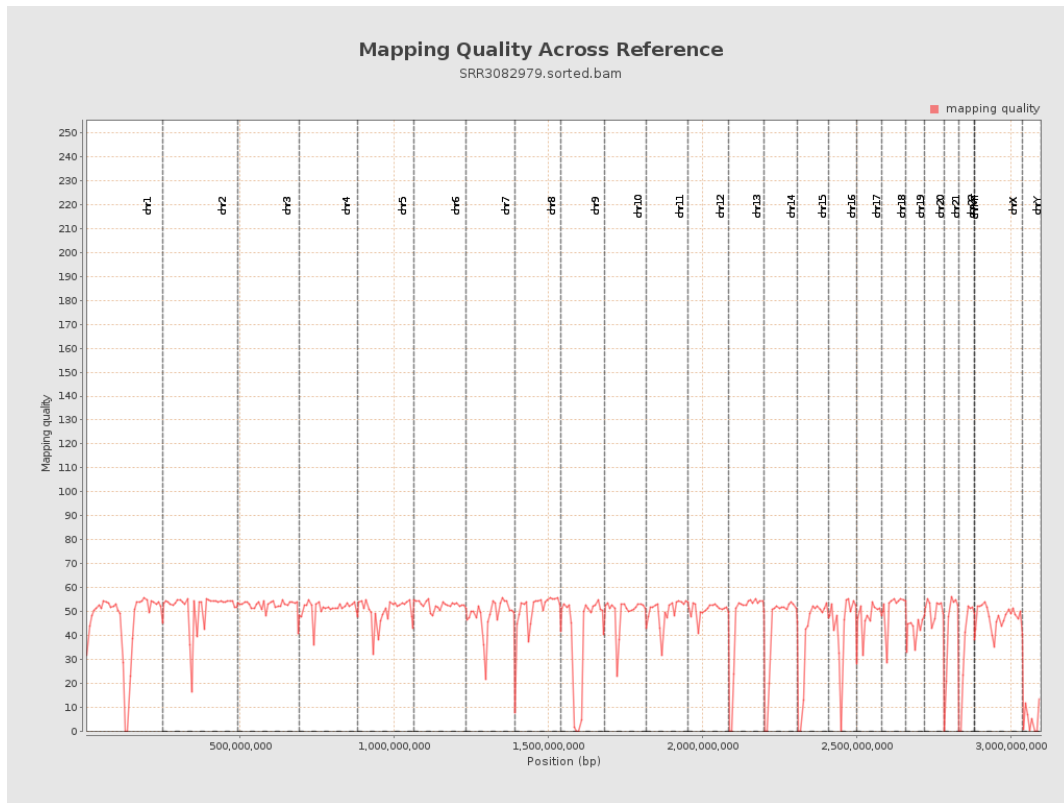
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

