

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/23 14:14:39

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR3080979.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_SRR3080979 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR3080979.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Fri Aug 23 14:14:38 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR3080979.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	4,118,703
Mapped reads	3,453,350 / 83.85%
Unmapped reads	665,353 / 16.15%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	24,105 / 0.59%
Read min/max/mean length	30 / 76 / 76.2
Duplicated reads (estimated)	239,556 / 5.82%
Duplication rate	5.59%
Clipped reads	1,377,324 / 33.44%

2.2. ACGT Content

Number/percentage of A's	65,782,361 / 28.23%
Number/percentage of C's	41,046,431 / 17.61%
Number/percentage of T's	76,188,261 / 32.69%
Number/percentage of G's	49,579,939 / 21.27%
Number/percentage of N's	464,093 / 0.2%
GC Percentage	38.89%

2.3. Coverage

Mean	0.0753

Standard Deviation	0.5847
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2.4. Mapping Quality

Mean Mapping Quality	47.31
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2.5. Mismatches and indels

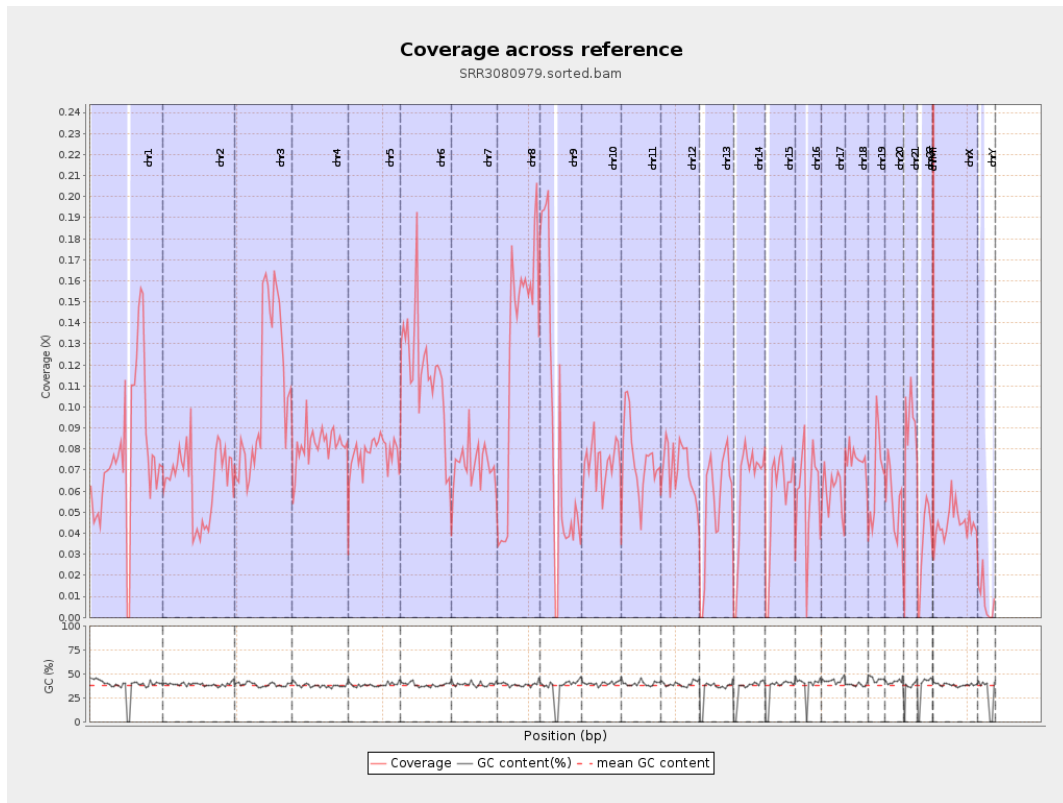
General error rate	0.99%
Mismatches	2,264,848
Insertions	17,474
Mapped reads with at least one insertion	0.5%
Deletions	46,762
Mapped reads with at least one deletion	1.34%
Homopolymer indels	49.27%

2.6. Chromosome stats

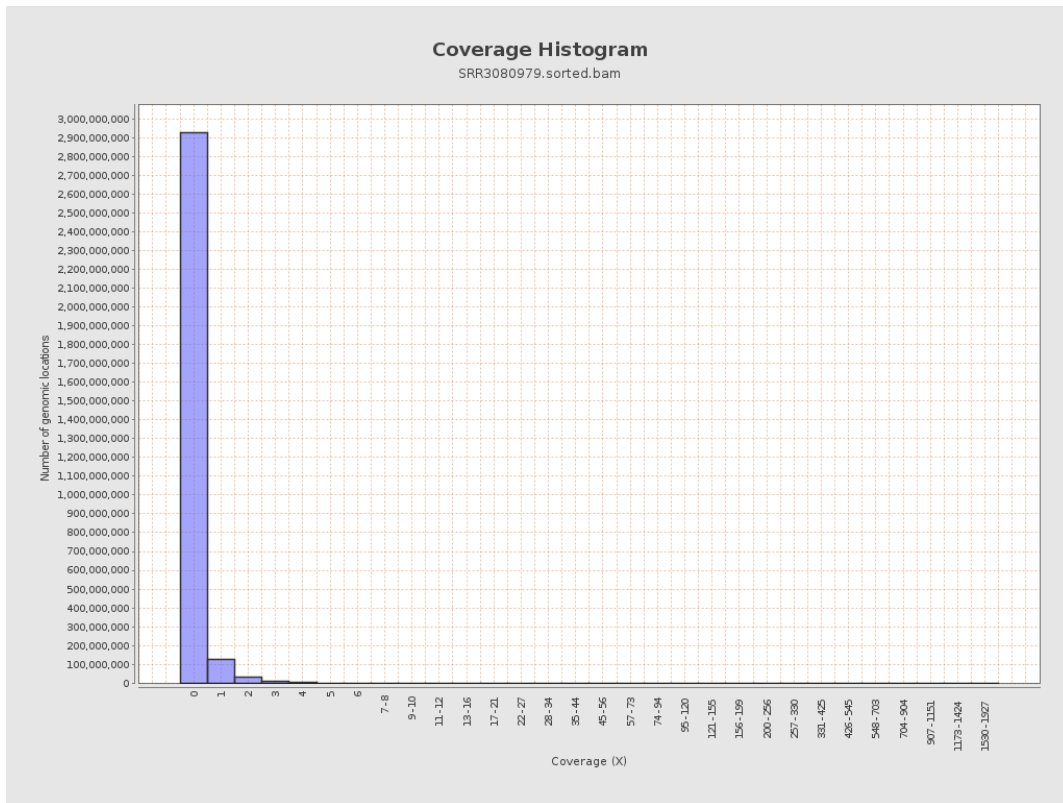
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	19152962	0.0768	0.8903
chr2	243199373	15724274	0.0647	0.488
chr3	198022430	21155376	0.1068	0.4315
chr4	191154276	15689718	0.0821	0.3921
chr5	180915260	14094719	0.0779	0.3699
chr6	171115067	20034601	0.1171	0.6213
chr7	159138663	11478063	0.0721	0.5424

chr8	146364022	18482023	0.1263	1.2517
chr9	141213431	11947119	0.0846	0.7101
chr10	135534747	9954358	0.0734	0.4766
chr11	135006516	10204966	0.0756	0.4399
chr12	133851895	9426401	0.0704	0.3717
chr13	115169878	6246740	0.0542	0.303
chr14	107349540	6658723	0.062	0.4019
chr15	102531392	5835488	0.0569	0.3096
chr16	90354753	5250460	0.0581	0.3768
chr17	81195210	4906505	0.0604	0.327
chr18	78077248	5869223	0.0752	1.1228
chr19	59128983	3981835	0.0673	0.6633
chr20	63025520	3515154	0.0558	0.3324
chr21	48129895	4152988	0.0863	0.4466
chr22	51304566	1833426	0.0357	0.2406
chrMT	16571	116221	7.0135	4.7825
chrX	155270560	6933611	0.0447	0.3145
chrY	59373566	498725	0.0084	0.1704

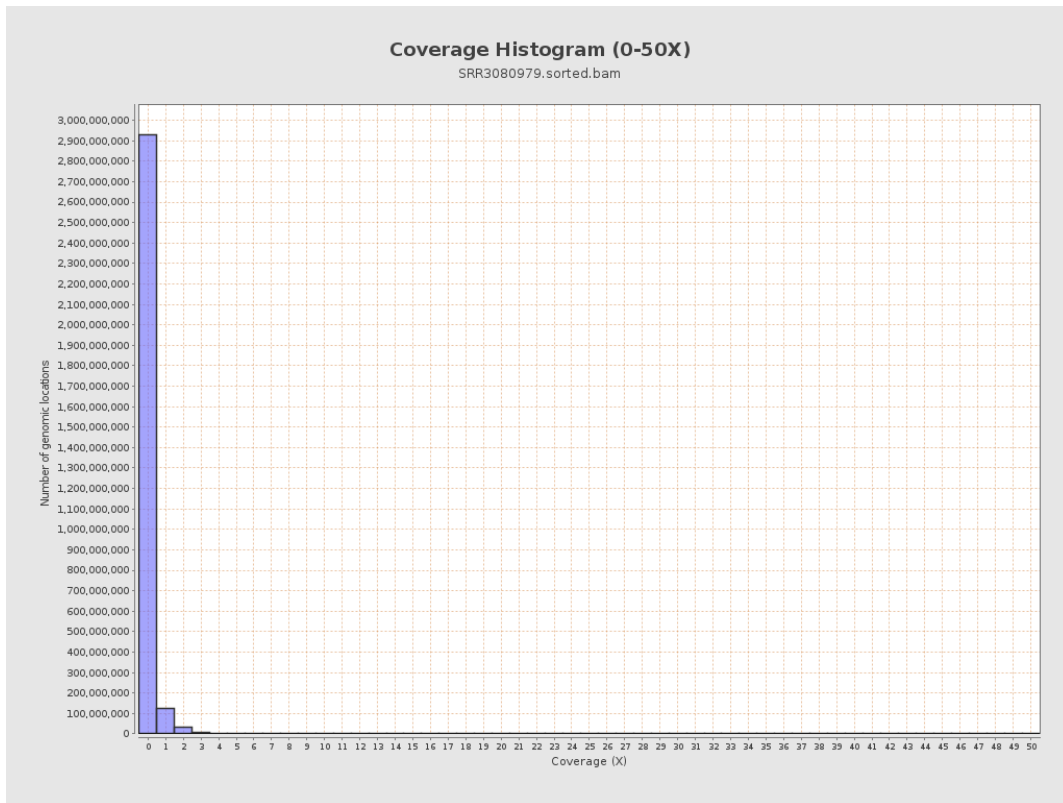
3. Results : Coverage across reference



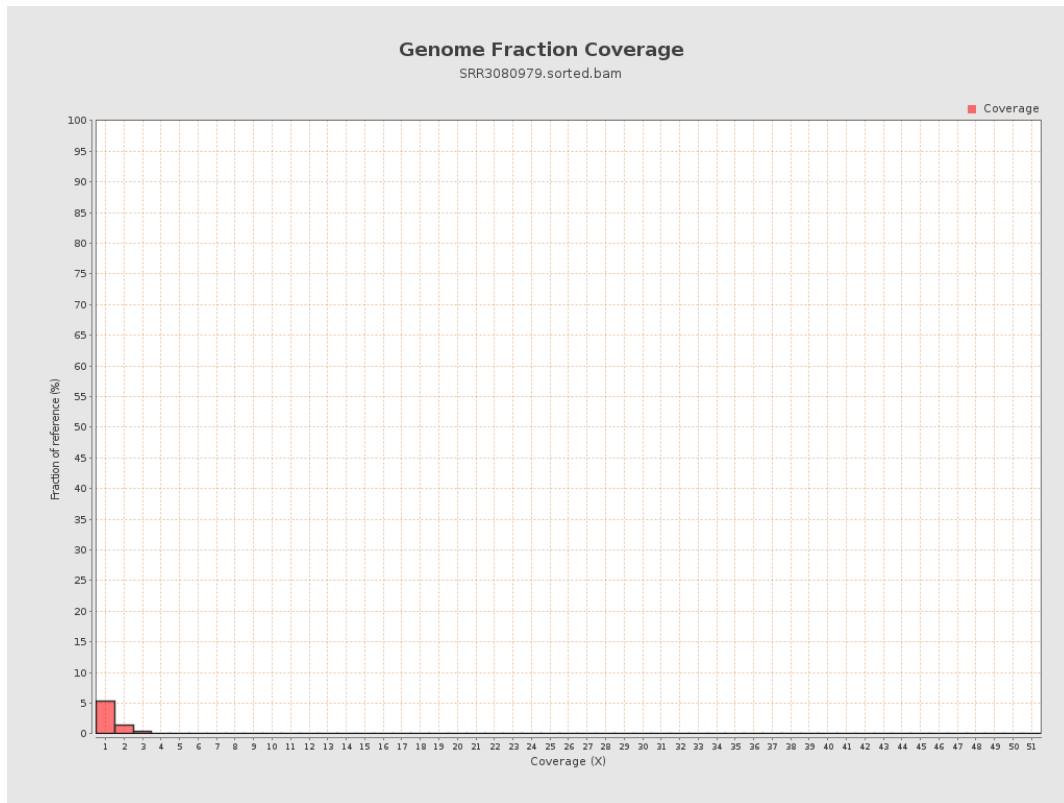
4. Results : Coverage Histogram



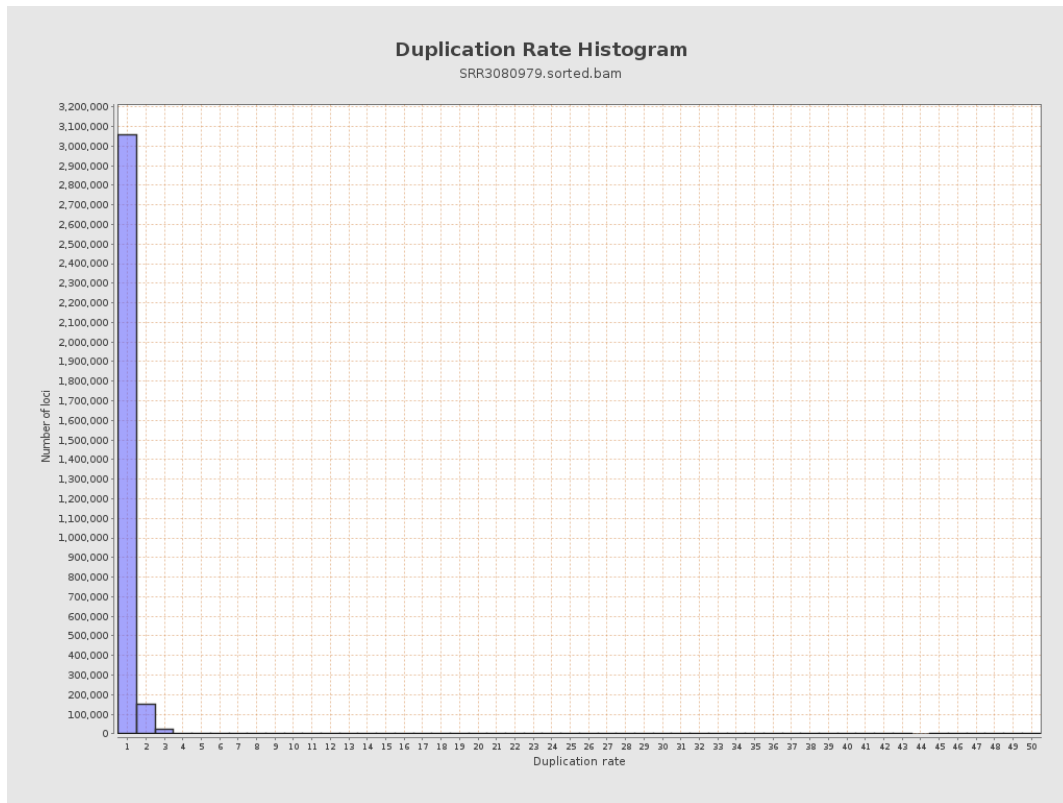
5. Results : Coverage Histogram (0-50X)



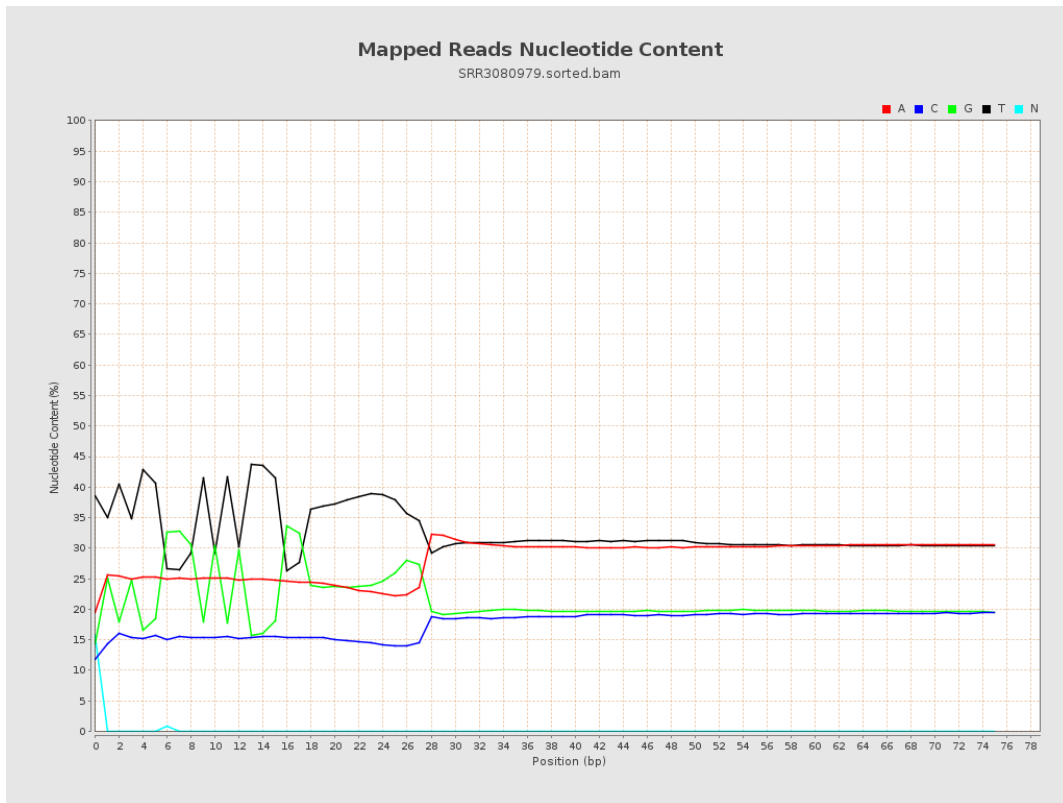
6. Results : Genome Fraction Coverage



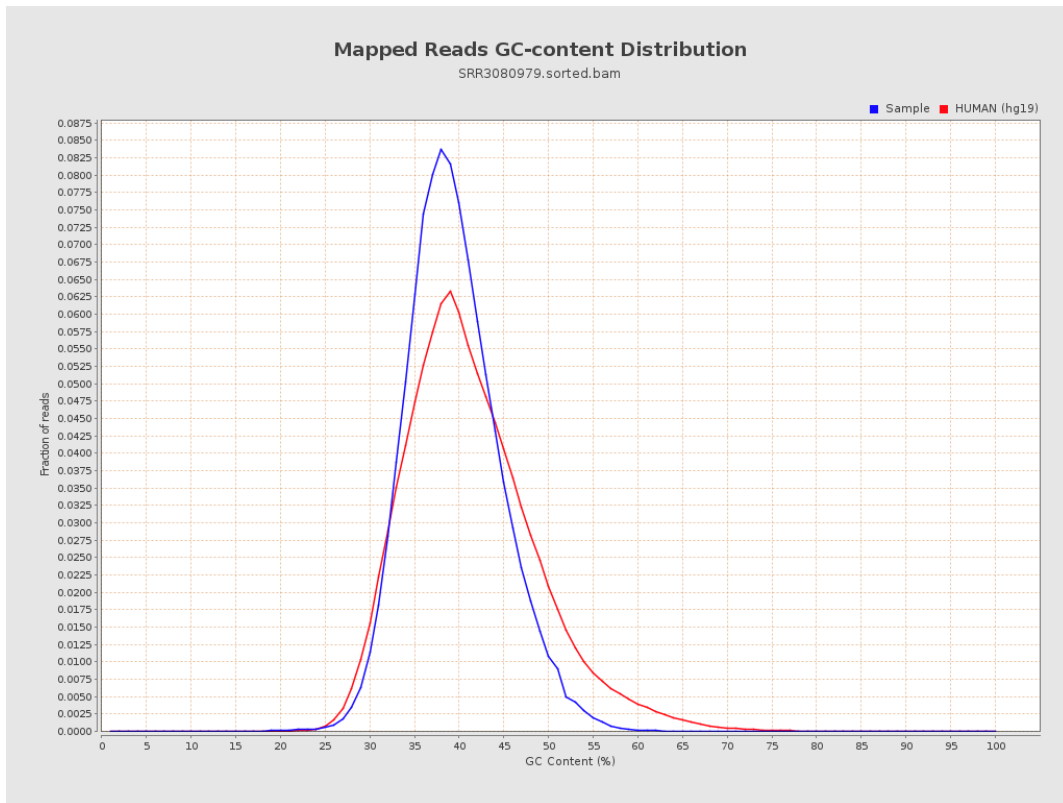
7. Results : Duplication Rate Histogram



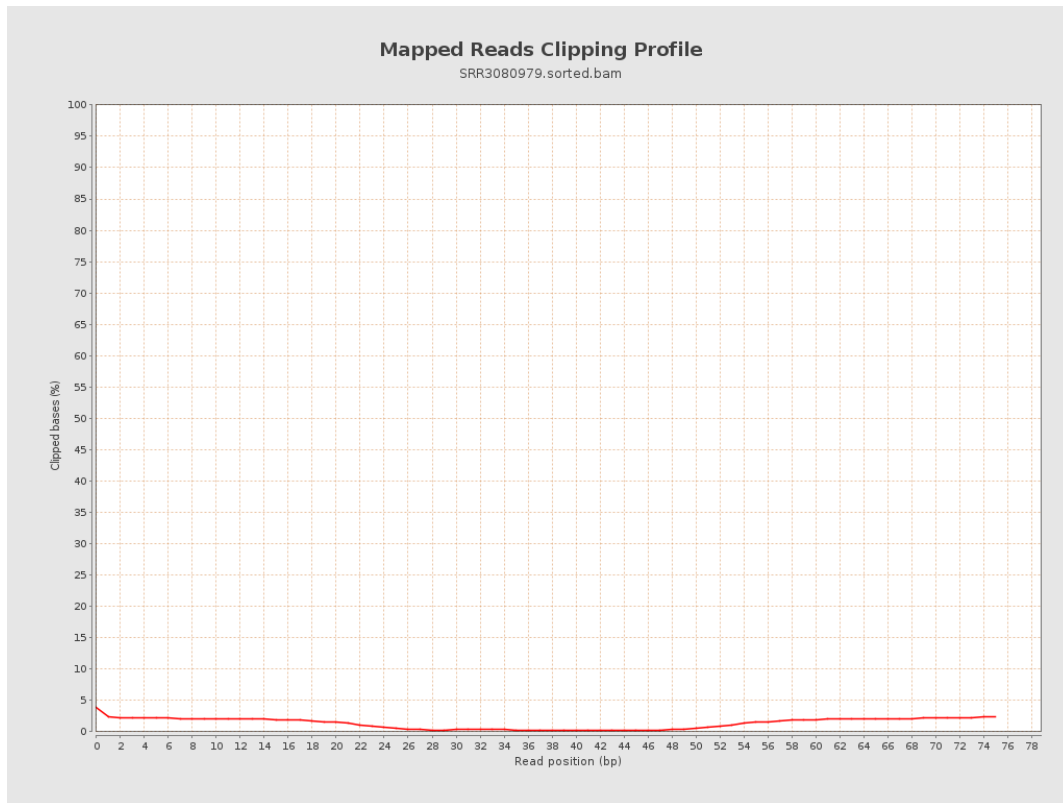
8. Results : Mapped Reads Nucleotide Content



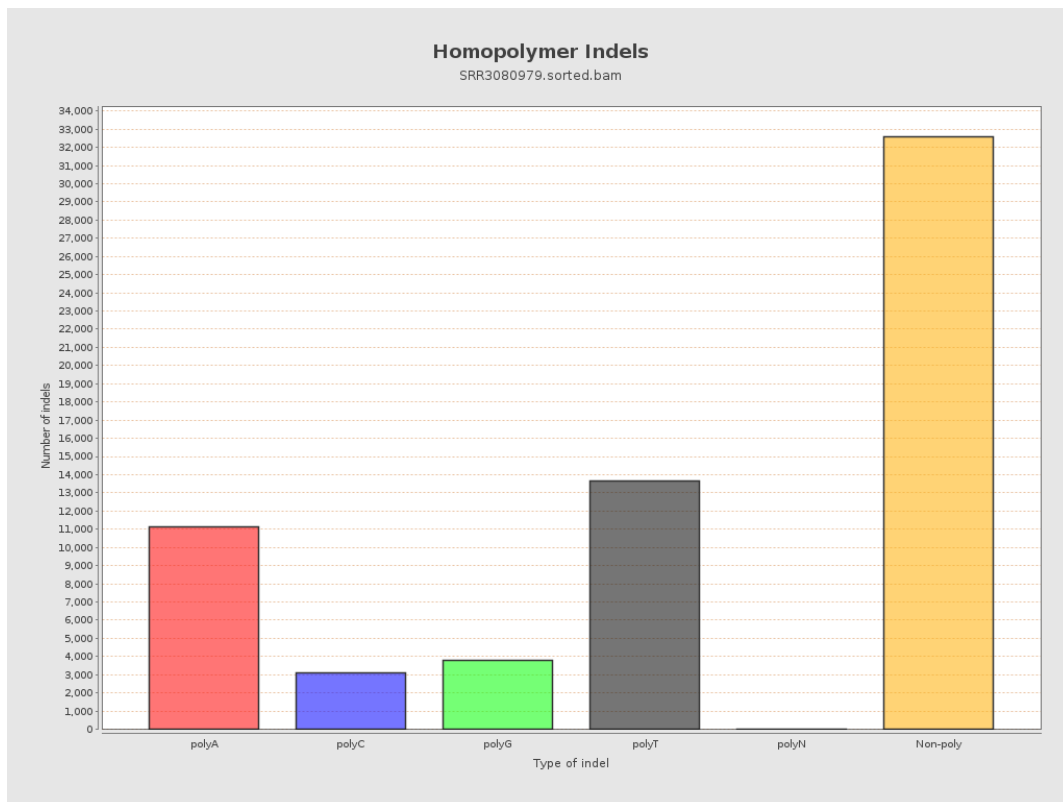
9. Results : Mapped Reads GC-content Distribution



10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

