

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/13 19:42:45

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6003995.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_SRR6003995 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6003995.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Fri Sep 13 19:42:44 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6003995.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,357,109
Mapped reads	3,030,829 / 90.28%
Unmapped reads	326,280 / 9.72%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	22,457 / 0.67%
Read min/max/mean length	30 / 76 / 76.23
Duplicated reads (estimated)	106,848 / 3.18%
Duplication rate	2.36%
Clipped reads	1,535,021 / 45.72%

2.2. ACGT Content

Number/percentage of A's	53,429,343 / 27.02%
Number/percentage of C's	38,532,442 / 19.49%
Number/percentage of T's	59,171,822 / 29.93%
Number/percentage of G's	46,551,650 / 23.54%
Number/percentage of N's	39,292 / 0.02%
GC Percentage	43.03%

2.3. Coverage

Mean	0.0639

Standard Deviation	0.546
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2.4. Mapping Quality

Mean Mapping Quality	41.97
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2.5. Mismatches and indels

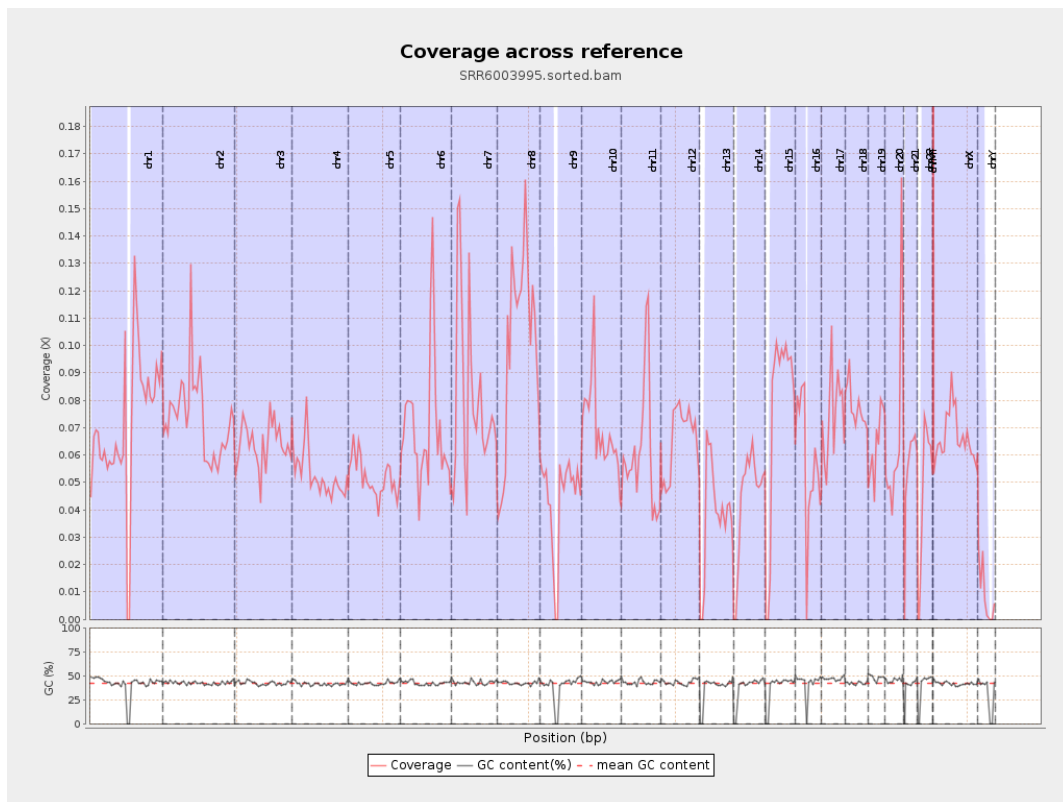
General error rate	0.85%
Mismatches	1,646,780
Insertions	16,207
Mapped reads with at least one insertion	0.53%
Deletions	49,581
Mapped reads with at least one deletion	1.62%
Homopolymer indels	44.73%

2.6. Chromosome stats

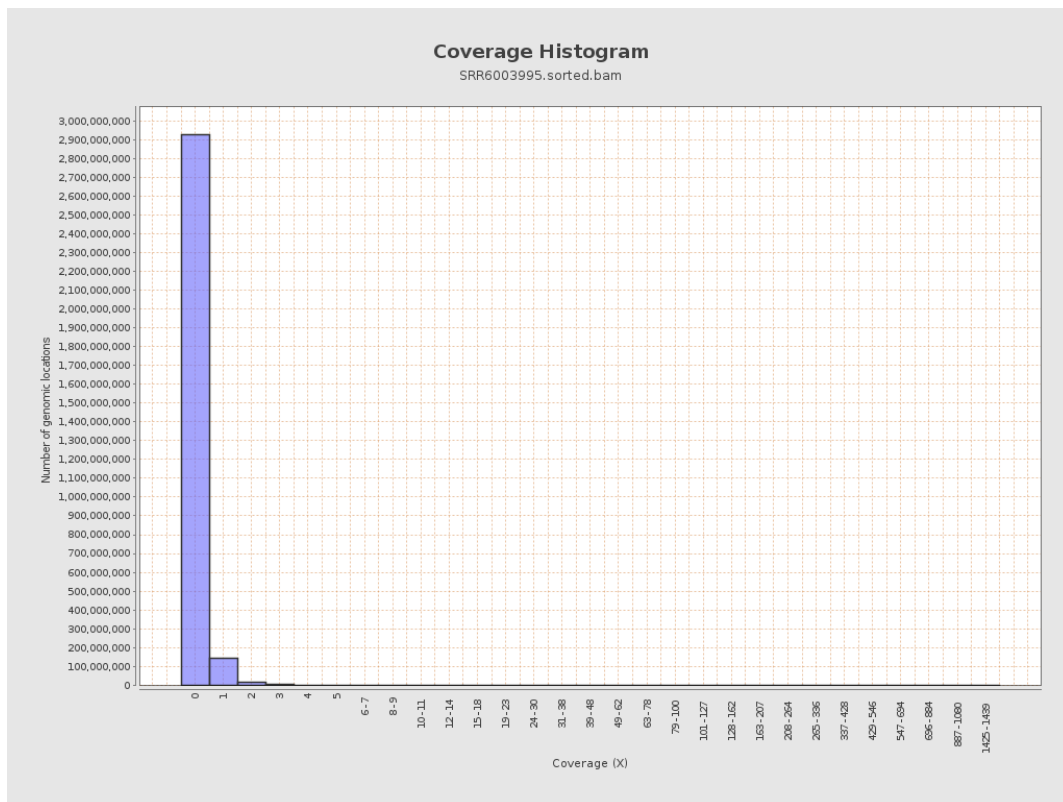
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	17793944	0.0714	0.9229
chr2	243199373	17810630	0.0732	0.8068
chr3	198022430	12720987	0.0642	0.295
chr4	191154276	10094604	0.0528	0.2933
chr5	180915260	9360386	0.0517	0.2605
chr6	171115067	11849366	0.0692	0.3633
chr7	159138663	12695415	0.0798	0.9699

chr8	146364022	14709616	0.1005	0.5261
chr9	141213431	6186895	0.0438	0.4498
chr10	135534747	9445349	0.0697	0.5556
chr11	135006516	8180405	0.0606	0.4163
chr12	133851895	8810174	0.0658	0.296
chr13	115169878	4413336	0.0383	0.2265
chr14	107349540	4789843	0.0446	0.2897
chr15	102531392	7697332	0.0751	0.3358
chr16	90354753	5132025	0.0568	0.338
chr17	81195210	6058885	0.0746	0.345
chr18	78077248	6078121	0.0778	0.9805
chr19	59128983	3764275	0.0637	0.7271
chr20	63025520	4328559	0.0687	0.3193
chr21	48129895	2603177	0.0541	0.2906
chr22	51304566	2421629	0.0472	0.2499
chrMT	16571	12636	0.7625	1.1462
chrX	155270560	10322324	0.0665	0.3488
chrY	59373566	526831	0.0089	0.1793

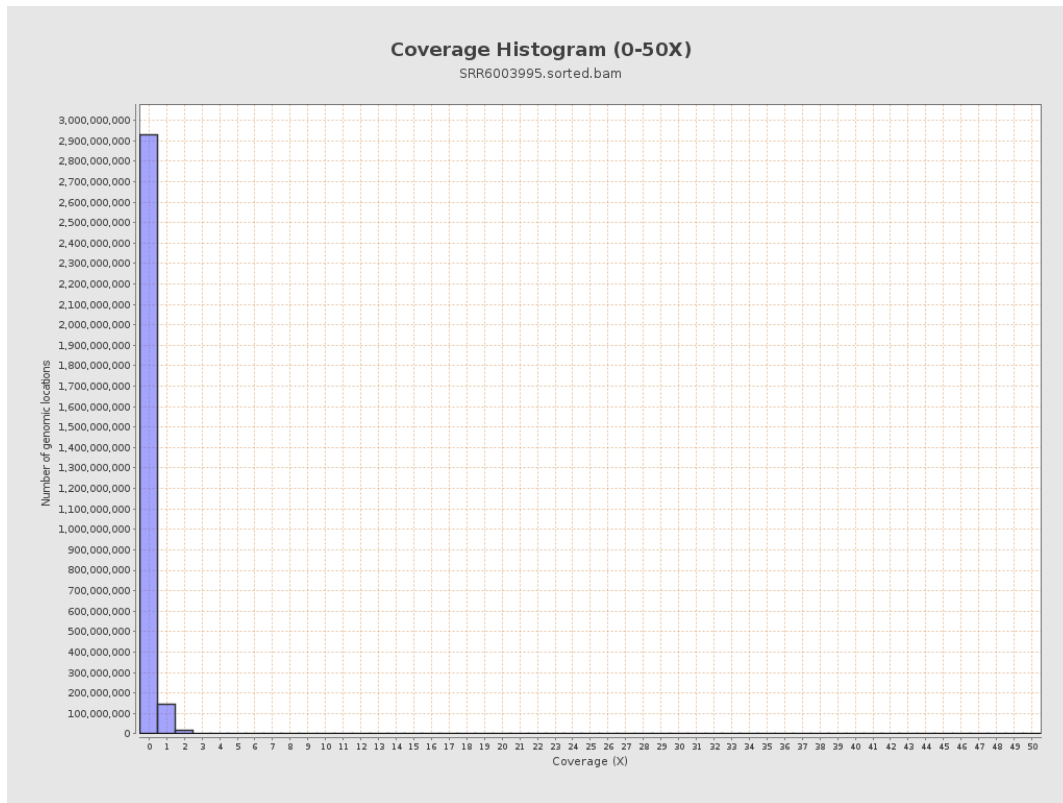
3. Results : Coverage across reference



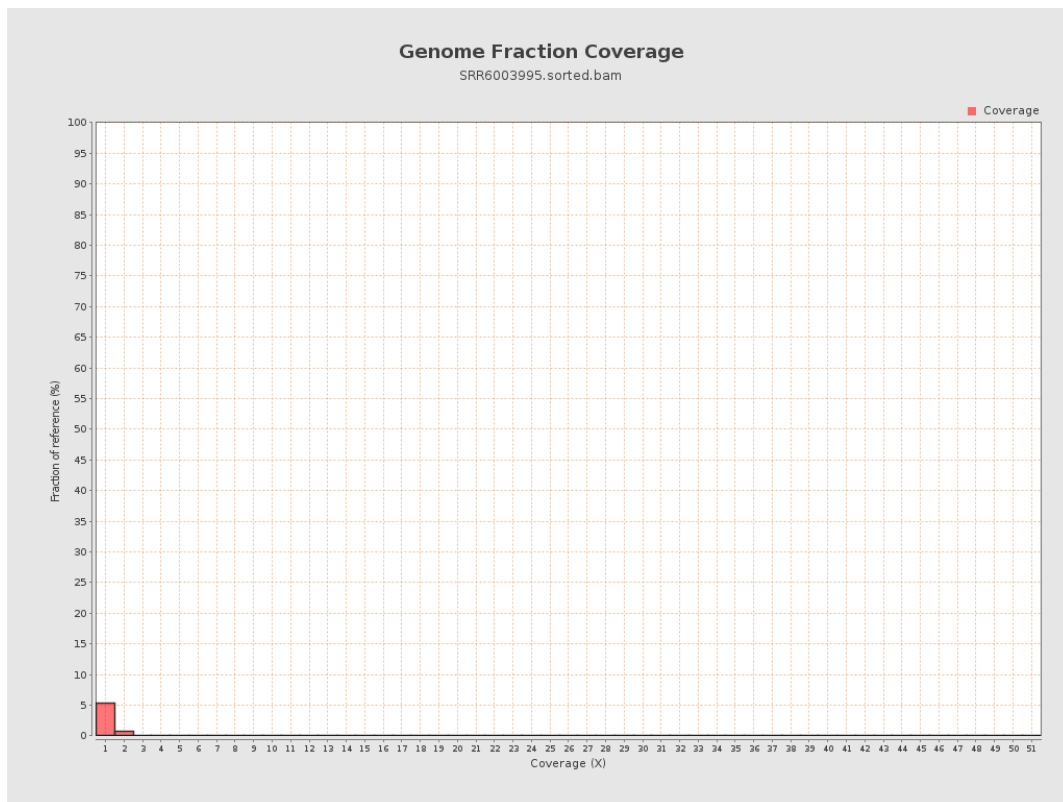
4. Results : Coverage Histogram



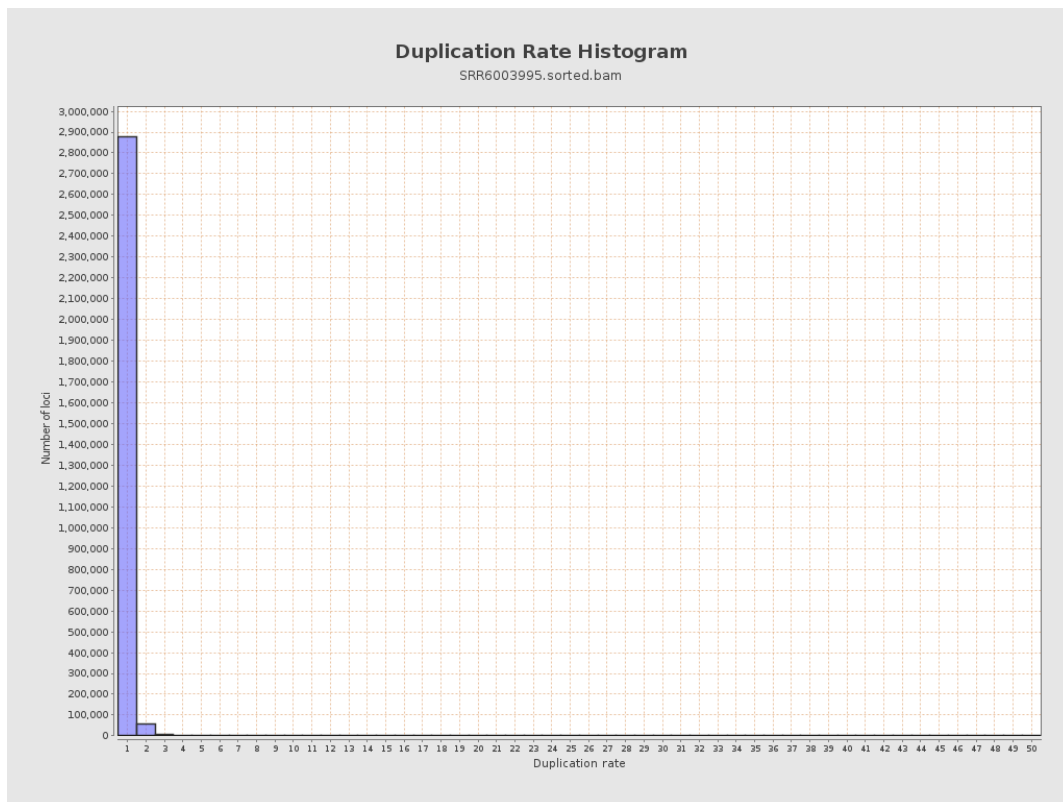
5. Results : Coverage Histogram (0-50X)



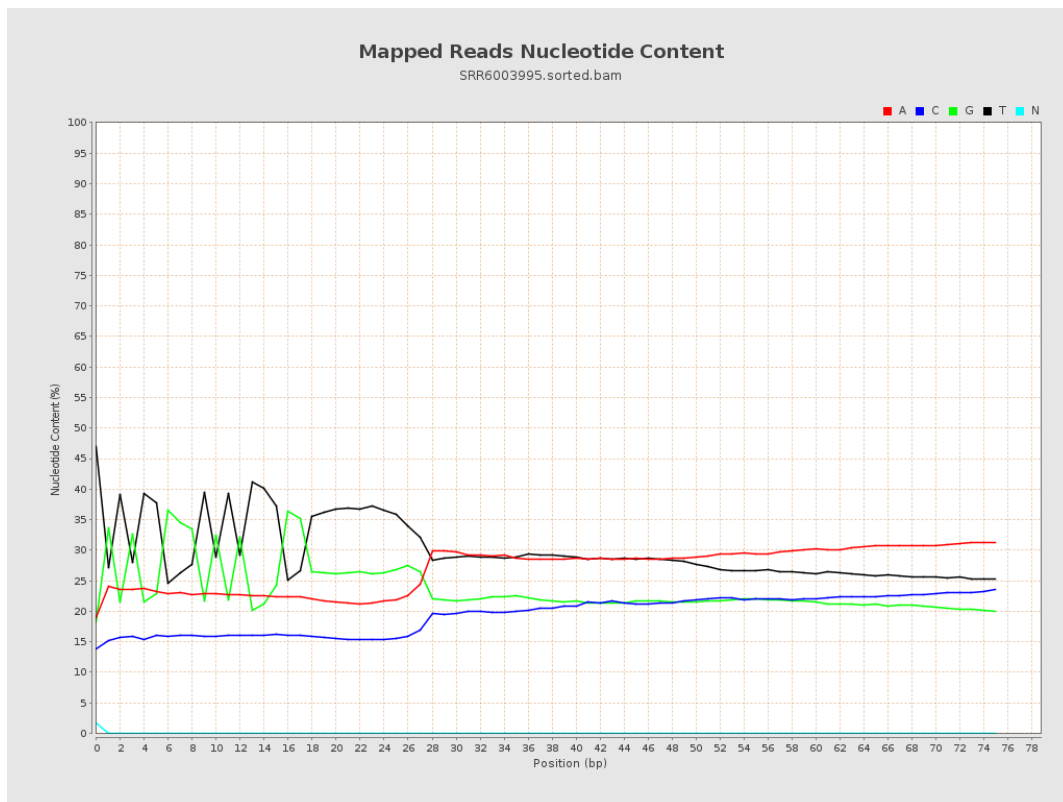
6. Results : Genome Fraction Coverage



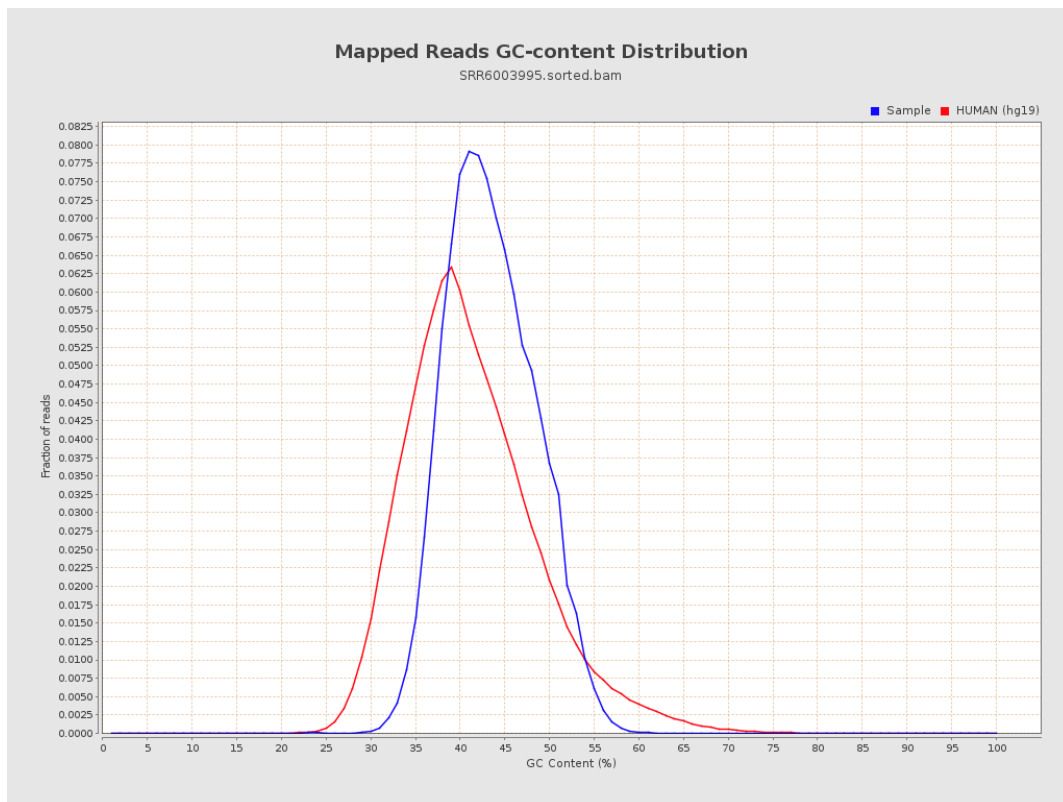
7. Results : Duplication Rate Histogram



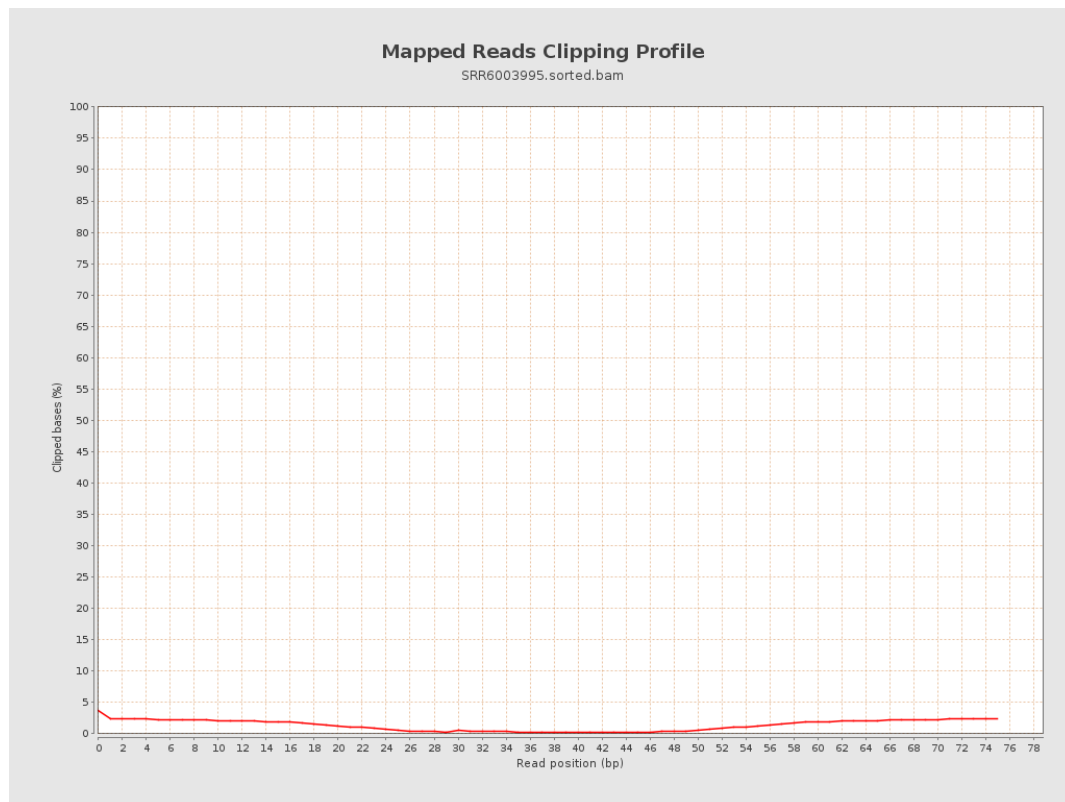
8. Results : Mapped Reads Nucleotide Content



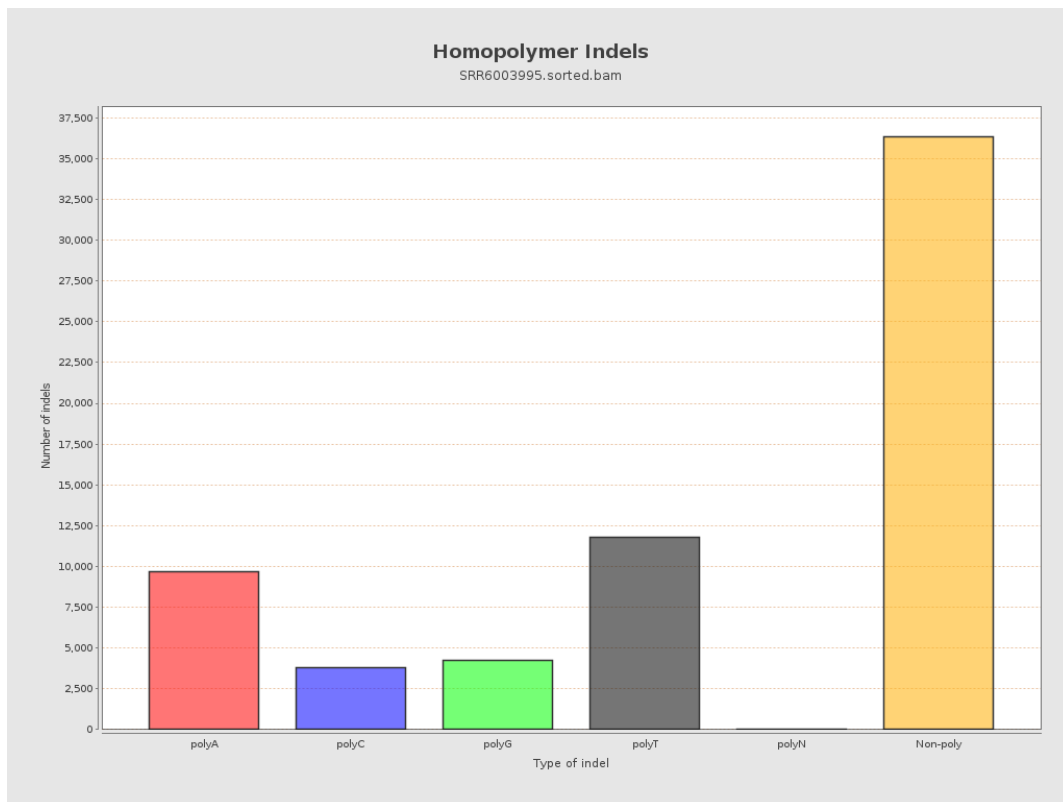
9. Results : Mapped Reads GC-content Distribution



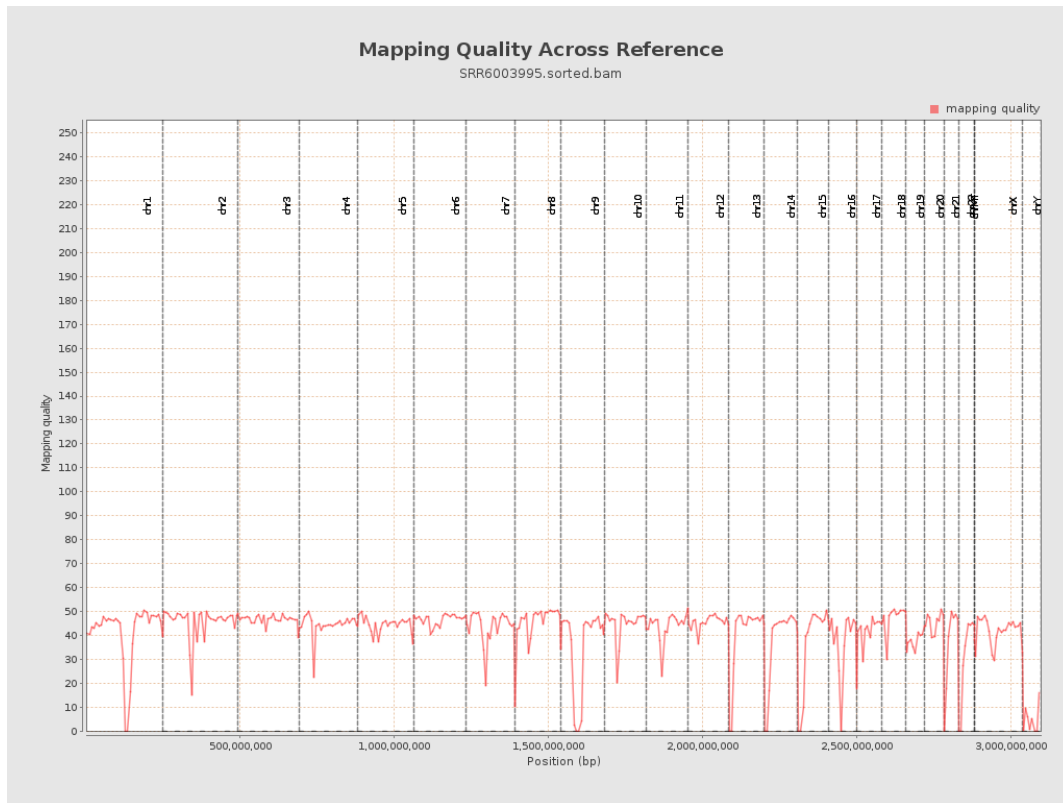
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

