

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/27 23:17:32

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR10524334.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_SRR10524334 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR10524334.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Aug 27 23:17:31 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR10524334.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,309,231
Mapped reads	3,044,021 / 91.99%
Unmapped reads	265,210 / 8.01%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	14,633 / 0.44%
Read min/max/mean length	30 / 76 / 76.15
Duplicated reads (estimated)	156,149 / 4.72%
Duplication rate	3.71%
Clipped reads	3,050,648 / 92.19%

2.2. ACGT Content

Number/percentage of A's	43,282,363 / 24.34%
Number/percentage of C's	34,011,498 / 19.13%
Number/percentage of T's	56,479,401 / 31.76%
Number/percentage of G's	44,036,674 / 24.76%
Number/percentage of N's	26,114 / 0.01%
GC Percentage	43.89%

2.3. Coverage

Mean	0.0575

Standard Deviation	0.4726
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2.4. Mapping Quality

Mean Mapping Quality	44.22
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2.5. Mismatches and indels

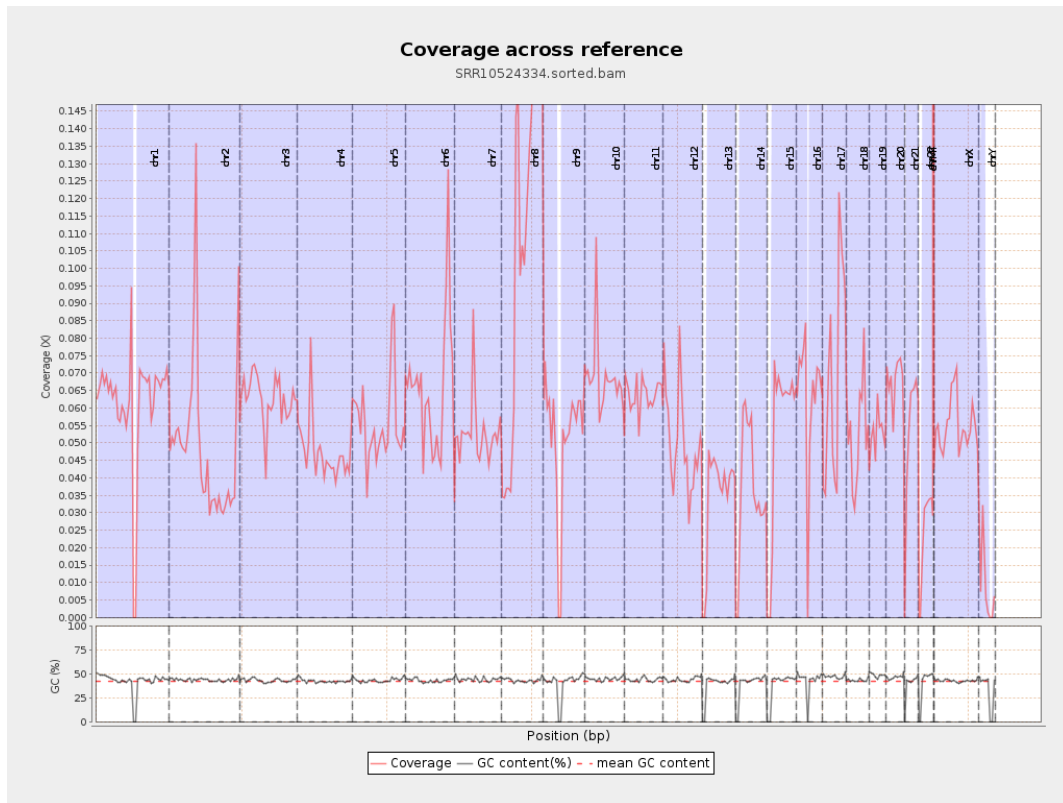
General error rate	0.52%
Mismatches	895,629
Insertions	11,562
Mapped reads with at least one insertion	0.38%
Deletions	29,367
Mapped reads with at least one deletion	0.96%
Homopolymer indels	41.09%

2.6. Chromosome stats

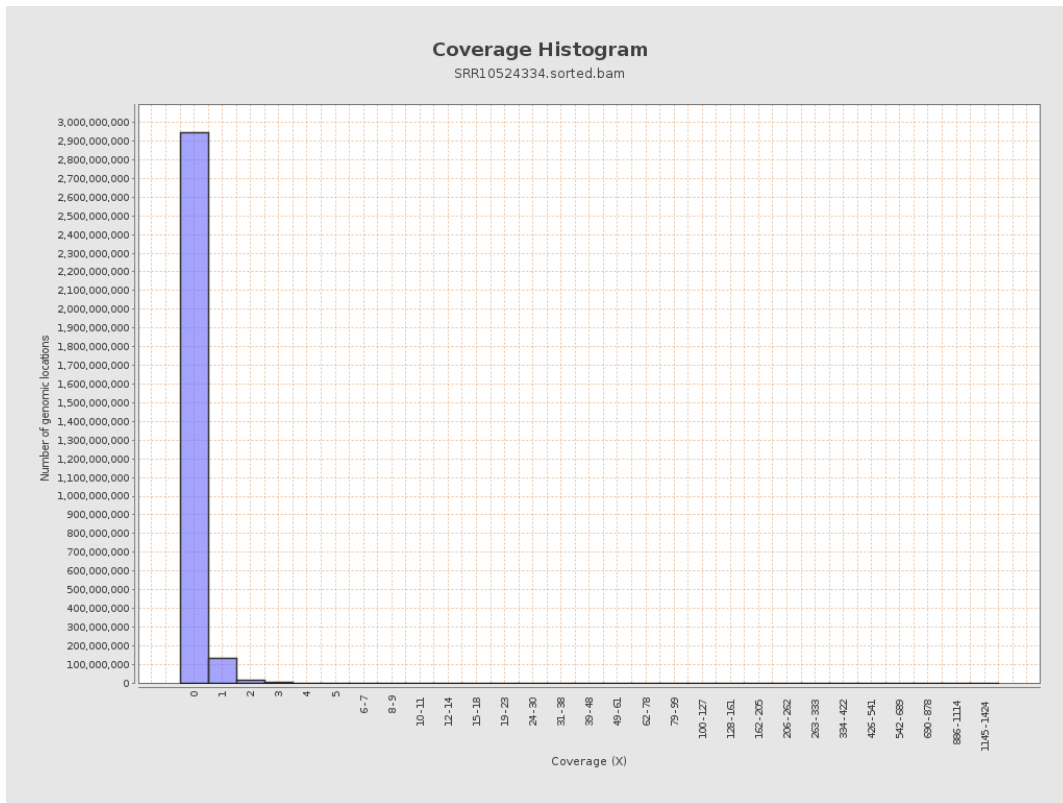
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	15286996	0.0613	0.9052
chr2	243199373	11870013	0.0488	0.6476
chr3	198022430	12403270	0.0626	0.2913
chr4	191154276	9014975	0.0472	0.2991
chr5	180915260	10123421	0.056	0.2736
chr6	171115067	11203876	0.0655	0.332
chr7	159138663	8431515	0.053	0.6031

chr8	146364022	18561085	0.1268	0.5873
chr9	141213431	7165894	0.0507	0.3461
chr10	135534747	9281756	0.0685	0.4642
chr11	135006516	8590576	0.0636	0.4013
chr12	133851895	6727285	0.0503	0.2643
chr13	115169878	3978771	0.0345	0.2153
chr14	107349540	3943406	0.0367	0.2379
chr15	102531392	5441892	0.0531	0.2707
chr16	90354753	5722331	0.0633	0.317
chr17	81195210	5597987	0.0689	0.3466
chr18	78077248	4157032	0.0532	0.6968
chr19	59128983	3136513	0.053	0.6201
chr20	63025520	4256183	0.0675	0.312
chr21	48129895	2546449	0.0529	0.3022
chr22	51304566	1215882	0.0237	0.1802
chrMT	16571	198484	11.9778	7.172
chrX	155270560	8501262	0.0548	0.3081
chrY	59373566	526154	0.0089	0.249

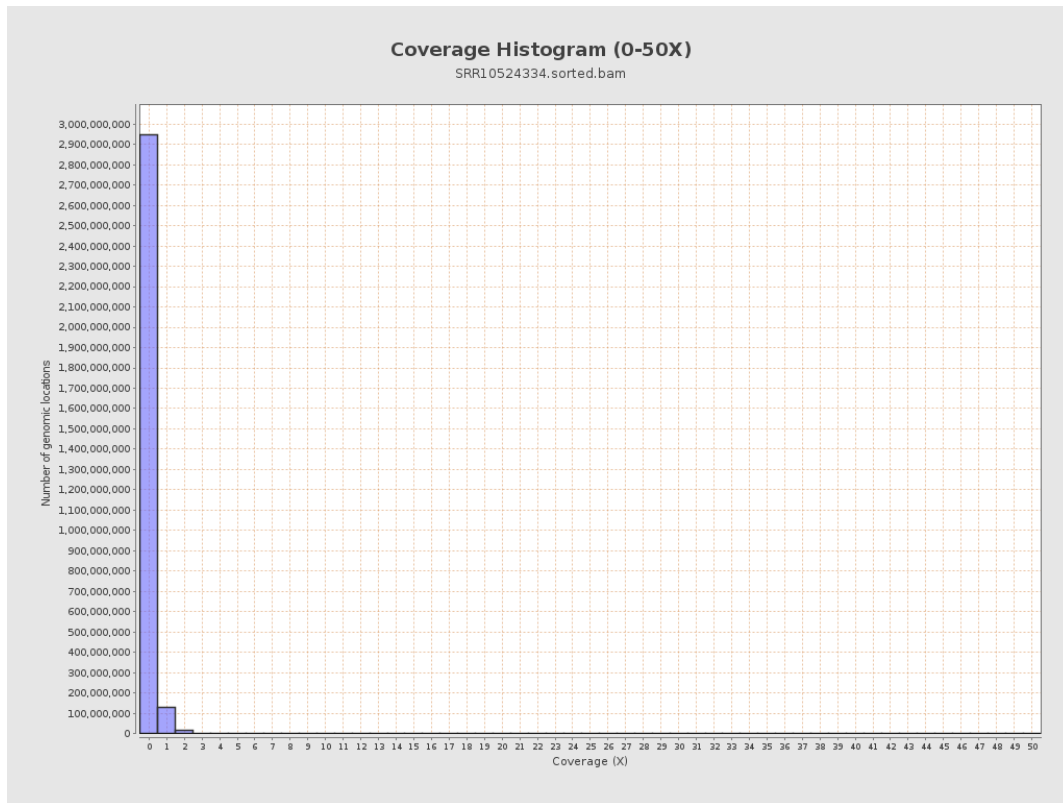
3. Results : Coverage across reference



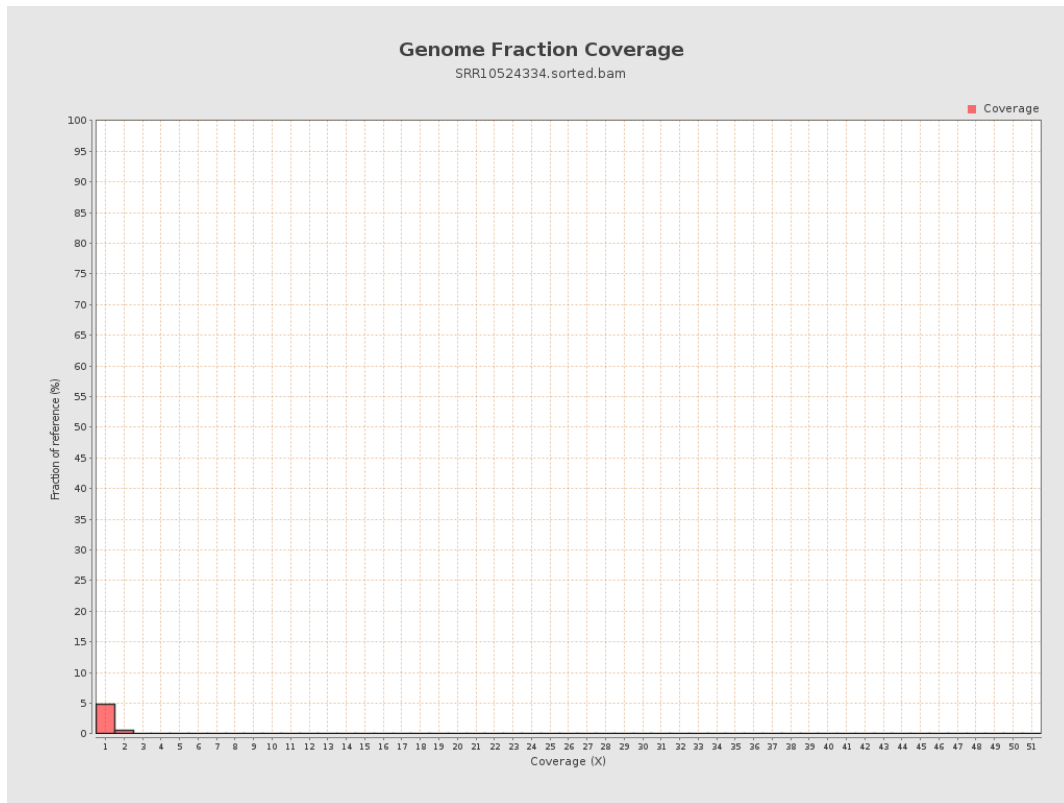
4. Results : Coverage Histogram



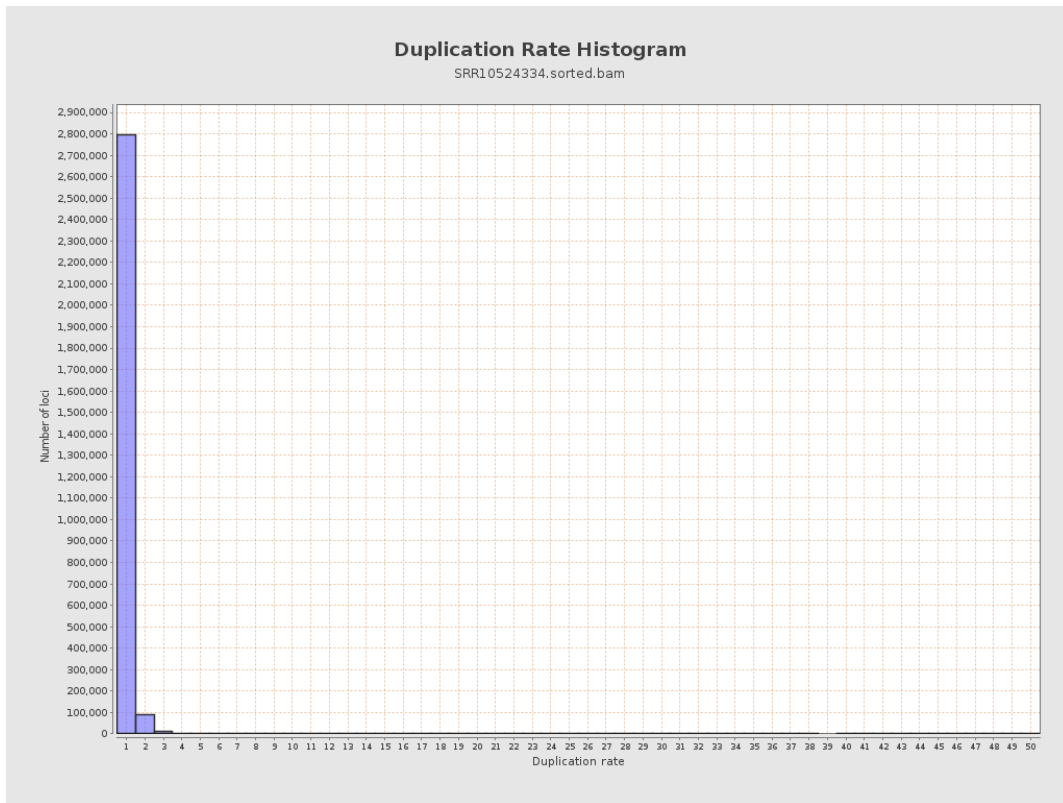
5. Results : Coverage Histogram (0-50X)



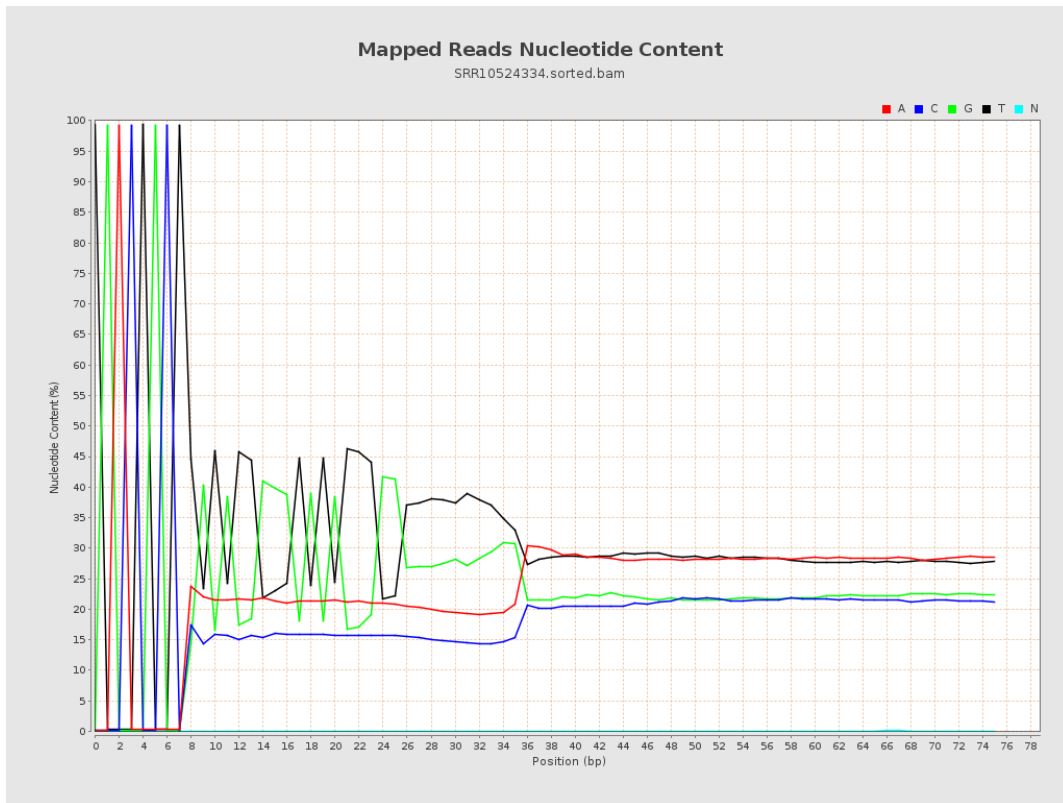
6. Results : Genome Fraction Coverage



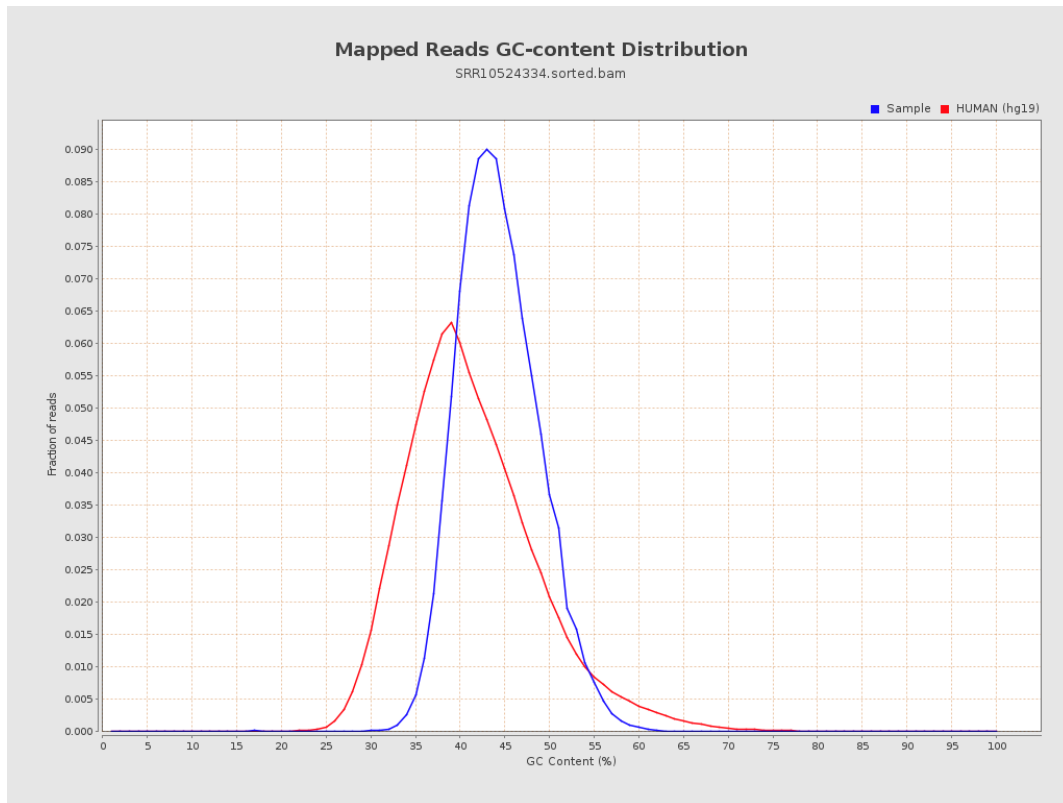
7. Results : Duplication Rate Histogram



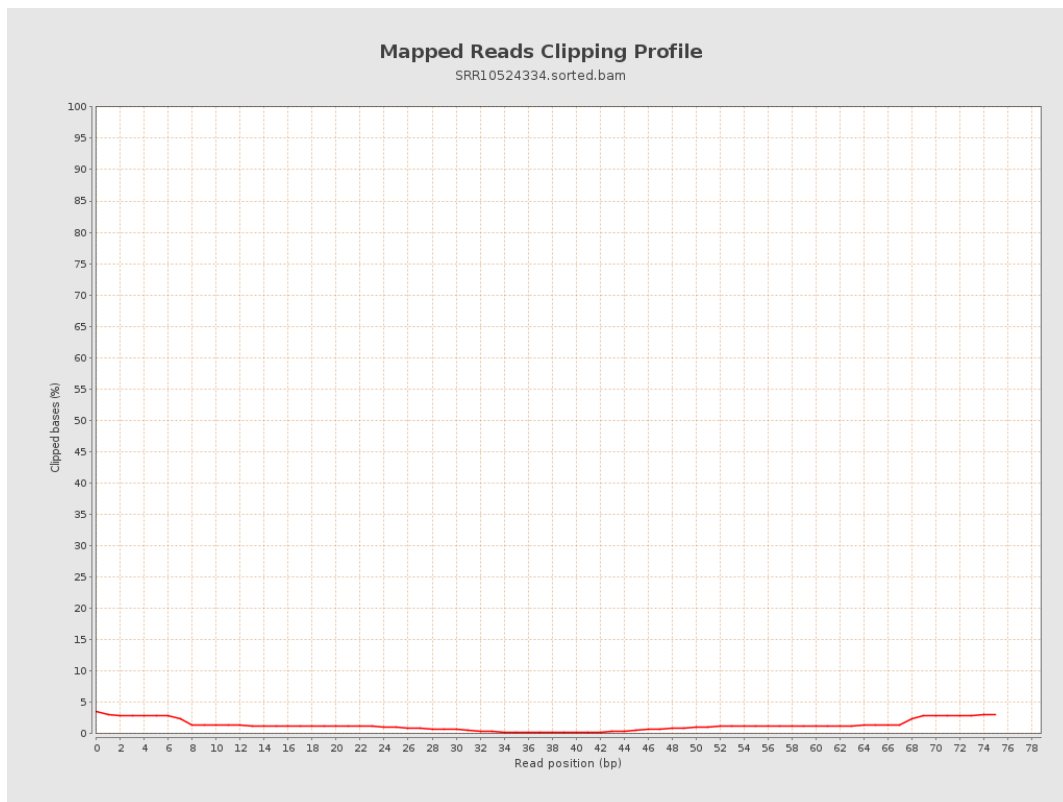
8. Results : Mapped Reads Nucleotide Content



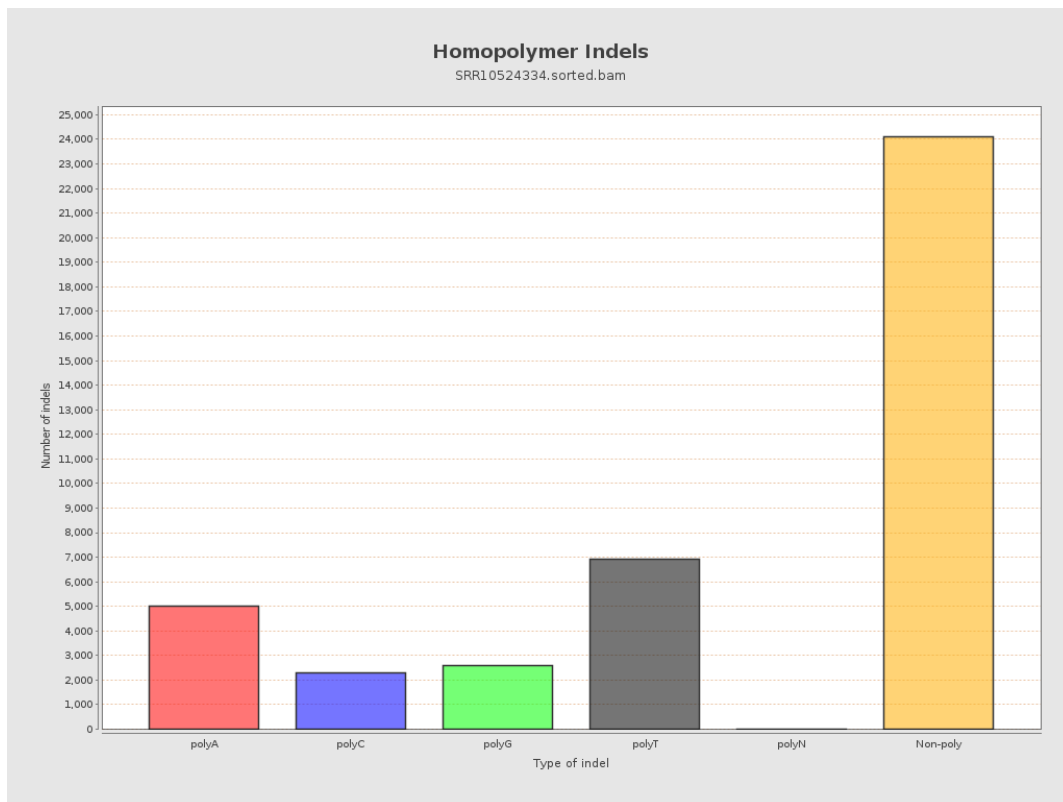
9. Results : Mapped Reads GC-content Distribution



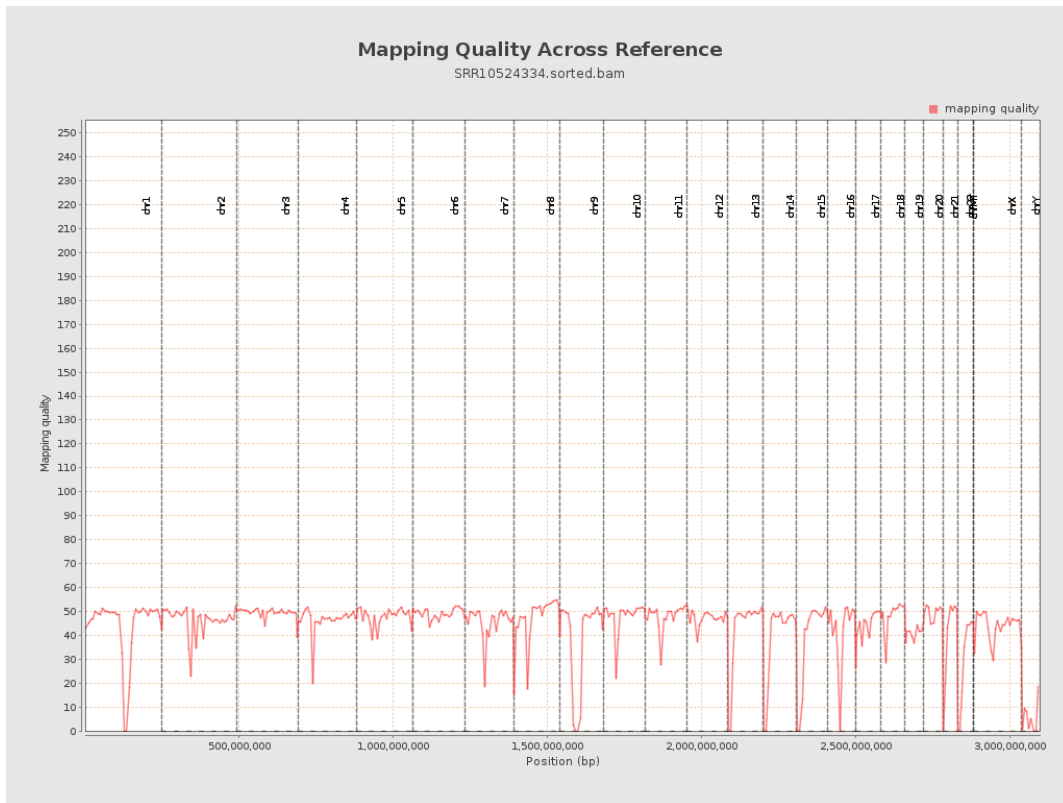
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

