

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/02 08:41:28

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR9716350.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_SRR9716350 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR9716350.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Sep 02 08:41:27 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR9716350.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	478,504
Mapped reads	432,488 / 90.38%
Unmapped reads	46,016 / 9.62%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	9,039 / 1.89%
Read min/max/mean length	30 / 101 / 101.68
Duplicated reads (estimated)	7,571 / 1.58%
Duplication rate	1.22%
Clipped reads	440,729 / 92.11%

2.2. ACGT Content

Number/percentage of A's	8,841,855 / 26.23%
Number/percentage of C's	6,514,226 / 19.33%
Number/percentage of T's	10,450,979 / 31.01%
Number/percentage of G's	7,893,305 / 23.42%
Number/percentage of N's	2,402 / 0.01%
GC Percentage	42.75%

2.3. Coverage

Mean	0.0109

Standard Deviation	0.137
--------------------	-------

2.4. Mapping Quality

Mean Mapping Quality	47.04
----------------------	-------

2.5. Mismatches and indels

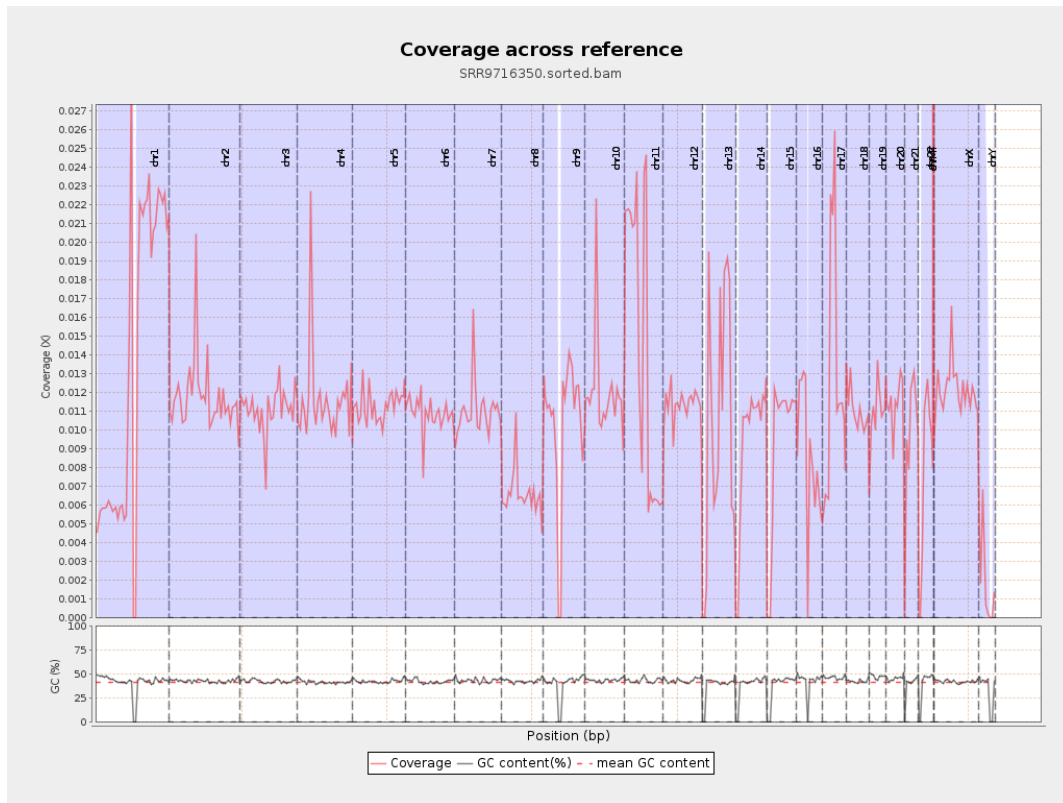
General error rate	0.72%
Mismatches	235,267
Insertions	2,807
Mapped reads with at least one insertion	0.64%
Deletions	7,844
Mapped reads with at least one deletion	1.79%
Homopolymer indels	42.15%

2.6. Chromosome stats

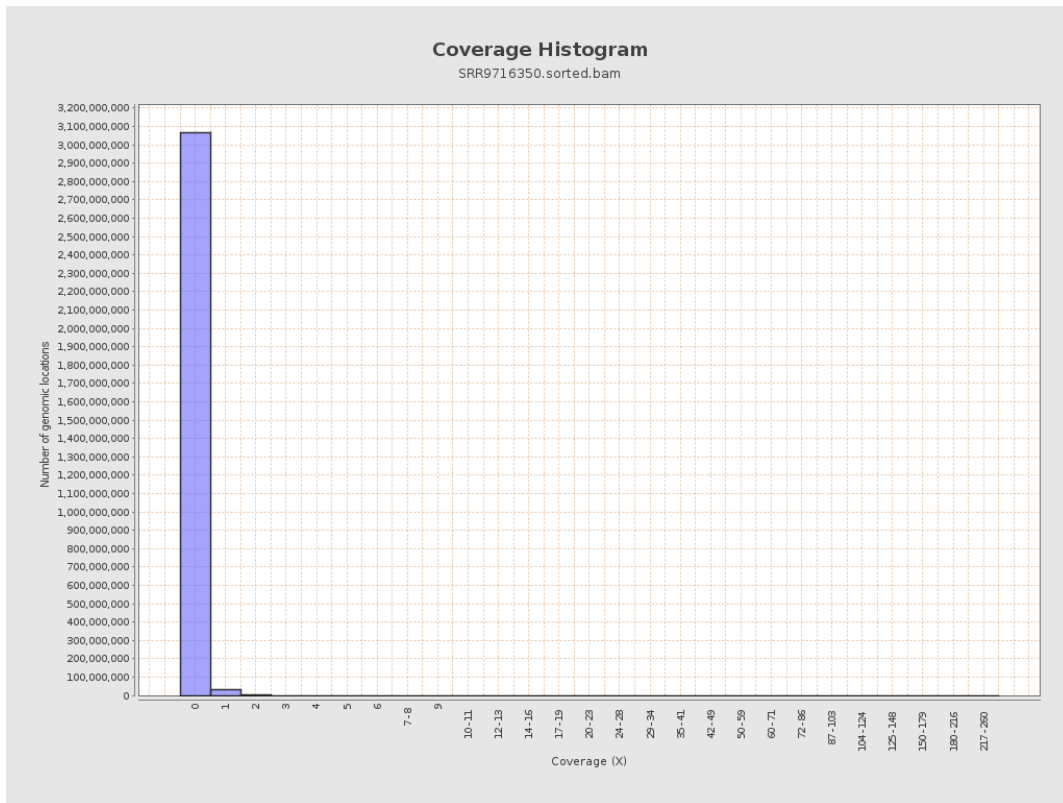
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	3319784	0.0133	0.2462
chr2	243199373	2867170	0.0118	0.1431
chr3	198022430	2196100	0.0111	0.1085
chr4	191154276	2209635	0.0116	0.1222
chr5	180915260	2035408	0.0113	0.1098
chr6	171115067	1859455	0.0109	0.1108
chr7	159138663	1757489	0.011	0.1514

chr8	146364022	960341	0.0066	0.1208
chr9	141213431	1440050	0.0102	0.1208
chr10	135534747	1620803	0.012	0.1424
chr11	135006516	1915338	0.0142	0.1451
chr12	133851895	1515350	0.0113	0.11
chr13	115169878	1173128	0.0102	0.1045
chr14	107349540	994212	0.0093	0.1015
chr15	102531392	955609	0.0093	0.0995
chr16	90354753	771048	0.0085	0.0973
chr17	81195210	1056077	0.013	0.1253
chr18	78077248	862941	0.0111	0.1622
chr19	59128983	656818	0.0111	0.1657
chr20	63025520	708605	0.0112	0.1119
chr21	48129895	466435	0.0097	0.1064
chr22	51304566	385605	0.0075	0.0896
chrMT	16571	3608	0.2177	0.4519
chrX	155270560	1881819	0.0121	0.1201
chrY	59373566	105049	0.0018	0.0749

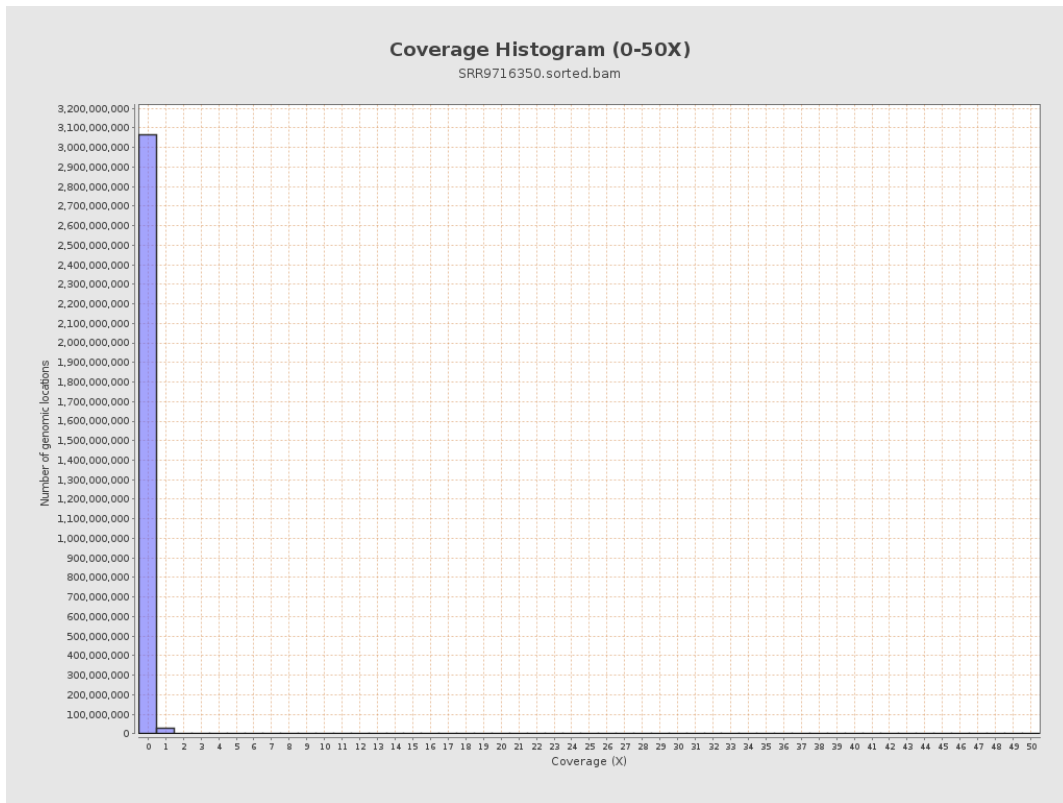
3. Results : Coverage across reference



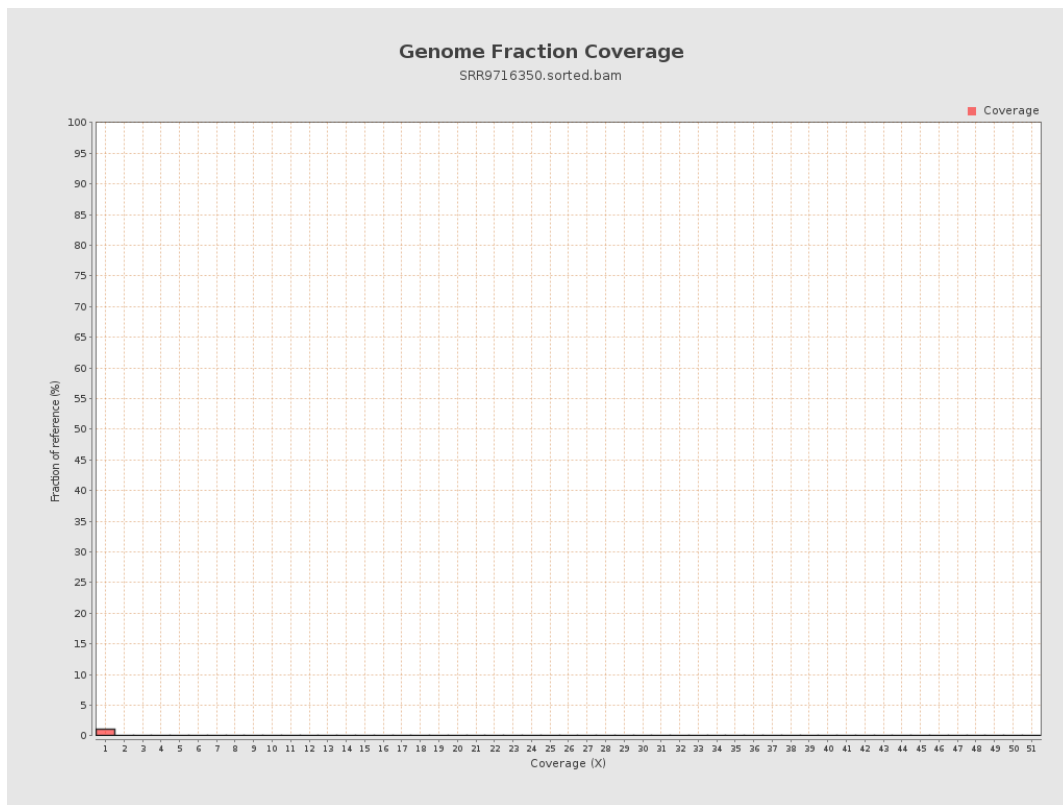
4. Results : Coverage Histogram



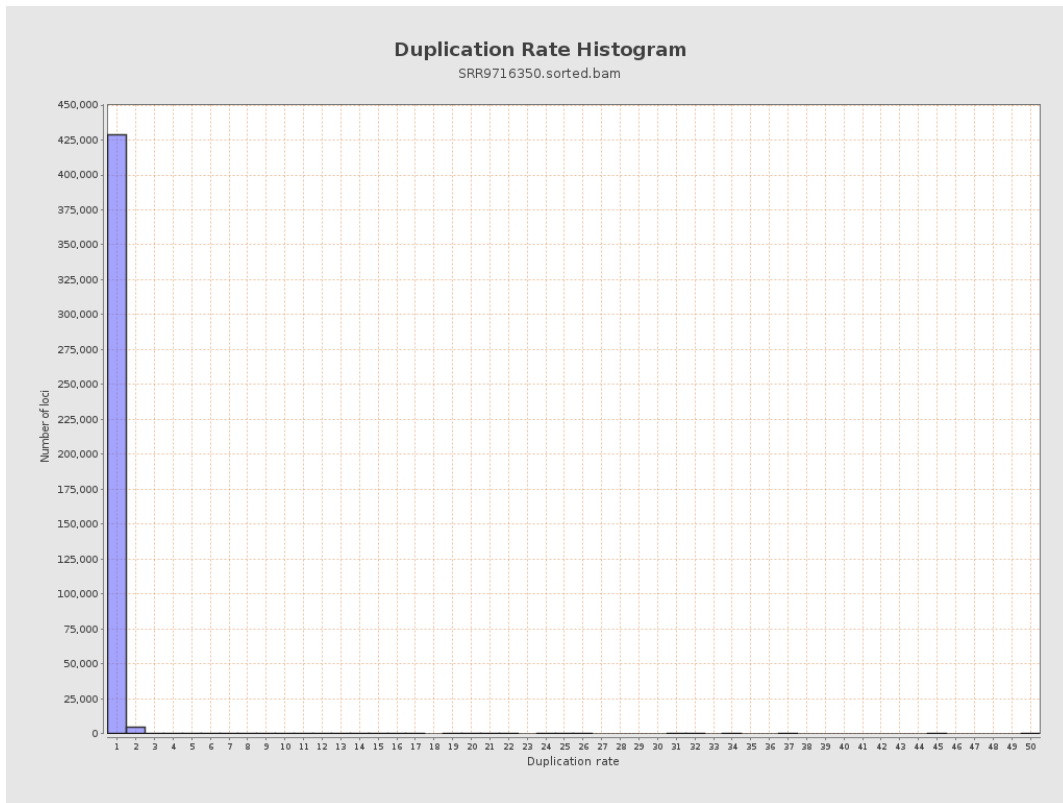
5. Results : Coverage Histogram (0-50X)



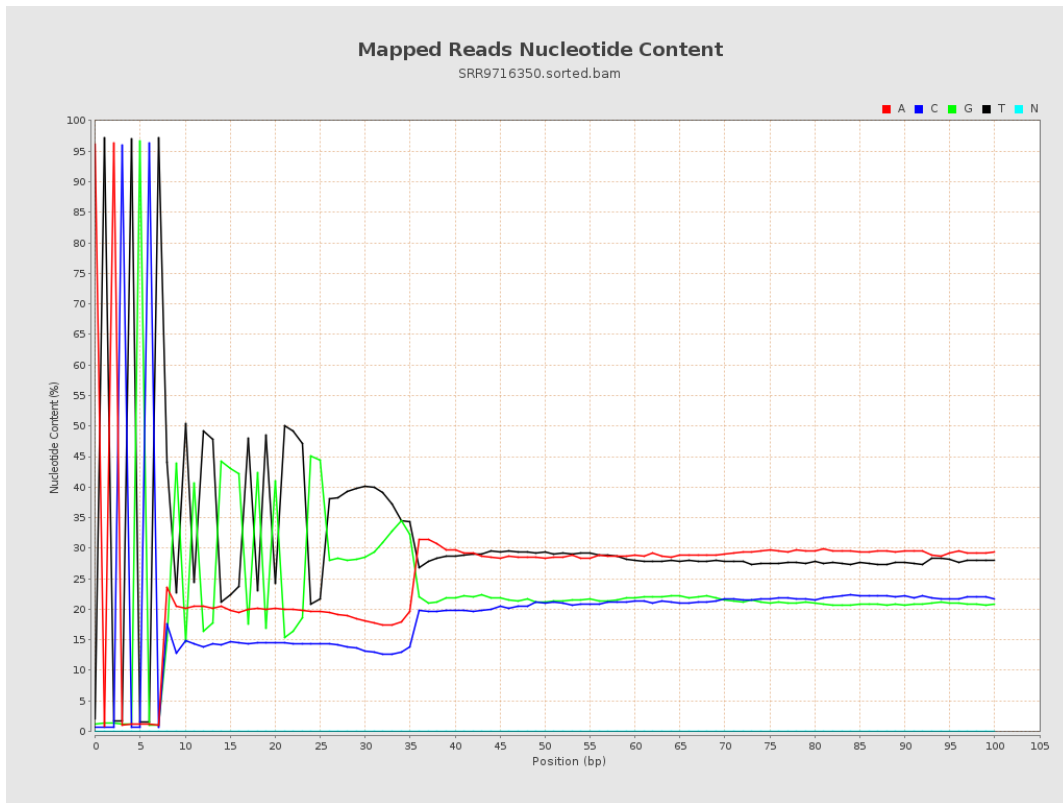
6. Results : Genome Fraction Coverage



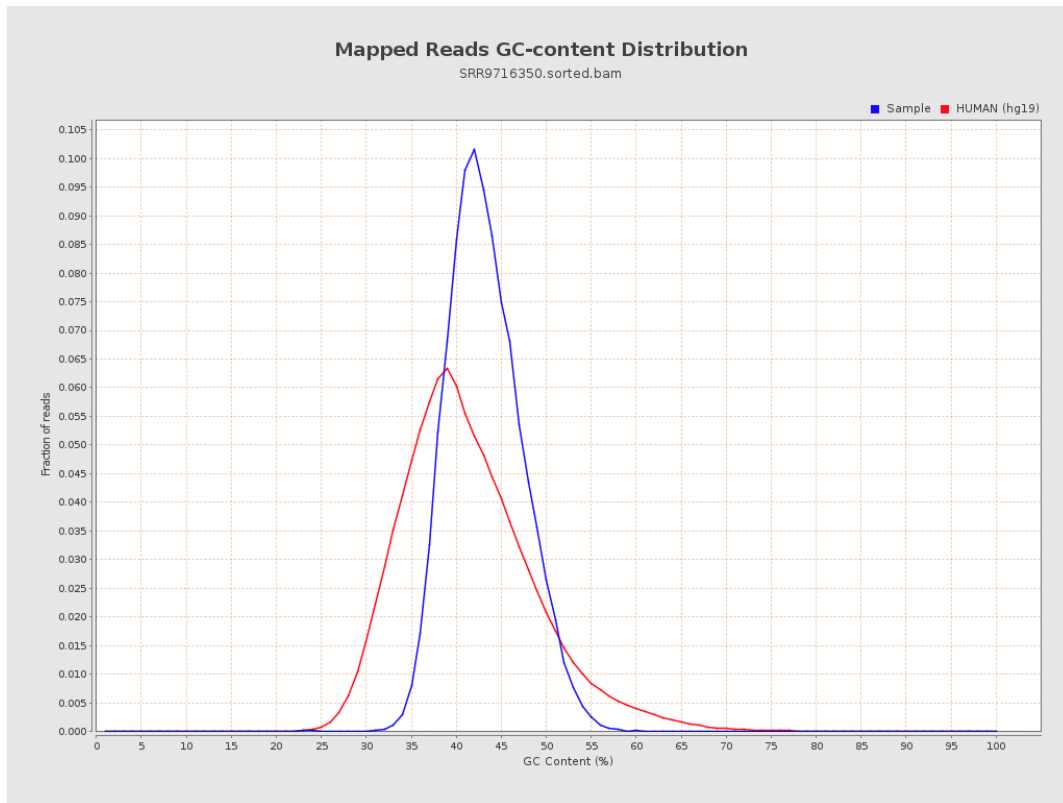
7. Results : Duplication Rate Histogram



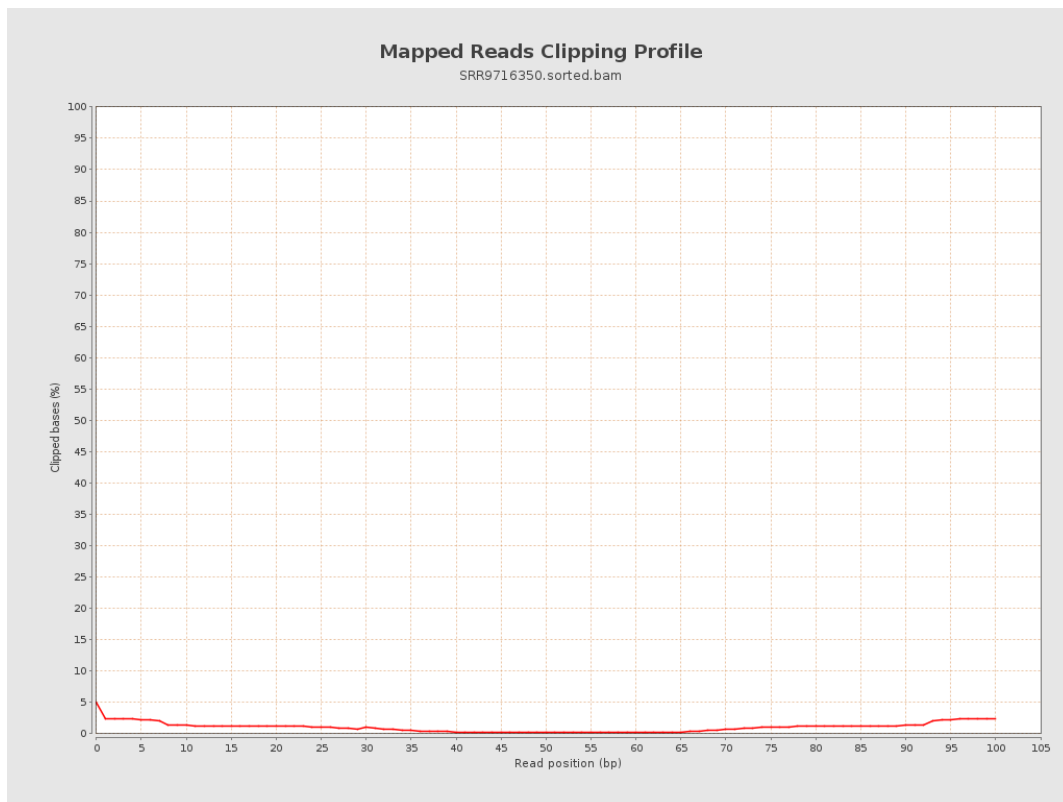
8. Results : Mapped Reads Nucleotide Content



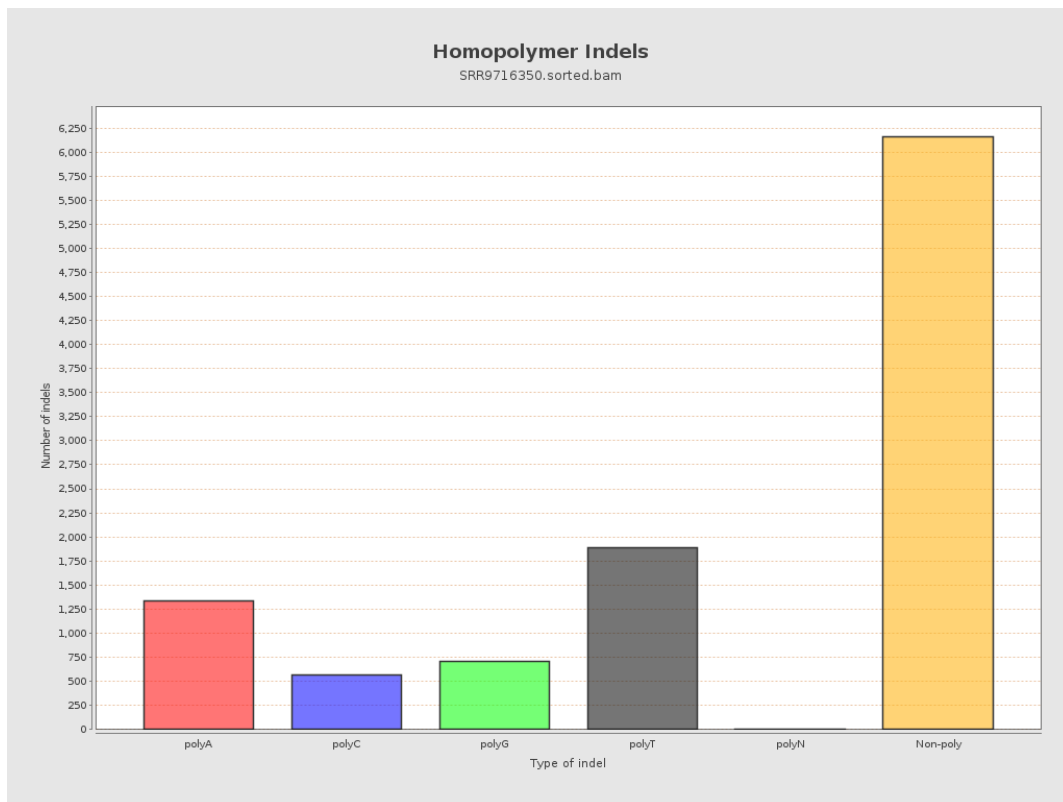
9. Results : Mapped Reads GC-content Distribution



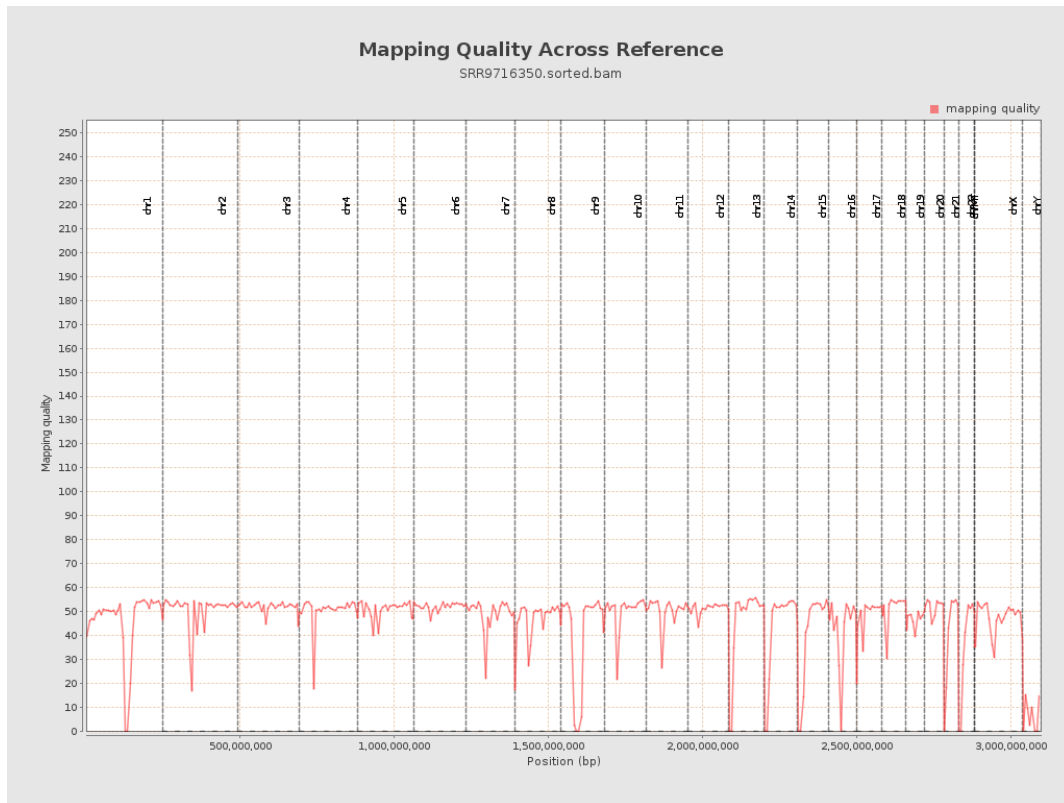
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

