

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/02 04:49:57

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR9716273.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_SRR9716273 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR9716273.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Sep 02 04:49:57 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR9716273.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	475,712
Mapped reads	403,503 / 84.82%
Unmapped reads	72,209 / 15.18%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	1,623 / 0.34%
Read min/max/mean length	30 / 76 / 76.11
Duplicated reads (estimated)	7,113 / 1.5%
Duplication rate	1.37%
Clipped reads	403,882 / 84.9%

2.2. ACGT Content

Number/percentage of A's	6,117,846 / 26.03%
Number/percentage of C's	4,014,364 / 17.08%
Number/percentage of T's	7,539,976 / 32.09%
Number/percentage of G's	5,825,884 / 24.79%
Number/percentage of N's	657 / 0%
GC Percentage	41.88%

2.3. Coverage

Mean	0.0076

Standard Deviation	0.1038
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2.4. Mapping Quality

Mean Mapping Quality	45.37
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2.5. Mismatches and indels

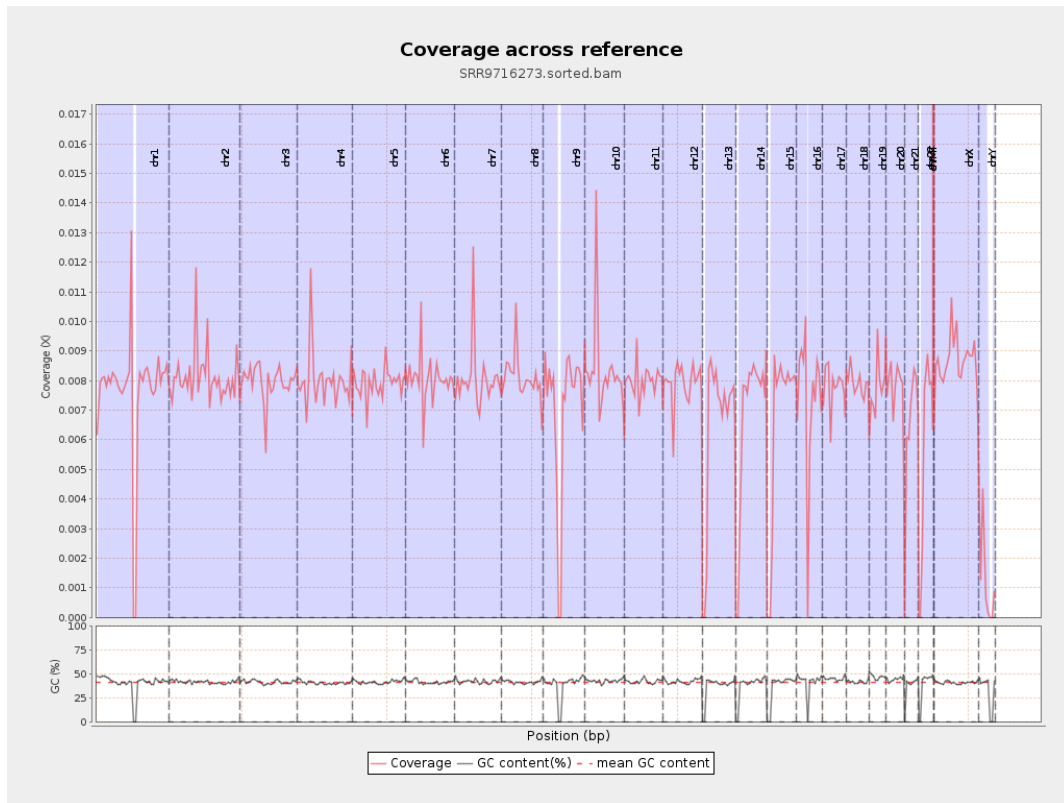
General error rate	0.52%
Mismatches	119,174
Insertions	1,772
Mapped reads with at least one insertion	0.44%
Deletions	4,620
Mapped reads with at least one deletion	1.14%
Homopolymer indels	42.32%

2.6. Chromosome stats

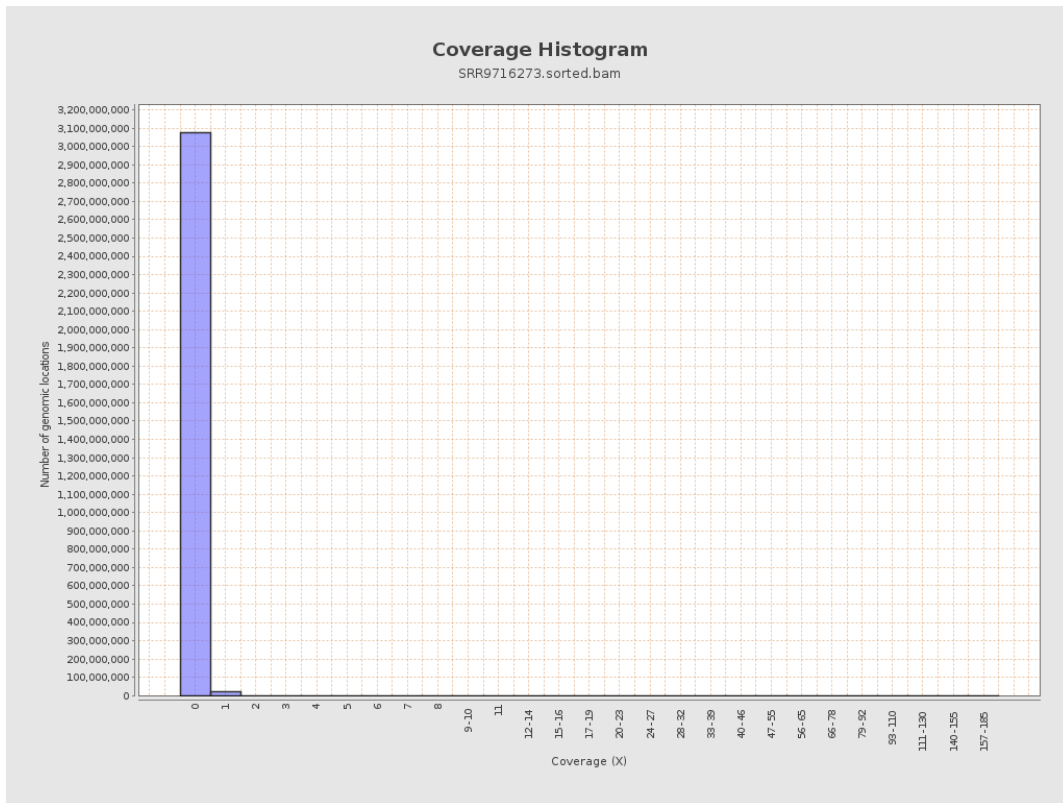
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	1898995	0.0076	0.1564
chr2	243199373	1975990	0.0081	0.1162
chr3	198022430	1574528	0.008	0.0922
chr4	191154276	1525581	0.008	0.0945
chr5	180915260	1431980	0.0079	0.0917
chr6	171115067	1378272	0.0081	0.0986
chr7	159138663	1282868	0.0081	0.1134

chr8	146364022	1183334	0.0081	0.1059
chr9	141213431	983618	0.007	0.0931
chr10	135534747	1126031	0.0083	0.1058
chr11	135006516	1076476	0.008	0.0975
chr12	133851895	1058634	0.0079	0.0921
chr13	115169878	738886	0.0064	0.0827
chr14	107349540	715751	0.0067	0.0847
chr15	102531392	676283	0.0066	0.0838
chr16	90354753	666110	0.0074	0.0901
chr17	81195210	625592	0.0077	0.091
chr18	78077248	620766	0.008	0.1224
chr19	59128983	460801	0.0078	0.122
chr20	63025520	490736	0.0078	0.0907
chr21	48129895	311995	0.0065	0.0867
chr22	51304566	282398	0.0055	0.0765
chrMT	16571	1658	0.1001	0.3156
chrX	155270560	1346082	0.0087	0.0987
chrY	59373566	72801	0.0012	0.0465

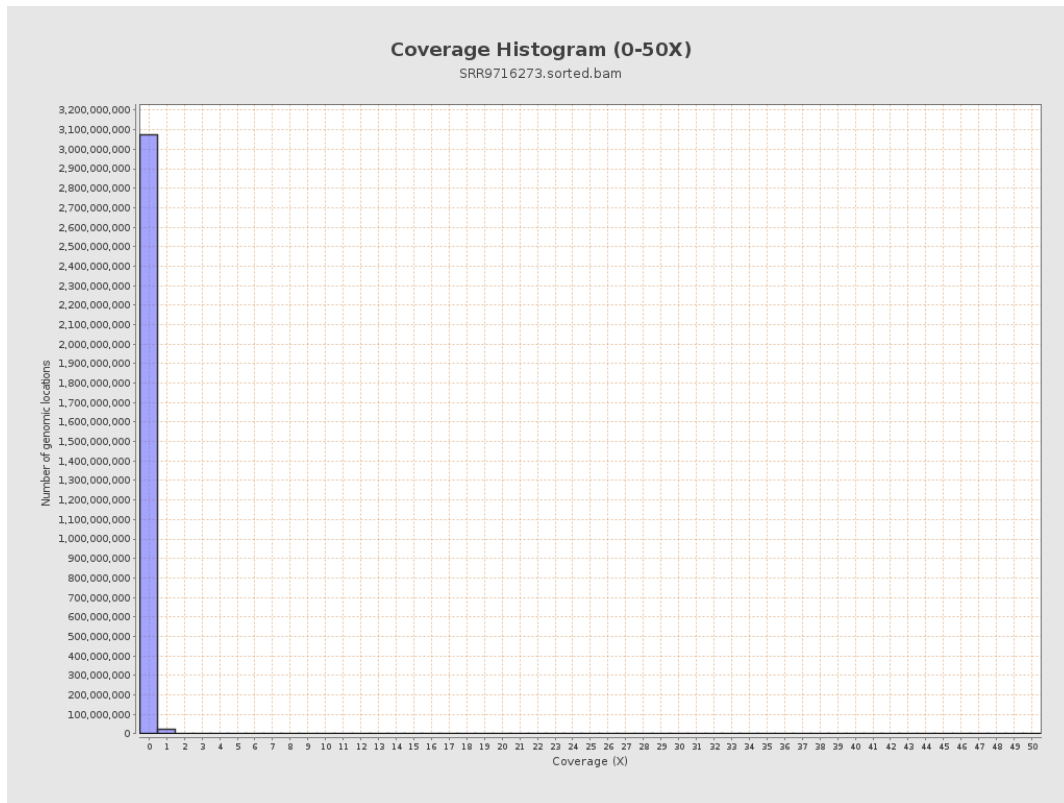
3. Results : Coverage across reference



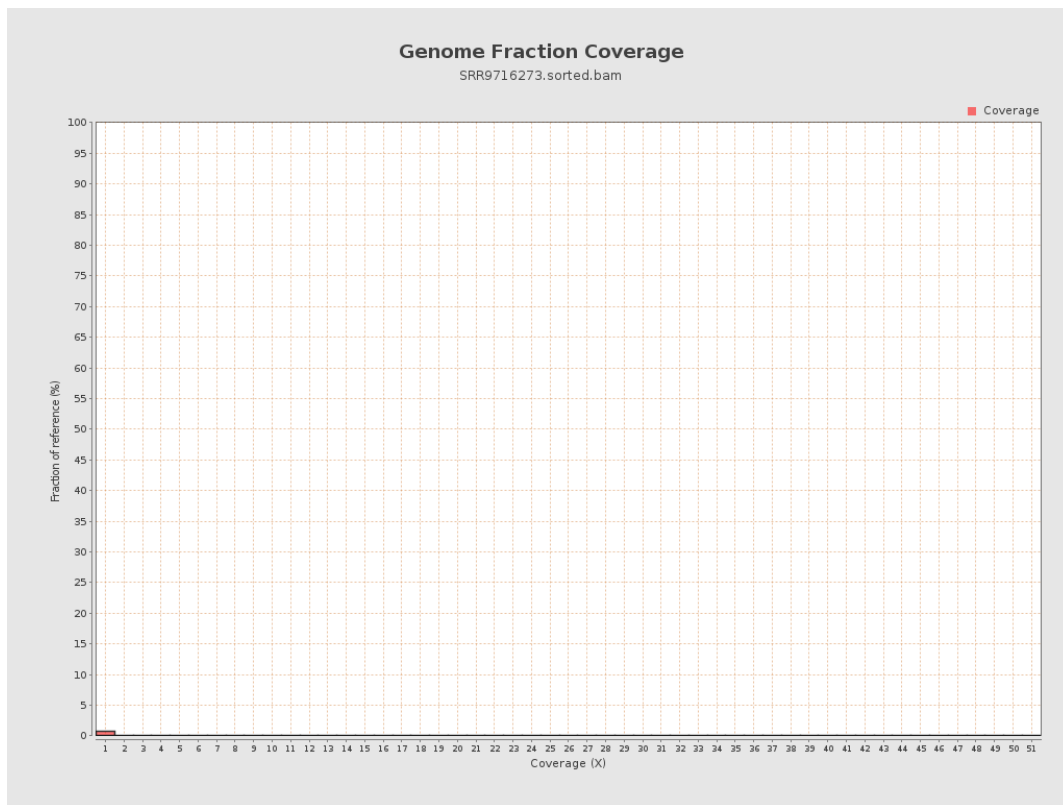
4. Results : Coverage Histogram



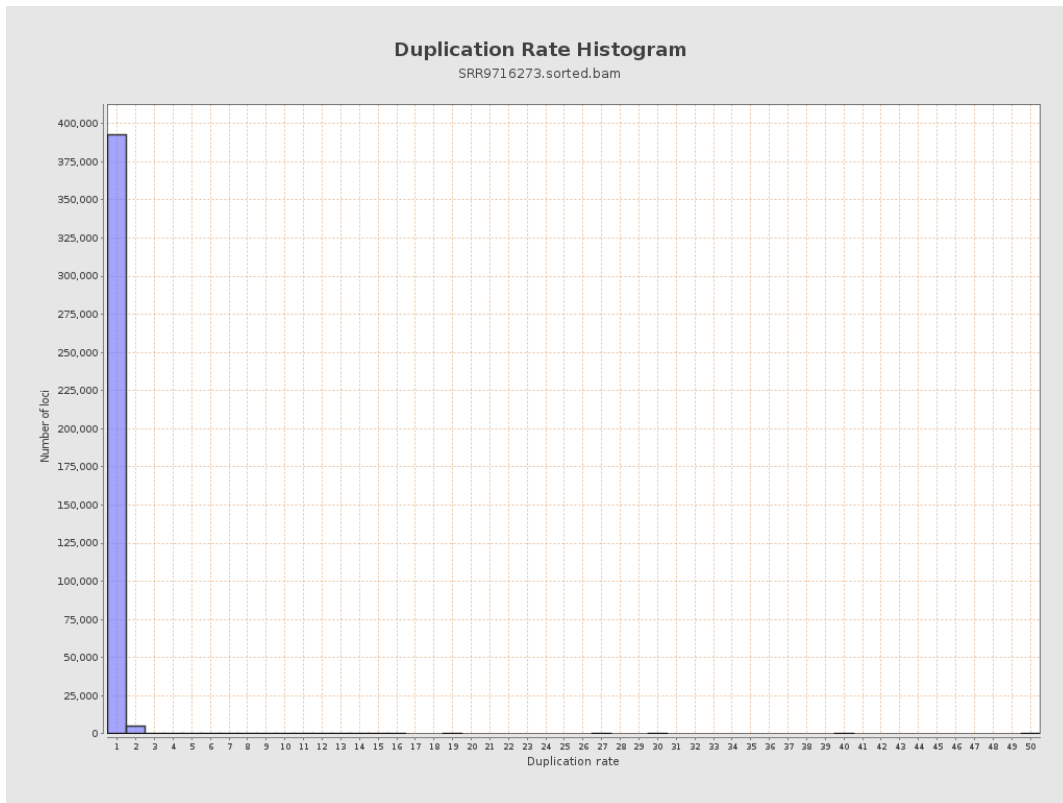
5. Results : Coverage Histogram (0-50X)



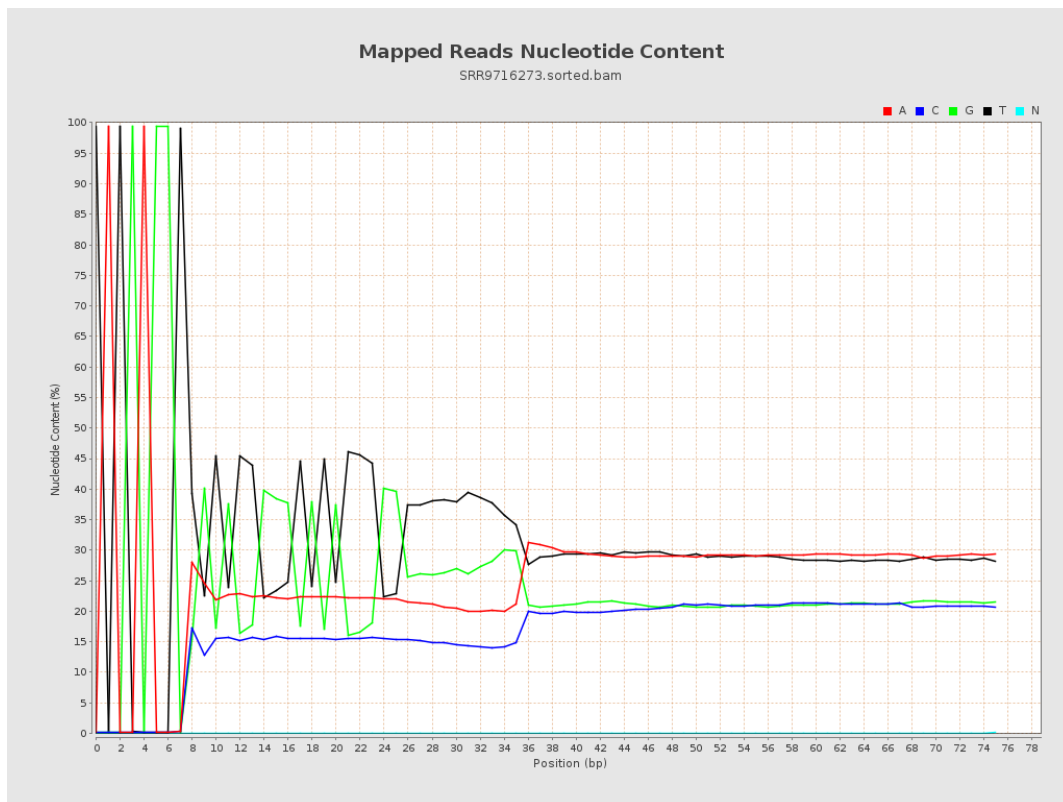
6. Results : Genome Fraction Coverage



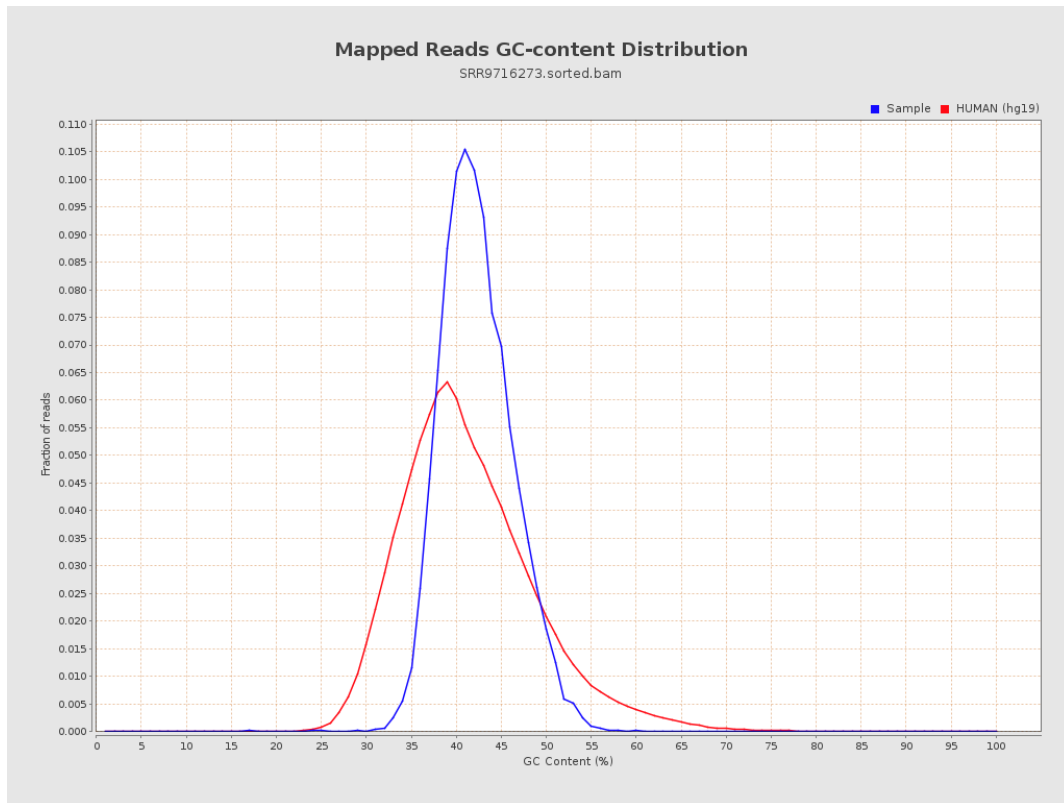
7. Results : Duplication Rate Histogram



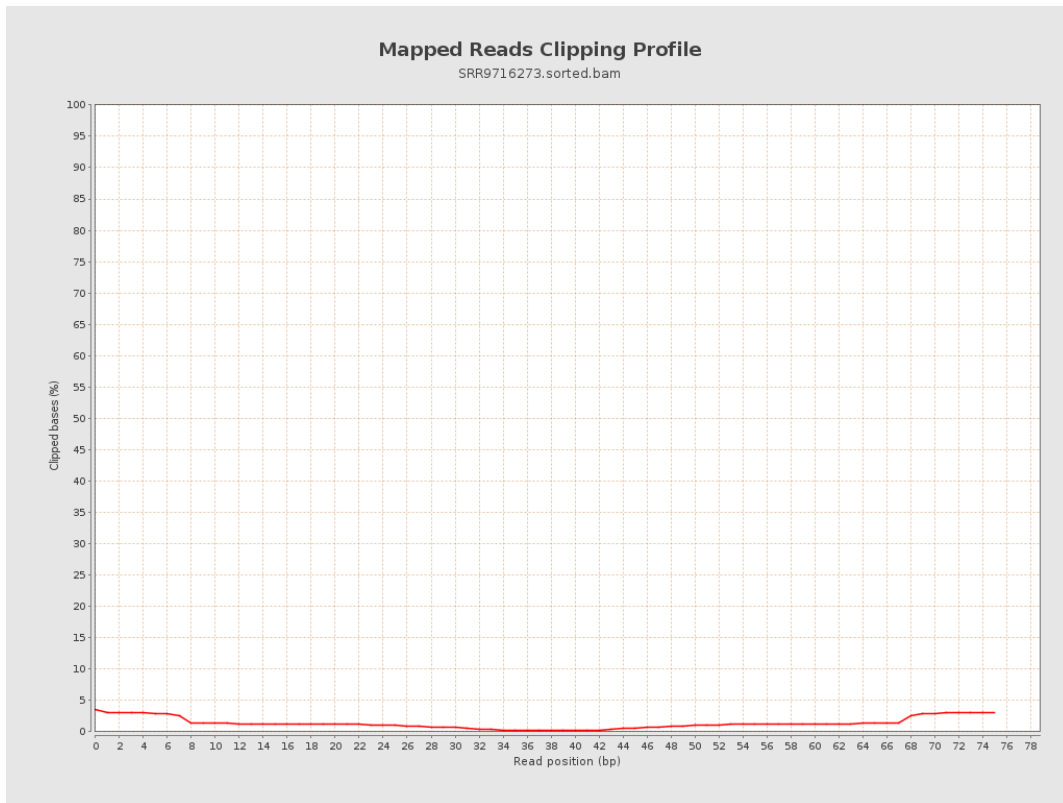
8. Results : Mapped Reads Nucleotide Content



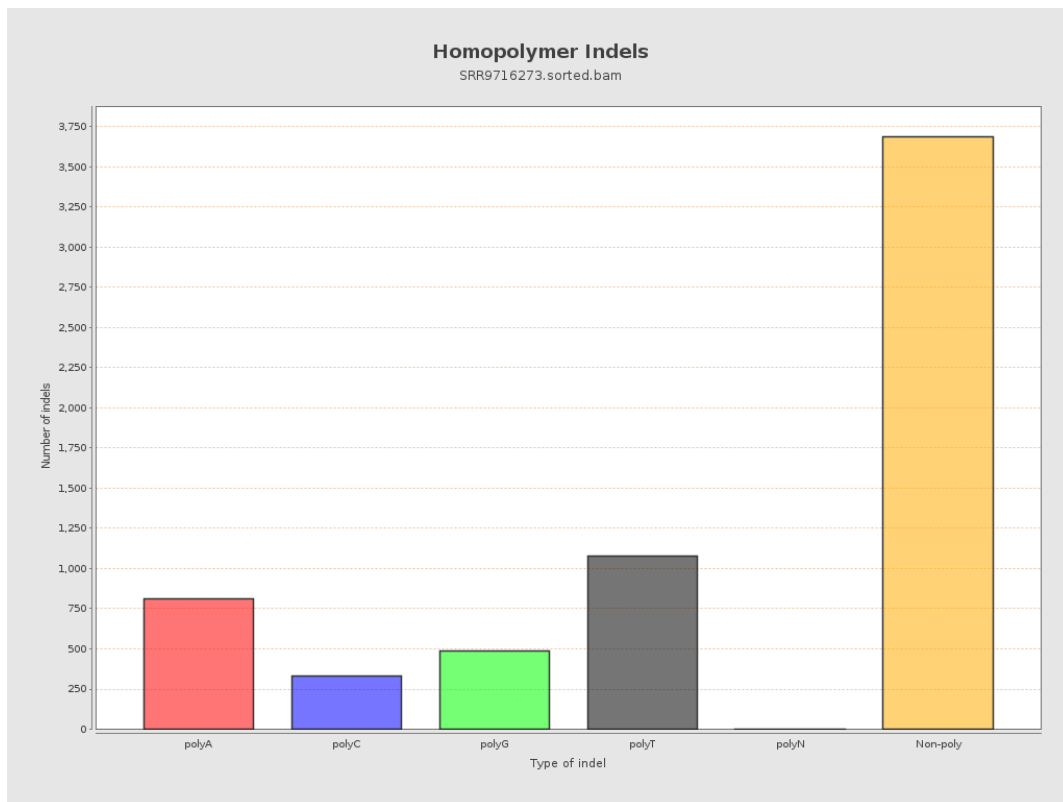
9. Results : Mapped Reads GC-content Distribution



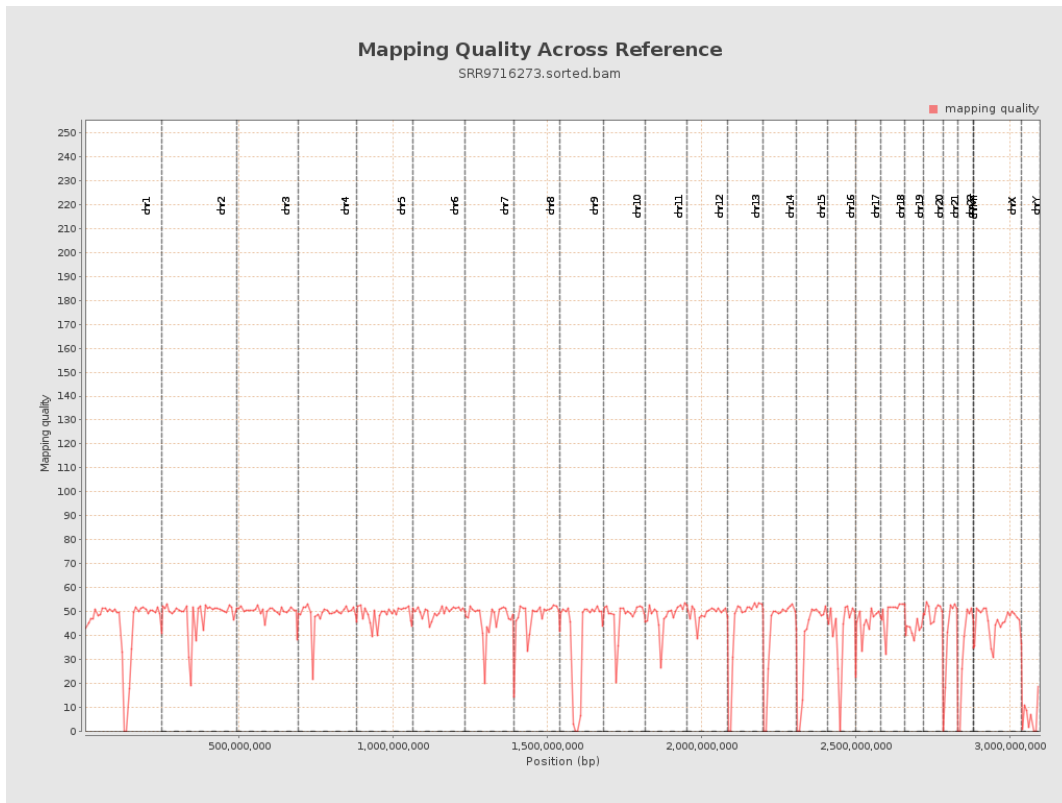
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

