

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/02 17:27:17

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR9716504.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_SRR9716504 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR9716504.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Sep 02 17:27:16 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR9716504.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	2,828,423
Mapped reads	2,491,659 / 88.09%
Unmapped reads	336,764 / 11.91%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	9,984 / 0.35%
Read min/max/mean length	30 / 76 / 76.12
Duplicated reads (estimated)	115,272 / 4.08%
Duplication rate	3.45%
Clipped reads	2,494,444 / 88.19%

2.2. ACGT Content

Number/percentage of A's	34,919,451 / 24.57%
Number/percentage of C's	28,190,497 / 19.84%
Number/percentage of T's	46,320,458 / 32.59%
Number/percentage of G's	32,679,205 / 23%
Number/percentage of N's	1,034 / 0%
GC Percentage	42.83%

2.3. Coverage

Mean	0.0459

Standard Deviation	0.3611
--------------------	--------

2.4. Mapping Quality

Mean Mapping Quality	43.89
----------------------	-------

2.5. Mismatches and indels

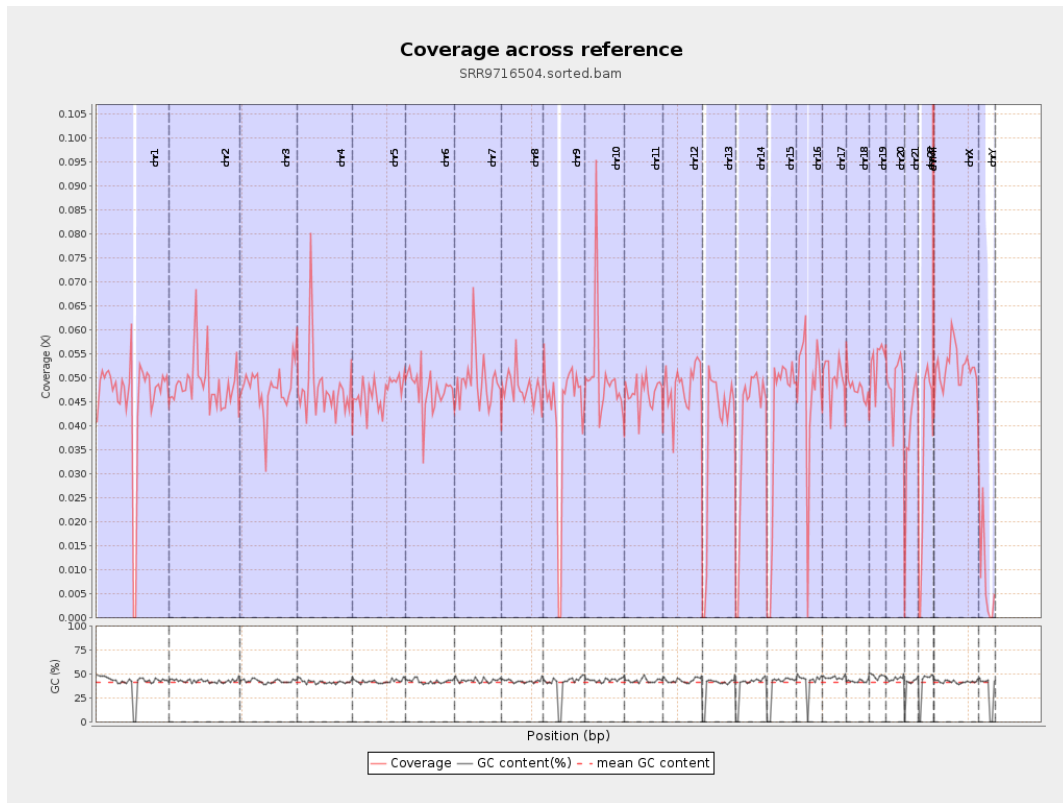
General error rate	0.54%
Mismatches	755,259
Insertions	9,424
Mapped reads with at least one insertion	0.38%
Deletions	27,180
Mapped reads with at least one deletion	1.08%
Homopolymer indels	41.41%

2.6. Chromosome stats

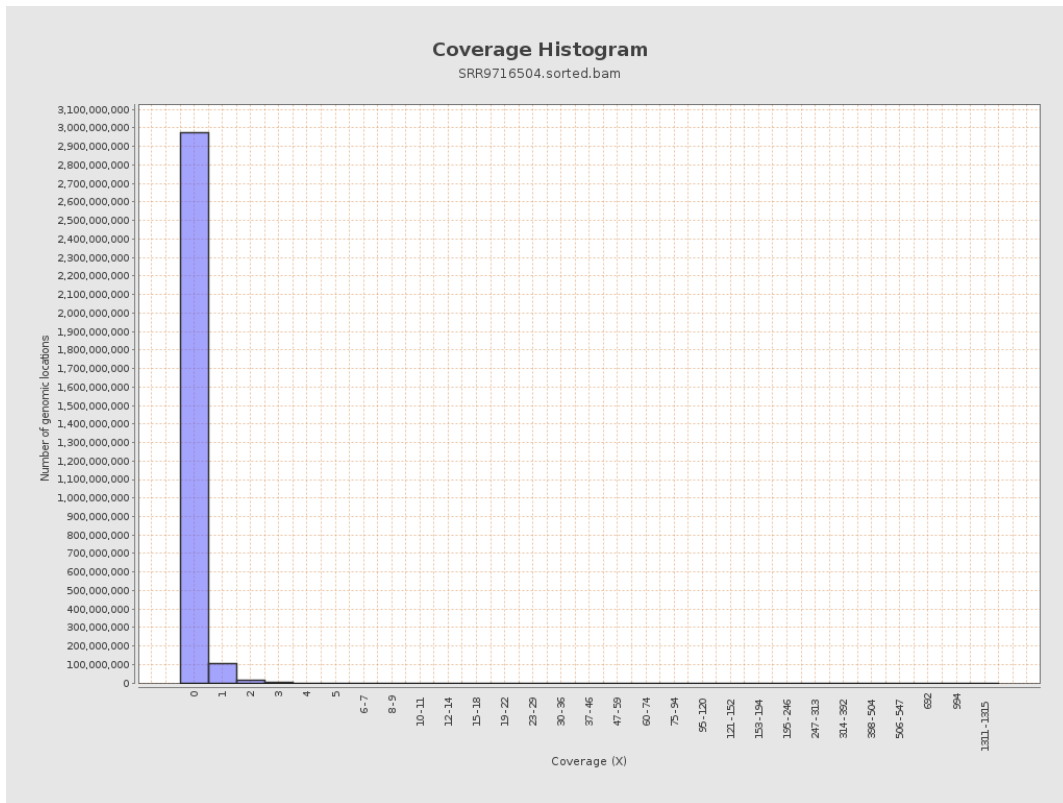
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	11367276	0.0456	0.4803
chr2	243199373	11856848	0.0488	0.616
chr3	198022430	9430997	0.0476	0.2561
chr4	191154276	9099983	0.0476	0.2959
chr5	180915260	8483003	0.0469	0.25
chr6	171115067	8110927	0.0474	0.3014
chr7	159138663	7890765	0.0496	0.4342

chr8	146364022	7002812	0.0478	0.3585
chr9	141213431	5919863	0.0419	0.2952
chr10	135534747	6789693	0.0501	0.4493
chr11	135006516	6317814	0.0468	0.3137
chr12	133851895	6451036	0.0482	0.2578
chr13	115169878	4431106	0.0385	0.2257
chr14	107349540	4275869	0.0398	0.2425
chr15	102531392	4172336	0.0407	0.2374
chr16	90354753	4305867	0.0477	0.2844
chr17	81195210	4010508	0.0494	0.2707
chr18	78077248	3739599	0.0479	0.449
chr19	59128983	3147452	0.0532	0.4281
chr20	63025520	3051172	0.0484	0.2696
chr21	48129895	1843507	0.0383	0.2606
chr22	51304566	1749148	0.0341	0.2151
chrMT	16571	154722	9.3369	6.0391
chrX	155270560	8084661	0.0521	0.2931
chrY	59373566	467076	0.0079	0.2289

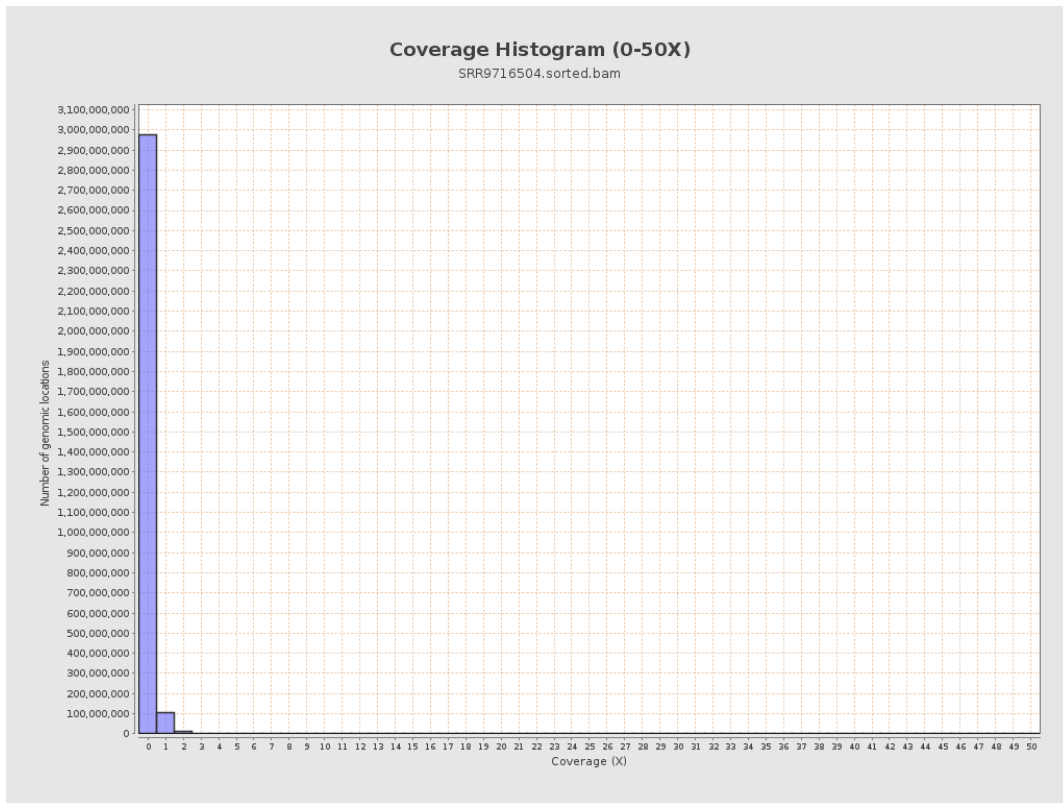
3. Results : Coverage across reference



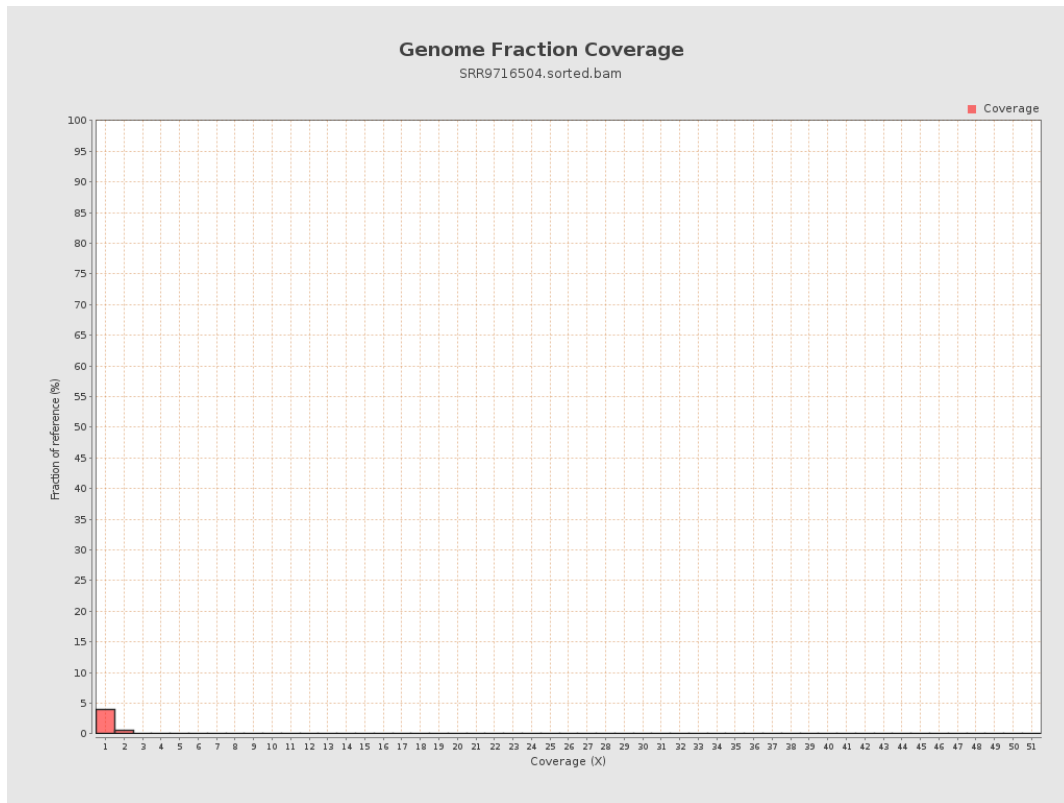
4. Results : Coverage Histogram



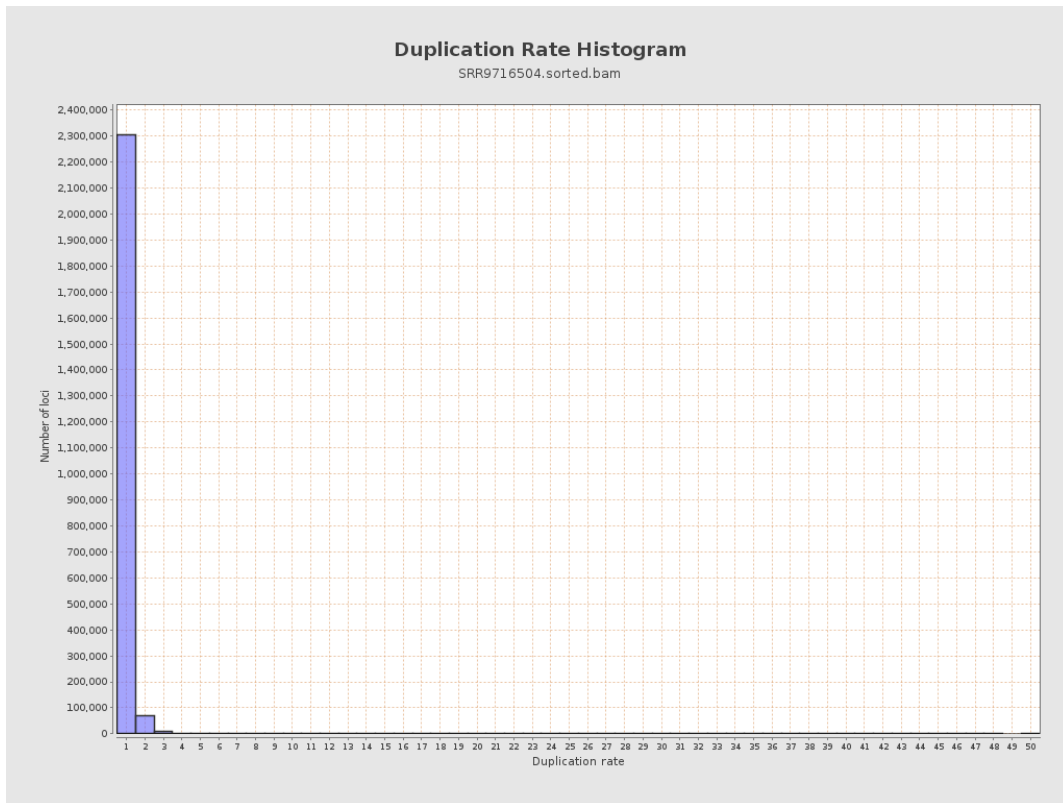
5. Results : Coverage Histogram (0-50X)



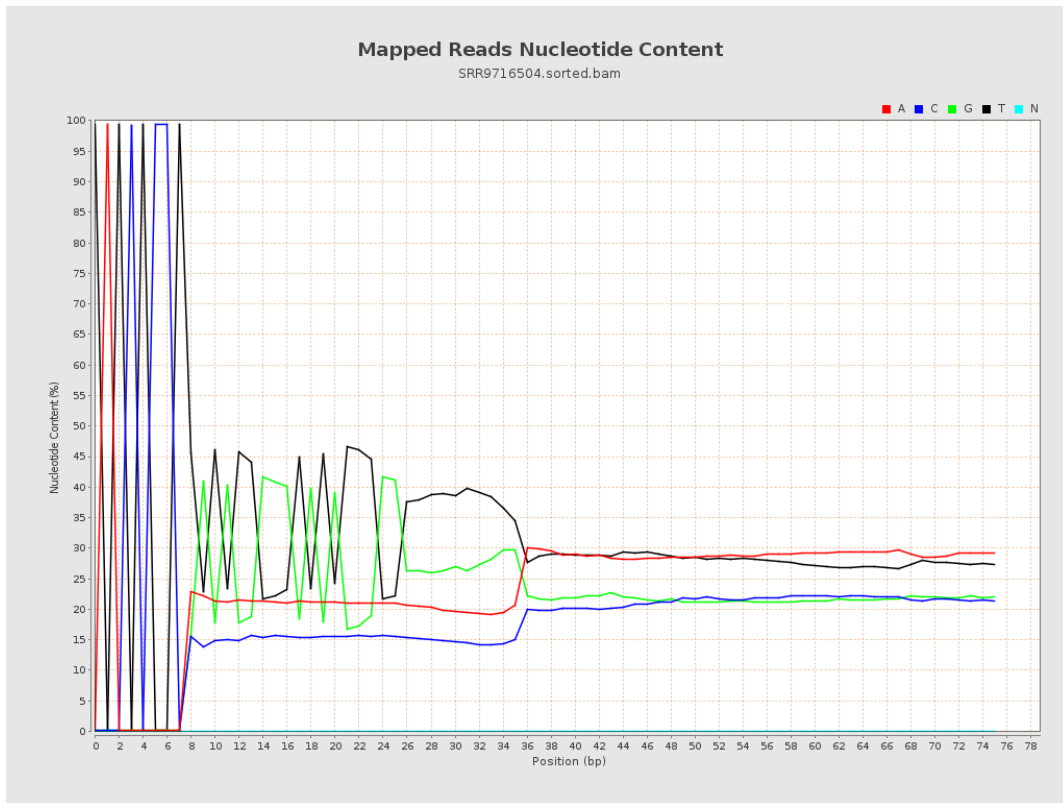
6. Results : Genome Fraction Coverage



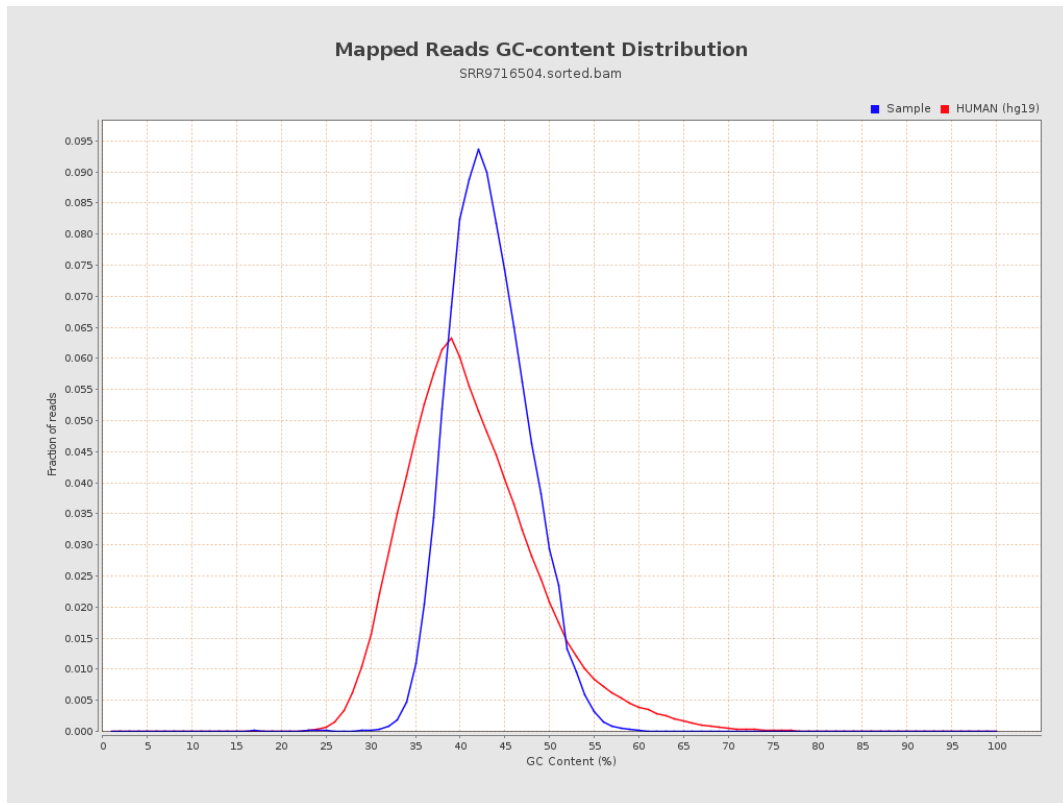
7. Results : Duplication Rate Histogram



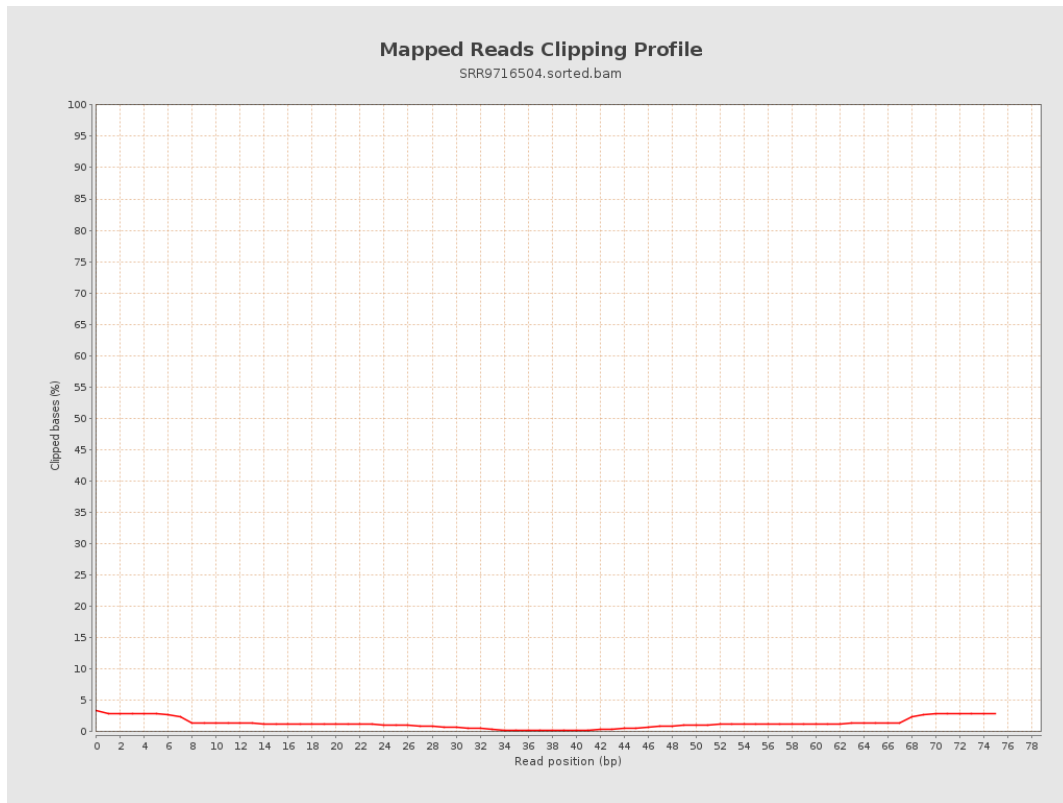
8. Results : Mapped Reads Nucleotide Content



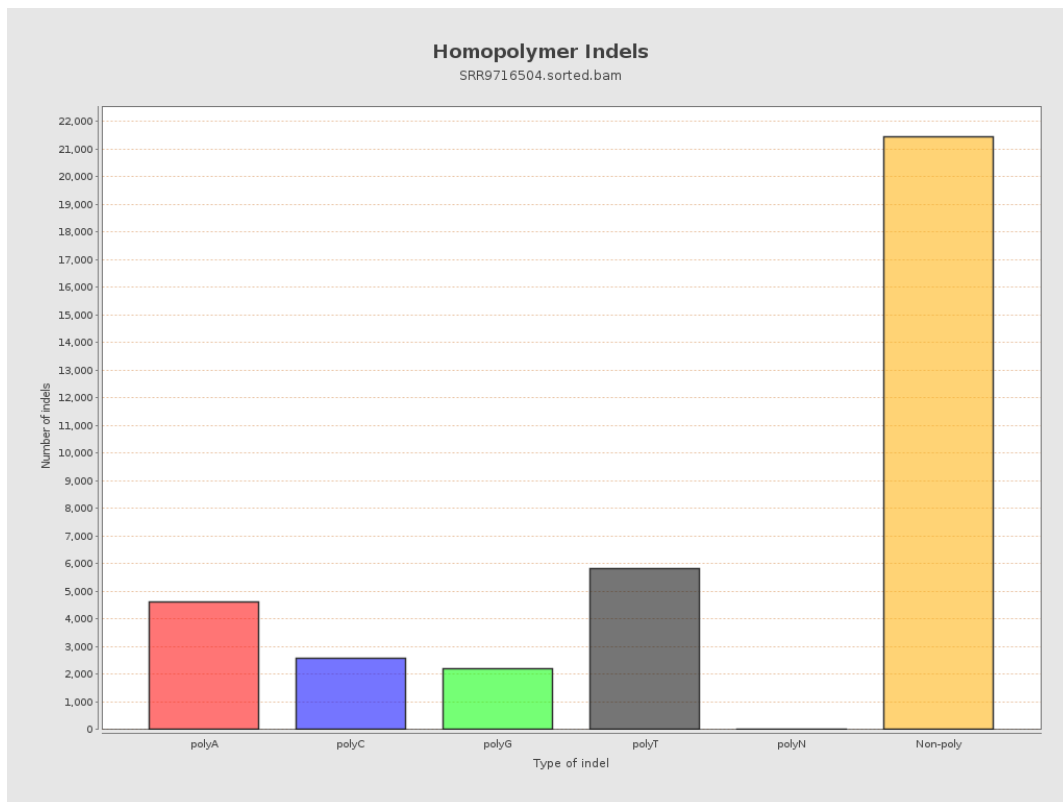
9. Results : Mapped Reads GC-content Distribution



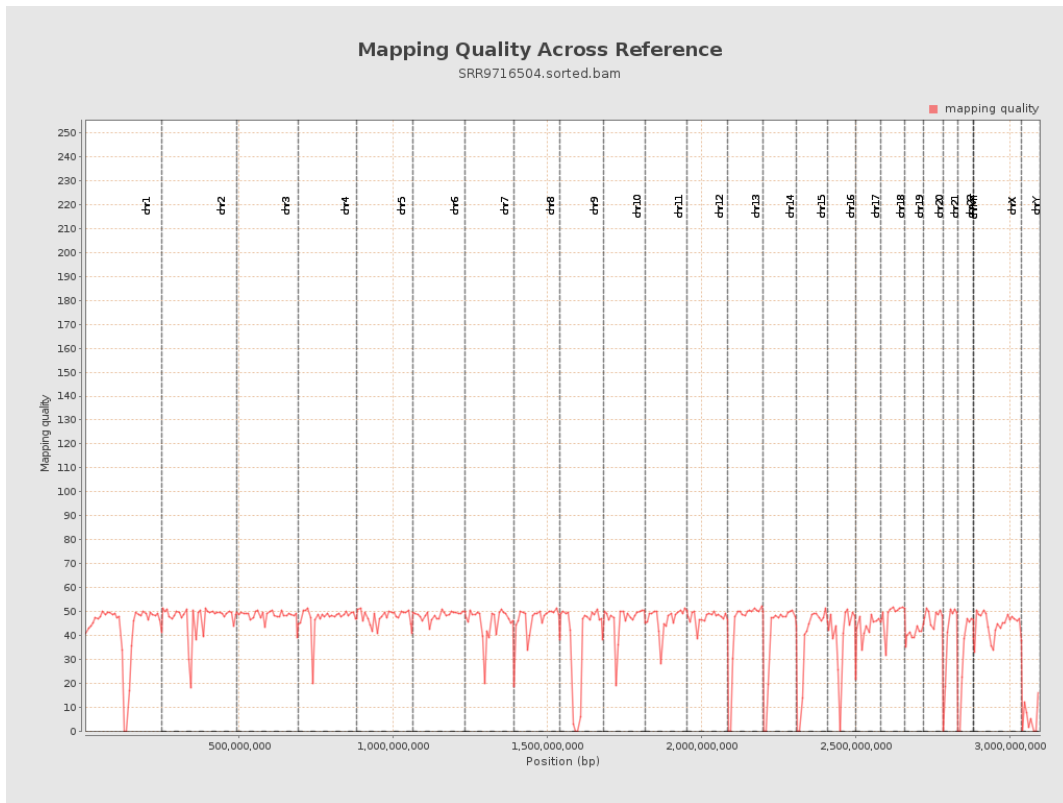
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

