

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/14 04:39:59

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6004149.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_SRR6004149 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6004149.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sat Sep 14 04:39:58 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6004149.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,286,238
Mapped reads	2,625,408 / 79.89%
Unmapped reads	660,830 / 20.11%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	11,626 / 0.35%
Read min/max/mean length	30 / 76 / 76.12
Duplicated reads (estimated)	233,053 / 7.09%
Duplication rate	7.01%
Clipped reads	1,624,884 / 49.45%

2.2. ACGT Content

Number/percentage of A's	42,270,757 / 25.68%
Number/percentage of C's	28,428,007 / 17.27%
Number/percentage of T's	54,868,094 / 33.34%
Number/percentage of G's	39,004,172 / 23.7%
Number/percentage of N's	16,493 / 0.01%
GC Percentage	40.97%

2.3. Coverage

Mean	0.0532

Standard Deviation	0.5335
--------------------	--------

2.4. Mapping Quality

Mean Mapping Quality	43.33
----------------------	-------

2.5. Mismatches and indels

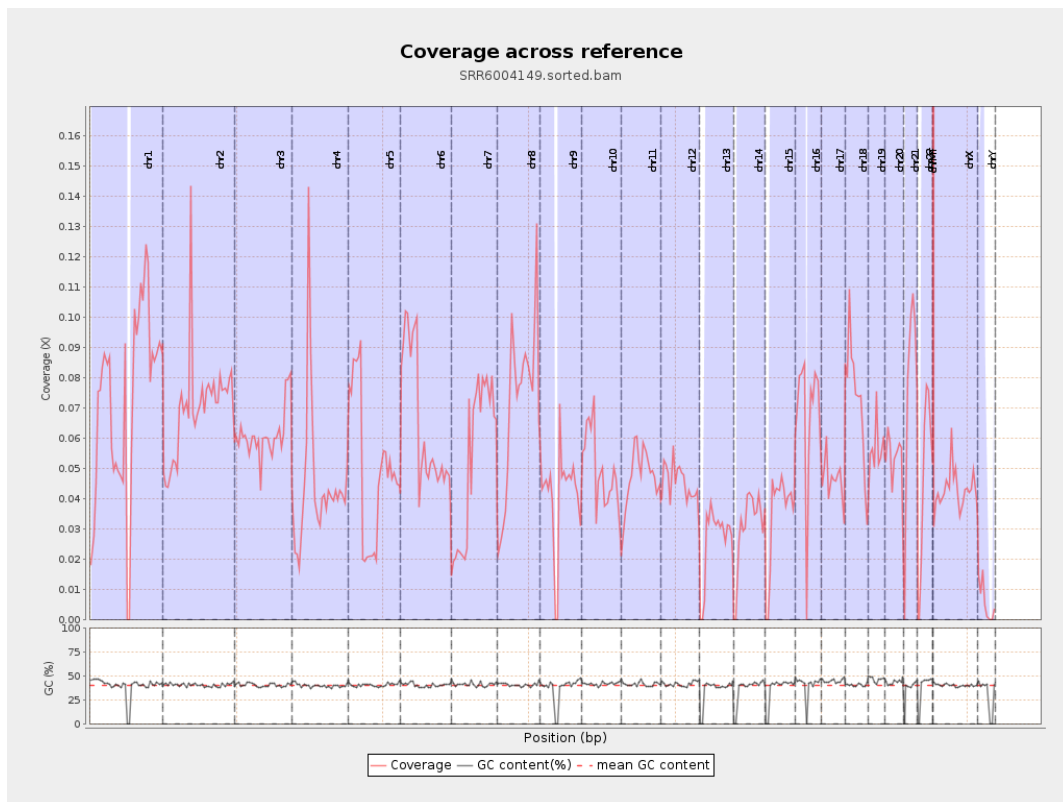
General error rate	1.03%
Mismatches	1,670,258
Insertions	14,690
Mapped reads with at least one insertion	0.56%
Deletions	50,896
Mapped reads with at least one deletion	1.92%
Homopolymer indels	49.46%

2.6. Chromosome stats

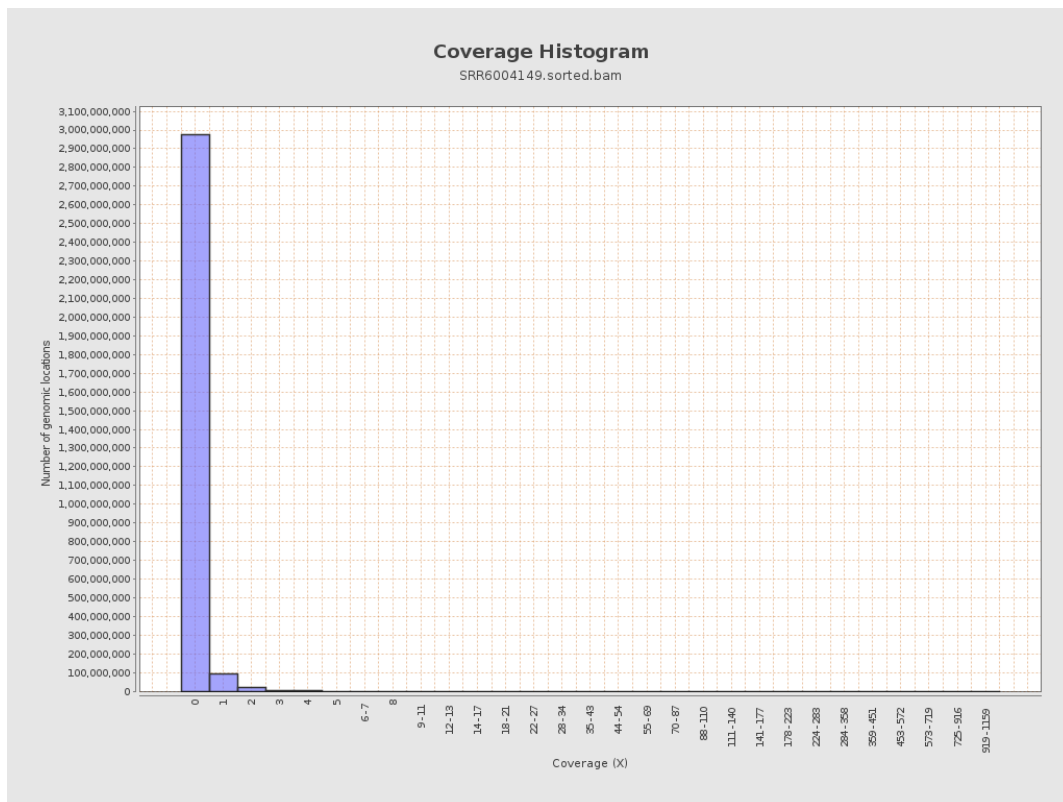
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	17978075	0.0721	0.8329
chr2	243199373	17016187	0.07	0.873
chr3	198022430	12059208	0.0609	0.3203
chr4	191154276	8570902	0.0448	0.2846
chr5	180915260	8907676	0.0492	0.2937
chr6	171115067	11060768	0.0646	0.4273
chr7	159138663	8413920	0.0529	0.5879

chr8	146364022	10463854	0.0715	0.5808
chr9	141213431	5816212	0.0412	0.5661
chr10	135534747	6690890	0.0494	0.3959
chr11	135006516	6381026	0.0473	0.4691
chr12	133851895	6046641	0.0452	0.2954
chr13	115169878	3032701	0.0263	0.2092
chr14	107349540	3274145	0.0305	0.3322
chr15	102531392	3498718	0.0341	0.2491
chr16	90354753	6043174	0.0669	0.3669
chr17	81195210	3779208	0.0465	0.3454
chr18	78077248	5757888	0.0737	1.1835
chr19	59128983	3331661	0.0563	0.7064
chr20	63025520	3405695	0.054	0.3516
chr21	48129895	3618944	0.0752	0.3795
chr22	51304566	2424742	0.0473	0.279
chrMT	16571	117828	7.1105	8.6539
chrX	155270560	6641477	0.0428	0.3727
chrY	59373566	340256	0.0057	0.1214

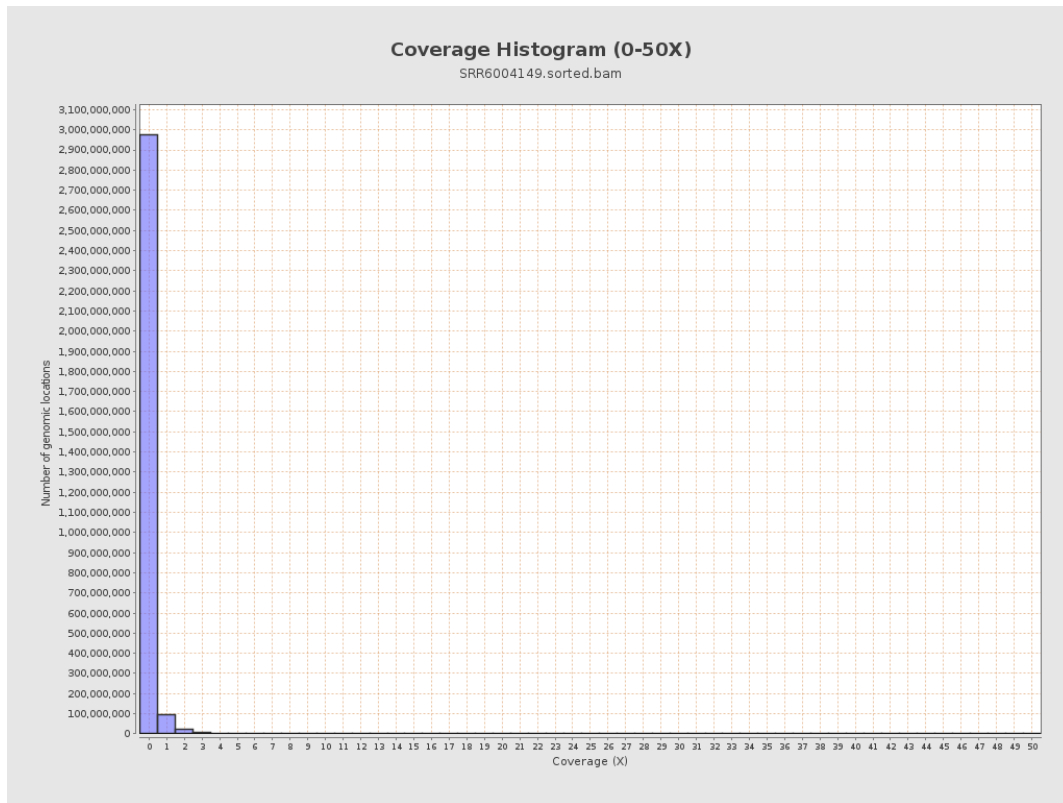
3. Results : Coverage across reference



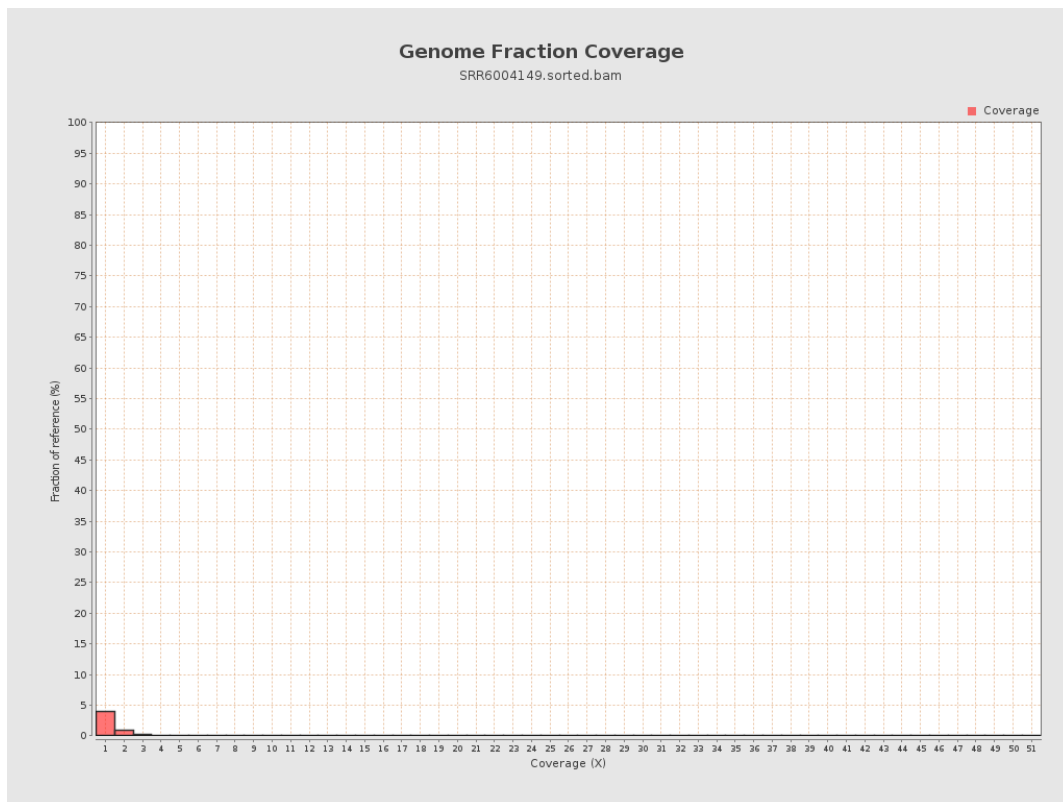
4. Results : Coverage Histogram



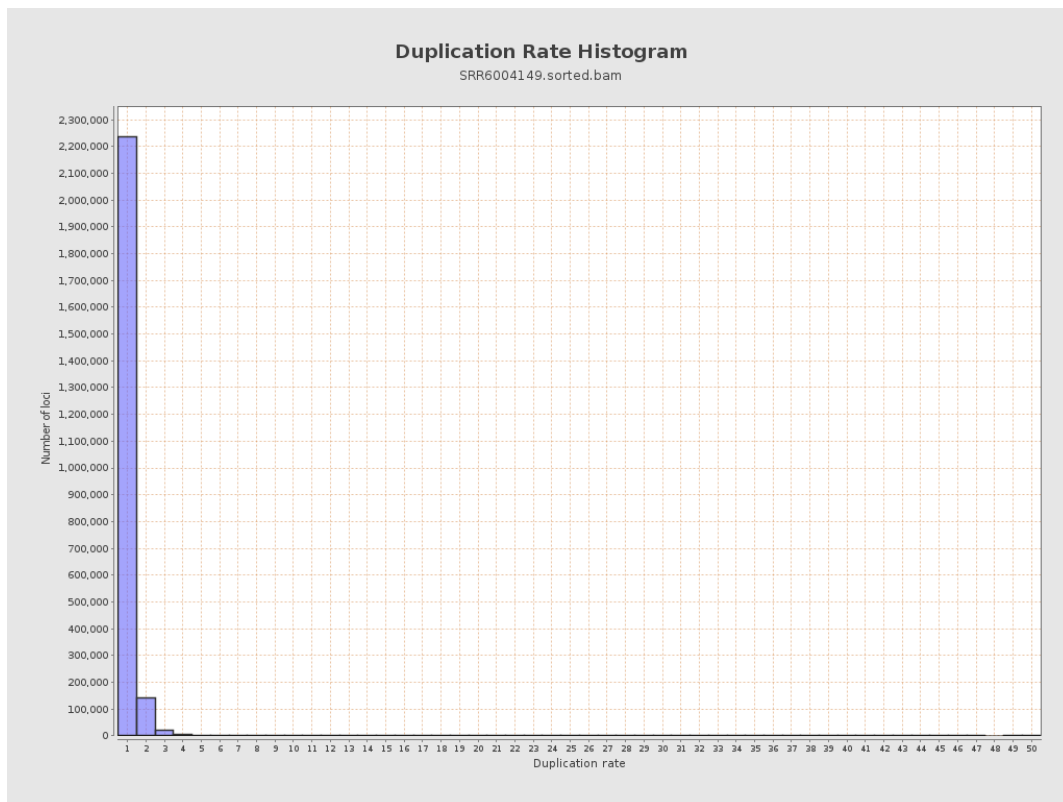
5. Results : Coverage Histogram (0-50X)



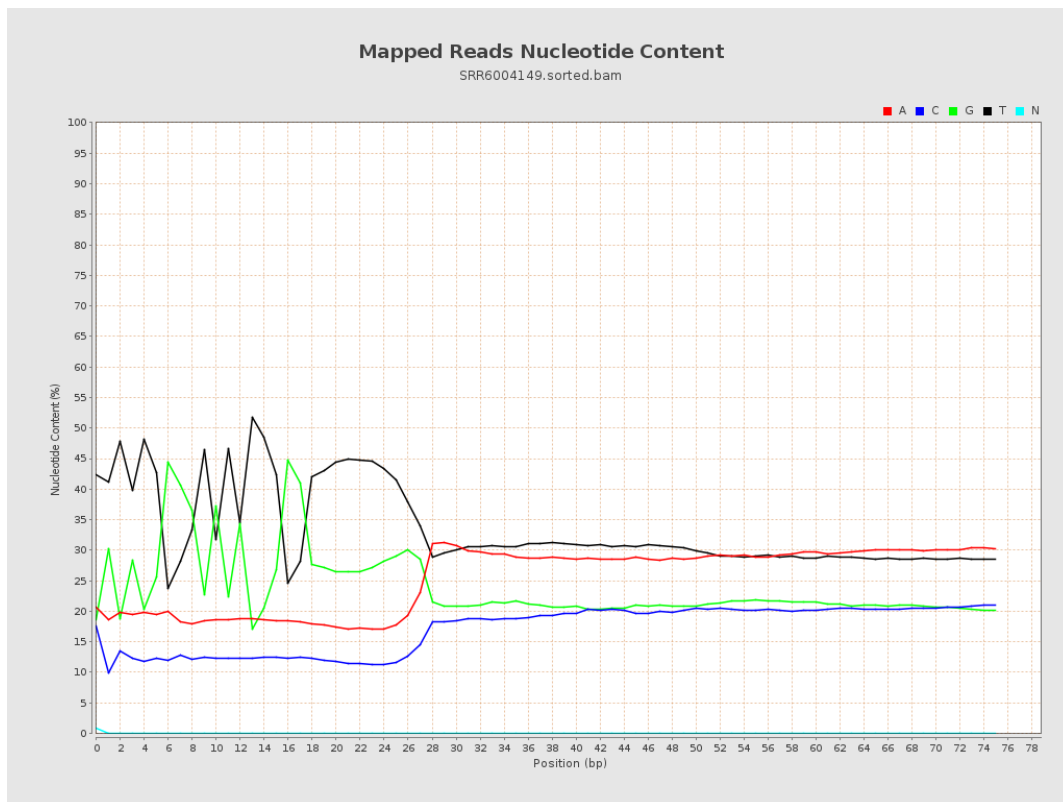
6. Results : Genome Fraction Coverage



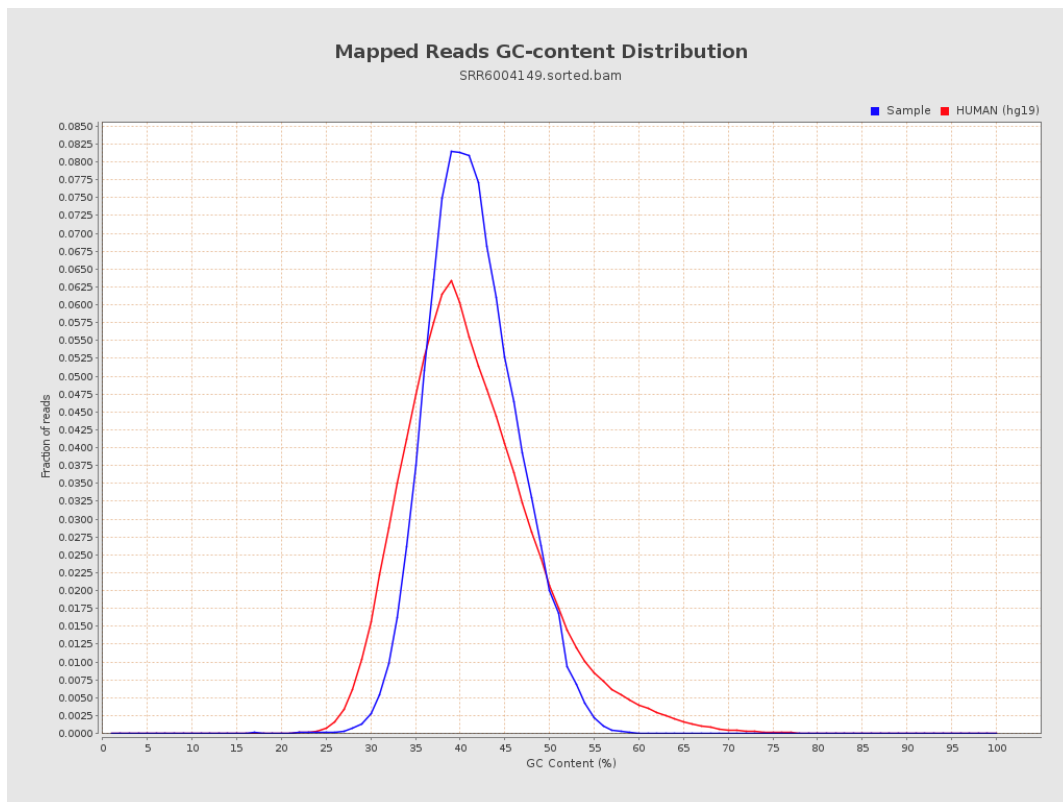
7. Results : Duplication Rate Histogram



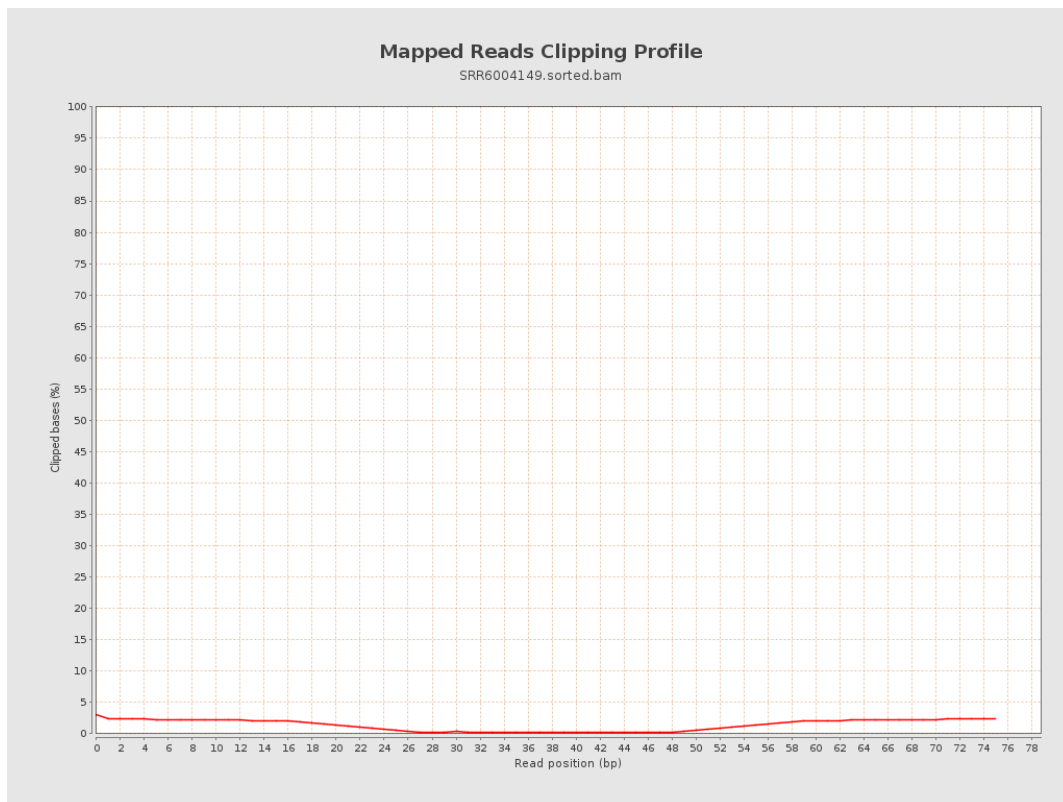
8. Results : Mapped Reads Nucleotide Content



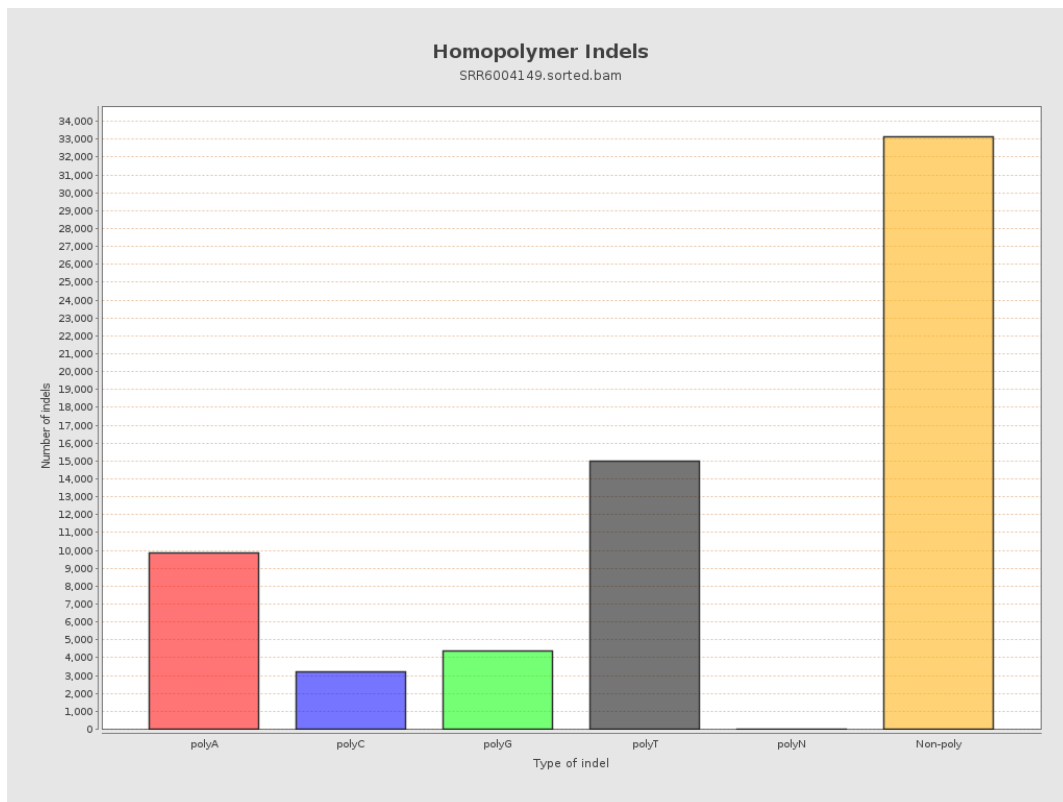
9. Results : Mapped Reads GC-content Distribution



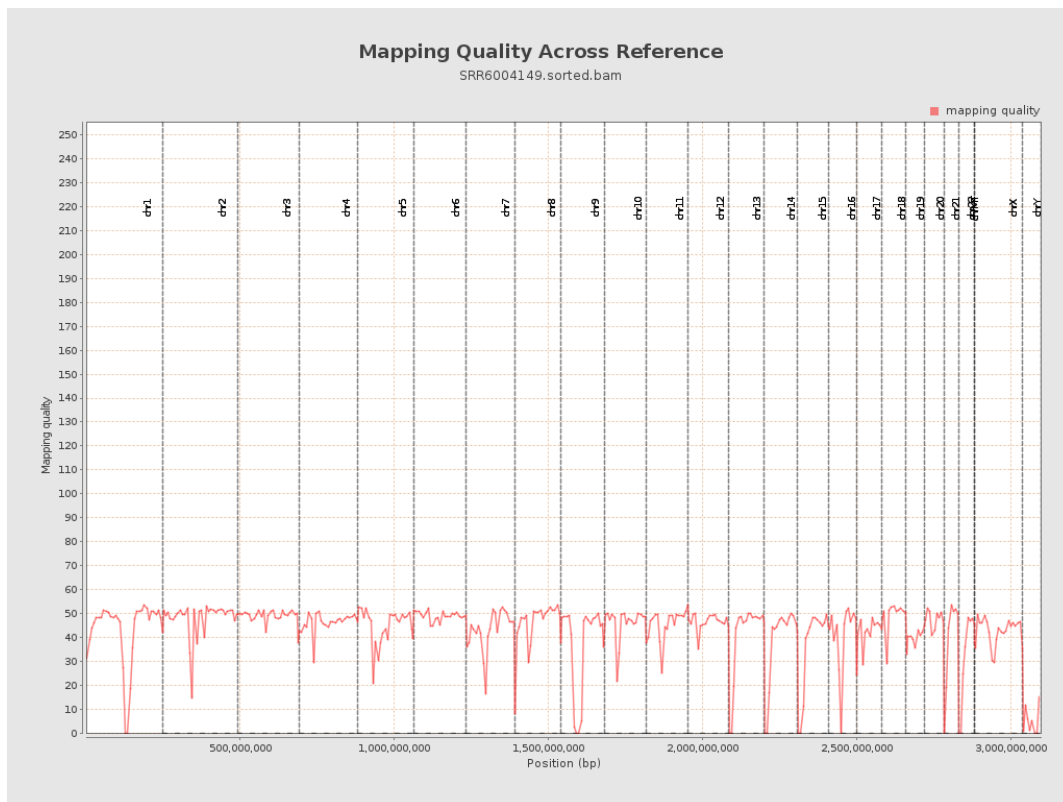
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

