

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/16 08:51:28

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6231867.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_SRR6231867 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6231867.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Sep 16 08:51:28 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6231867.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	5,754,586
Mapped reads	5,132,548 / 89.19%
Unmapped reads	622,038 / 10.81%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	46,498 / 0.81%
Read min/max/mean length	30 / 76 / 76.28
Duplicated reads (estimated)	314,815 / 5.47%
Duplication rate	4.13%
Clipped reads	2,465,414 / 42.84%

2.2. ACGT Content

Number/percentage of A's	99,450,675 / 29.39%
Number/percentage of C's	64,589,769 / 19.09%
Number/percentage of T's	101,585,491 / 30.02%
Number/percentage of G's	72,690,434 / 21.48%
Number/percentage of N's	26,105 / 0.01%
GC Percentage	40.57%

2.3. Coverage

Mean	0.1093

Standard Deviation	3.89
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2.4. Mapping Quality

Mean Mapping Quality	45.75
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2.5. Mismatches and indels

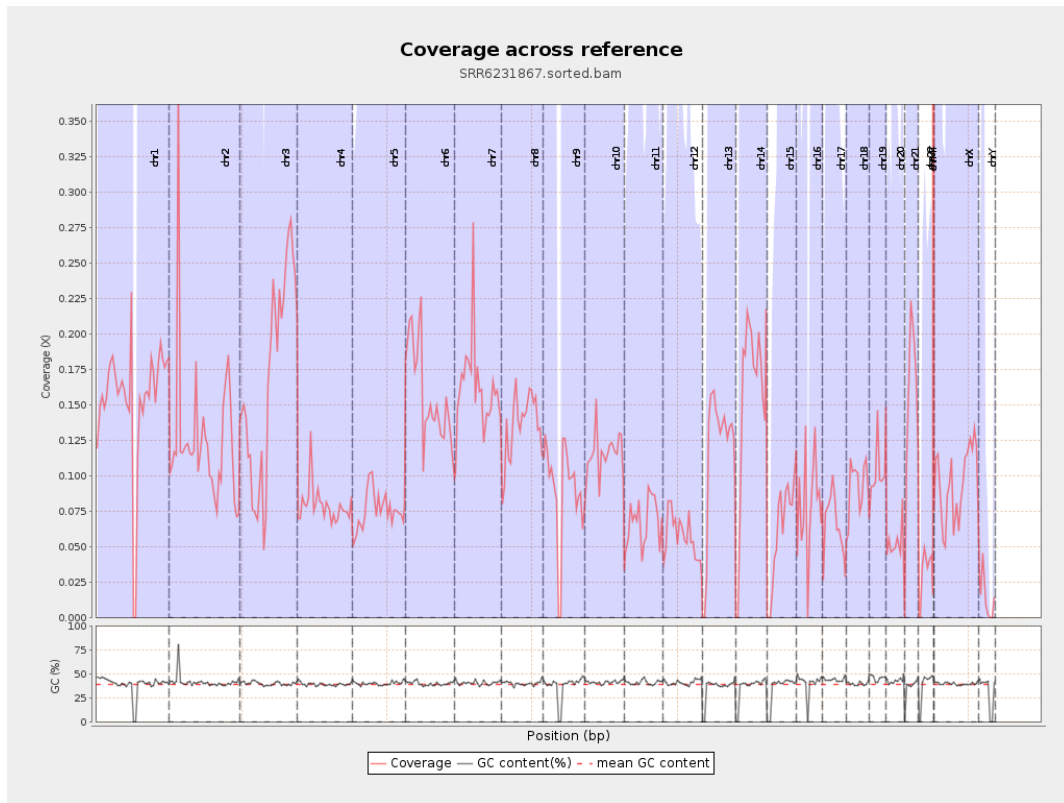
General error rate	0.63%
Mismatches	2,080,505
Insertions	23,615
Mapped reads with at least one insertion	0.46%
Deletions	85,730
Mapped reads with at least one deletion	1.65%
Homopolymer indels	45.67%

2.6. Chromosome stats

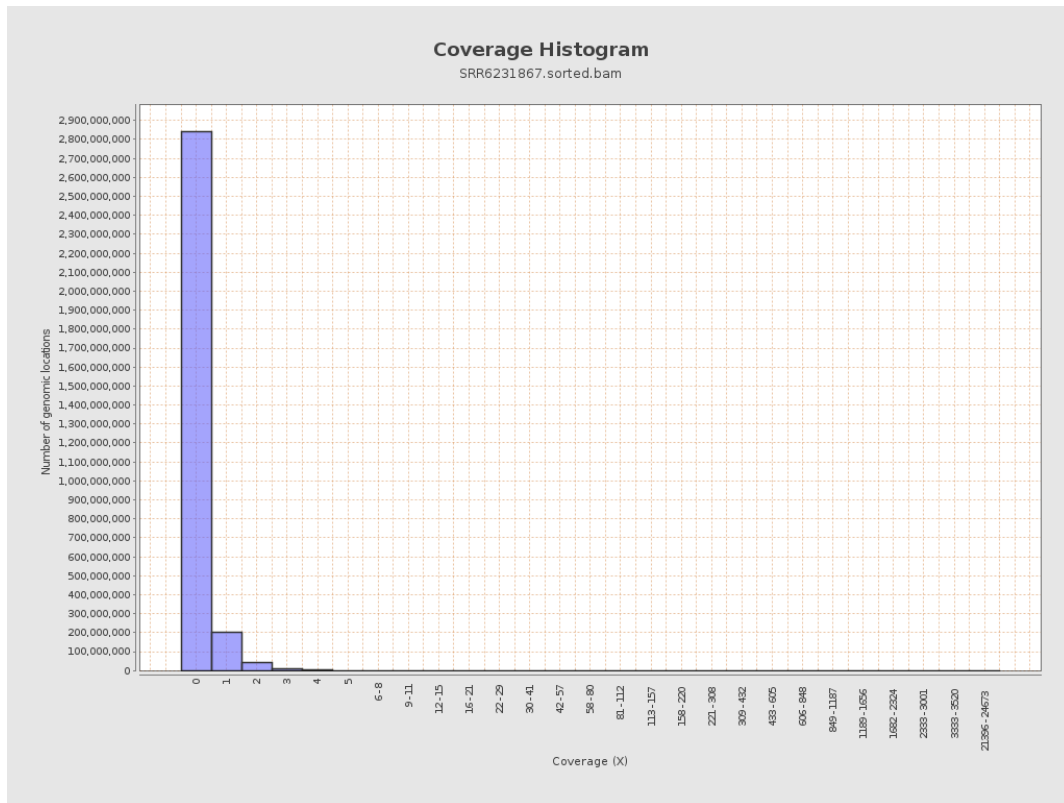
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	38351581	0.1539	2.293
chr2	243199373	30521192	0.1255	13.4145
chr3	198022430	33217395	0.1677	0.512
chr4	191154276	15068057	0.0788	0.3898
chr5	180915260	13638441	0.0754	0.3524
chr6	171115067	26898587	0.1572	0.7203
chr7	159138663	25950104	0.1631	1.9059

chr8	146364022	19776509	0.1351	1.4745
chr9	141213431	12647328	0.0896	0.667
chr10	135534747	15680628	0.1157	0.7518
chr11	135006516	9146952	0.0678	0.5854
chr12	133851895	7682924	0.0574	0.3139
chr13	115169878	13353289	0.1159	0.4192
chr14	107349540	16422997	0.153	0.5641
chr15	102531392	6348938	0.0619	0.3093
chr16	90354753	7155061	0.0792	0.4888
chr17	81195210	5273416	0.0649	0.3735
chr18	78077248	7252239	0.0929	1.6016
chr19	59128983	6082495	0.1029	1.389
chr20	63025520	3345543	0.0531	0.3445
chr21	48129895	7286735	0.1514	0.5151
chr22	51304566	1617962	0.0315	0.212
chrMT	16571	253701	15.3099	8.4131
chrX	155270560	14716465	0.0948	0.4681
chrY	59373566	799939	0.0135	0.258

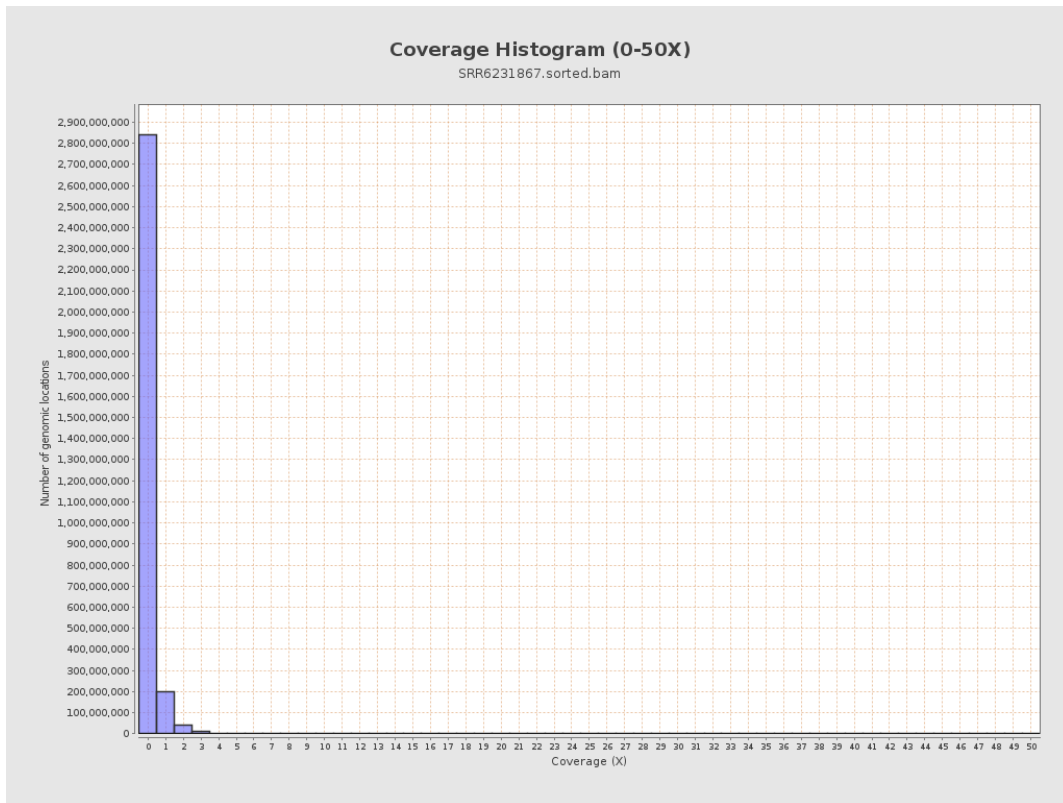
3. Results : Coverage across reference



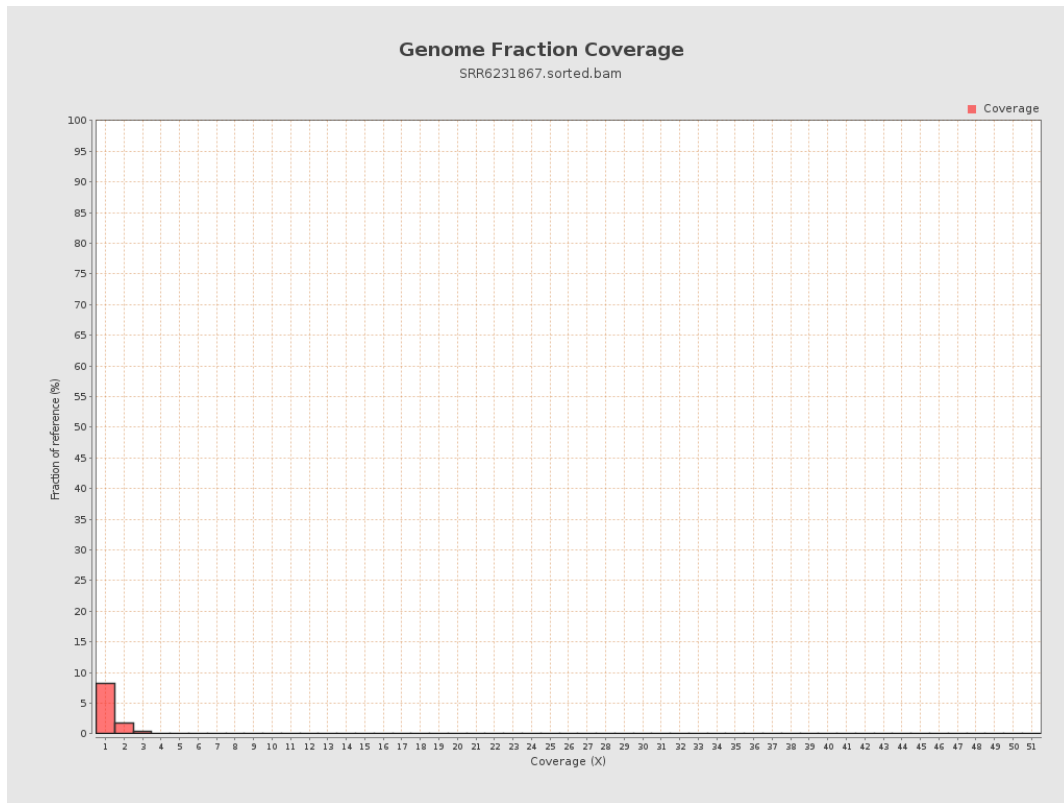
4. Results : Coverage Histogram



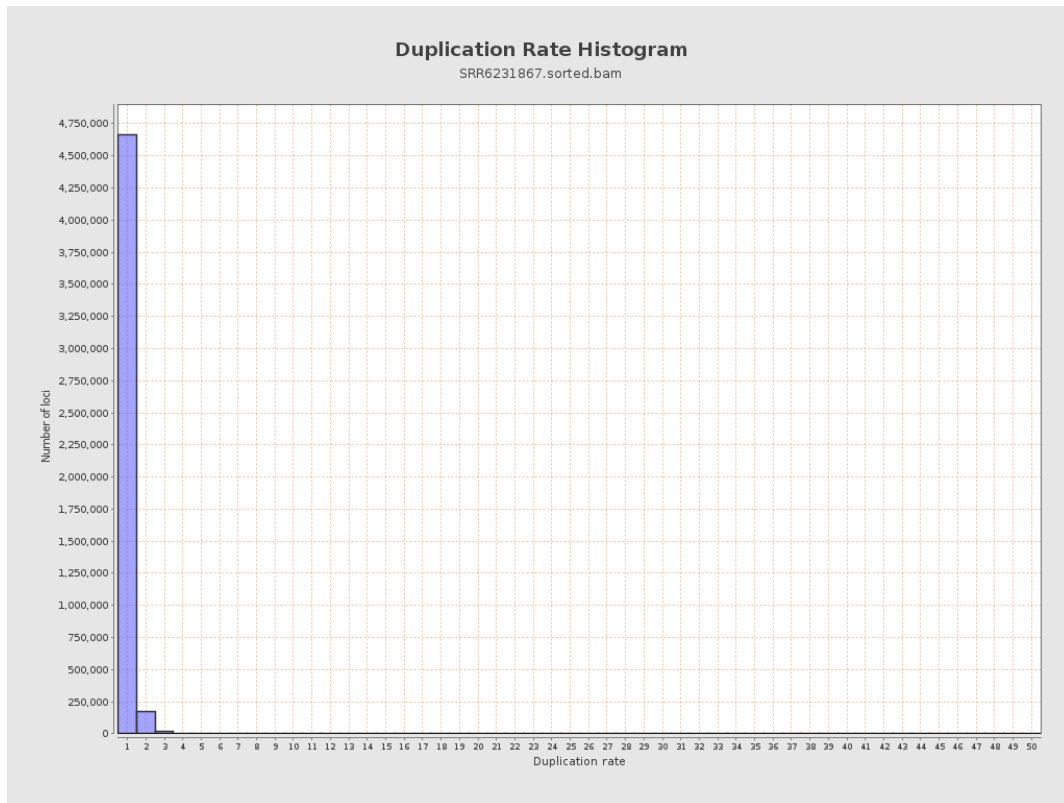
5. Results : Coverage Histogram (0-50X)



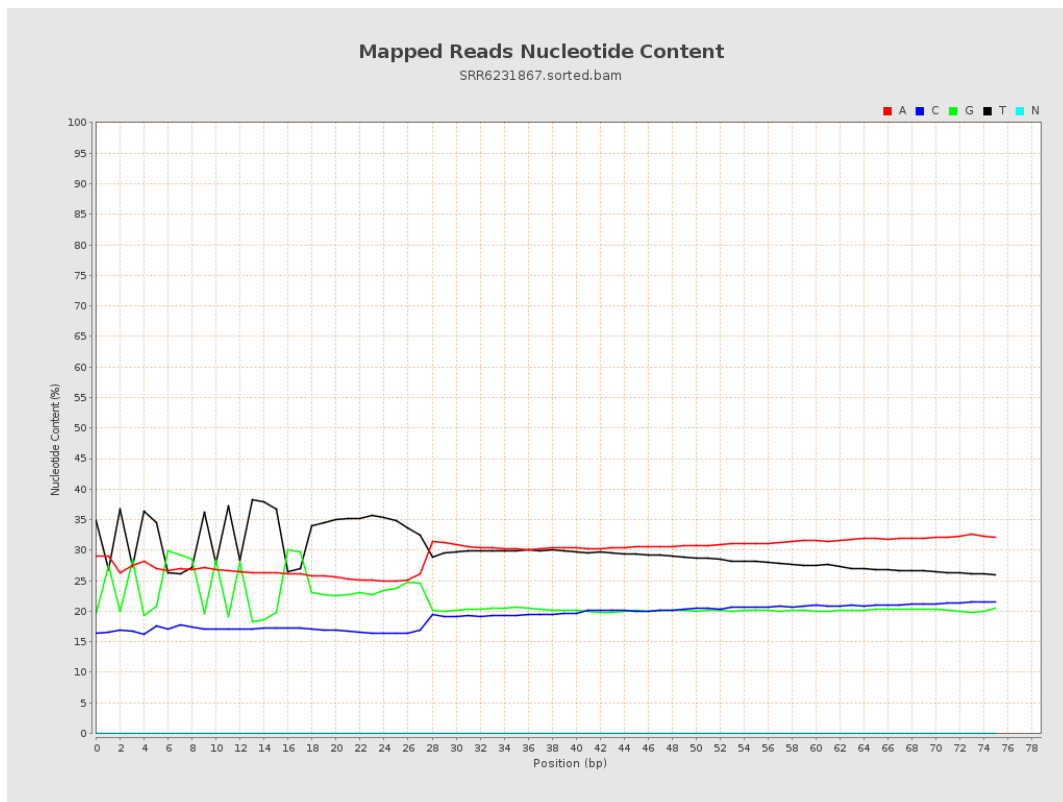
6. Results : Genome Fraction Coverage



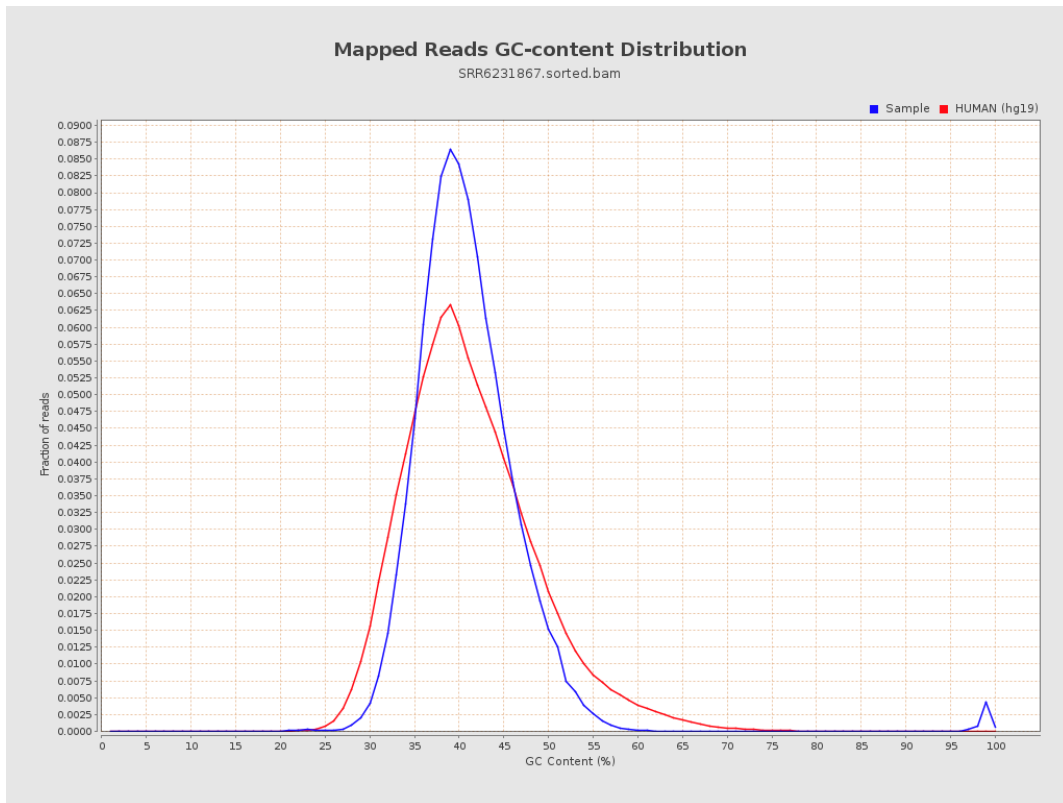
7. Results : Duplication Rate Histogram



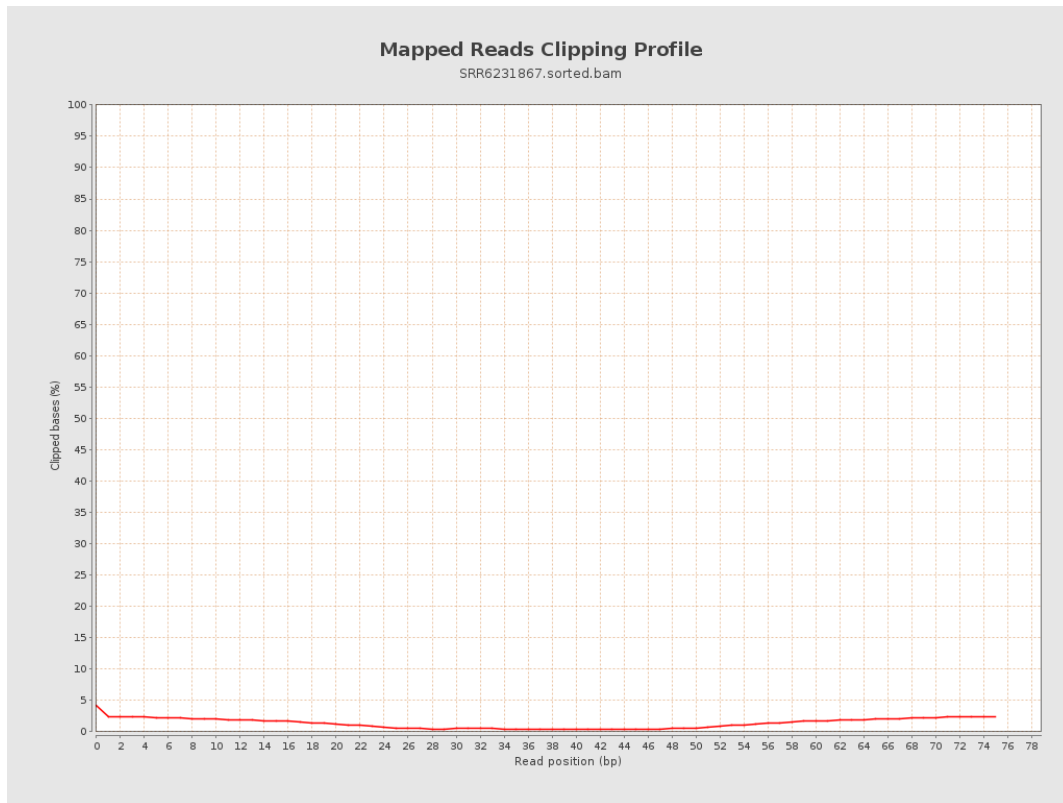
8. Results : Mapped Reads Nucleotide Content



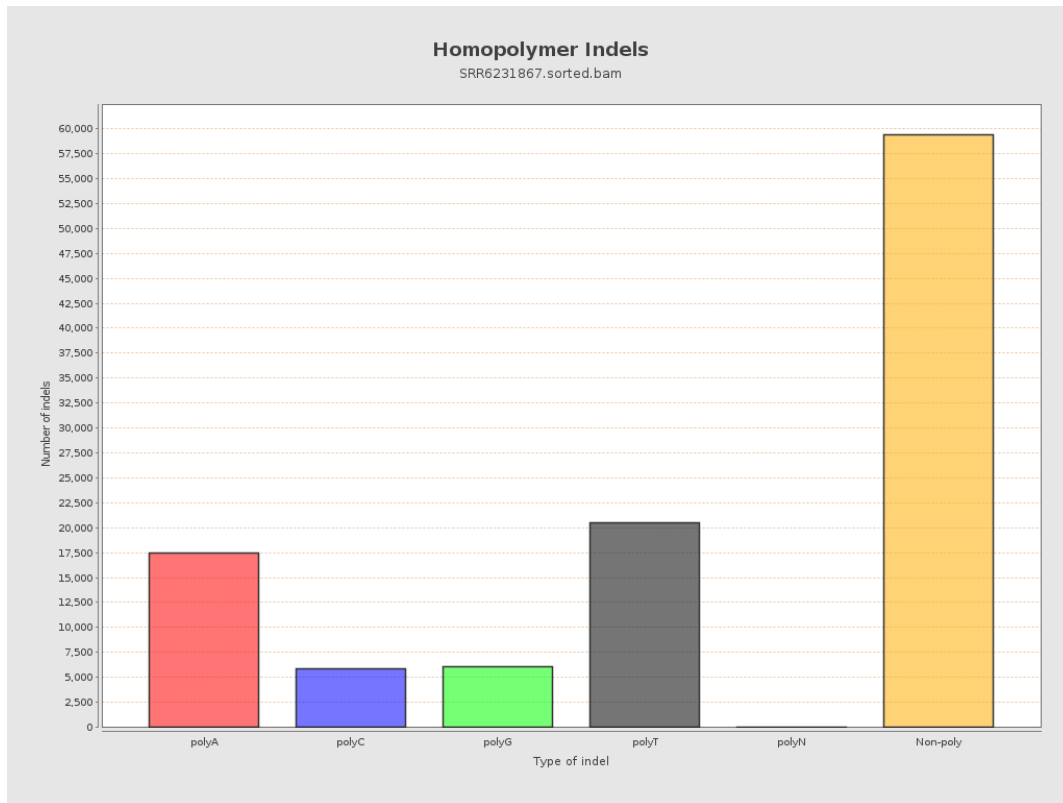
9. Results : Mapped Reads GC-content Distribution



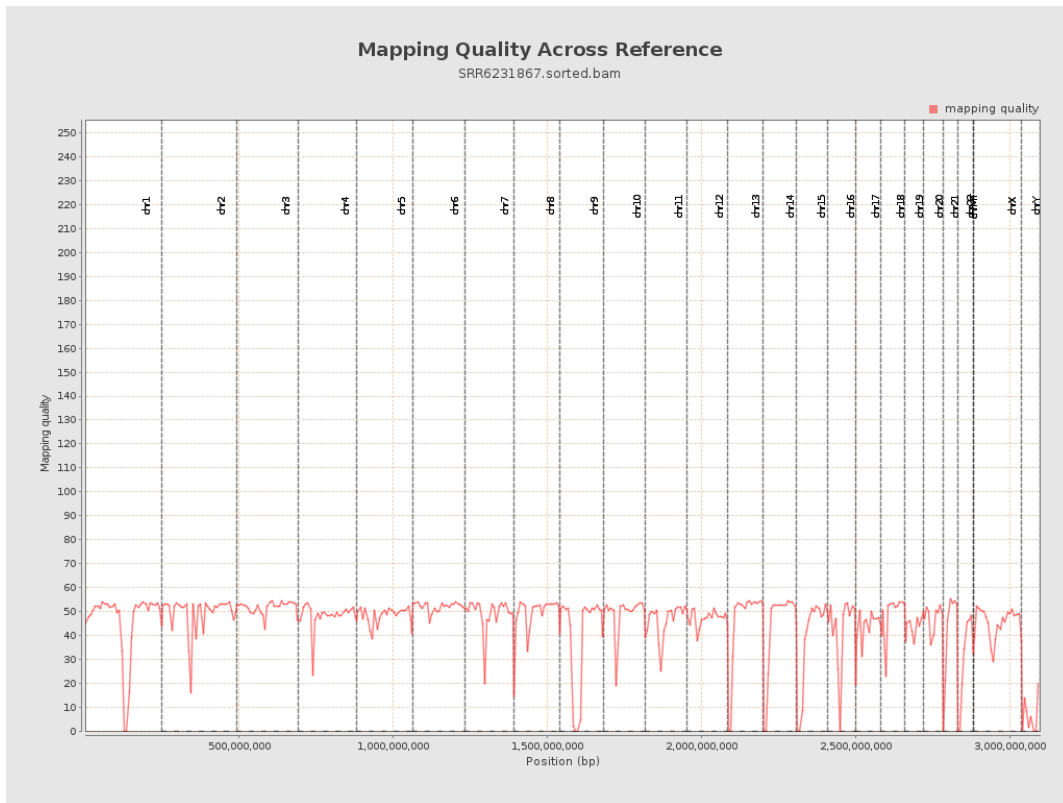
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

