

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/03 12:05:36

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR9716785.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_SRR9716785 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR9716785.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Sep 03 12:05:35 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR9716785.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,432,719
Mapped reads	3,139,951 / 91.47%
Unmapped reads	292,768 / 8.53%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	65,546 / 1.91%
Read min/max/mean length	30 / 101 / 101.69
Duplicated reads (estimated)	151,555 / 4.42%
Duplication rate	3.6%
Clipped reads	3,203,617 / 93.33%

2.2. ACGT Content

Number/percentage of A's	60,005,818 / 25.92%
Number/percentage of C's	47,268,499 / 20.42%
Number/percentage of T's	69,218,555 / 29.9%
Number/percentage of G's	54,974,874 / 23.75%
Number/percentage of N's	27,040 / 0.01%
GC Percentage	44.17%

2.3. Coverage

Mean	0.0748

Standard Deviation	0.508
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2.4. Mapping Quality

Mean Mapping Quality	46.41
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2.5. Mismatches and indels

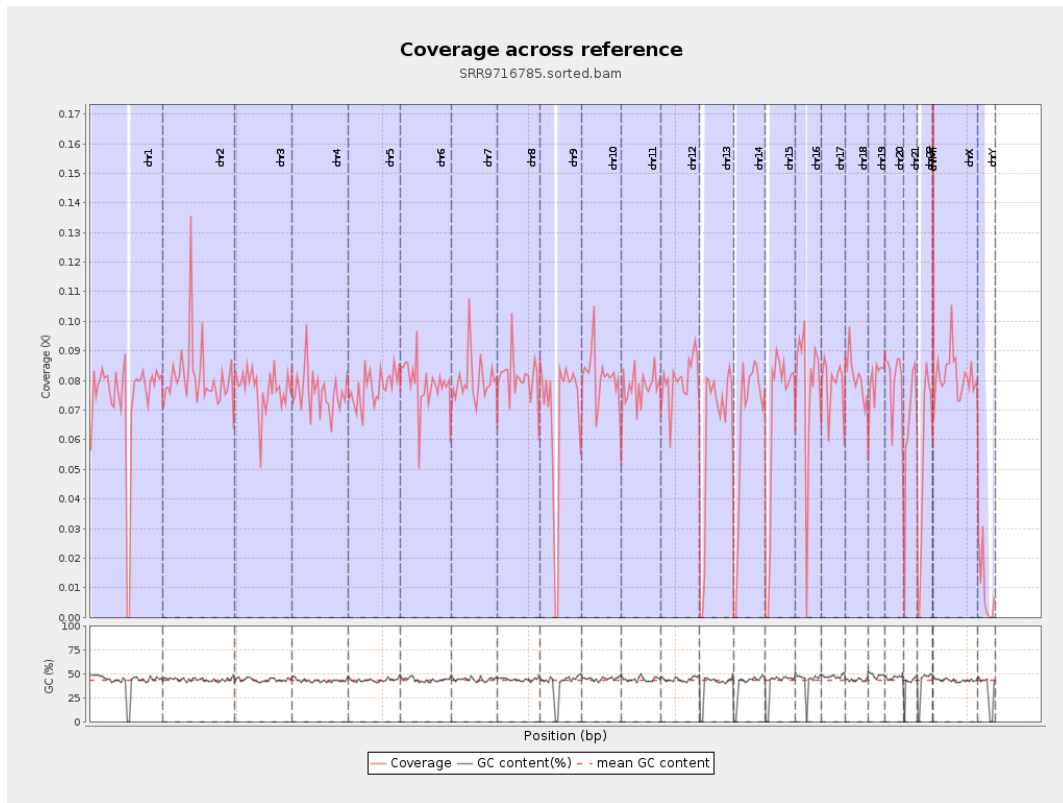
General error rate	0.71%
Mismatches	1,617,038
Insertions	16,722
Mapped reads with at least one insertion	0.53%
Deletions	45,560
Mapped reads with at least one deletion	1.43%
Homopolymer indels	42.34%

2.6. Chromosome stats

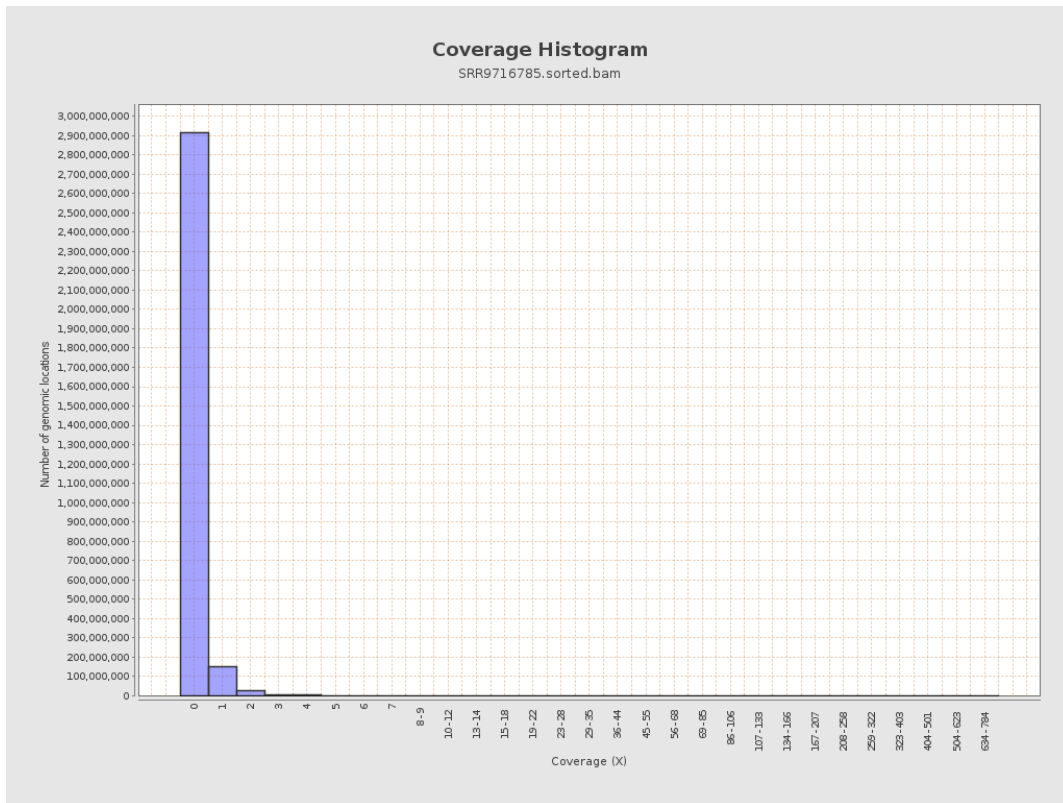
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	18296548	0.0734	0.5579
chr2	243199373	19777465	0.0813	0.7194
chr3	198022430	15263373	0.0771	0.3373
chr4	191154276	14560766	0.0762	0.3739
chr5	180915260	14034064	0.0776	0.3444
chr6	171115067	13462747	0.0787	0.4113
chr7	159138663	12785665	0.0803	0.6832

chr8	146364022	11807263	0.0807	0.6859
chr9	141213431	9616293	0.0681	0.5431
chr10	135534747	11000955	0.0812	0.5149
chr11	135006516	10464262	0.0775	0.5413
chr12	133851895	10776216	0.0805	0.3549
chr13	115169878	7288480	0.0633	0.3052
chr14	107349540	7036549	0.0655	0.4022
chr15	102531392	6818457	0.0665	0.322
chr16	90354753	6901576	0.0764	0.3871
chr17	81195210	6434606	0.0792	0.3777
chr18	78077248	6366317	0.0815	1.0295
chr19	59128983	4754811	0.0804	0.5351
chr20	63025520	5029693	0.0798	0.364
chr21	48129895	3107447	0.0646	0.3521
chr22	51304566	2756533	0.0537	0.2853
chrMT	16571	7597	0.4585	0.9626
chrX	155270560	12661463	0.0815	0.4557
chrY	59373566	563494	0.0095	0.258

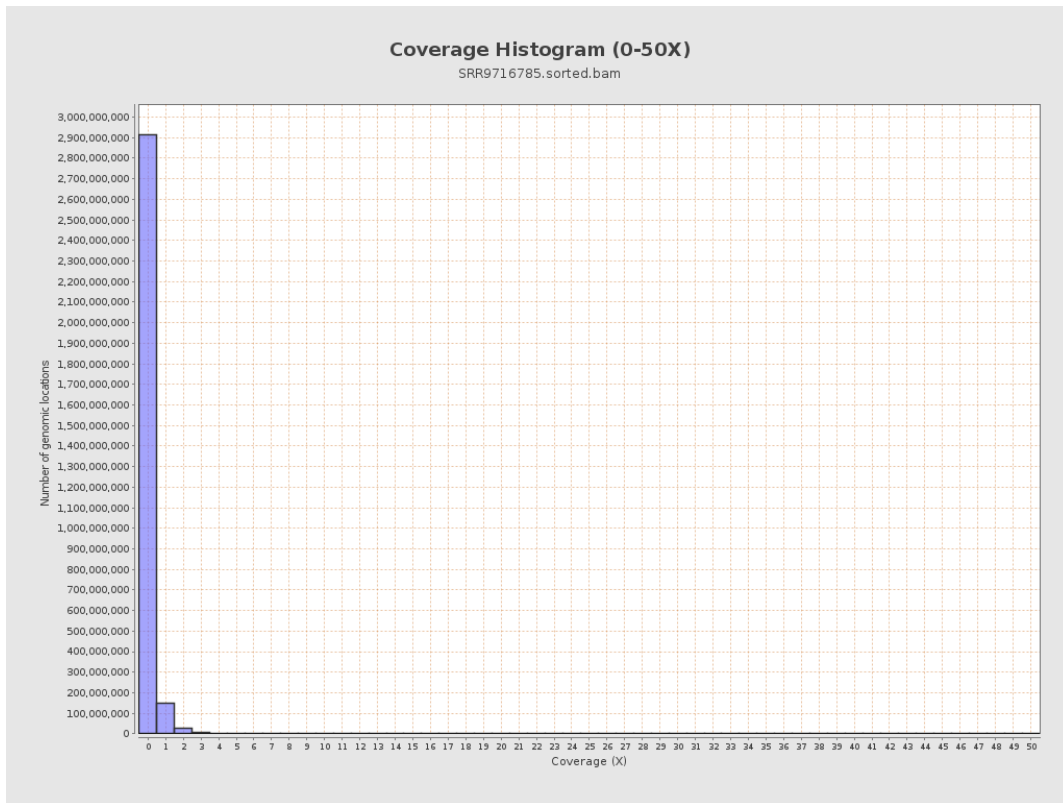
3. Results : Coverage across reference



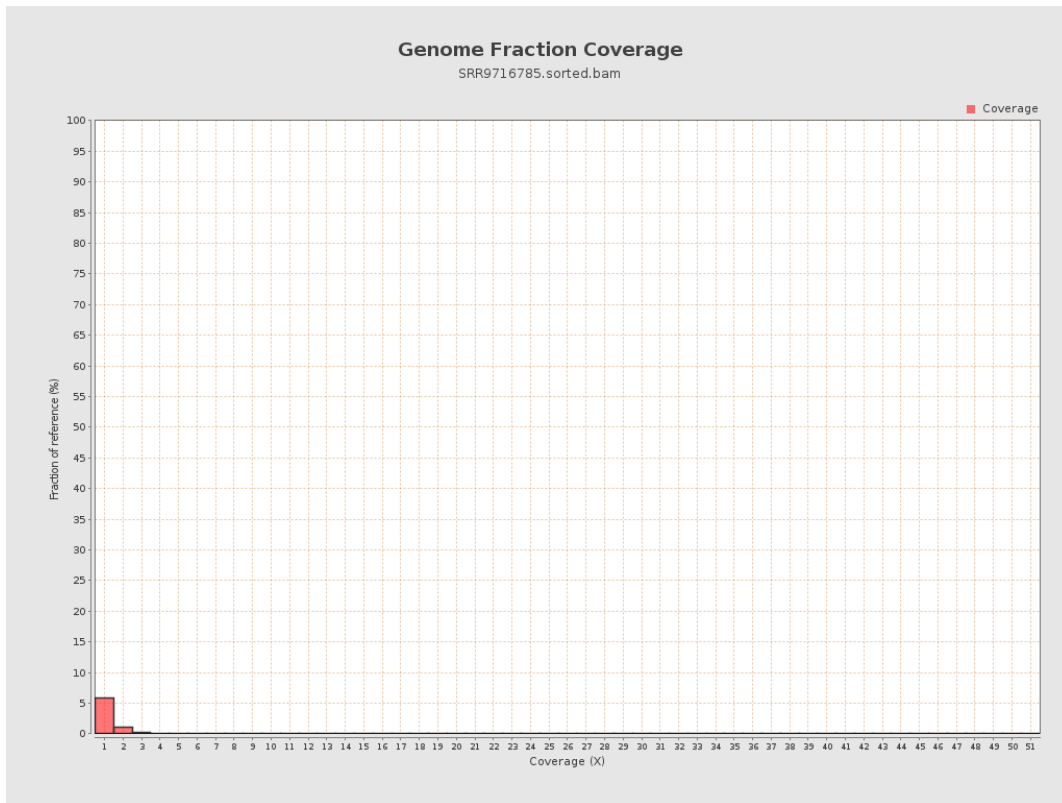
4. Results : Coverage Histogram



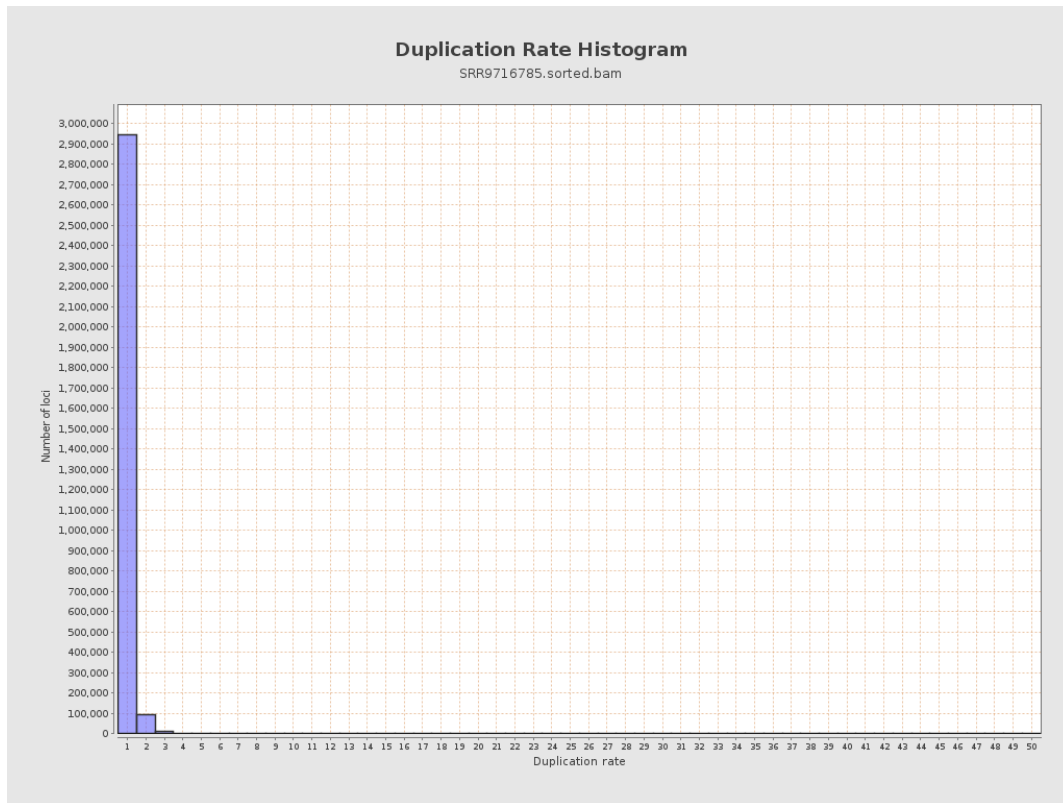
5. Results : Coverage Histogram (0-50X)



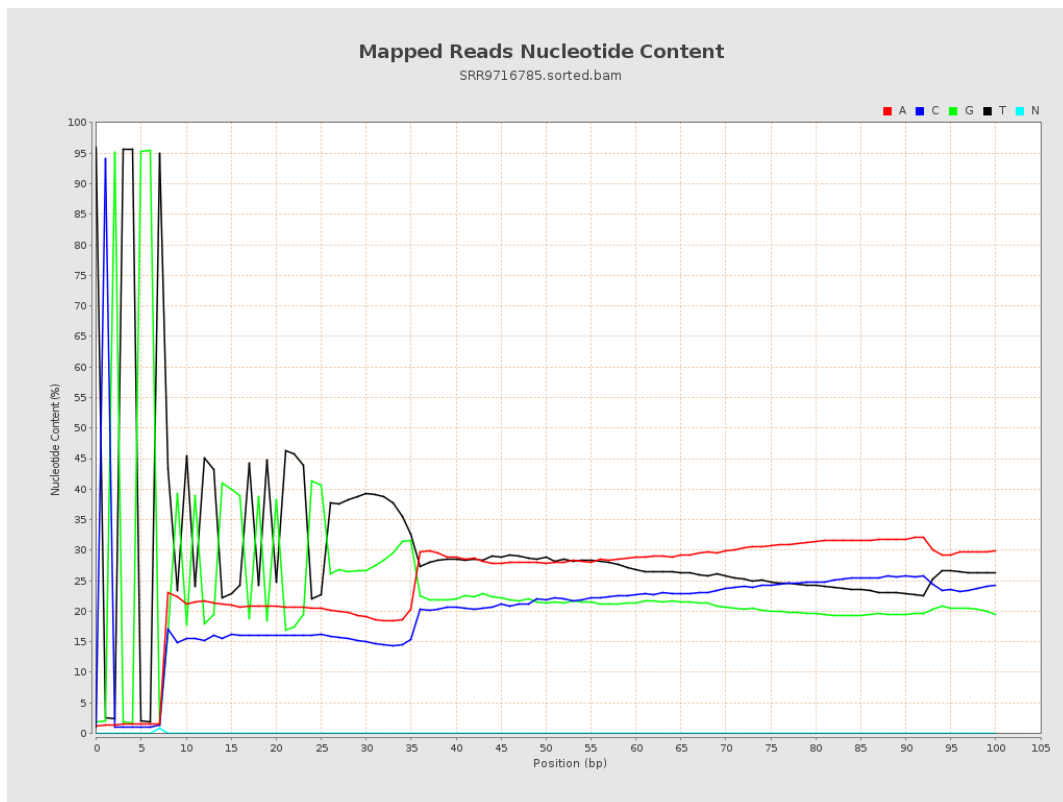
6. Results : Genome Fraction Coverage



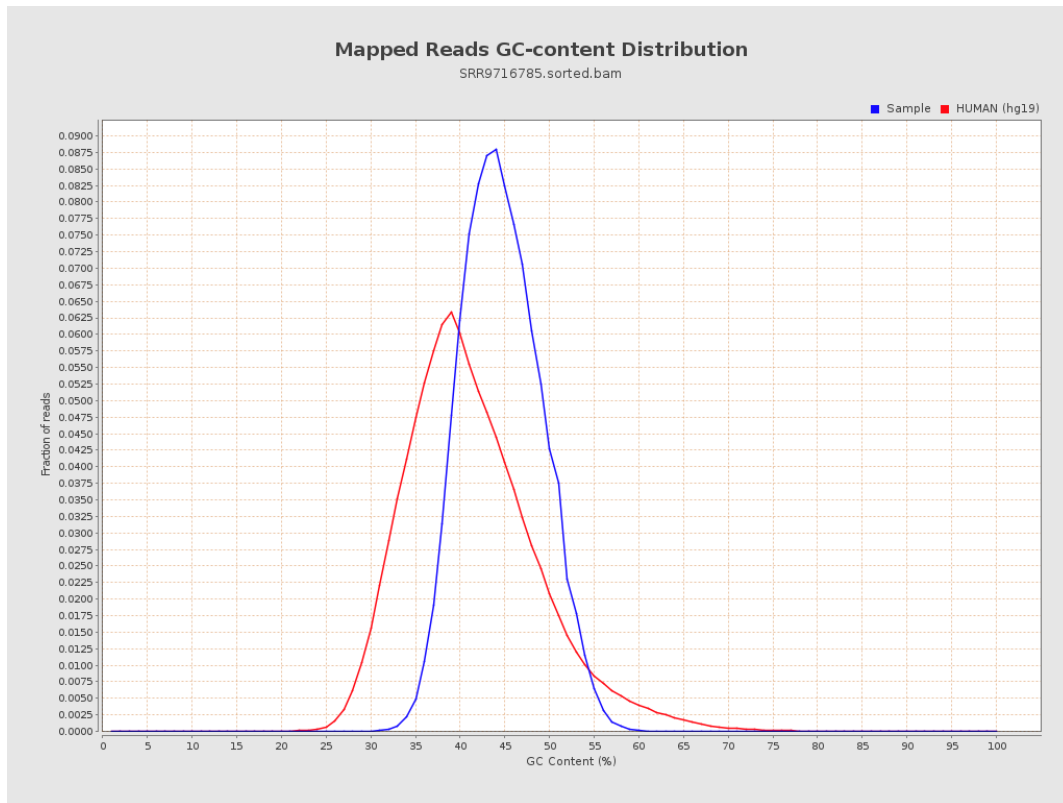
7. Results : Duplication Rate Histogram



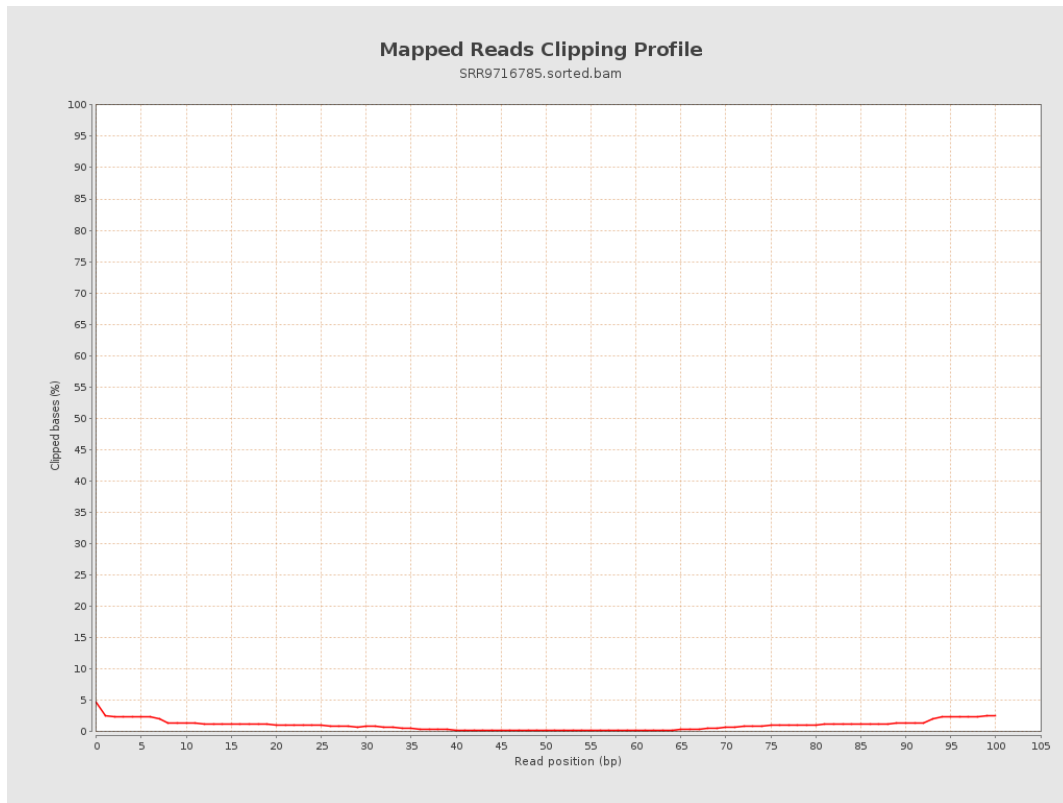
8. Results : Mapped Reads Nucleotide Content



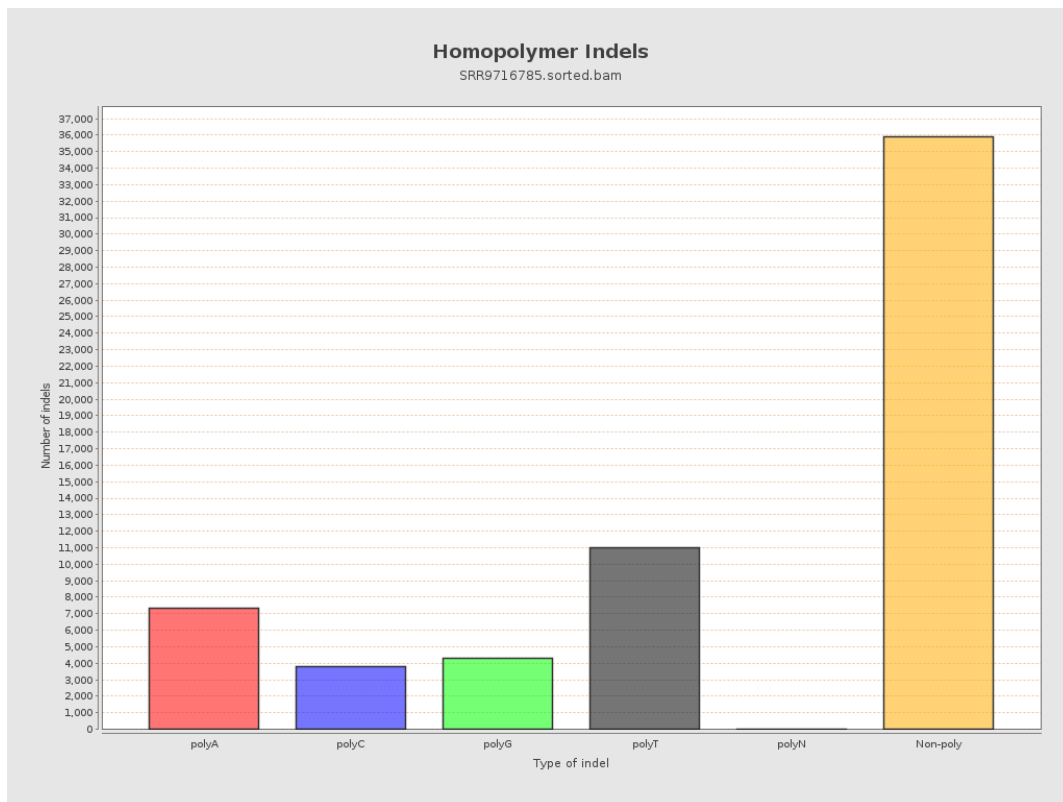
9. Results : Mapped Reads GC-content Distribution



10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

