

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/27 20:58:40

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR10524283.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_SRR10524283 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR10524283.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Aug 27 20:58:39 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR10524283.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	1,396,093
Mapped reads	1,285,565 / 92.08%
Unmapped reads	110,528 / 7.92%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	4,246 / 0.3%
Read min/max/mean length	30 / 76 / 76.1
Duplicated reads (estimated)	49,219 / 3.53%
Duplication rate	2.75%
Clipped reads	1,286,316 / 92.14%

2.2. ACGT Content

Number/percentage of A's	19,349,999 / 25.65%
Number/percentage of C's	13,017,863 / 17.26%
Number/percentage of T's	25,241,519 / 33.46%
Number/percentage of G's	17,811,511 / 23.61%
Number/percentage of N's	10,521 / 0.01%
GC Percentage	40.87%

2.3. Coverage

Mean	0.0244

Standard Deviation	0.2728
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2.4. Mapping Quality

Mean Mapping Quality	46.72
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2.5. Mismatches and indels

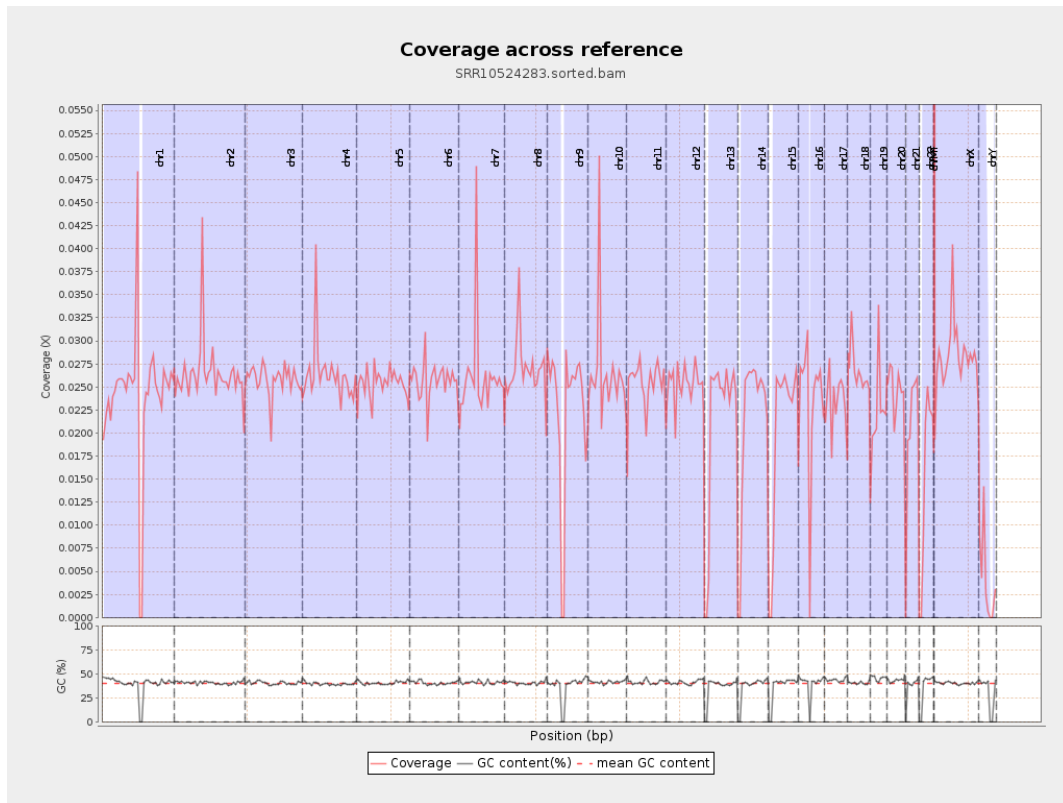
General error rate	0.51%
Mismatches	371,447
Insertions	5,188
Mapped reads with at least one insertion	0.4%
Deletions	14,421
Mapped reads with at least one deletion	1.11%
Homopolymer indels	44.21%

2.6. Chromosome stats

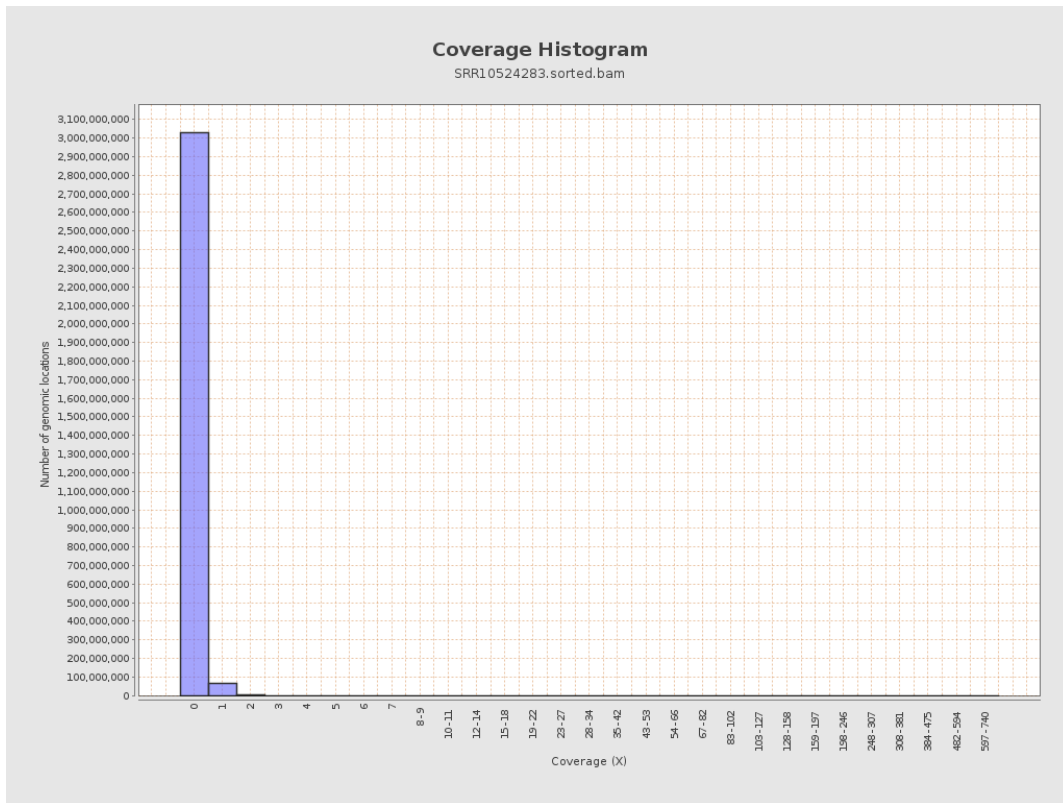
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	5984114	0.024	0.5601
chr2	243199373	6430458	0.0264	0.2722
chr3	198022430	5106583	0.0258	0.1731
chr4	191154276	5025128	0.0263	0.1938
chr5	180915260	4619012	0.0255	0.1728
chr6	171115067	4397958	0.0257	0.1866
chr7	159138663	4164777	0.0262	0.3784

chr8	146364022	3946088	0.027	0.2809
chr9	141213431	3168258	0.0224	0.2346
chr10	135534747	3582661	0.0264	0.2524
chr11	135006516	3447121	0.0255	0.2267
chr12	133851895	3389873	0.0253	0.1764
chr13	115169878	2430991	0.0211	0.1557
chr14	107349540	2287263	0.0213	0.1645
chr15	102531392	2082218	0.0203	0.1526
chr16	90354753	2086780	0.0231	0.178
chr17	81195210	1896703	0.0234	0.1805
chr18	78077248	2089259	0.0268	0.4422
chr19	59128983	1333039	0.0225	0.3507
chr20	63025520	1530900	0.0243	0.1717
chr21	48129895	989566	0.0206	0.1729
chr22	51304566	806250	0.0157	0.1341
chrMT	16571	11347	0.6848	0.9405
chrX	155270560	4411107	0.0284	0.2082
chrY	59373566	236908	0.004	0.1152

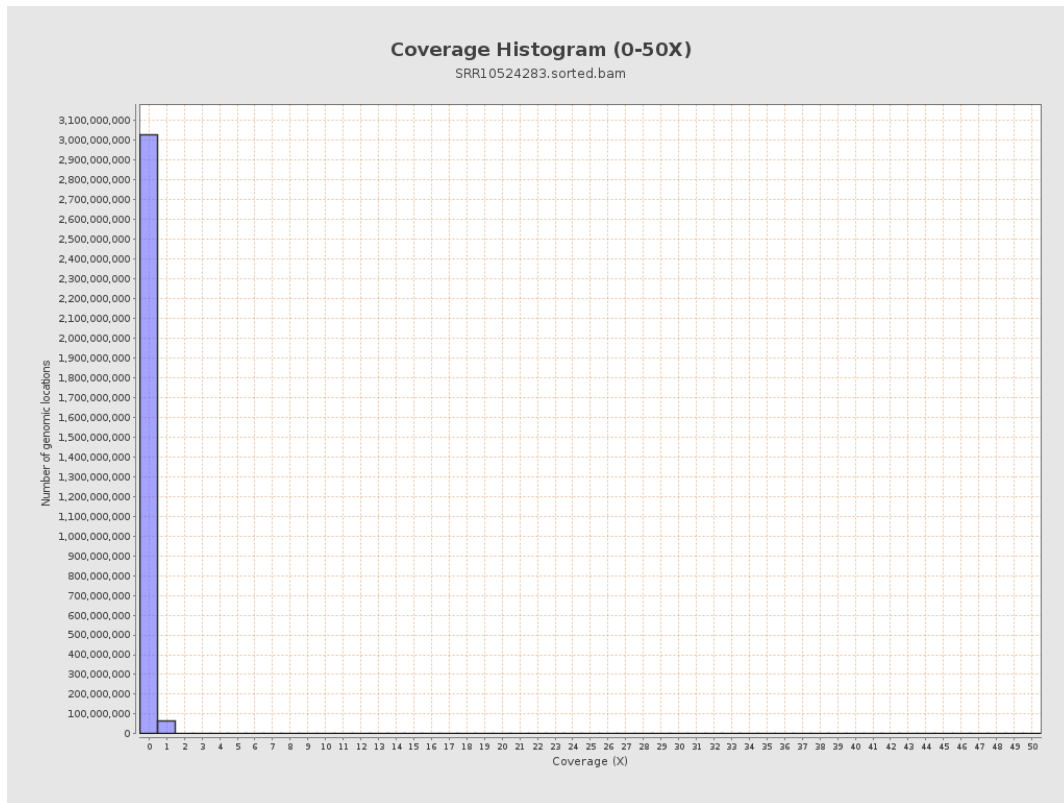
3. Results : Coverage across reference



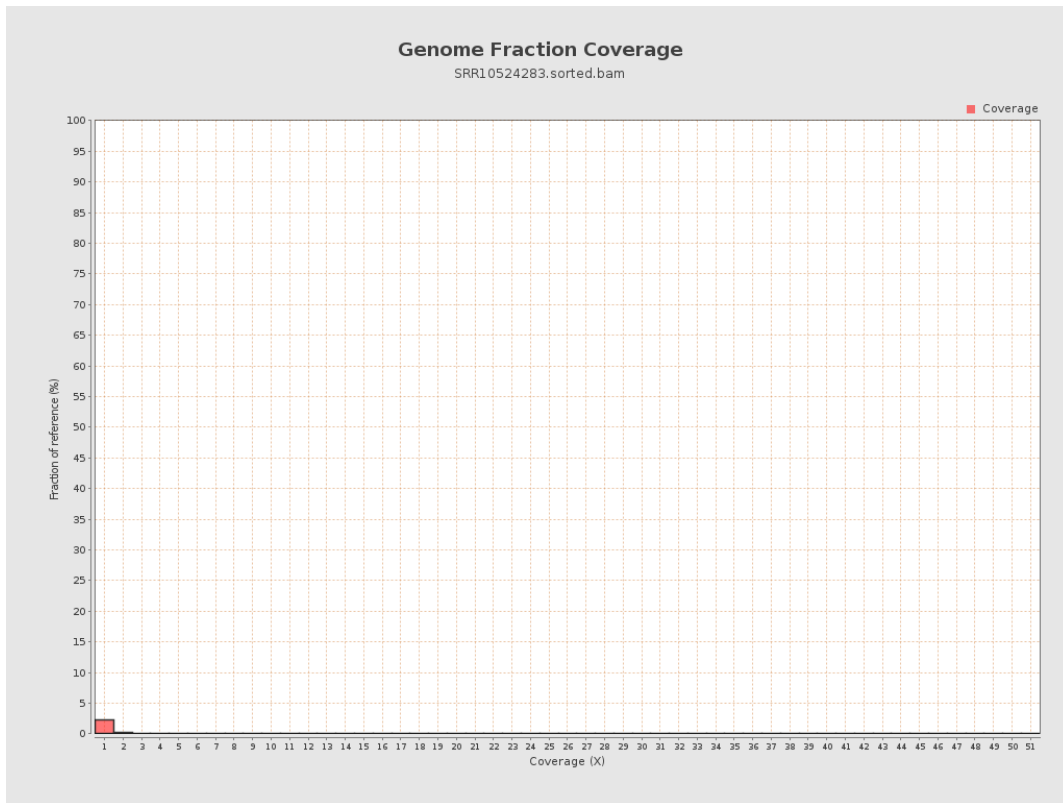
4. Results : Coverage Histogram



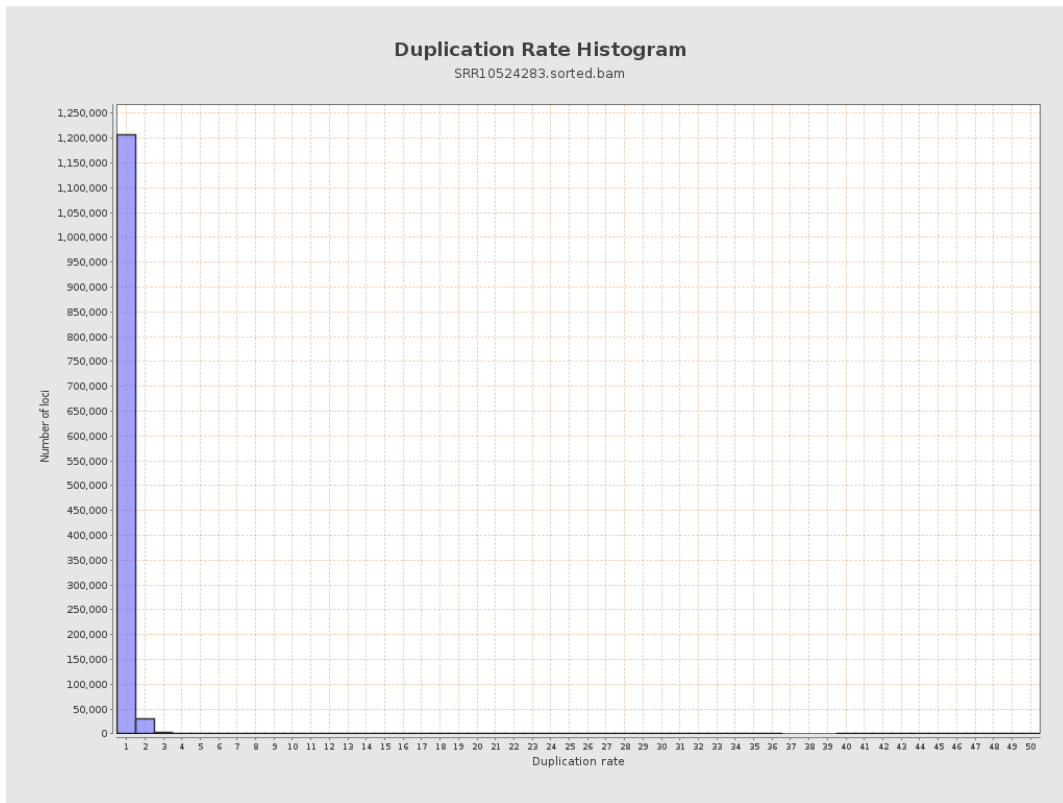
5. Results : Coverage Histogram (0-50X)



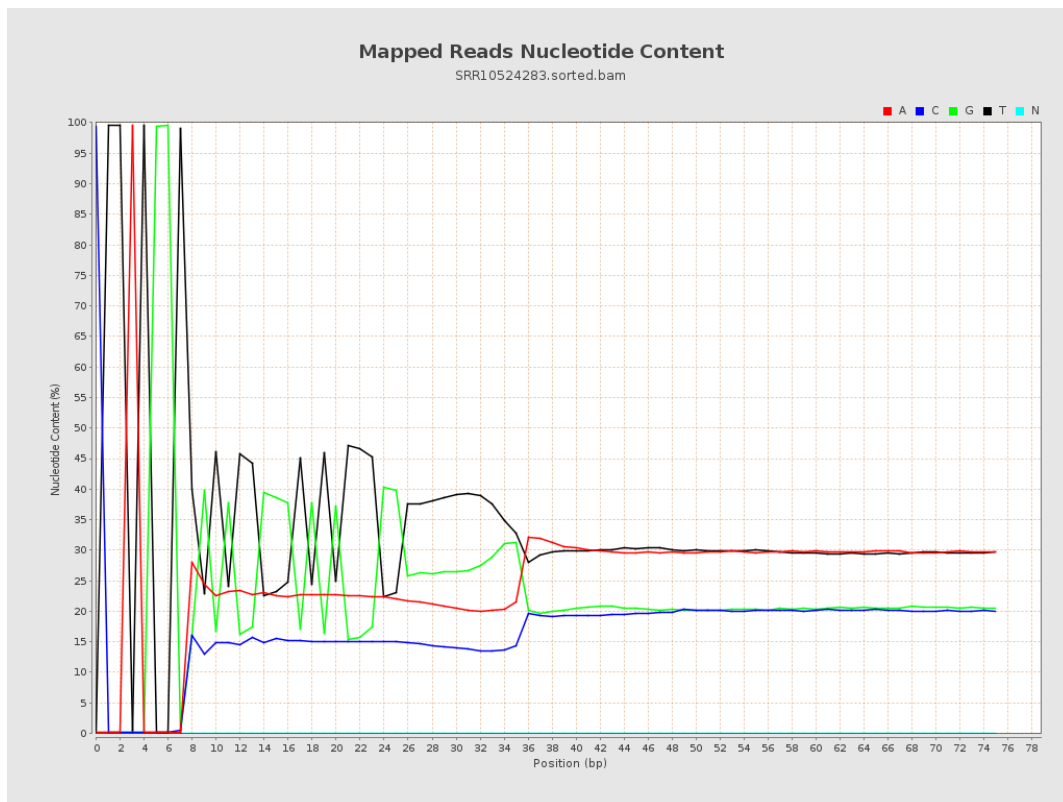
6. Results : Genome Fraction Coverage



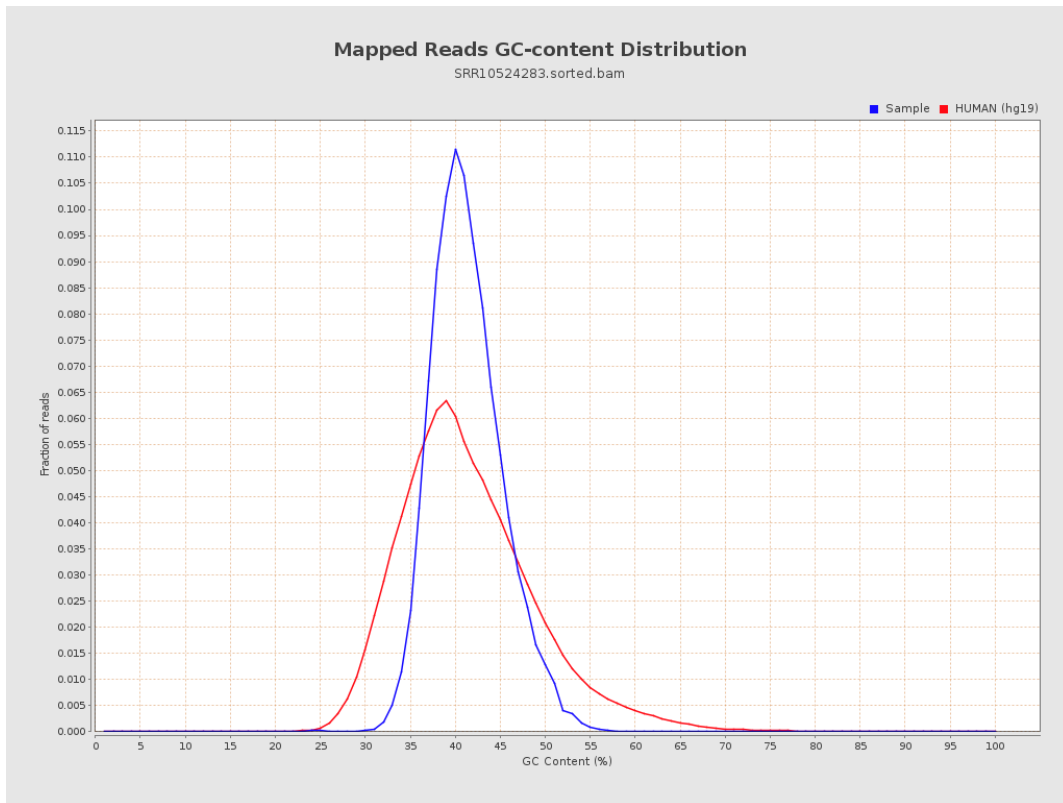
7. Results : Duplication Rate Histogram



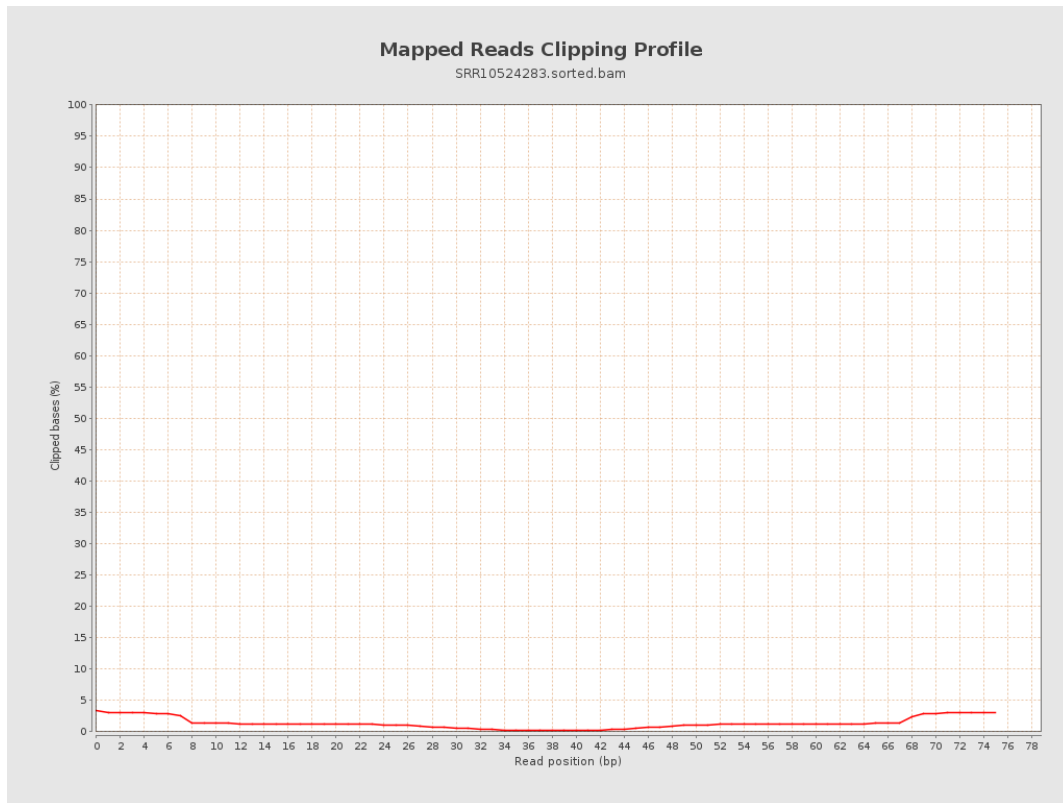
8. Results : Mapped Reads Nucleotide Content



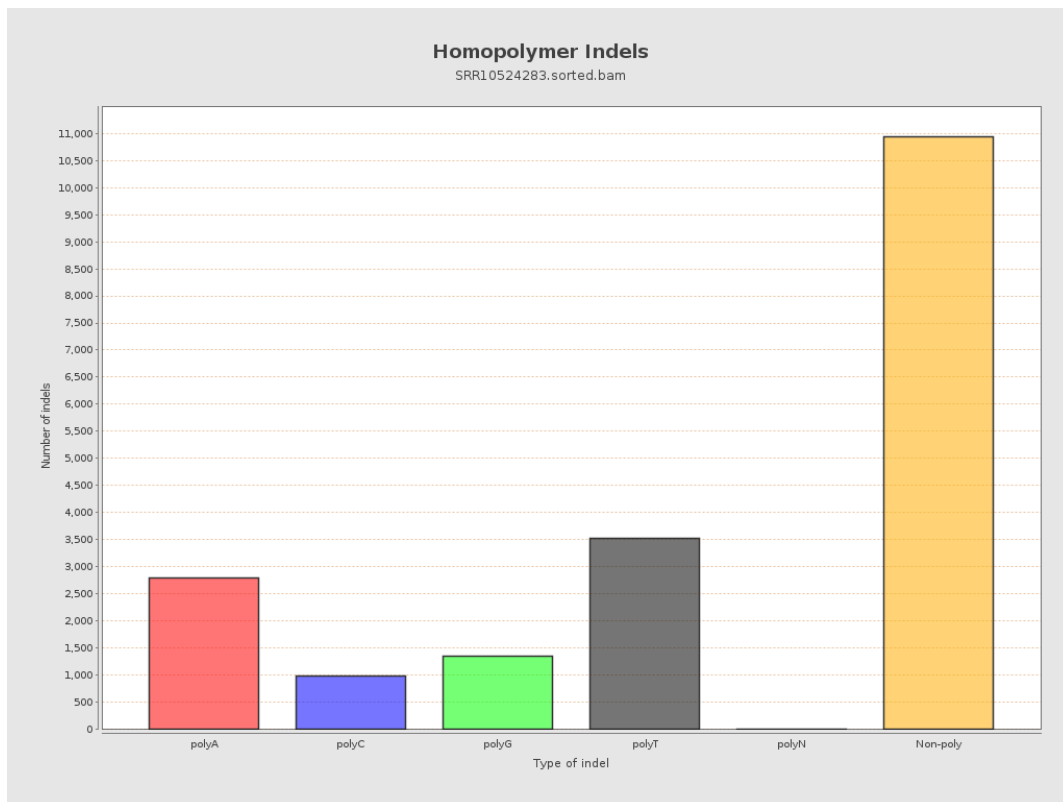
9. Results : Mapped Reads GC-content Distribution



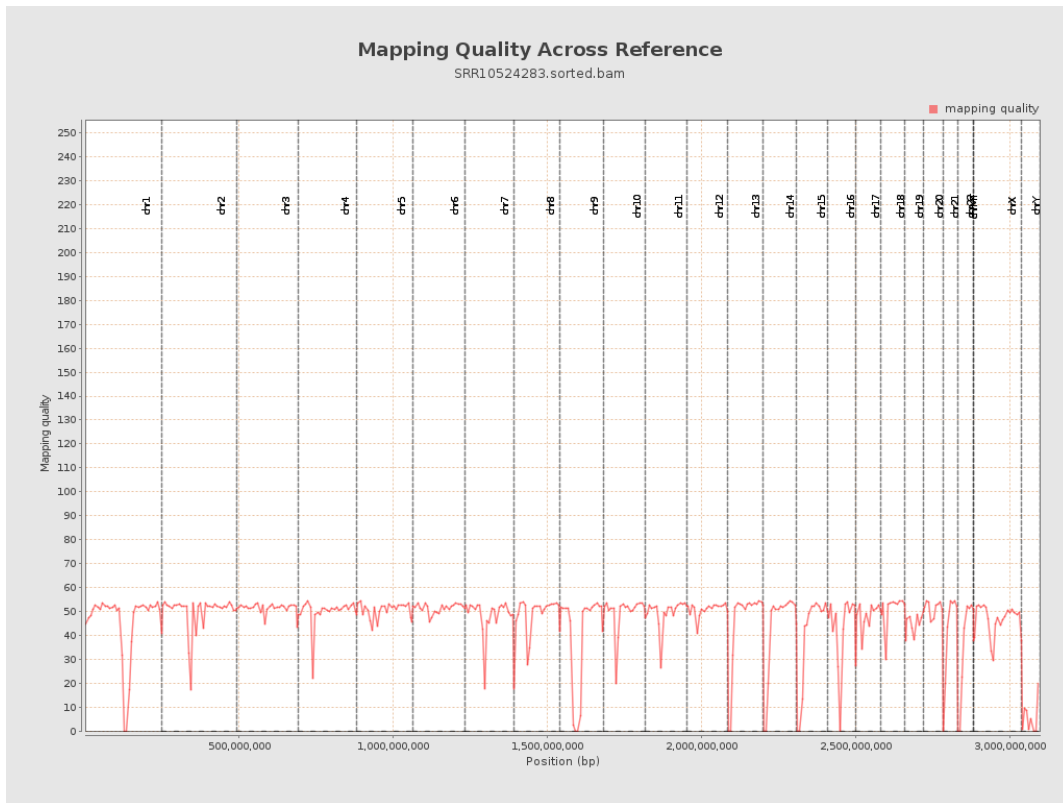
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

