

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/12/20 01:31:33

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR1506315.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_SRR1506315 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR1506315.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Fri Dec 20 01:31:33 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR1506315.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	9,889,590
Mapped reads	7,237,427 / 73.18%
Unmapped reads	2,652,163 / 26.82%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	116,627 / 1.18%
Read min/max/mean length	30 / 94 / 94.42
Duplicated reads (estimated)	1,462,154 / 14.78%
Duplication rate	14.97%
Clipped reads	4,380,889 / 44.3%

2.2. ACGT Content

Number/percentage of A's	166,536,237 / 28.95%
Number/percentage of C's	102,423,192 / 17.81%
Number/percentage of T's	161,127,666 / 28.01%
Number/percentage of G's	144,976,871 / 25.21%
Number/percentage of N's	96,070 / 0.02%
GC Percentage	43.01%

2.3. Coverage

Mean	0.1859

Standard Deviation	1.2386
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2.4. Mapping Quality

Mean Mapping Quality	47.51
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2.5. Mismatches and indels

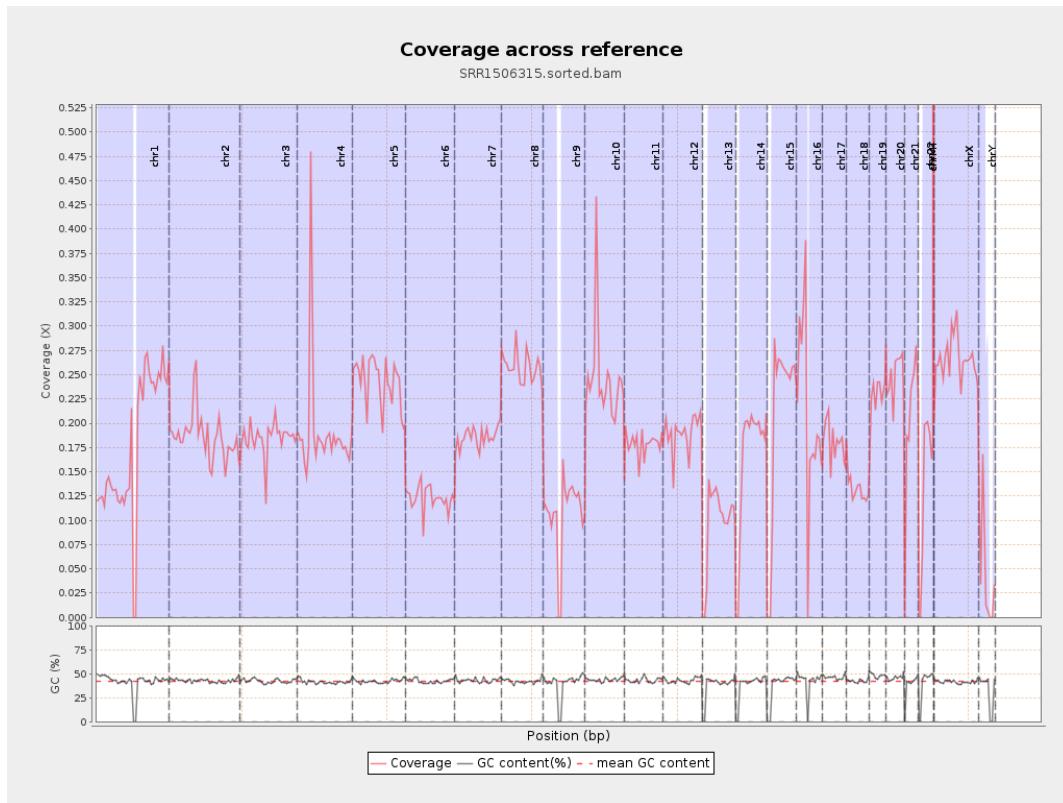
General error rate	0.88%
Mismatches	4,929,041
Insertions	53,003
Mapped reads with at least one insertion	0.71%
Deletions	128,439
Mapped reads with at least one deletion	1.74%
Homopolymer indels	43.19%

2.6. Chromosome stats

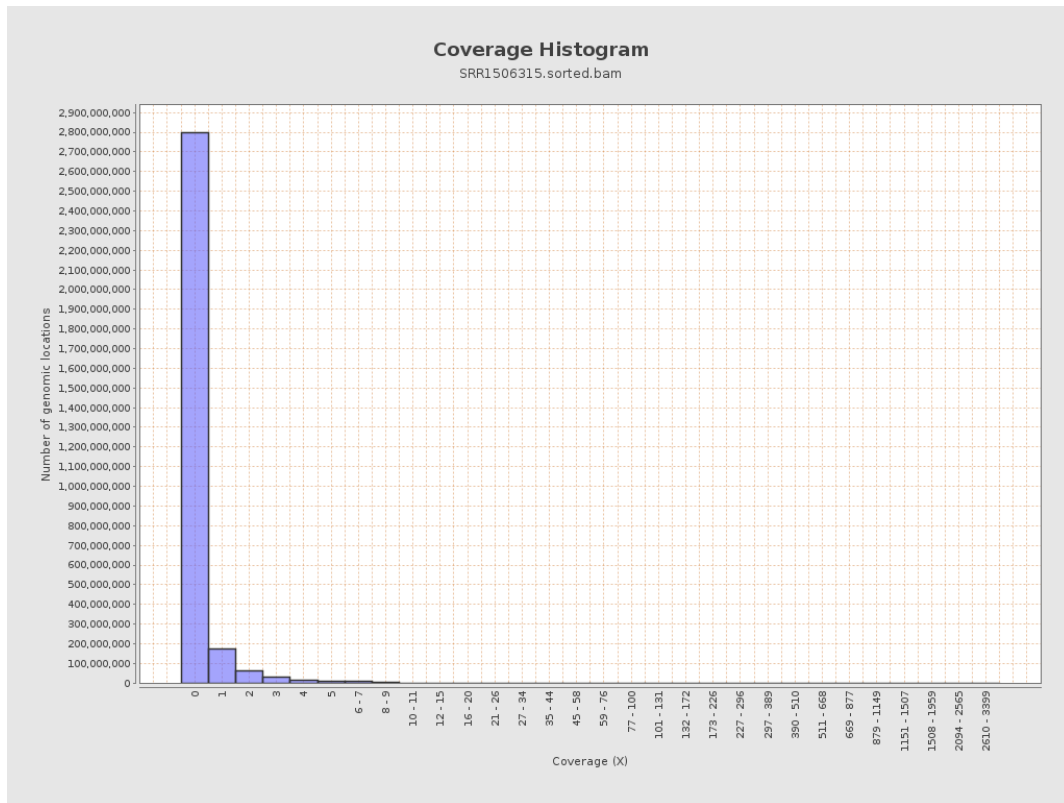
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	43663608	0.1752	1.8242
chr2	243199373	45438844	0.1868	1.2374
chr3	198022430	36494210	0.1843	0.7469
chr4	191154276	36174067	0.1892	1.313
chr5	180915260	43731786	0.2417	0.8844
chr6	171115067	21098546	0.1233	0.6533
chr7	159138663	29549290	0.1857	1.1341

chr8	146364022	37640445	0.2572	2.4371
chr9	141213431	15016991	0.1063	0.9528
chr10	135534747	33223569	0.2451	1.6208
chr11	135006516	24127597	0.1787	1.0784
chr12	133851895	25338832	0.1893	0.7835
chr13	115169878	11236715	0.0976	0.5366
chr14	107349540	17594746	0.1639	0.7716
chr15	102531392	21525408	0.2099	0.8214
chr16	90354753	19181816	0.2123	1.1239
chr17	81195210	14578387	0.1795	0.869
chr18	78077248	10320908	0.1322	1.5805
chr19	59128983	13629604	0.2305	1.4601
chr20	63025520	15636112	0.2481	0.9793
chr21	48129895	9932421	0.2064	1.0896
chr22	51304566	6702643	0.1306	0.6536
chrMT	16571	55034	3.3211	3.9846
chrX	155270560	41098064	0.2647	0.9975
chrY	59373566	2435061	0.041	1.4861

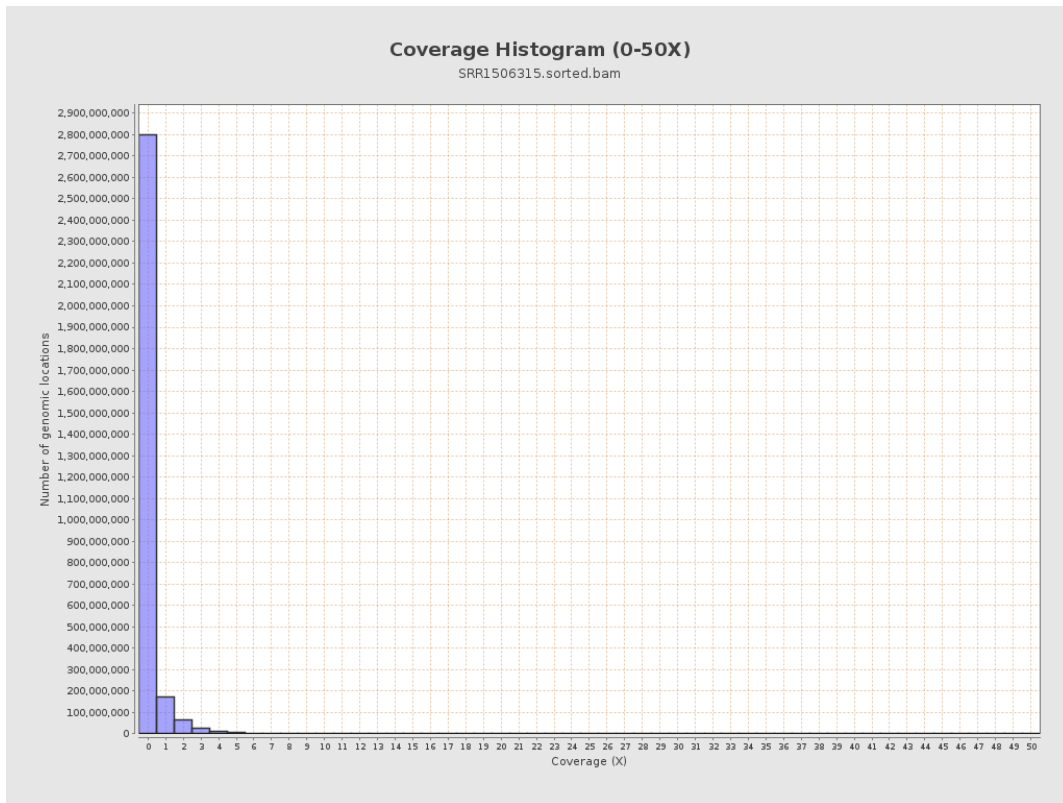
3. Results : Coverage across reference



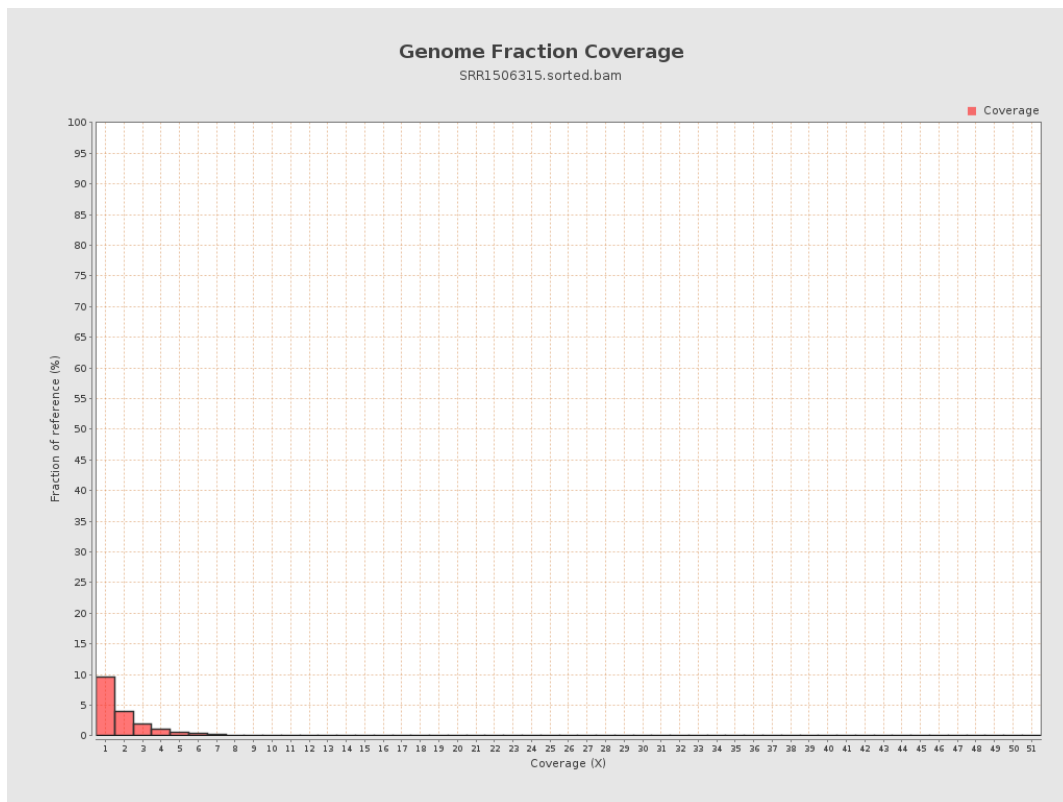
4. Results : Coverage Histogram



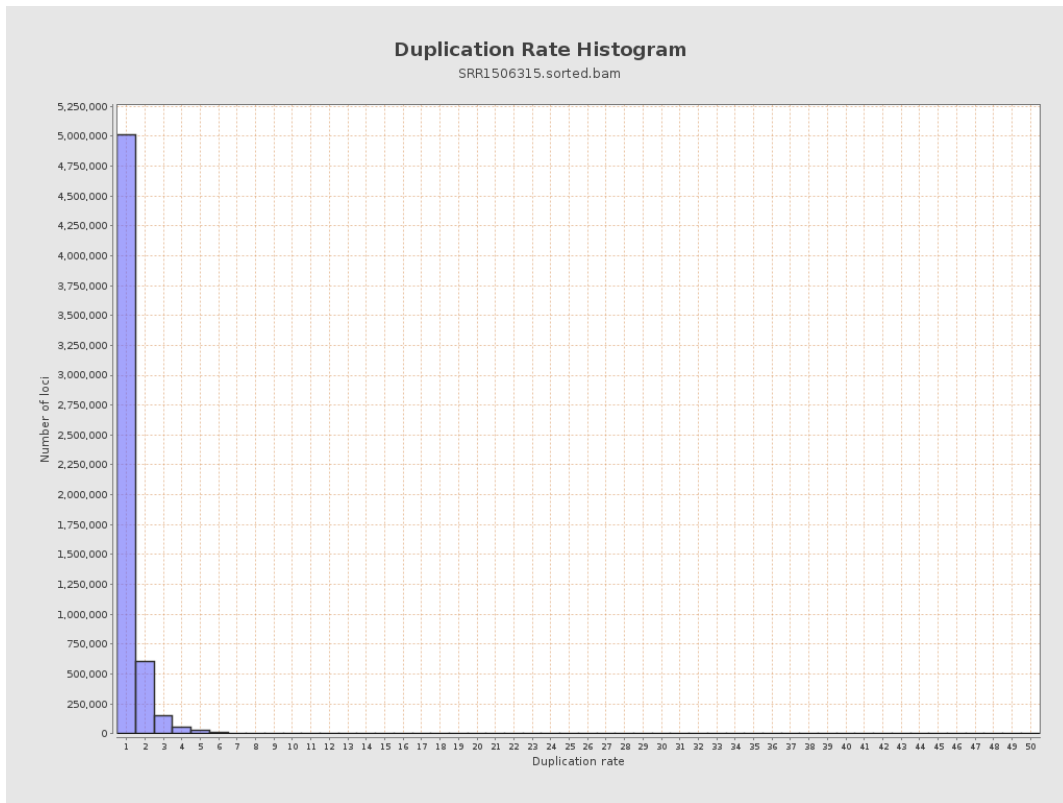
5. Results : Coverage Histogram (0-50X)



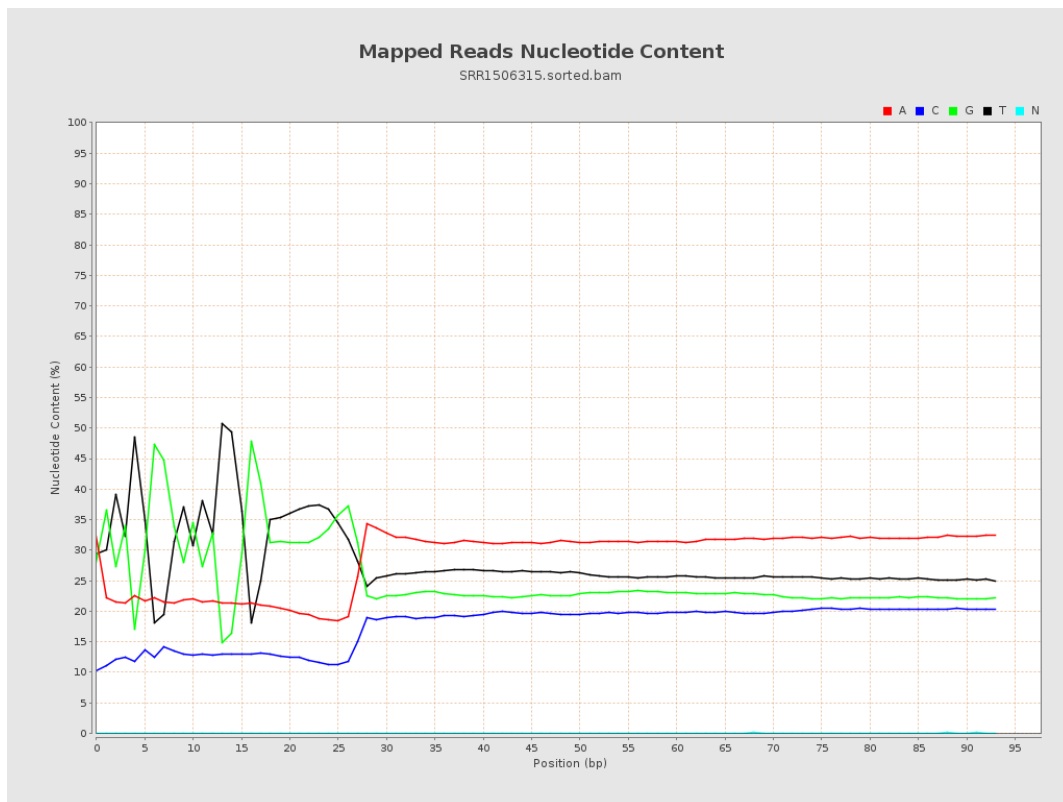
6. Results : Genome Fraction Coverage



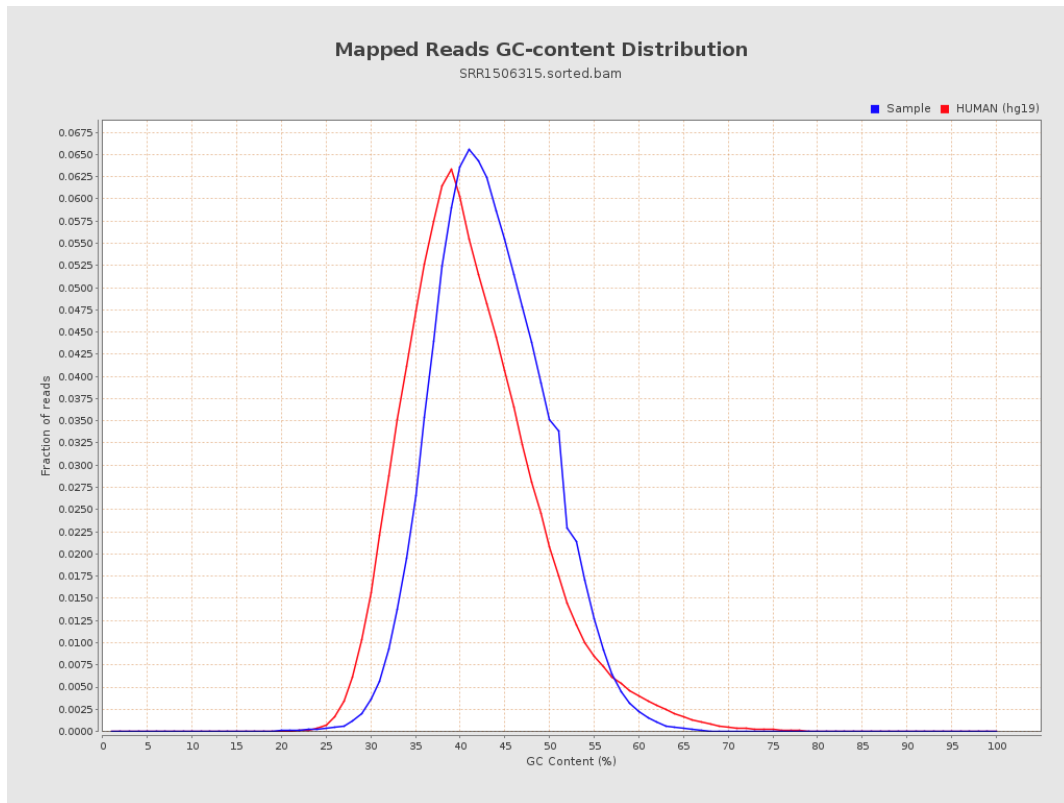
7. Results : Duplication Rate Histogram



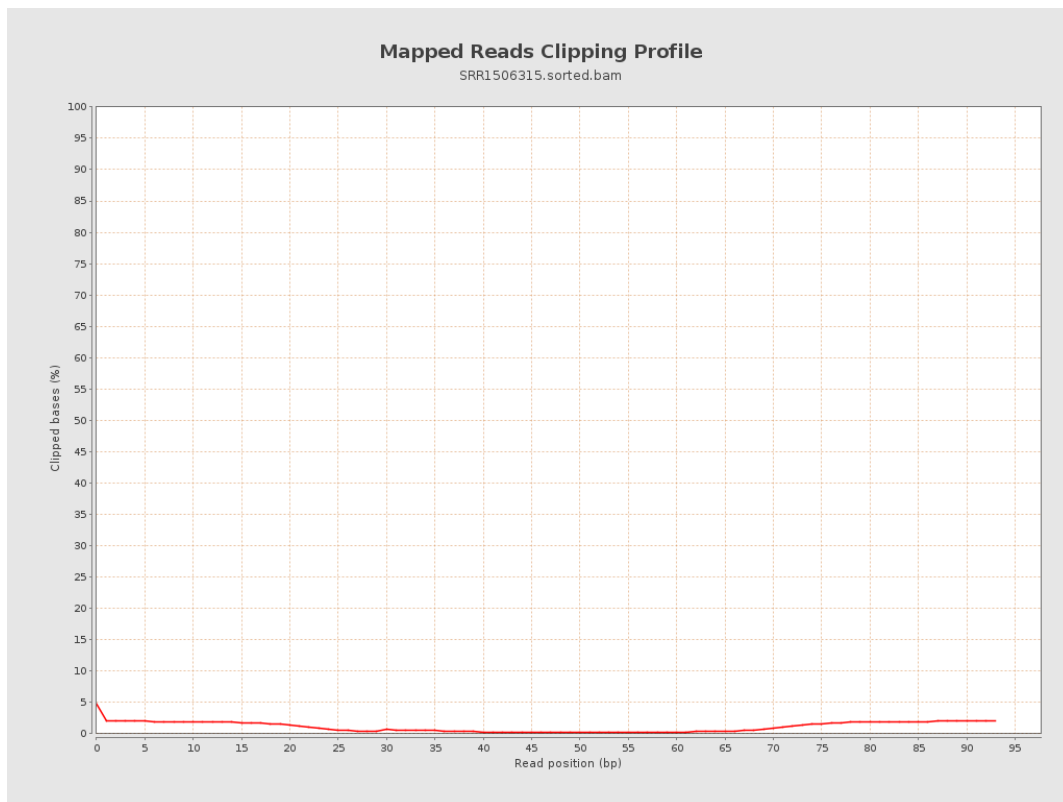
8. Results : Mapped Reads Nucleotide Content



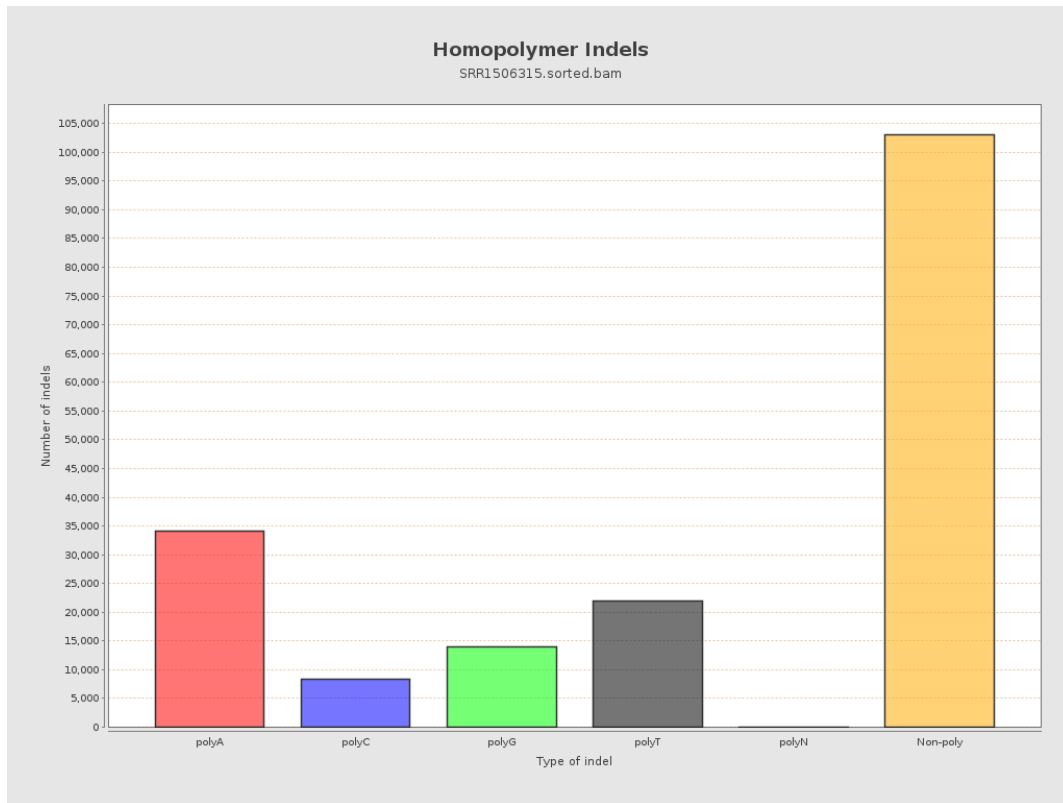
9. Results : Mapped Reads GC-content Distribution



10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

