

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/12/16 10:24:59

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR1153365.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_SRR1153365 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR1153365.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Dec 16 10:24:59 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR1153365.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	46,735,180
Mapped reads	46,432,066 / 99.35%
Unmapped reads	303,114 / 0.65%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	256,510 / 0.55%
Read min/max/mean length	30 / 100 / 100.23
Duplicated reads (estimated)	31,140,868 / 66.63%
Duplication rate	57.48%
Clipped reads	5,996,487 / 12.83%

2.2. ACGT Content

Number/percentage of A's	1,204,642,041 / 26.65%
Number/percentage of C's	1,047,110,873 / 23.17%
Number/percentage of T's	1,239,249,302 / 27.42%
Number/percentage of G's	1,027,316,256 / 22.73%
Number/percentage of N's	1,317,681 / 0.03%
GC Percentage	45.9%

2.3. Coverage

Mean	1.4602

Standard Deviation	16.9997
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2.4. Mapping Quality

Mean Mapping Quality	50.51
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2.5. Mismatches and indels

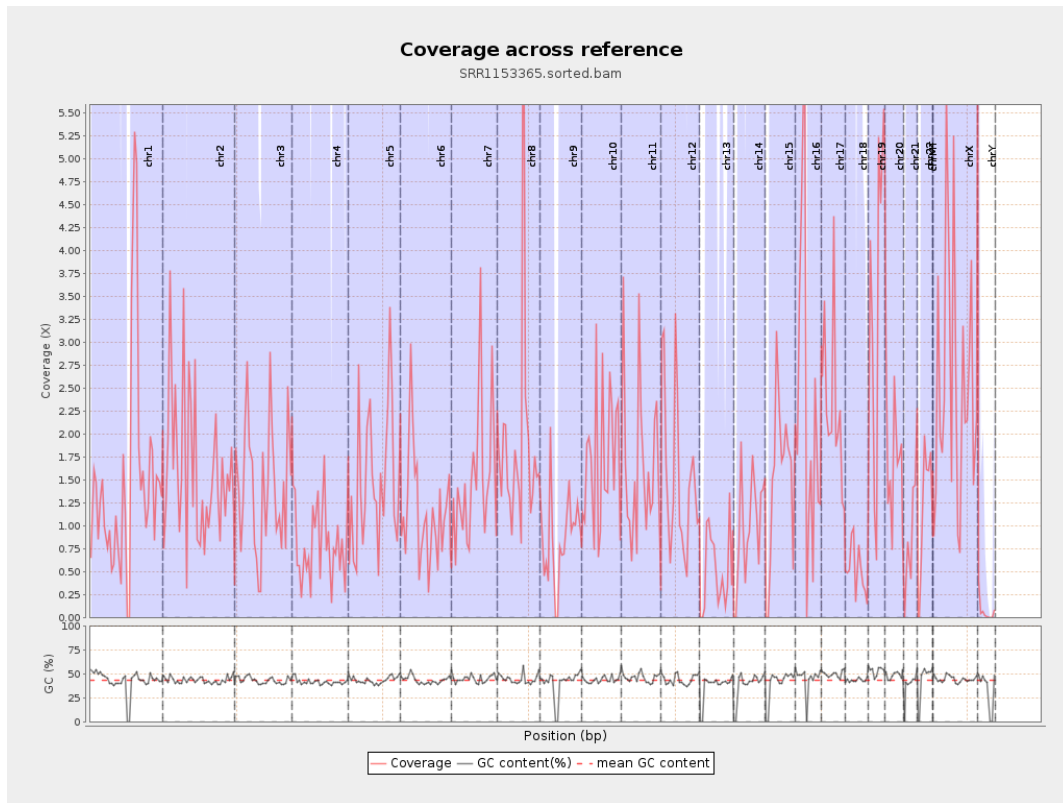
General error rate	0.26%
Mismatches	11,195,672
Insertions	319,798
Mapped reads with at least one insertion	0.68%
Deletions	280,804
Mapped reads with at least one deletion	0.6%
Homopolymer indels	49.89%

2.6. Chromosome stats

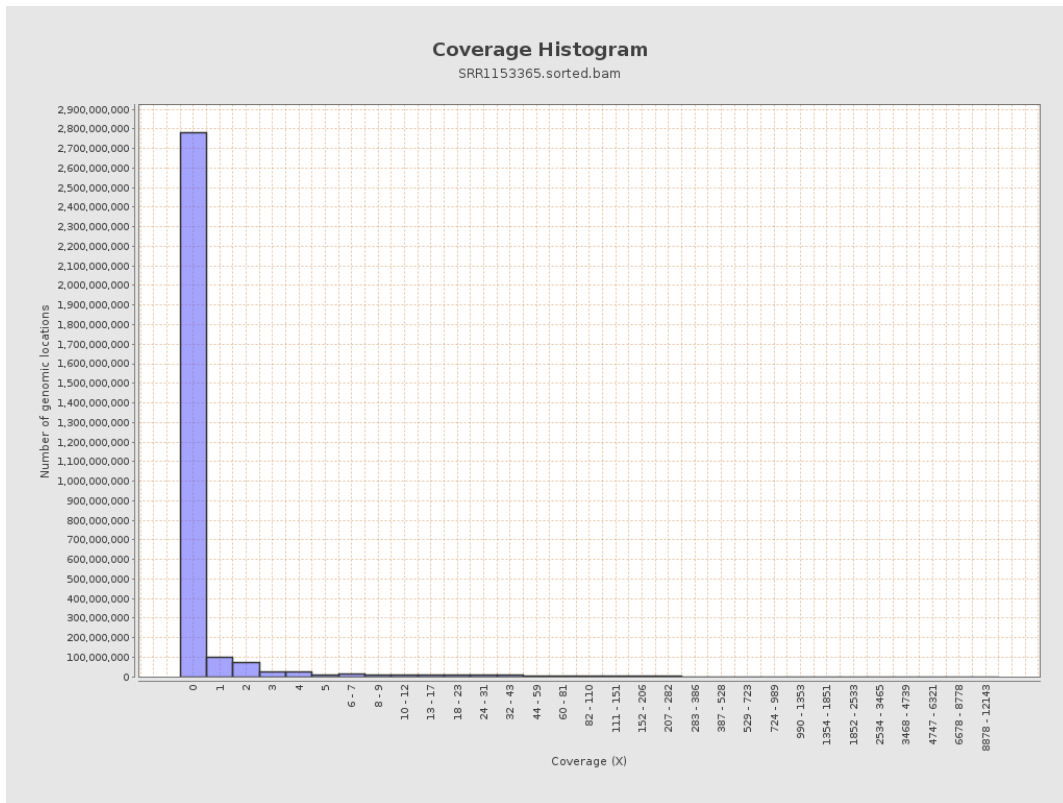
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	359765860	1.4434	14.3302
chr2	243199373	390578273	1.606	16.4966
chr3	198022430	278456419	1.4062	14.5216
chr4	191154276	154016222	0.8057	10.0218
chr5	180915260	271790024	1.5023	17.0013
chr6	171115067	194251829	1.1352	12.8251
chr7	159138663	238563583	1.4991	16.7116

chr8	146364022	279015685	1.9063	32.39
chr9	141213431	119033086	0.8429	10.1386
chr10	135534747	234869707	1.7329	17.8007
chr11	135006516	221168419	1.6382	15.3191
chr12	133851895	208664414	1.5589	14.9786
chr13	115169878	58607896	0.5089	8.0692
chr14	107349540	105616875	0.9839	10.5082
chr15	102531392	149708365	1.4601	13.4883
chr16	90354753	201574649	2.2309	22.5101
chr17	81195210	189985955	2.3399	18.6454
chr18	78077248	41439802	0.5308	7.5365
chr19	59128983	201696485	3.4111	28.0203
chr20	63025520	108243784	1.7175	16.8642
chr21	48129895	45368573	0.9426	11.1229
chr22	51304566	61217256	1.1932	12.7913
chrMT	16571	26517	1.6002	5.6108
chrX	155270560	402950851	2.5952	28.0437
chrY	59373566	3585899	0.0604	2.4031

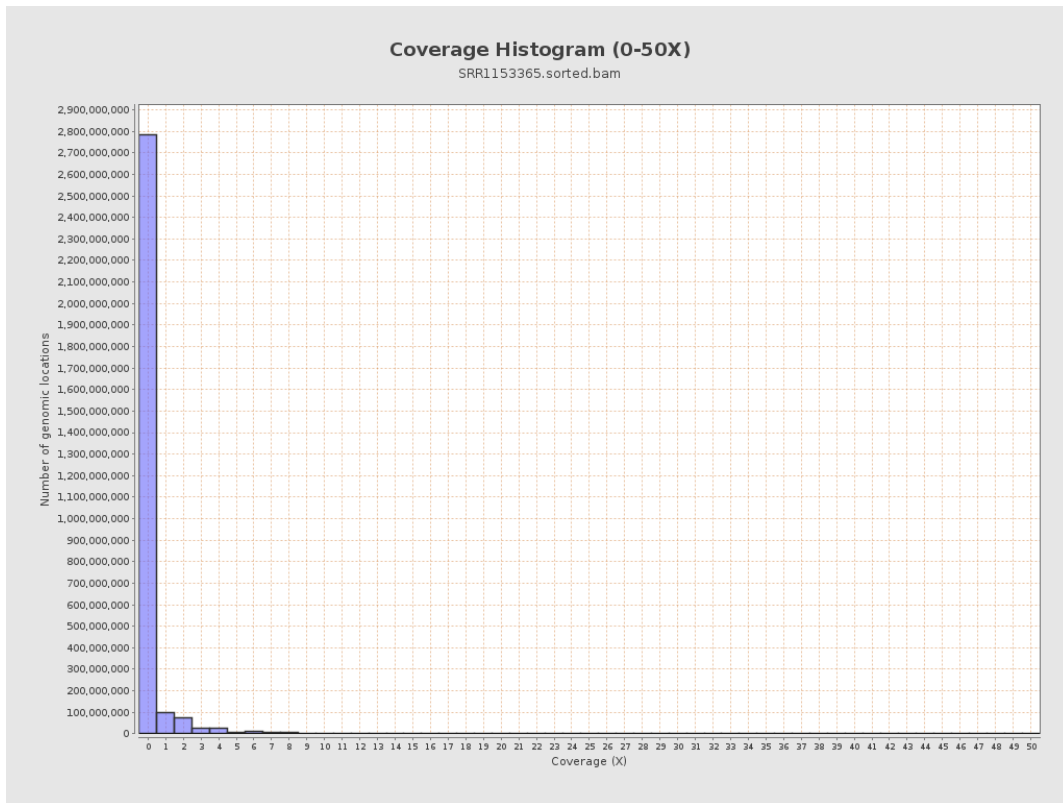
3. Results : Coverage across reference



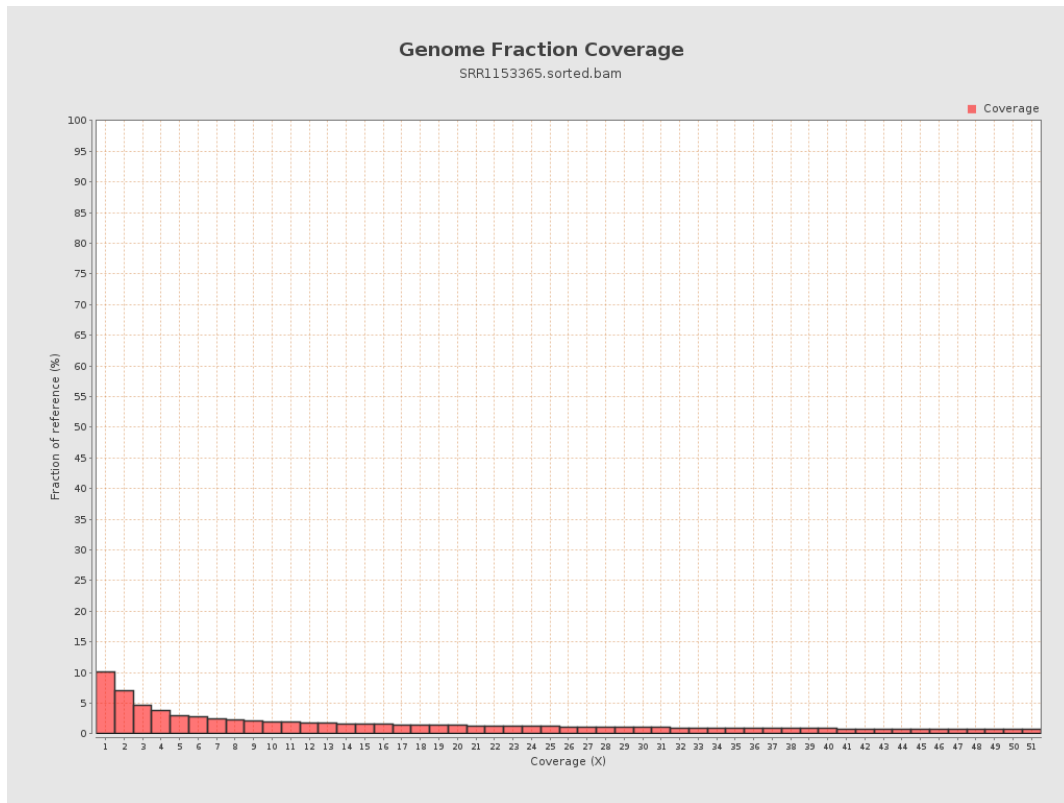
4. Results : Coverage Histogram



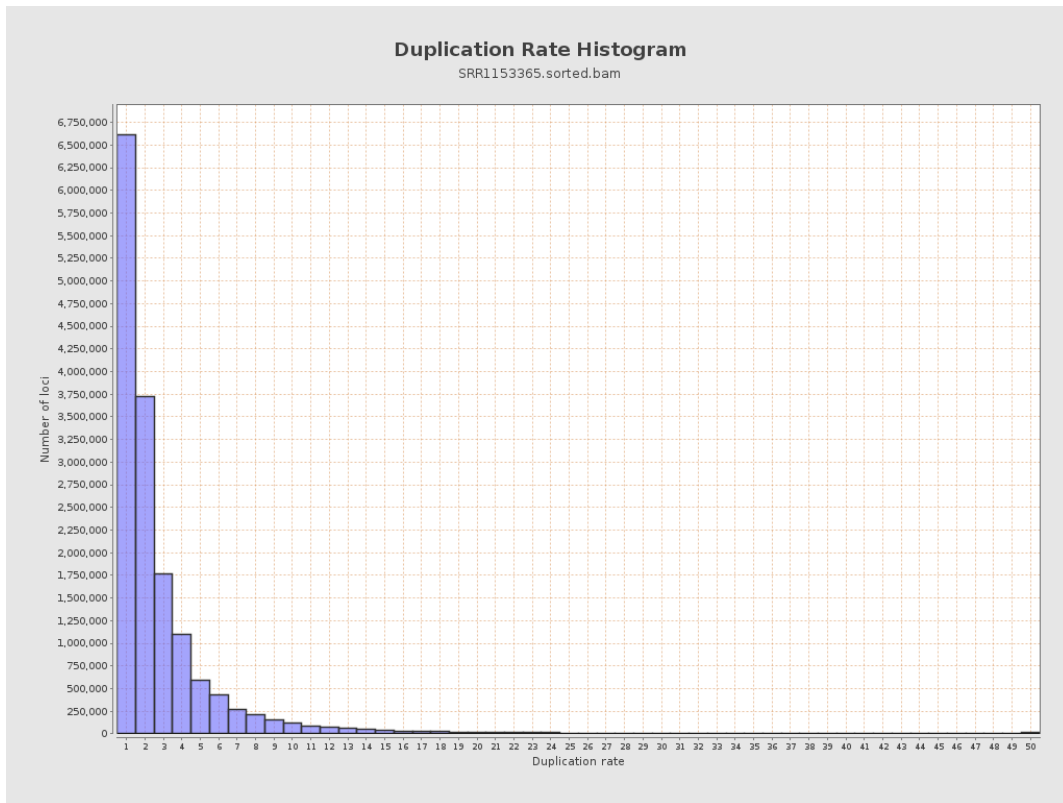
5. Results : Coverage Histogram (0-50X)



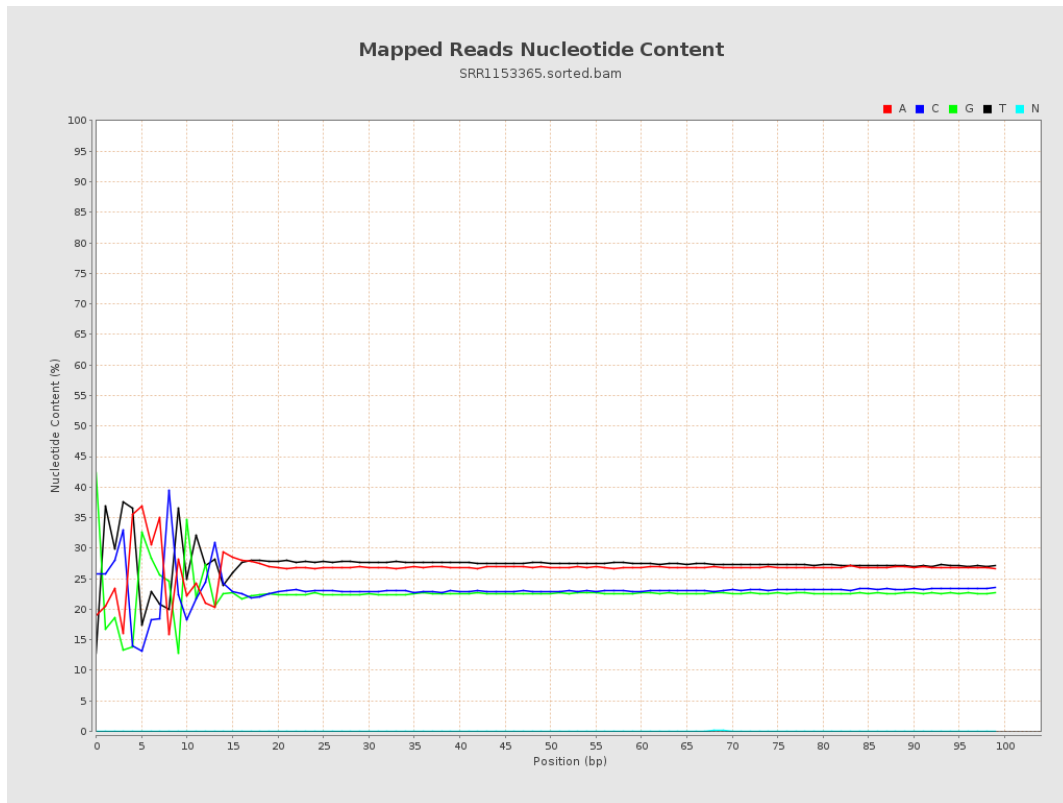
6. Results : Genome Fraction Coverage



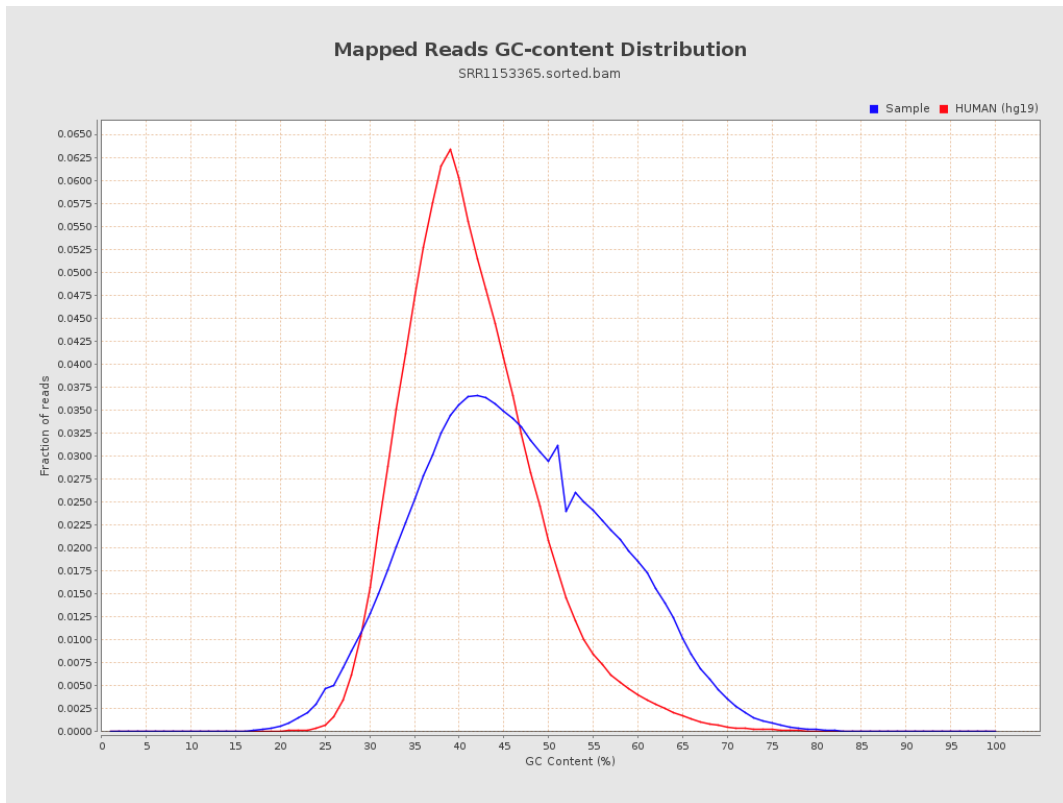
7. Results : Duplication Rate Histogram



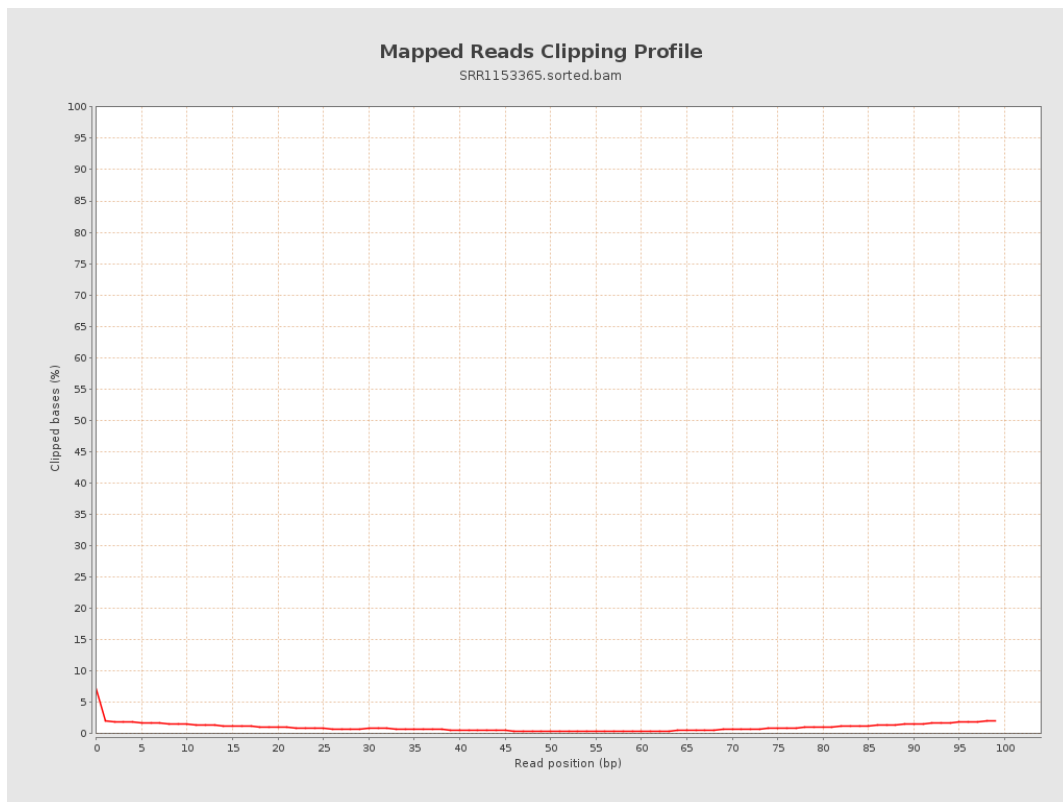
8. Results : Mapped Reads Nucleotide Content



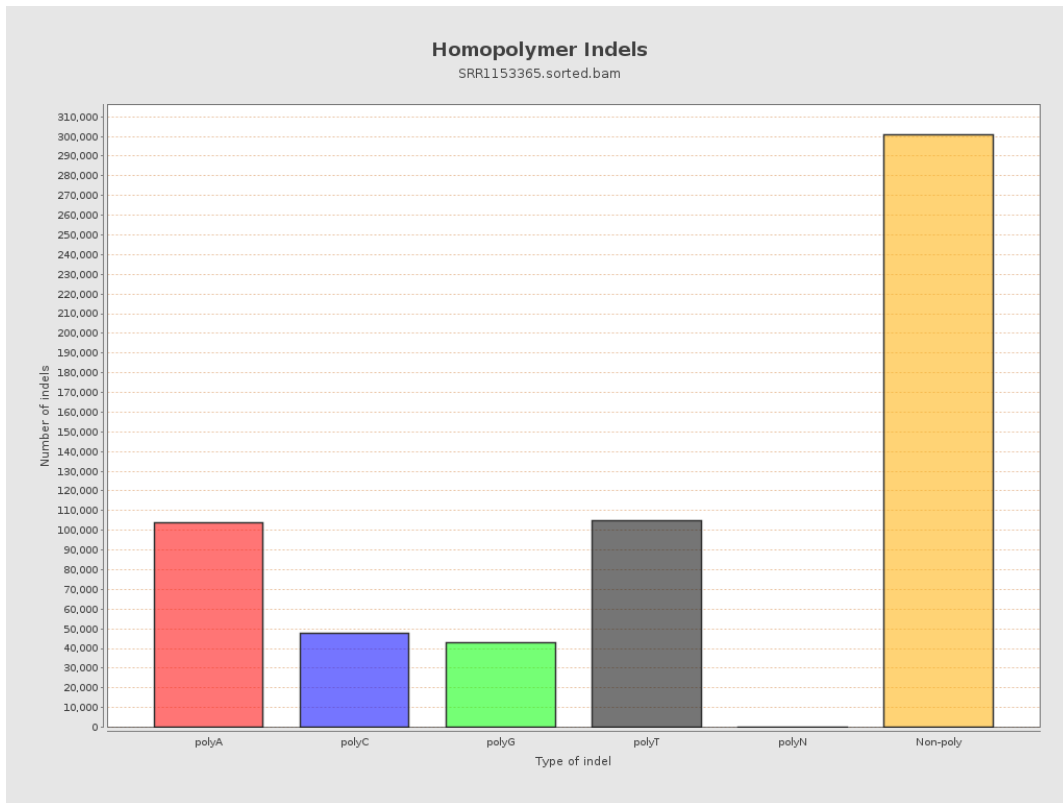
9. Results : Mapped Reads GC-content Distribution



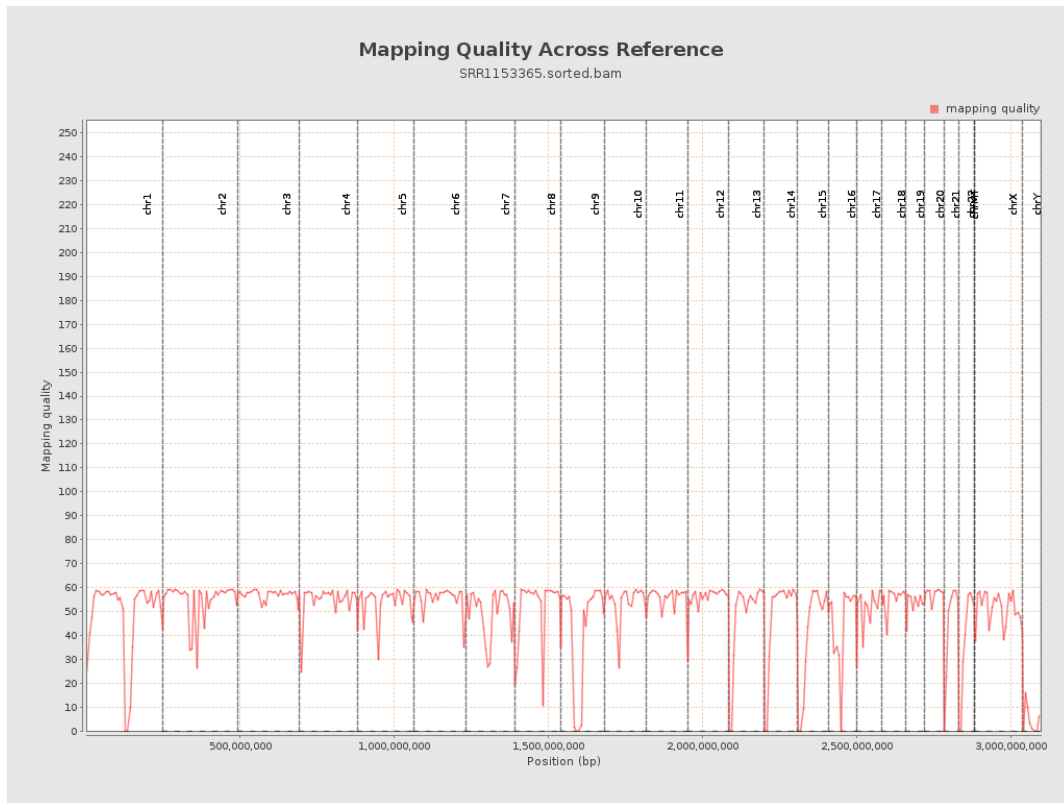
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

