

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/12/16 19:10:30

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR1153381.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_SRR1153381 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR1153381.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Dec 16 19:10:29 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR1153381.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	81,611,374
Mapped reads	81,064,562 / 99.33%
Unmapped reads	546,812 / 0.67%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	424,748 / 0.52%
Read min/max/mean length	30 / 100 / 100.21
Duplicated reads (estimated)	48,604,964 / 59.56%
Duplication rate	45.81%
Clipped reads	12,206,164 / 14.96%

2.2. ACGT Content

Number/percentage of A's	1,951,037,107 / 24.73%
Number/percentage of C's	1,992,757,243 / 25.25%
Number/percentage of T's	1,948,057,829 / 24.69%
Number/percentage of G's	1,997,858,961 / 25.32%
Number/percentage of N's	1,059,697 / 0.01%
GC Percentage	50.57%

2.3. Coverage

Mean	2.5492

Standard Deviation	41.3595
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2.4. Mapping Quality

Mean Mapping Quality	48.65
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2.5. Mismatches and indels

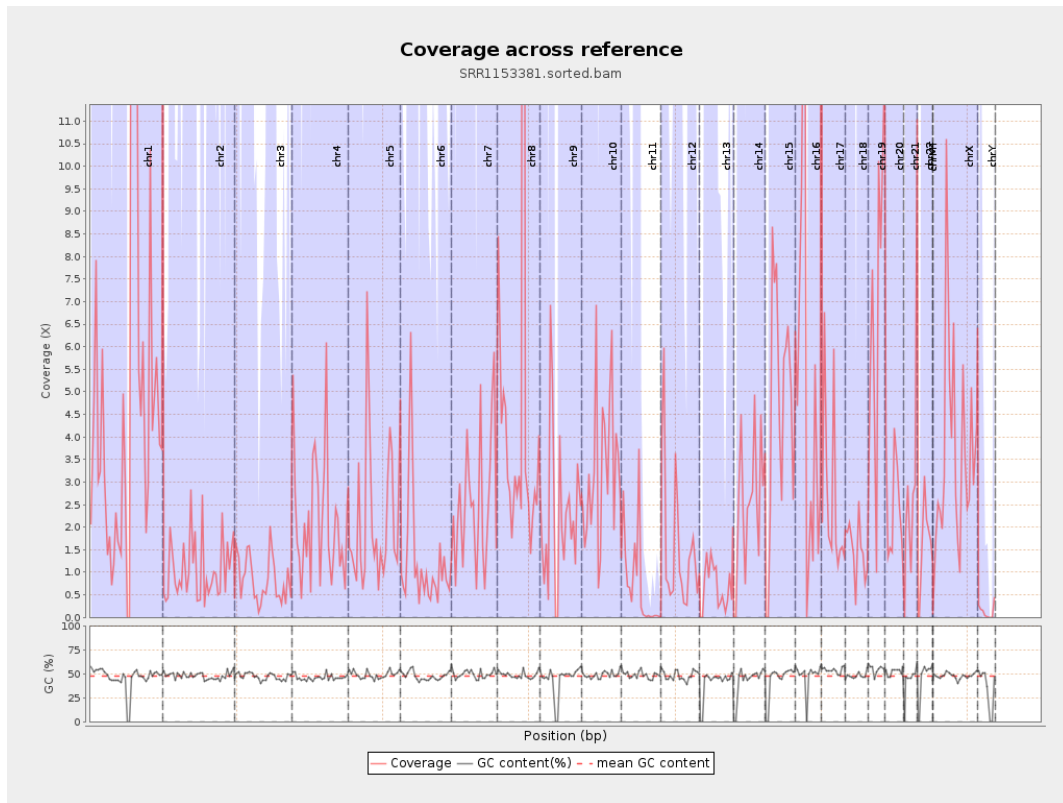
General error rate	0.3%
Mismatches	22,442,946
Insertions	560,909
Mapped reads with at least one insertion	0.68%
Deletions	359,862
Mapped reads with at least one deletion	0.44%
Homopolymer indels	45.55%

2.6. Chromosome stats

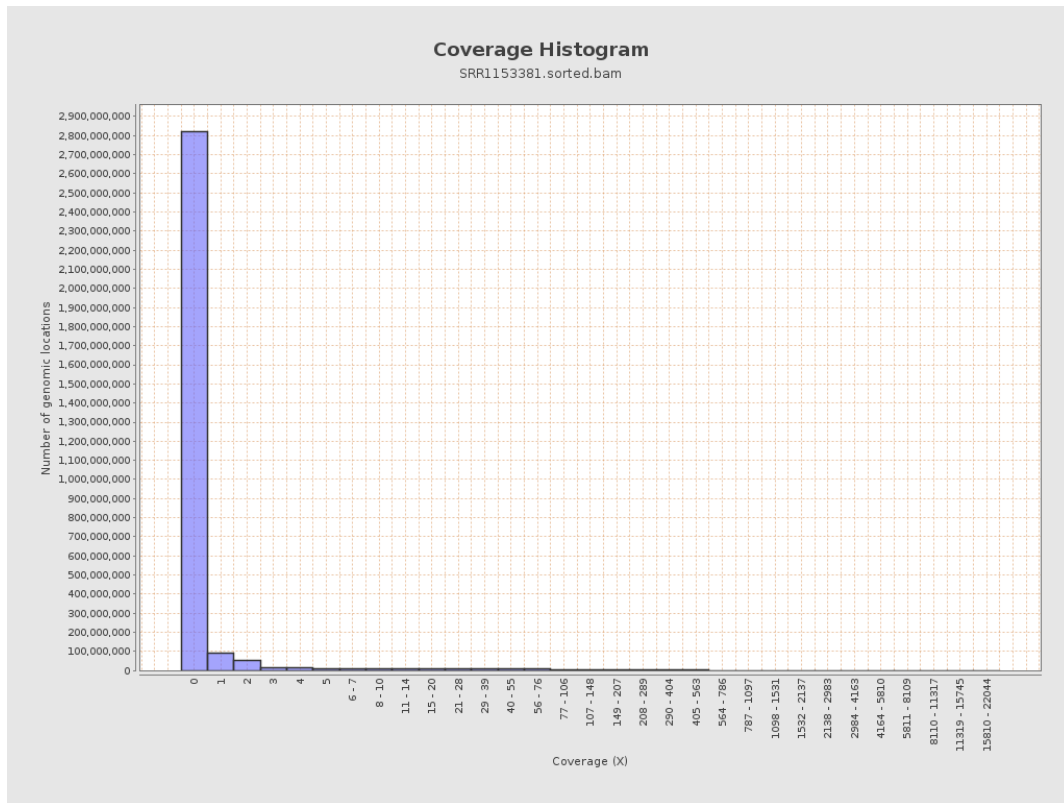
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	1269880140	5.0948	55.7836
chr2	243199373	259164839	1.0656	17.7424
chr3	198022430	170406368	0.8605	14.4616
chr4	191154276	412660222	2.1588	41.0335
chr5	180915260	393507119	2.1751	31.4794
chr6	171115067	206579138	1.2073	19.1239
chr7	159138663	403575225	2.536	40.7488

chr8	146364022	751213555	5.1325	111.3314
chr9	141213431	312135004	2.2104	27.8654
chr10	135534747	444231577	3.2776	34.7349
chr11	135006516	101590341	0.7525	13.0512
chr12	133851895	213131839	1.5923	25.7405
chr13	115169878	79013128	0.6861	13.1277
chr14	107349540	270269503	2.5177	27.0888
chr15	102531392	466582230	4.5506	39.2058
chr16	90354753	453633579	5.0206	47.4837
chr17	81195210	216255368	2.6634	37.8812
chr18	78077248	117800882	1.5088	21.3241
chr19	59128983	405079315	6.8508	57.1918
chr20	63025520	152727256	2.4233	24.8776
chr21	48129895	142880877	2.9687	65.115
chr22	51304566	80856259	1.576	18.8014
chrMT	16571	977	0.059	0.391
chrX	155270560	560643516	3.6108	43.765
chrY	59373566	7780224	0.131	7.3241

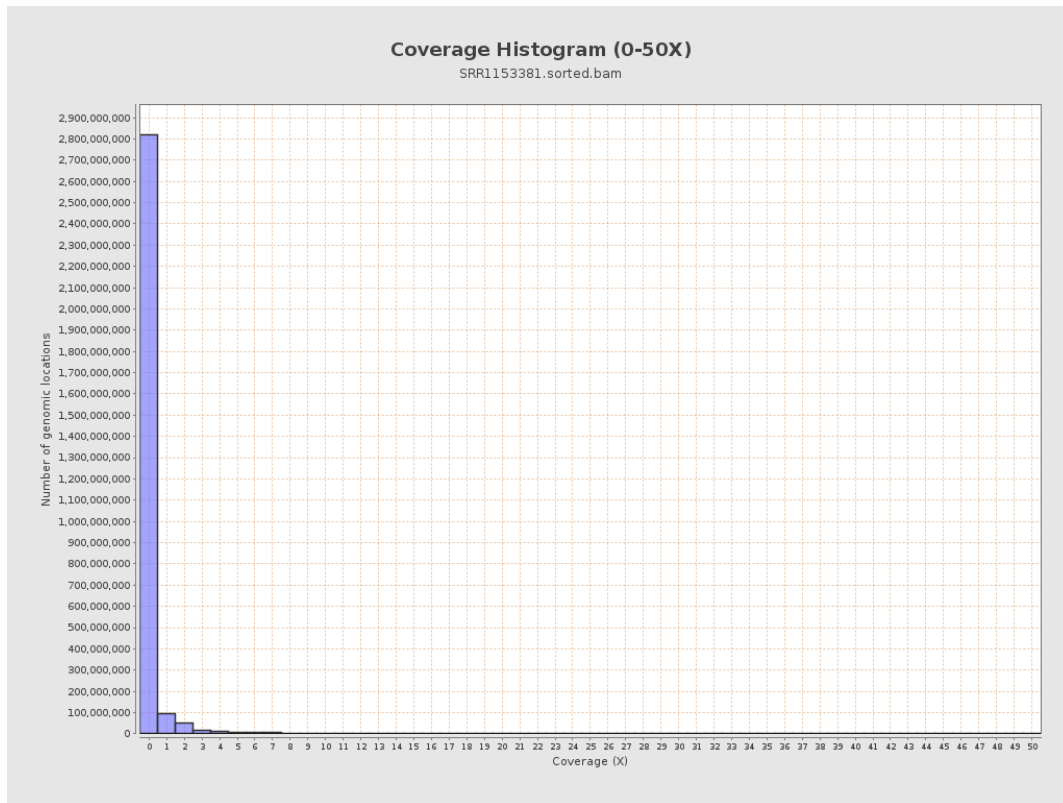
3. Results : Coverage across reference



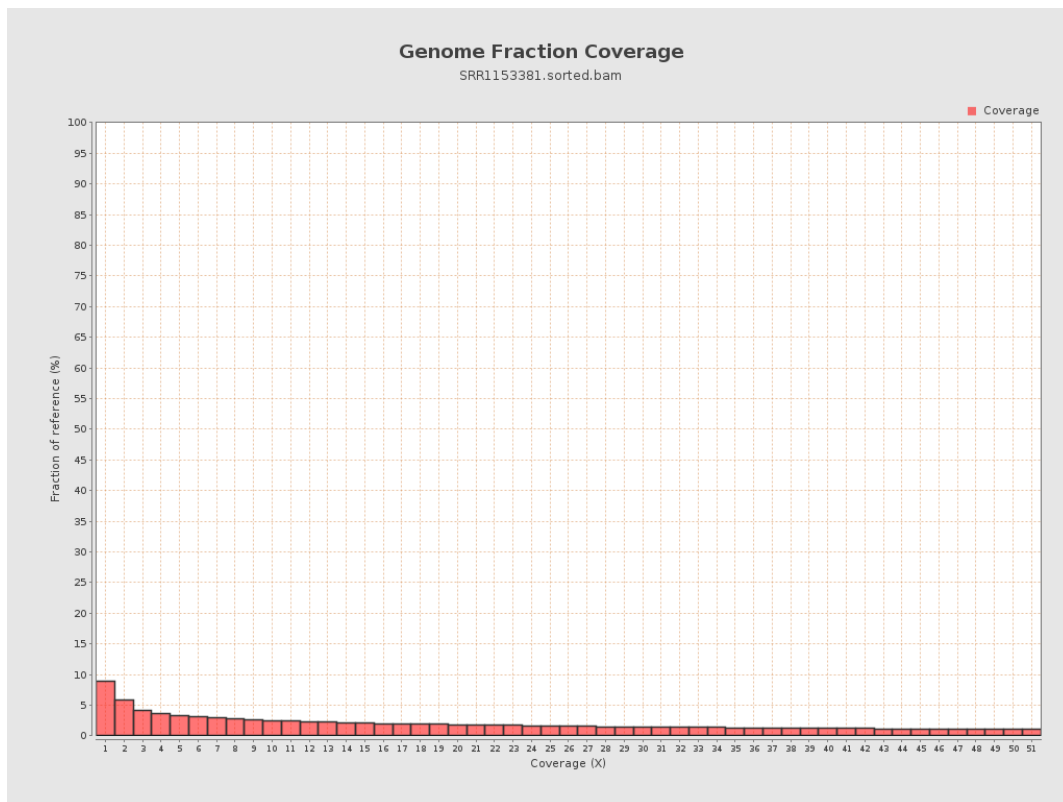
4. Results : Coverage Histogram



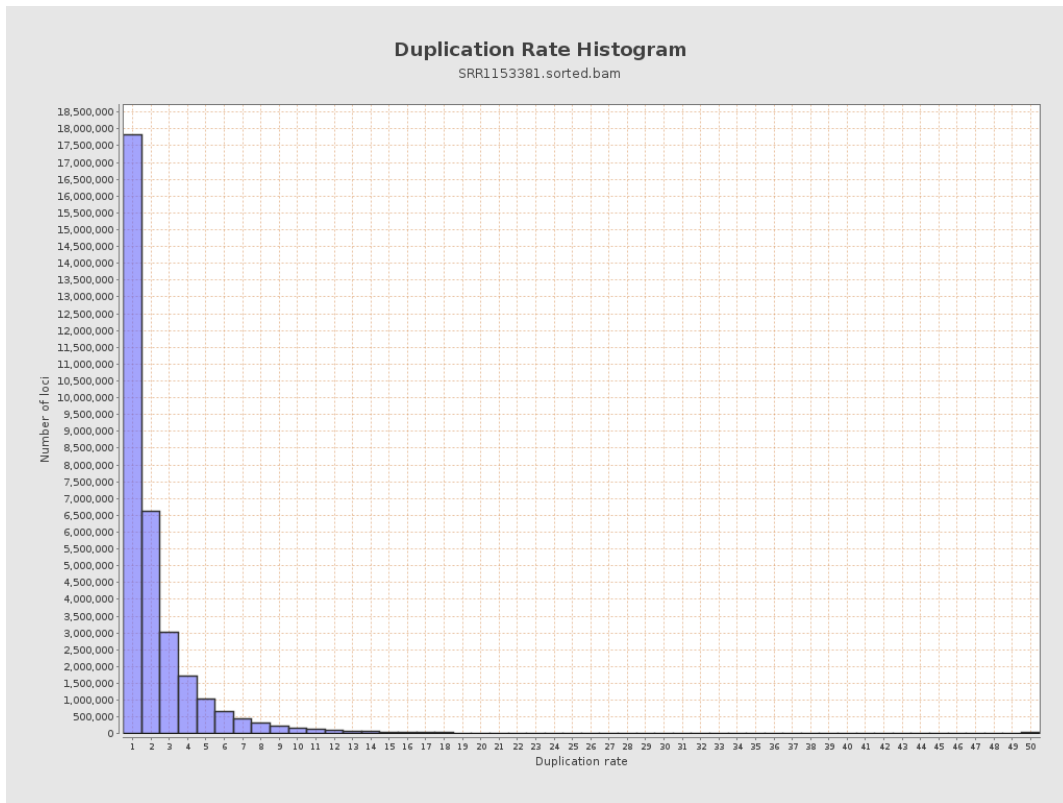
5. Results : Coverage Histogram (0-50X)



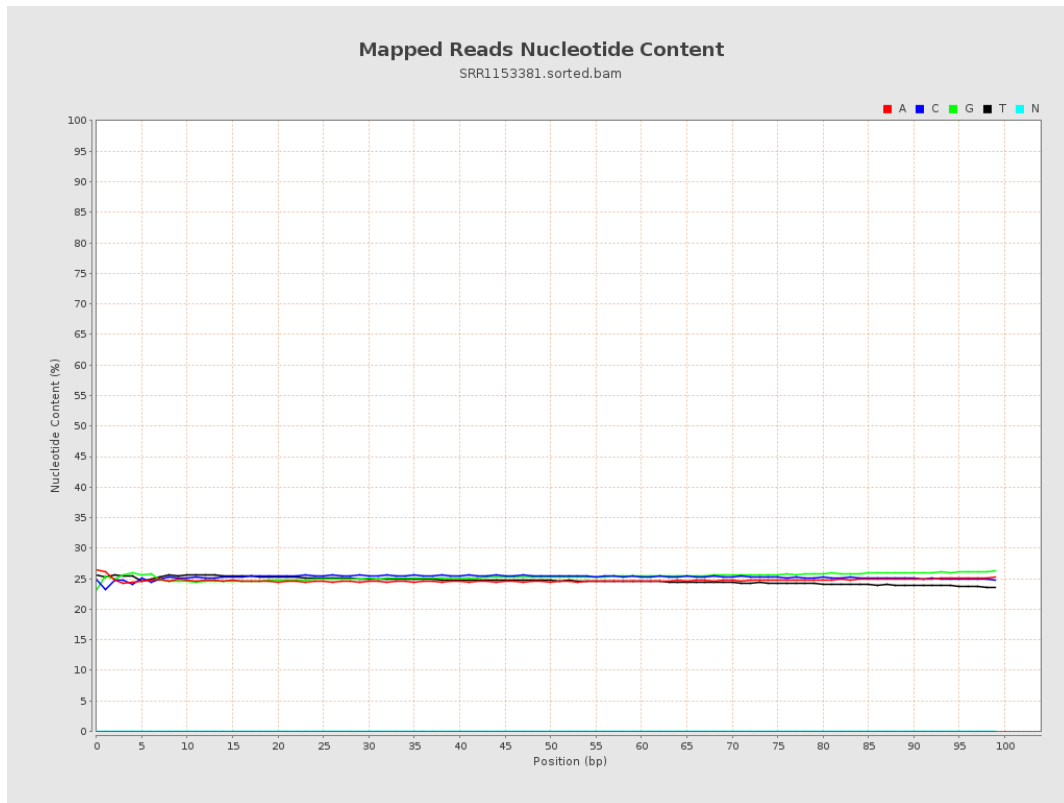
6. Results : Genome Fraction Coverage



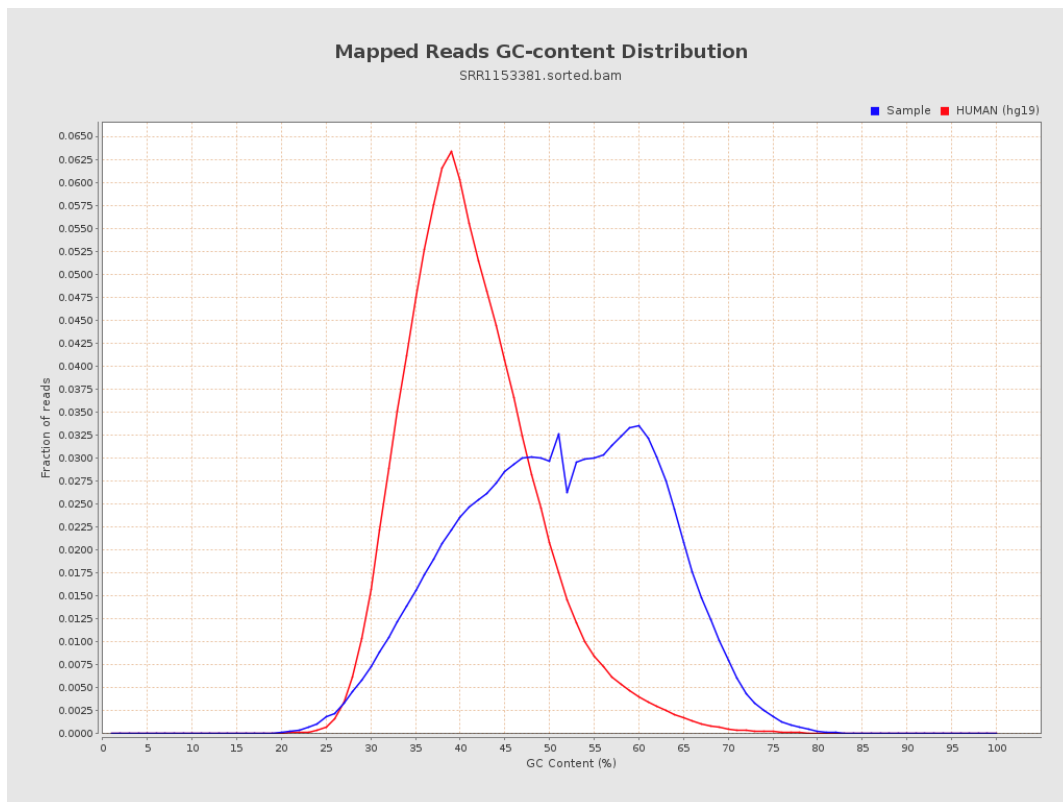
7. Results : Duplication Rate Histogram



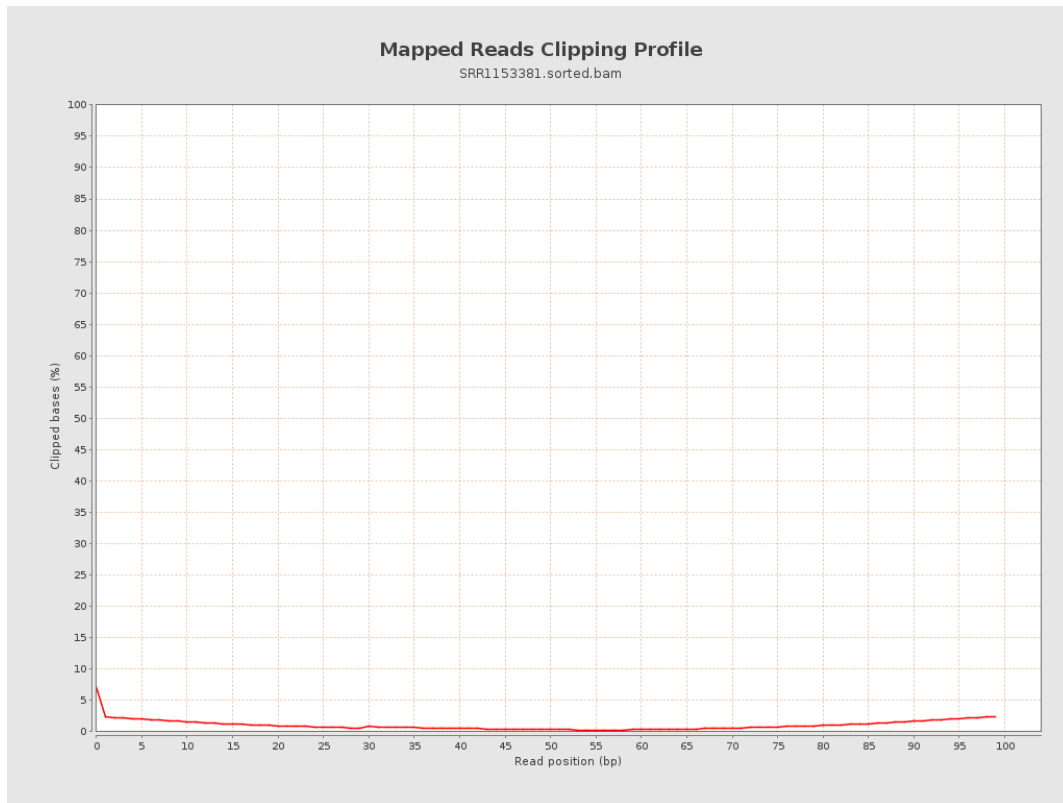
8. Results : Mapped Reads Nucleotide Content



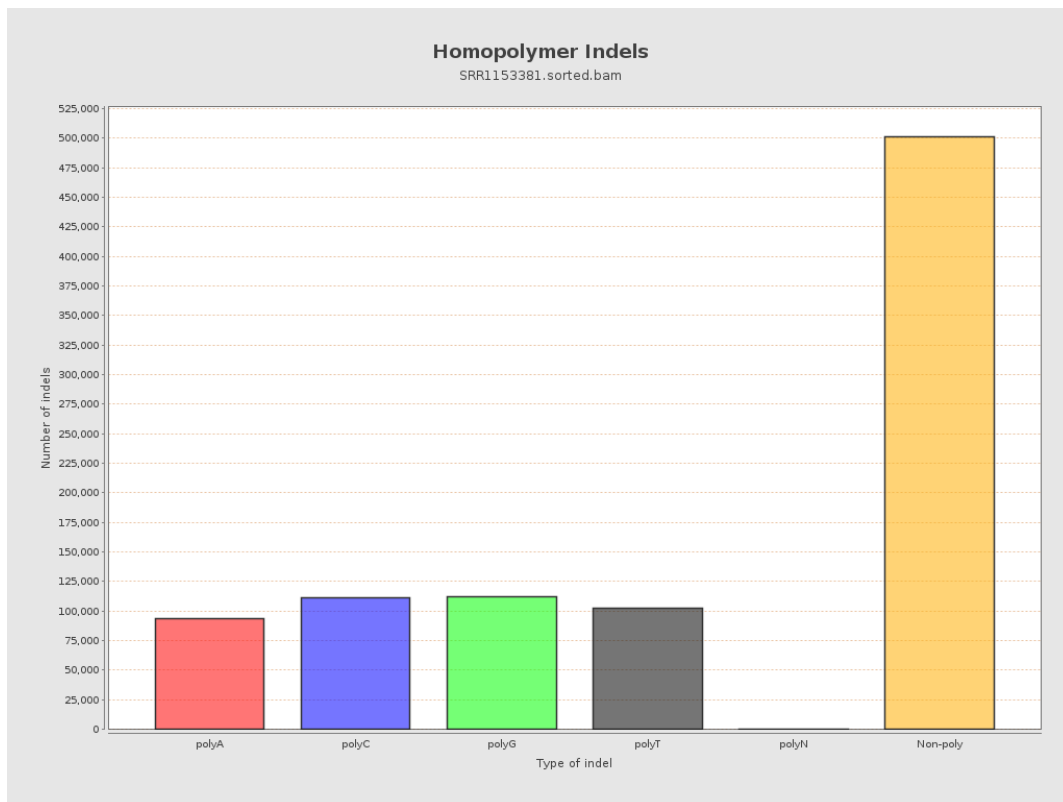
9. Results : Mapped Reads GC-content Distribution



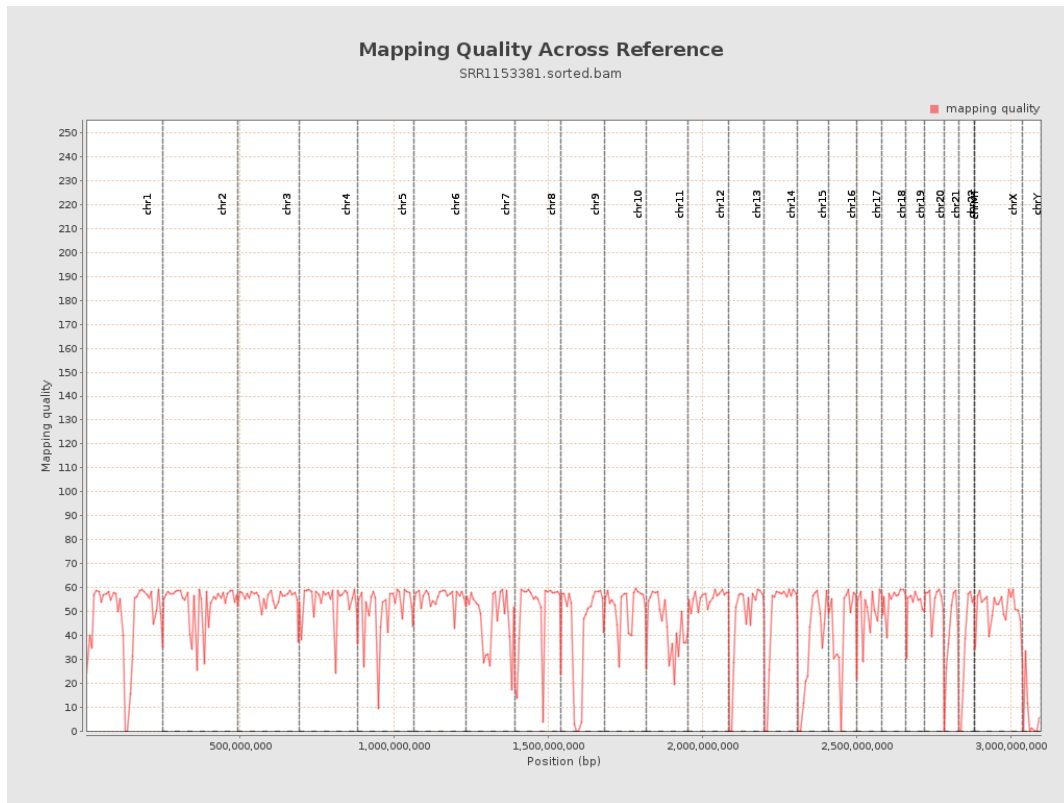
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

