

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/24 05:10:36

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR3081249.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_SRR3081249 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR3081249.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sat Aug 24 05:10:22 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR3081249.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	4,526,553
Mapped reads	3,904,857 / 86.27%
Unmapped reads	621,696 / 13.73%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	38,545 / 0.85%
Read min/max/mean length	30 / 76 / 76.3
Duplicated reads (estimated)	227,042 / 5.02%
Duplication rate	4.41%
Clipped reads	1,688,328 / 37.3%

2.2. ACGT Content

Number/percentage of A's	71,571,981 / 27.36%
Number/percentage of C's	47,865,959 / 18.3%
Number/percentage of T's	83,662,862 / 31.98%
Number/percentage of G's	58,455,946 / 22.34%
Number/percentage of N's	69,855 / 0.03%
GC Percentage	40.64%

2.3. Coverage

Mean	0.0845

Standard Deviation	0.6853
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2.4. Mapping Quality

Mean Mapping Quality	46.26
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2.5. Mismatches and indels

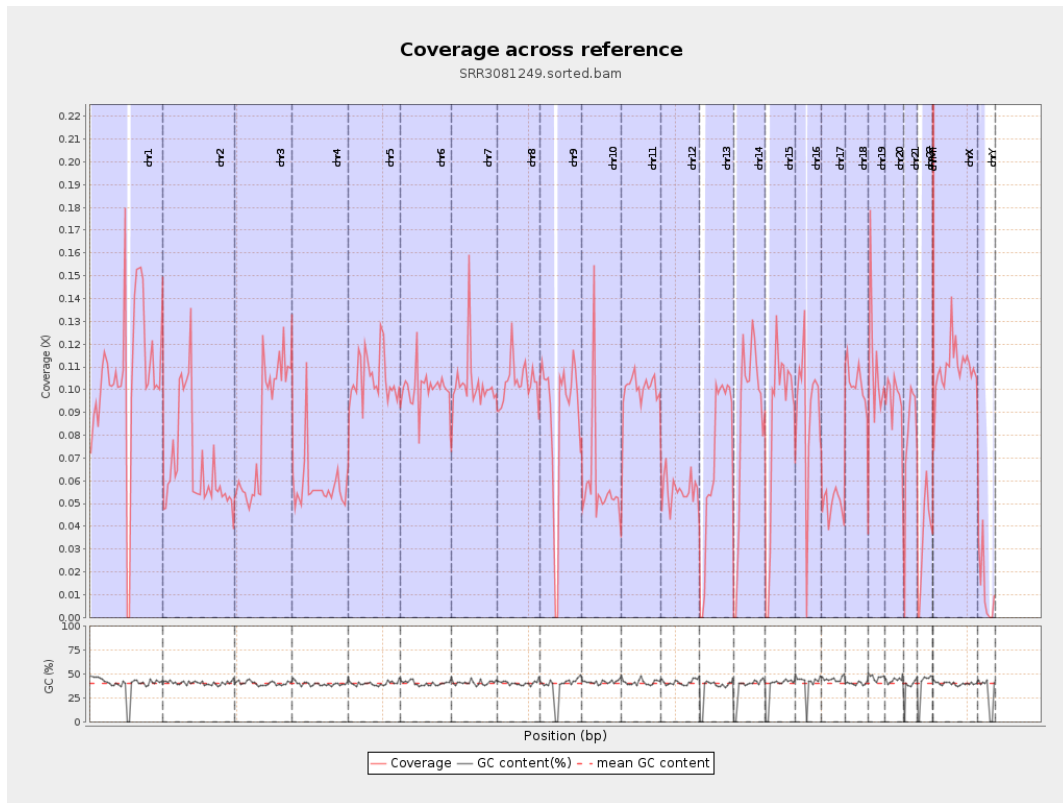
General error rate	0.86%
Mismatches	2,213,267
Insertions	21,578
Mapped reads with at least one insertion	0.55%
Deletions	65,147
Mapped reads with at least one deletion	1.65%
Homopolymer indels	49.47%

2.6. Chromosome stats

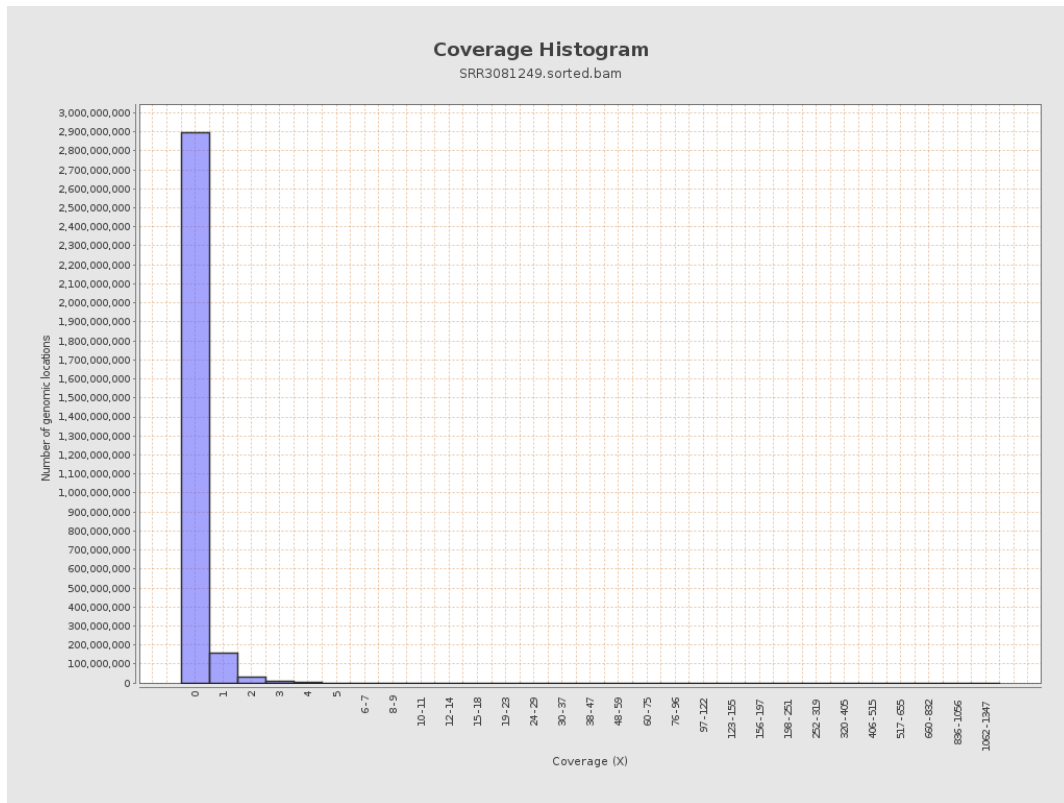
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	26511124	0.1064	1.2795
chr2	243199373	16329227	0.0671	0.7194
chr3	198022430	16407023	0.0829	0.3736
chr4	191154276	11073551	0.0579	0.3968
chr5	180915260	18930069	0.1046	0.4052
chr6	171115067	17209552	0.1006	0.5124
chr7	159138663	16280285	0.1023	1.0329

chr8	146364022	15073287	0.103	0.8324
chr9	141213431	12534418	0.0888	0.651
chr10	135534747	7873139	0.0581	0.8121
chr11	135006516	13464009	0.0997	0.7375
chr12	133851895	7467101	0.0558	0.3049
chr13	115169878	8083728	0.0702	0.3257
chr14	107349540	9524909	0.0887	0.4141
chr15	102531392	8697121	0.0848	0.3633
chr16	90354753	8122672	0.0899	0.4268
chr17	81195210	4075879	0.0502	0.3964
chr18	78077248	8046586	0.1031	0.9963
chr19	59128983	6451233	0.1091	1.0245
chr20	63025520	6017136	0.0955	0.4068
chr21	48129895	3796062	0.0789	0.4306
chr22	51304566	1802599	0.0351	0.2275
chrMT	16571	246590	14.8808	7.6848
chrX	155270560	16967269	0.1093	0.4632
chrY	59373566	753202	0.0127	0.3414

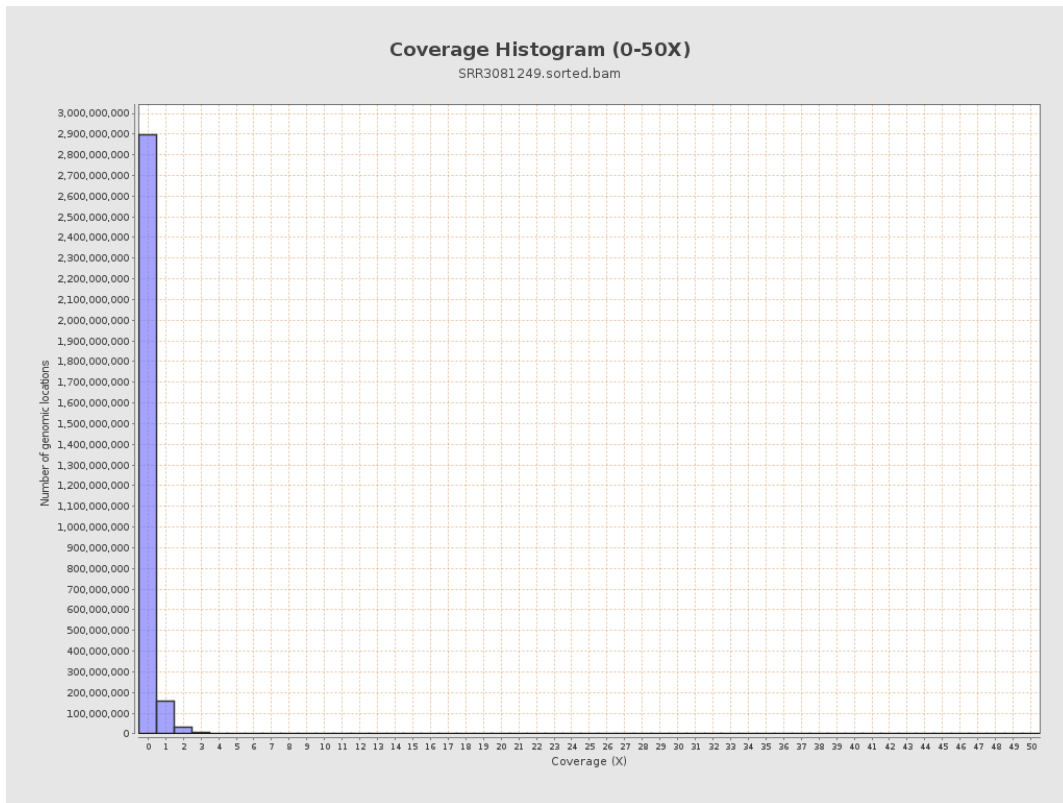
3. Results : Coverage across reference



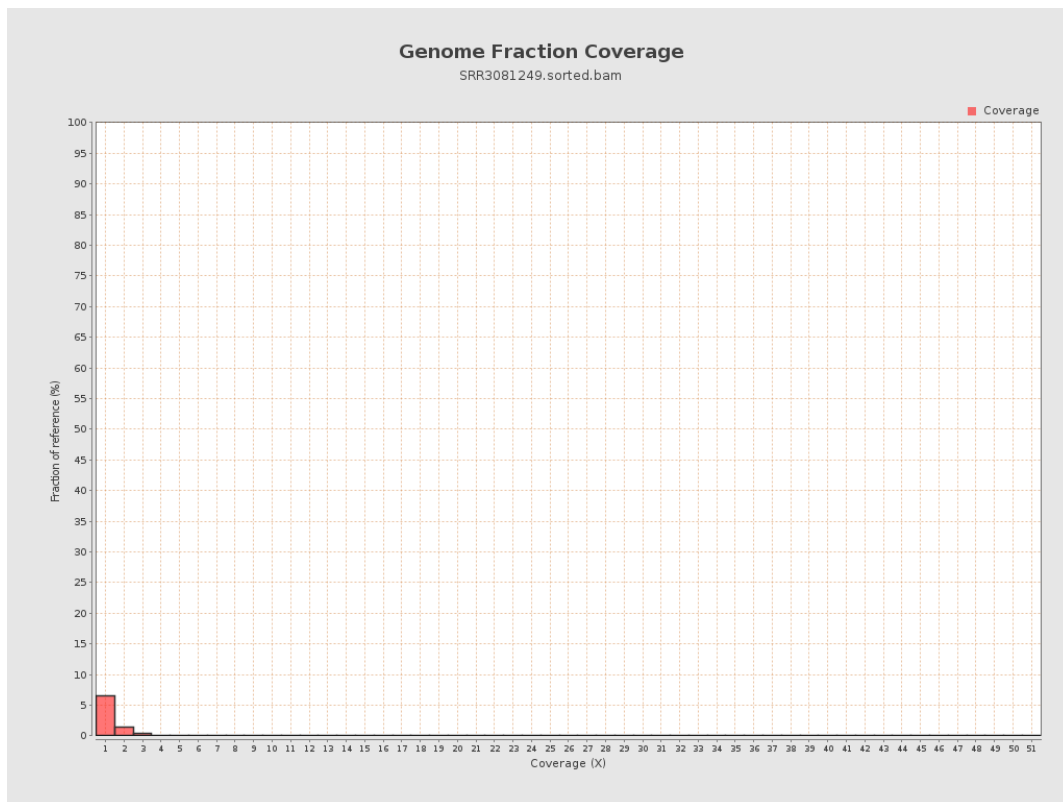
4. Results : Coverage Histogram



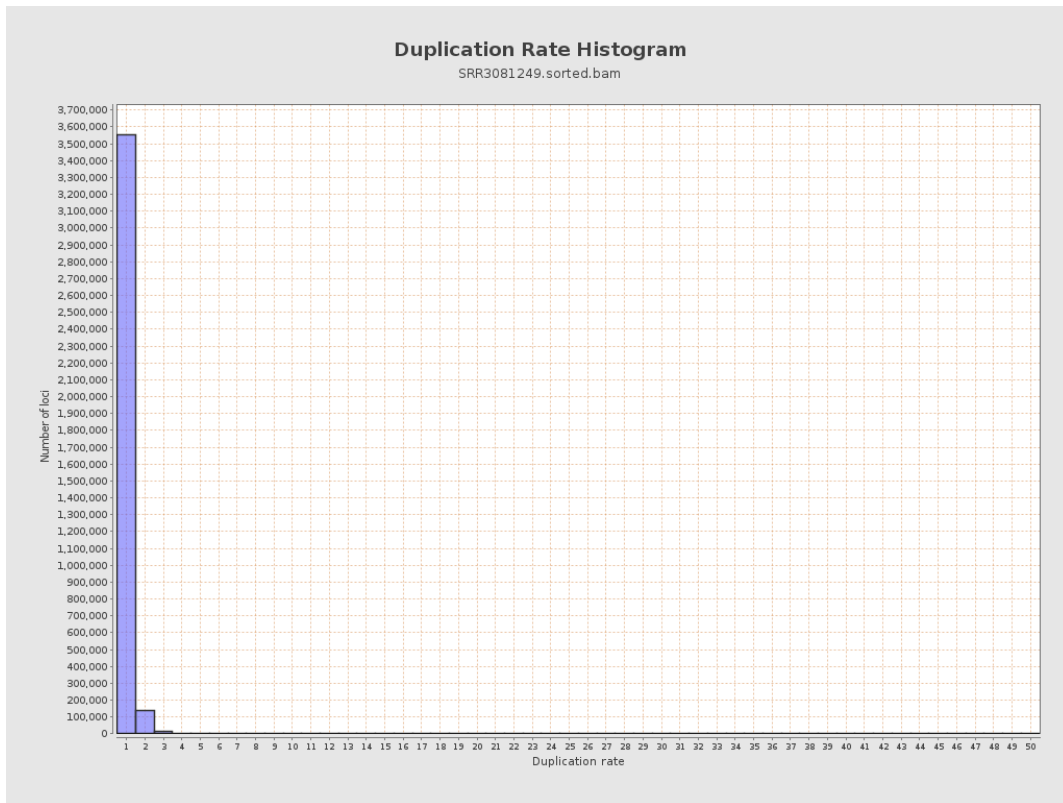
5. Results : Coverage Histogram (0-50X)



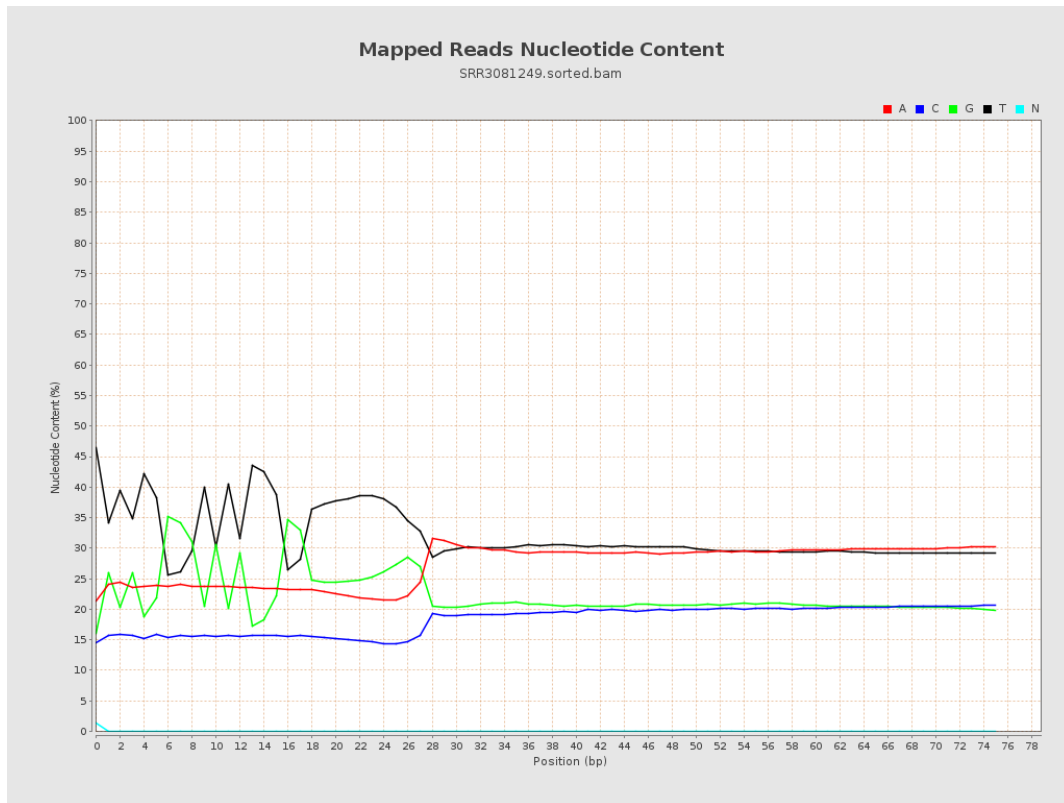
6. Results : Genome Fraction Coverage



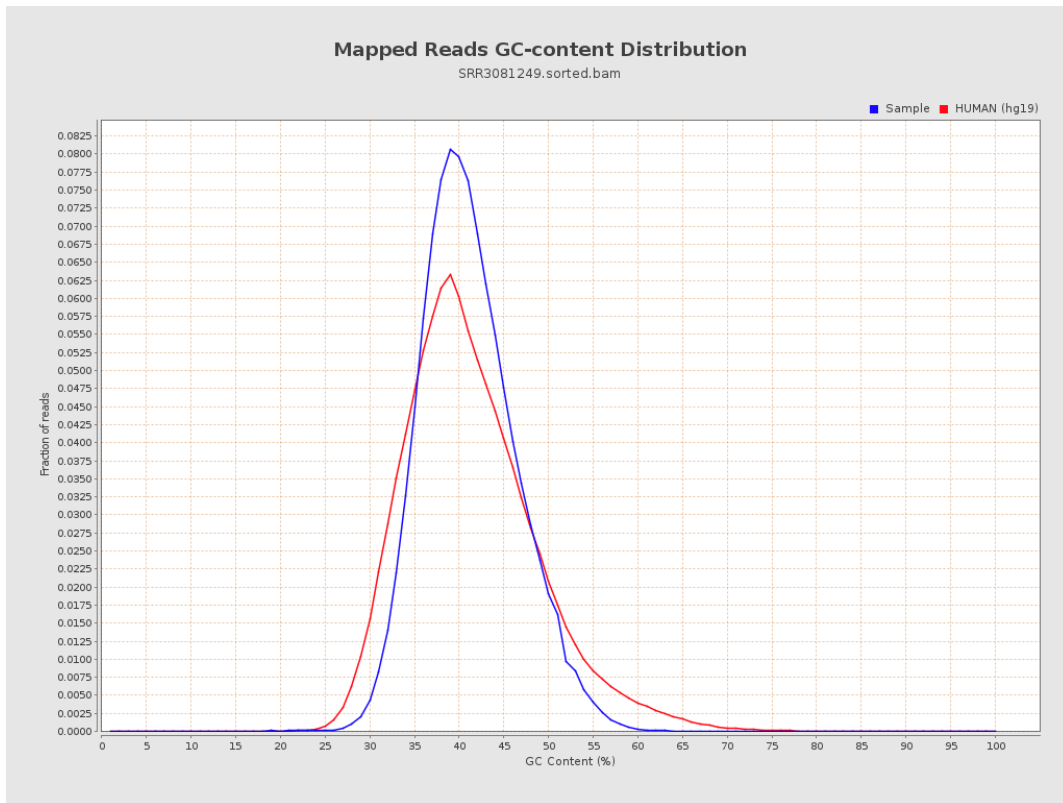
7. Results : Duplication Rate Histogram



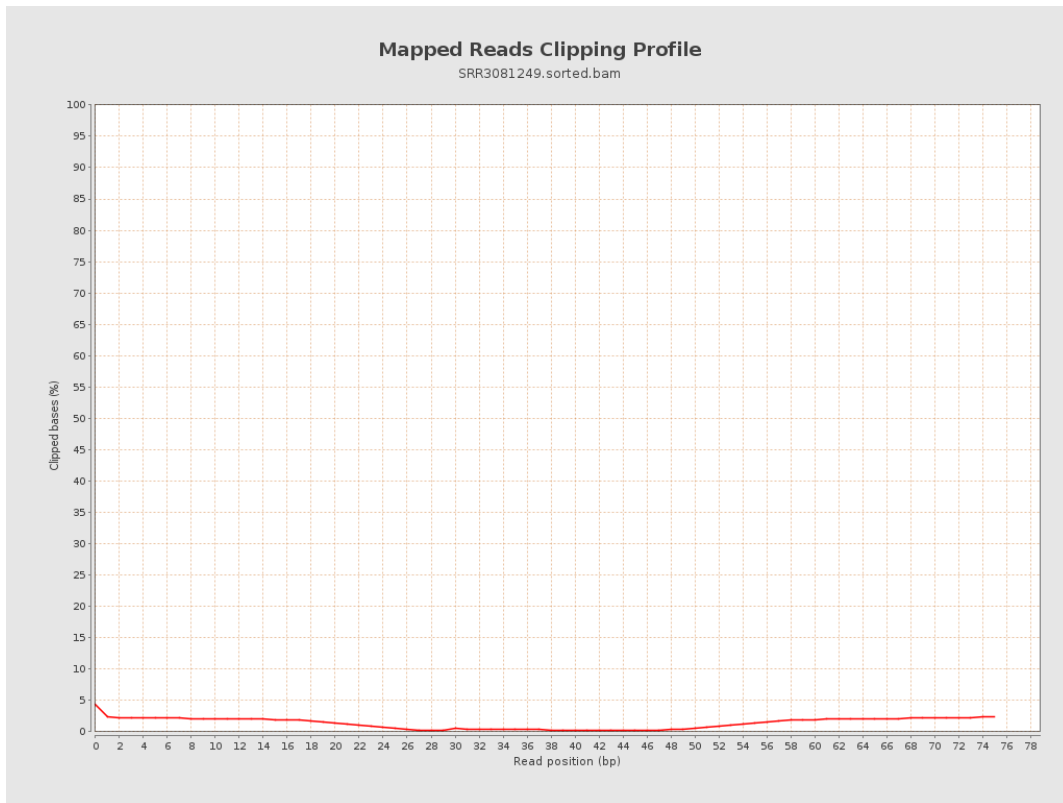
8. Results : Mapped Reads Nucleotide Content



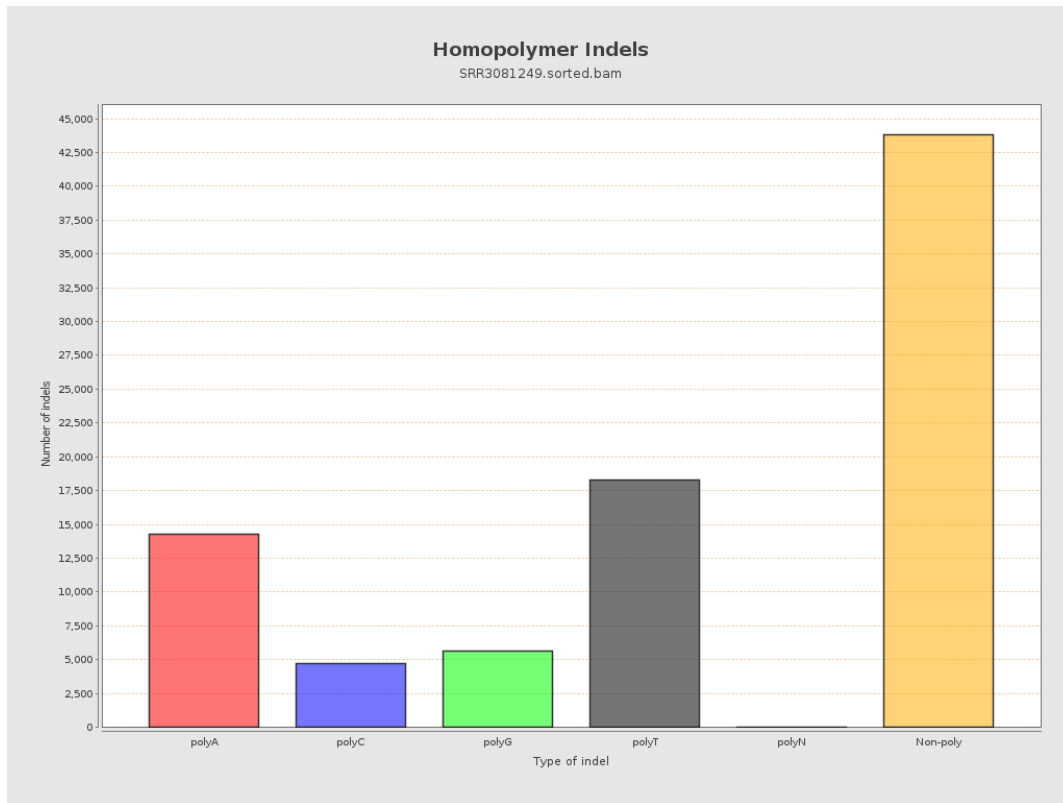
9. Results : Mapped Reads GC-content Distribution



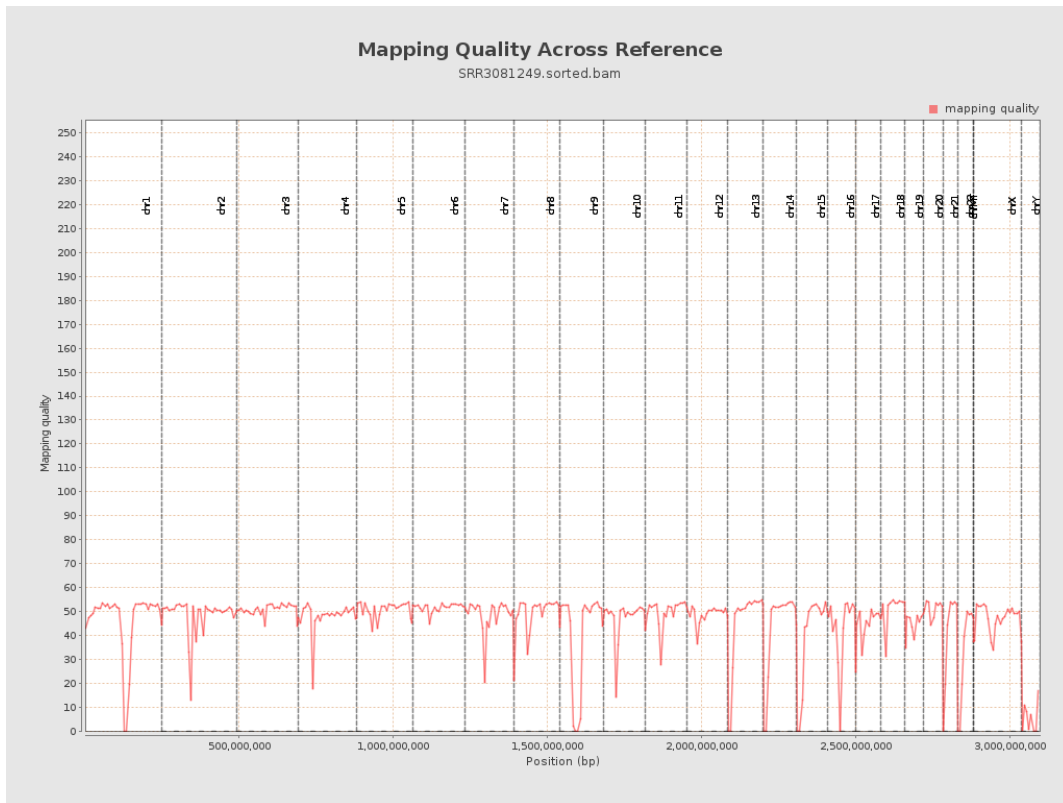
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

