

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/29 06:22:45

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR10524929.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_SRR10524929 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR10524929.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Thu Aug 29 06:22:41 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR10524929.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	710,203
Mapped reads	659,069 / 92.8%
Unmapped reads	51,134 / 7.2%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	2,492 / 0.35%
Read min/max/mean length	30 / 76 / 76.12
Duplicated reads (estimated)	20,177 / 2.84%
Duplication rate	2.41%
Clipped reads	660,254 / 92.97%

2.2. ACGT Content

Number/percentage of A's	9,399,262 / 24.53%
Number/percentage of C's	6,875,567 / 17.95%
Number/percentage of T's	11,998,141 / 31.32%
Number/percentage of G's	10,036,432 / 26.2%
Number/percentage of N's	322 / 0%
GC Percentage	44.15%

2.3. Coverage

Mean	0.0124

Standard Deviation	0.1447
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2.4. Mapping Quality

Mean Mapping Quality	44.78
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2.5. Mismatches and indels

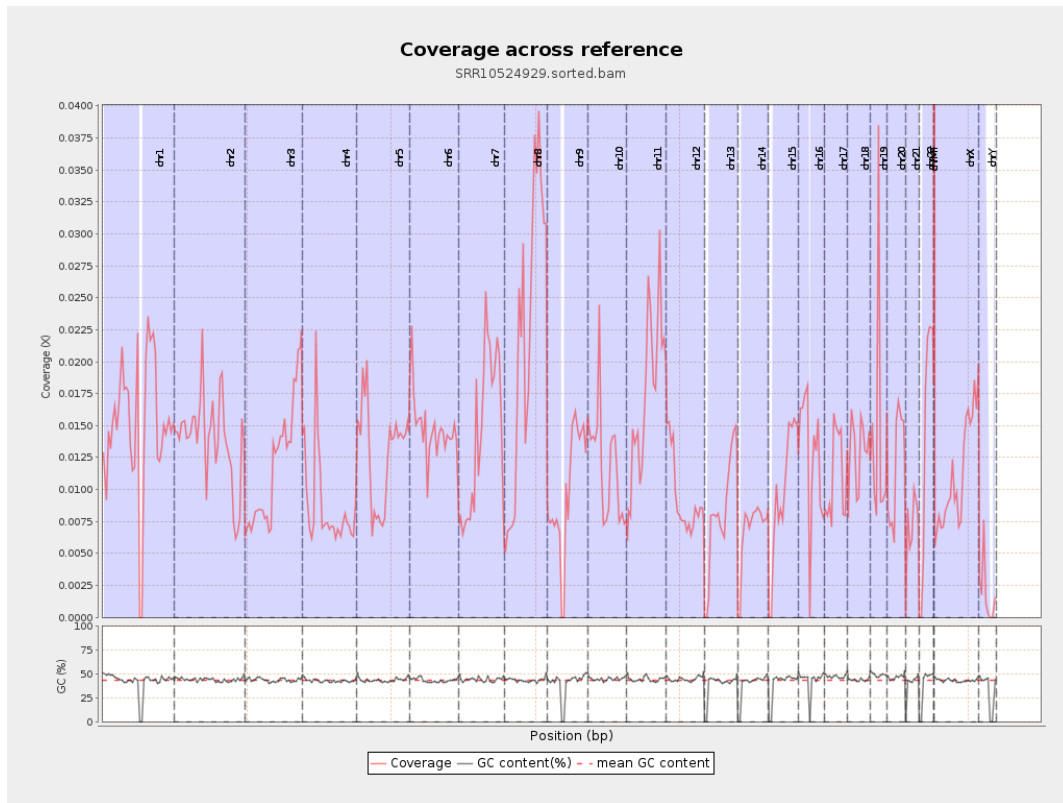
General error rate	0.51%
Mismatches	190,149
Insertions	2,474
Mapped reads with at least one insertion	0.37%
Deletions	7,259
Mapped reads with at least one deletion	1.09%
Homopolymer indels	43.73%

2.6. Chromosome stats

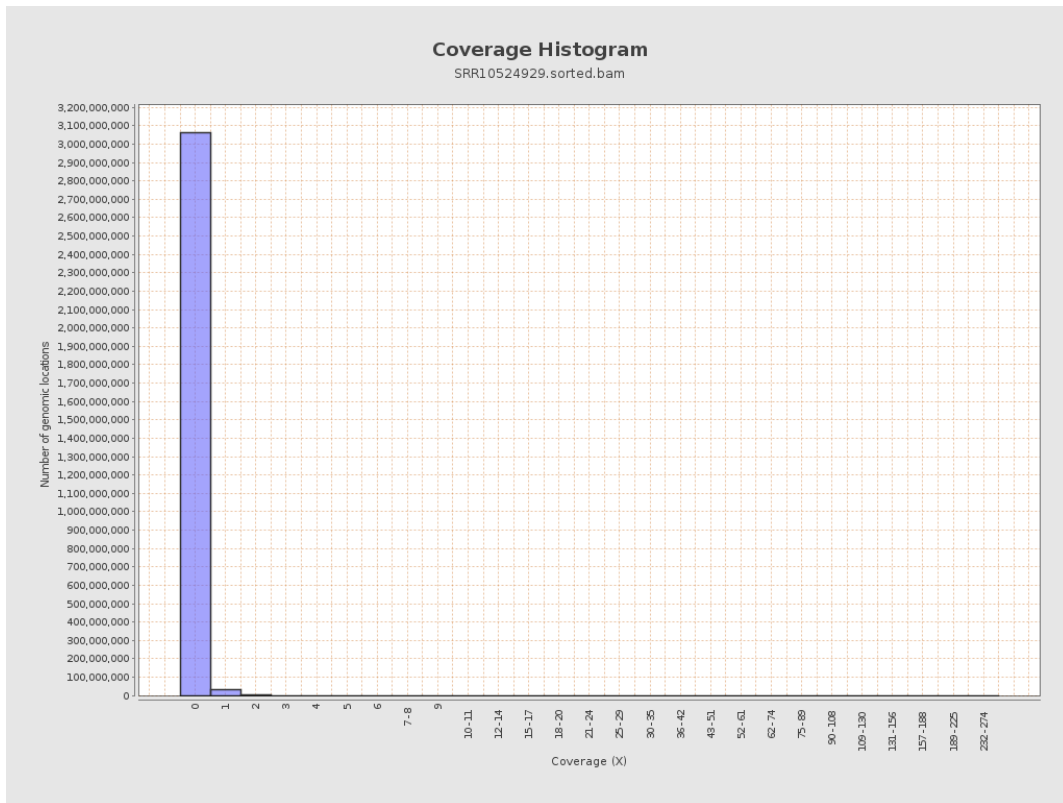
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	3743170	0.015	0.2371
chr2	243199373	3380664	0.0139	0.1686
chr3	198022430	2339479	0.0118	0.1149
chr4	191154276	1690596	0.0088	0.1148
chr5	180915260	2330653	0.0129	0.12
chr6	171115067	2518711	0.0147	0.1325
chr7	159138663	2279234	0.0143	0.1607

chr8	146364022	3236922	0.0221	0.1852
chr9	141213431	1389797	0.0098	0.1182
chr10	135534747	1627834	0.012	0.1491
chr11	135006516	2350453	0.0174	0.1508
chr12	133851895	1257560	0.0094	0.1048
chr13	115169878	942892	0.0082	0.0961
chr14	107349540	723754	0.0067	0.0883
chr15	102531392	996975	0.0097	0.1041
chr16	90354753	1132681	0.0125	0.1221
chr17	81195210	889258	0.011	0.1128
chr18	78077248	1006183	0.0129	0.1834
chr19	59128983	838415	0.0142	0.1747
chr20	63025520	734358	0.0117	0.1151
chr21	48129895	337832	0.007	0.1026
chr22	51304566	731074	0.0142	0.1269
chrMT	16571	3441	0.2077	0.5186
chrX	155270560	1728088	0.0111	0.1162
chrY	59373566	111816	0.0019	0.0803

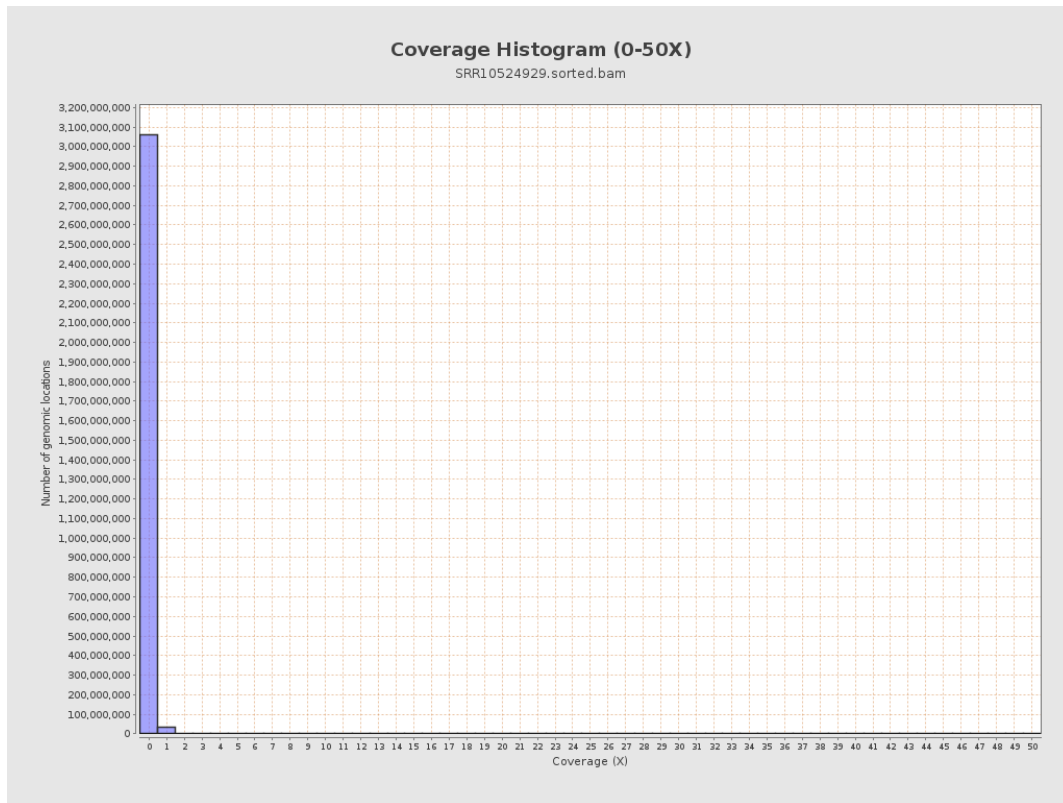
3. Results : Coverage across reference



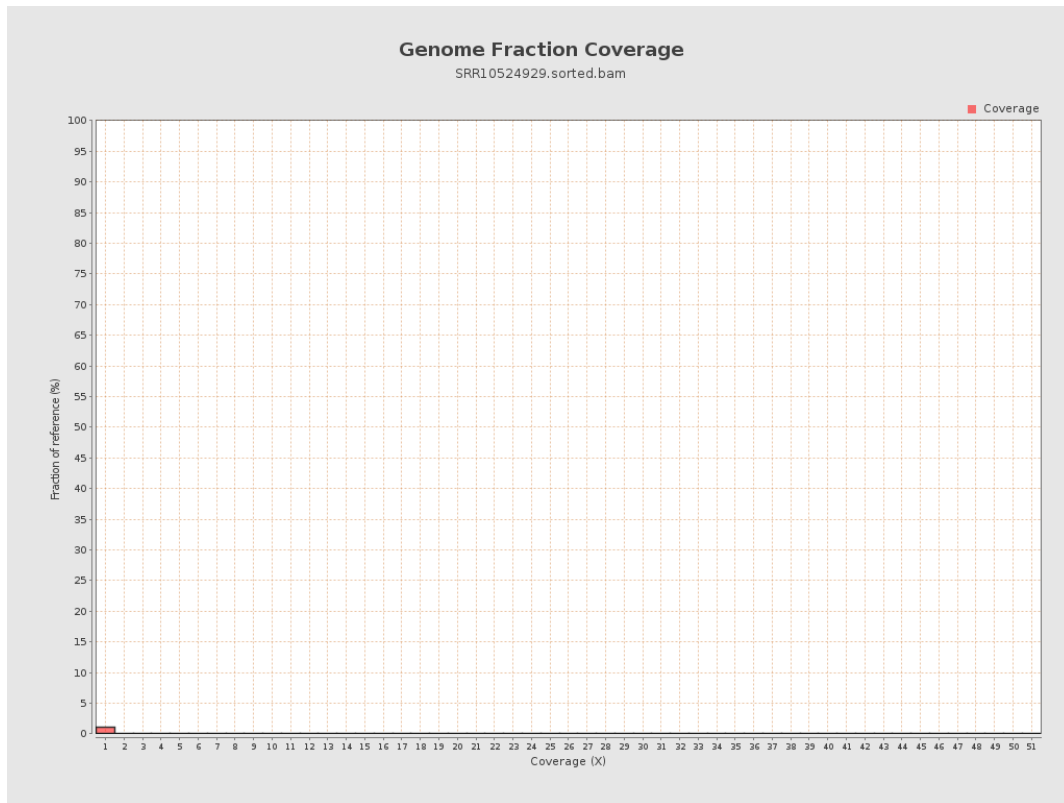
4. Results : Coverage Histogram



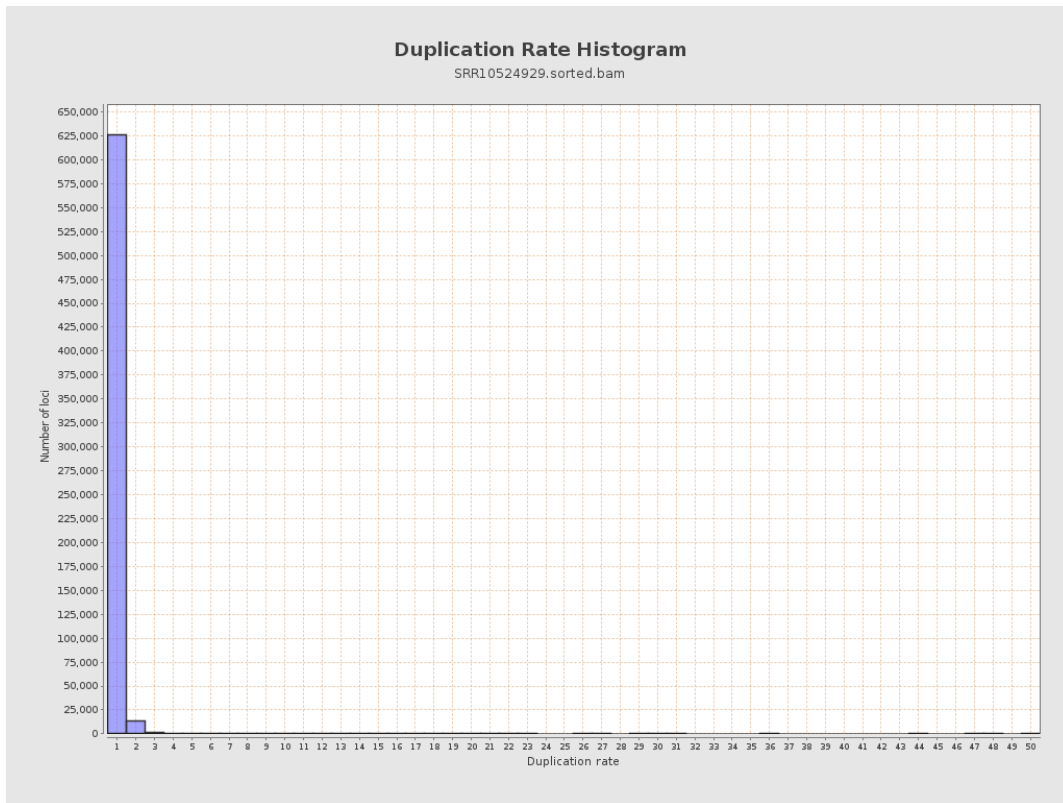
5. Results : Coverage Histogram (0-50X)



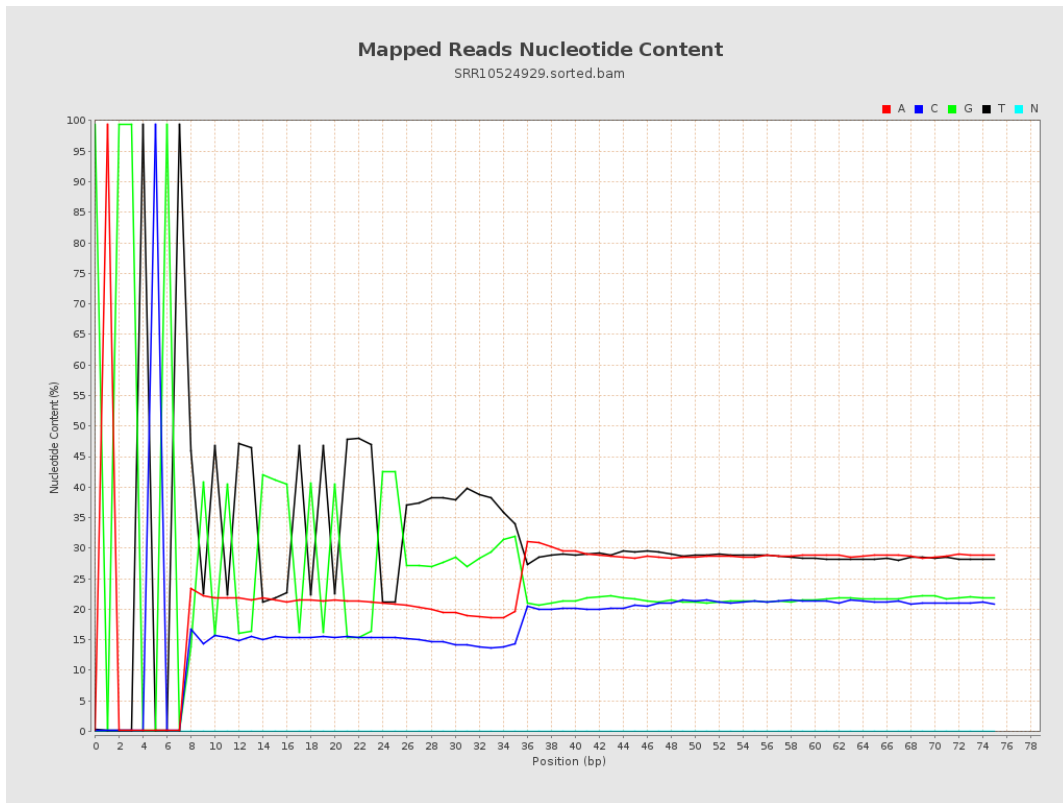
6. Results : Genome Fraction Coverage



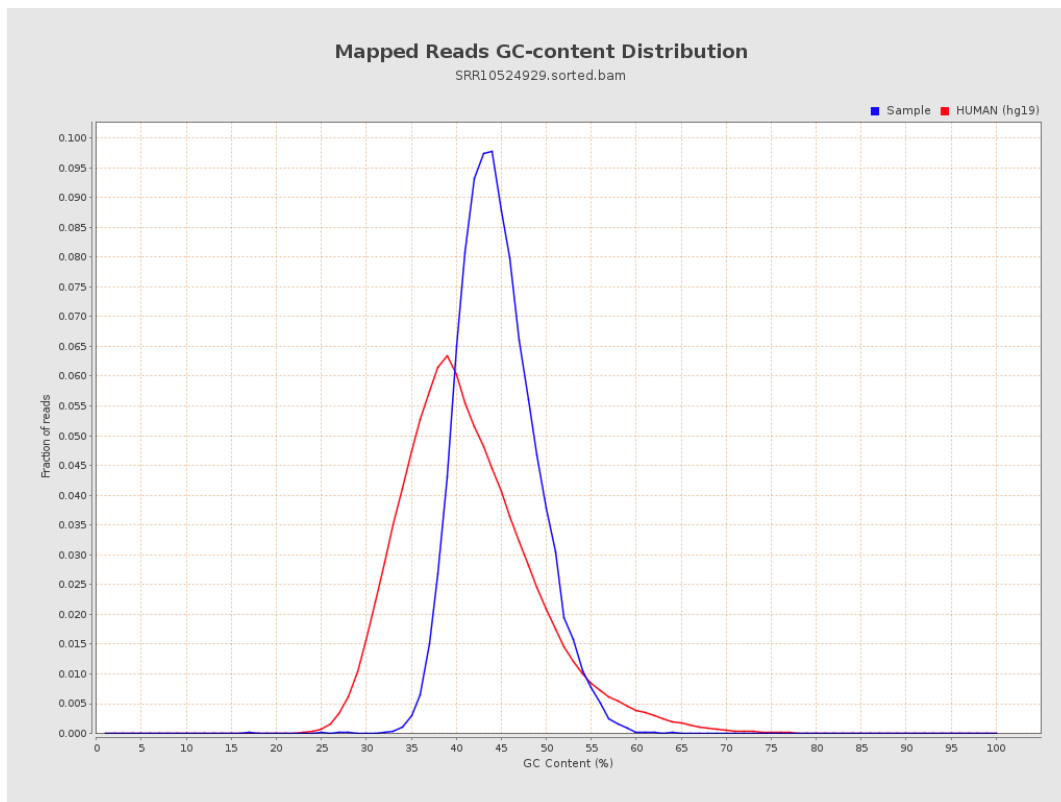
7. Results : Duplication Rate Histogram



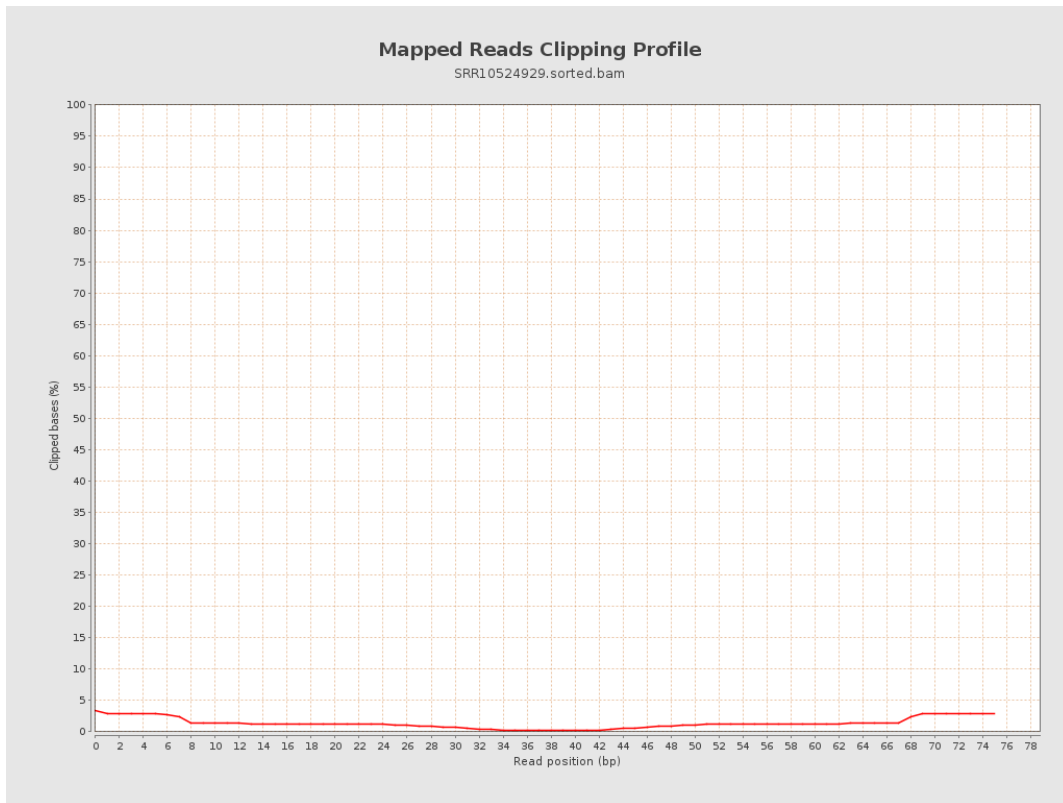
8. Results : Mapped Reads Nucleotide Content



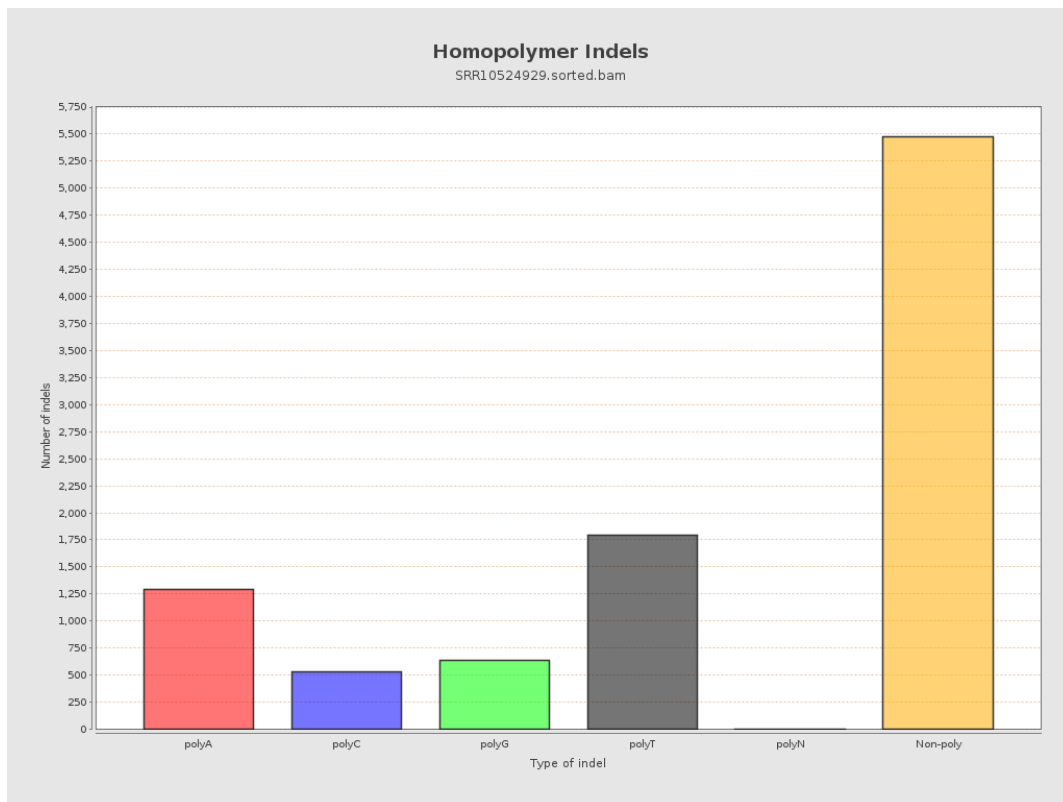
9. Results : Mapped Reads GC-content Distribution



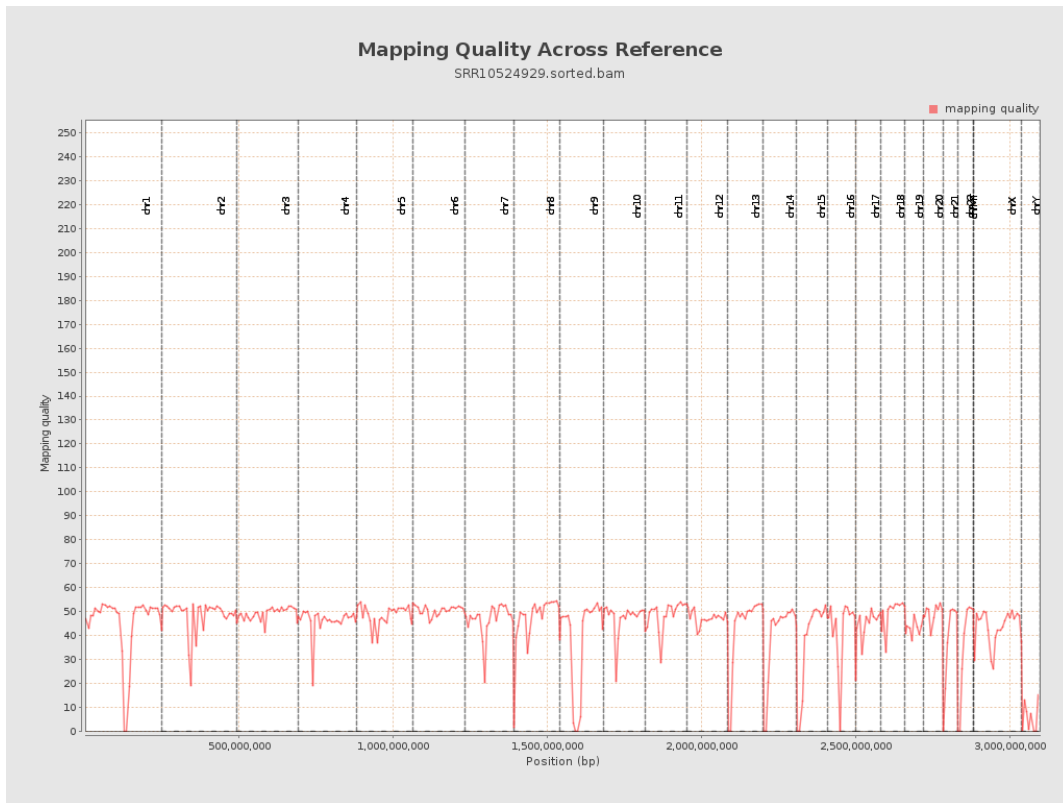
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

