

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/03 13:42:11

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR9716823.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_SRR9716823 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR9716823.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Sep 03 13:42:10 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR9716823.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	760,685
Mapped reads	598,497 / 78.68%
Unmapped reads	162,188 / 21.32%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	664 / 0.09%
Read min/max/mean length	30 / 76 / 76.03
Duplicated reads (estimated)	37,251 / 4.9%
Duplication rate	5.58%
Clipped reads	598,047 / 78.62%

2.2. ACGT Content

Number/percentage of A's	6,913,509 / 22.04%
Number/percentage of C's	6,051,547 / 19.29%
Number/percentage of T's	9,941,725 / 31.7%
Number/percentage of G's	8,458,044 / 26.97%
Number/percentage of N's	595 / 0%
GC Percentage	46.26%

2.3. Coverage

Mean	0.0101

Standard Deviation	0.12
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2.4. Mapping Quality

Mean Mapping Quality	43
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2.5. Mismatches and indels

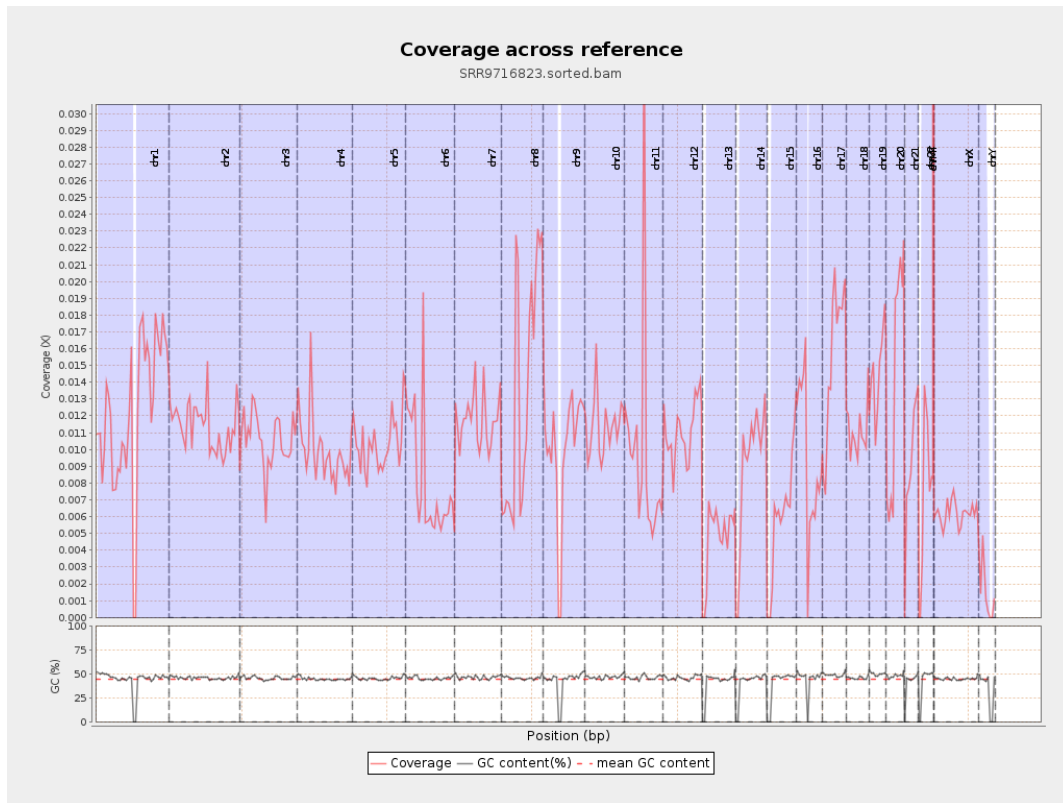
General error rate	0.65%
Mismatches	198,983
Insertions	1,940
Mapped reads with at least one insertion	0.32%
Deletions	4,398
Mapped reads with at least one deletion	0.73%
Homopolymer indels	41.51%

2.6. Chromosome stats

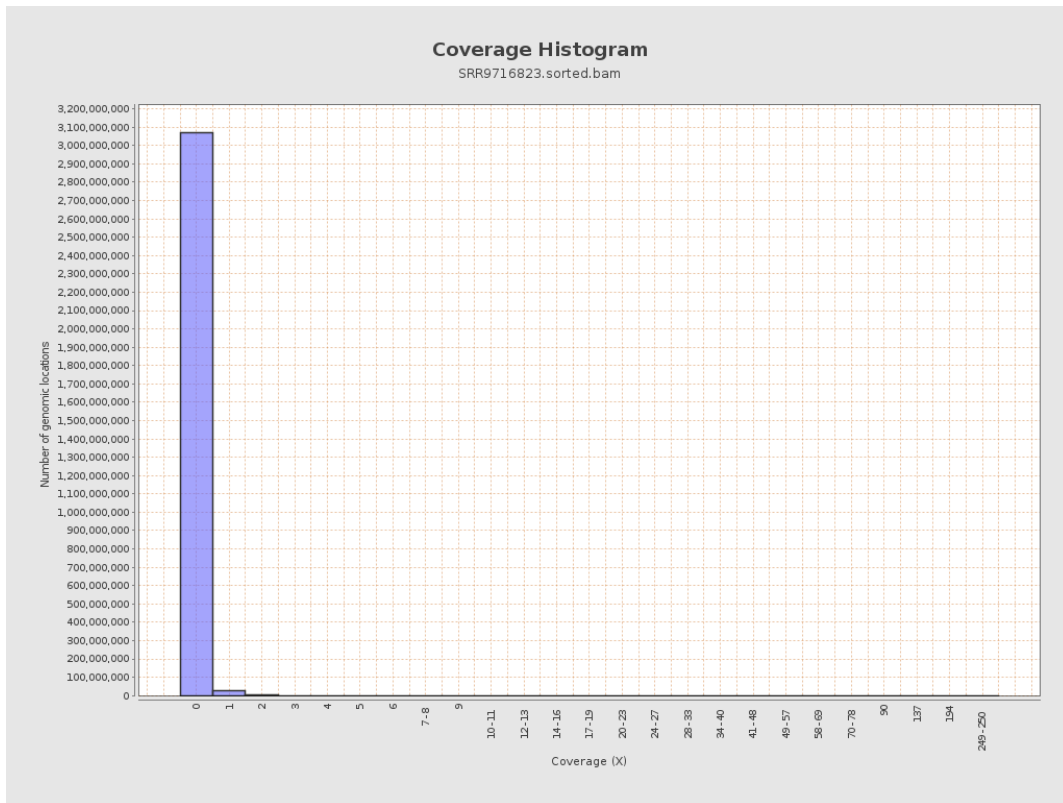
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	3042764	0.0122	0.1356
chr2	243199373	2770840	0.0114	0.156
chr3	198022430	2091294	0.0106	0.114
chr4	191154276	1877146	0.0098	0.1132
chr5	180915260	1872544	0.0104	0.1128
chr6	171115067	1387455	0.0081	0.1052
chr7	159138663	1866818	0.0117	0.1315

chr8	146364022	1976380	0.0135	0.1317
chr9	141213431	1392615	0.0099	0.1133
chr10	135534747	1533671	0.0113	0.1281
chr11	135006516	1323429	0.0098	0.1165
chr12	133851895	1479721	0.0111	0.117
chr13	115169878	543298	0.0047	0.0768
chr14	107349540	971042	0.009	0.1056
chr15	102531392	614309	0.006	0.0857
chr16	90354753	848582	0.0094	0.1111
chr17	81195210	1294596	0.0159	0.1428
chr18	78077248	857738	0.011	0.1268
chr19	59128983	860601	0.0146	0.1553
chr20	63025520	871735	0.0138	0.1355
chr21	48129895	446647	0.0093	0.1118
chr22	51304566	364565	0.0071	0.0959
chrMT	16571	36633	2.2107	2.2661
chrX	155270560	964054	0.0062	0.0889
chrY	59373566	84302	0.0014	0.049

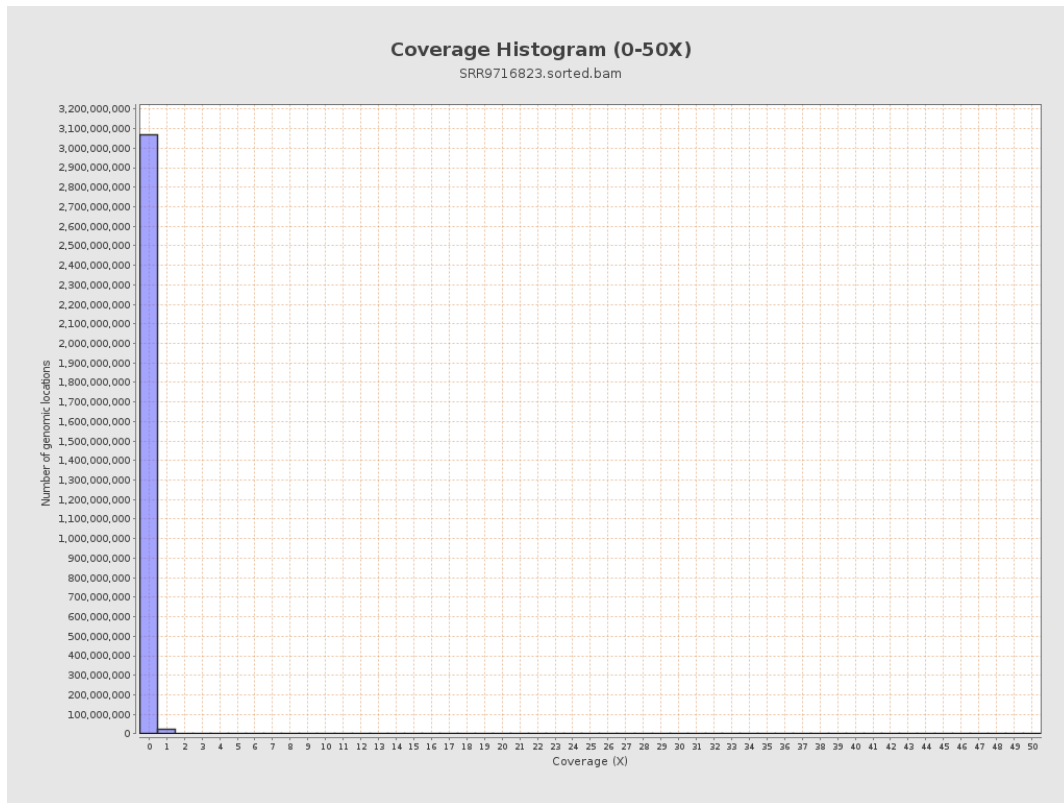
3. Results : Coverage across reference



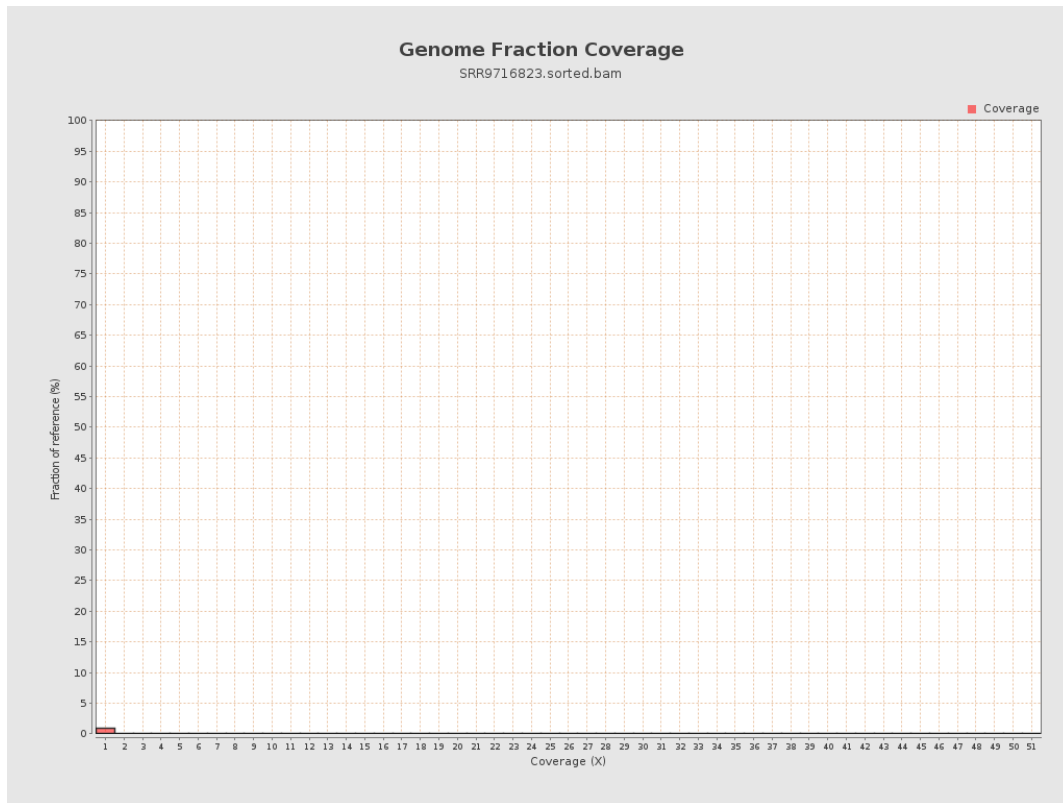
4. Results : Coverage Histogram



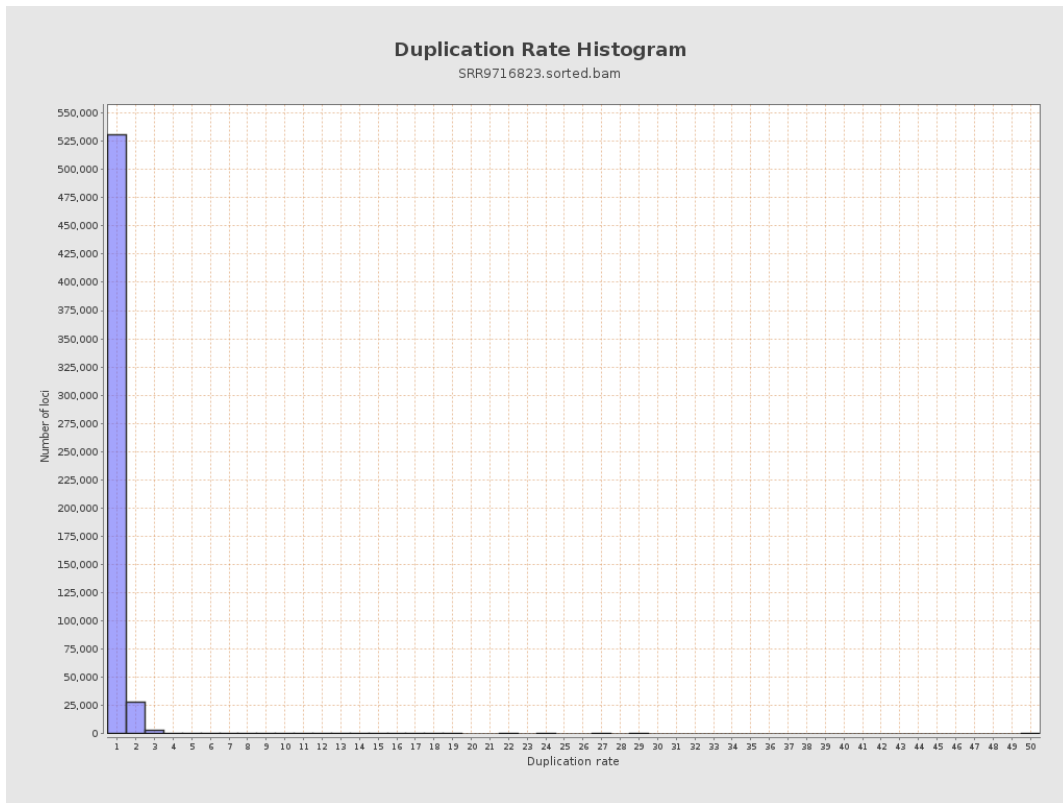
5. Results : Coverage Histogram (0-50X)



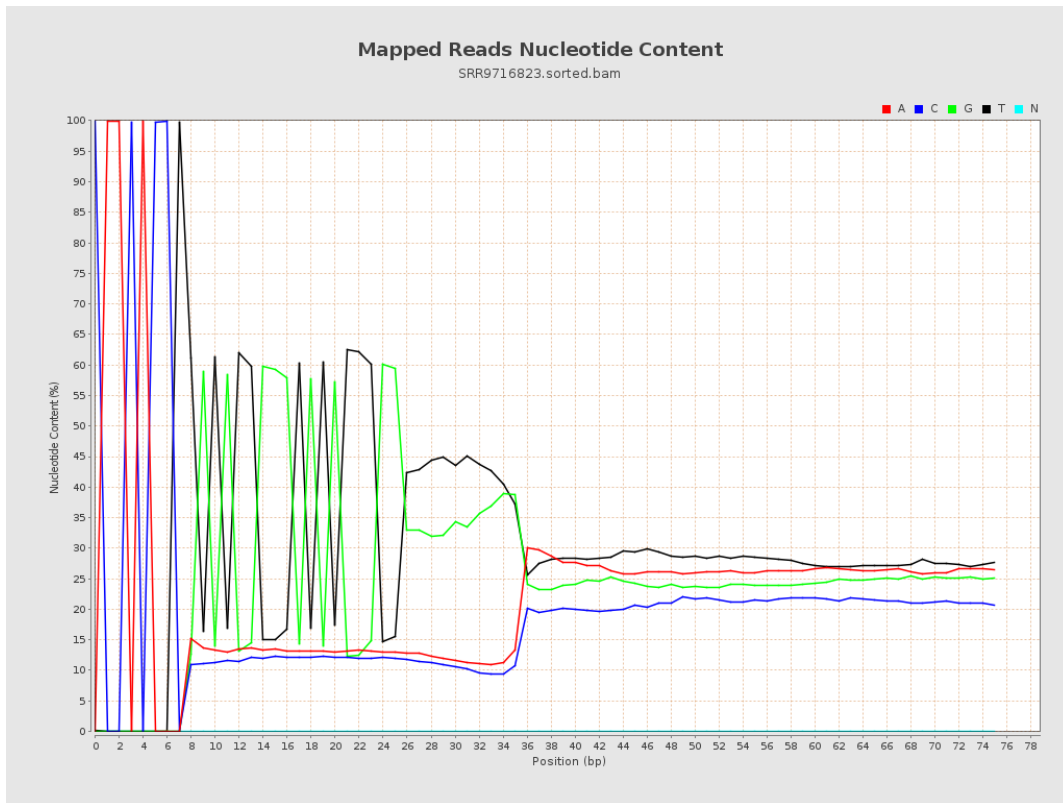
6. Results : Genome Fraction Coverage



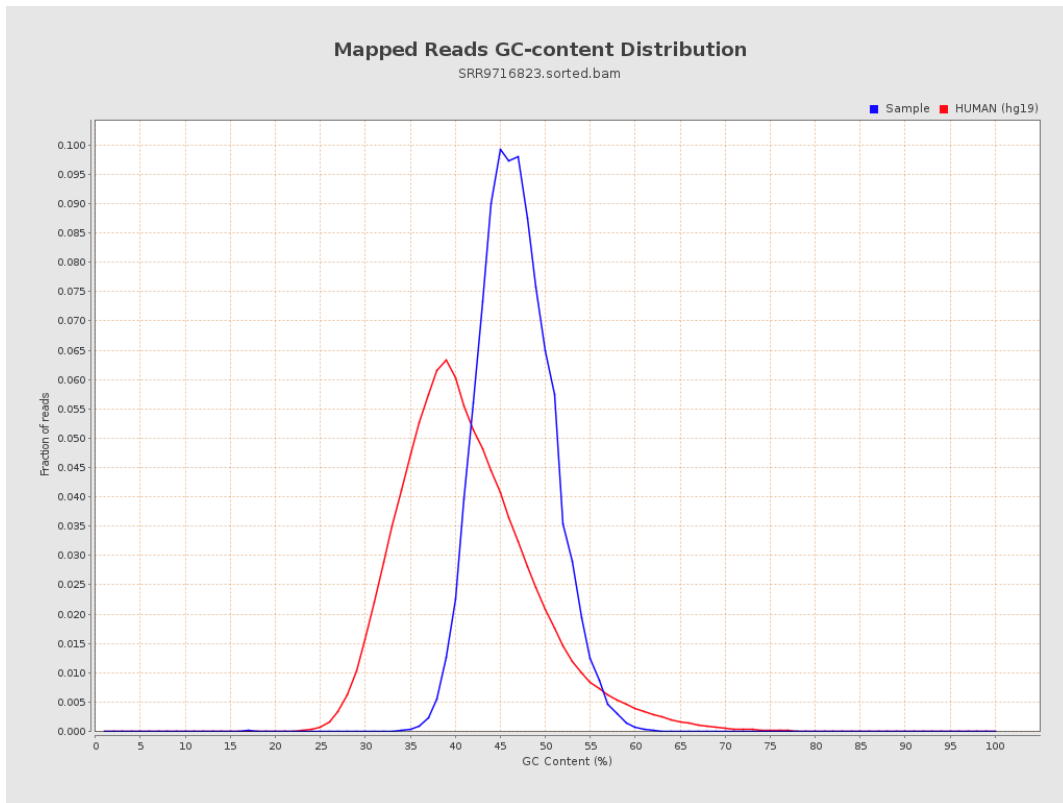
7. Results : Duplication Rate Histogram



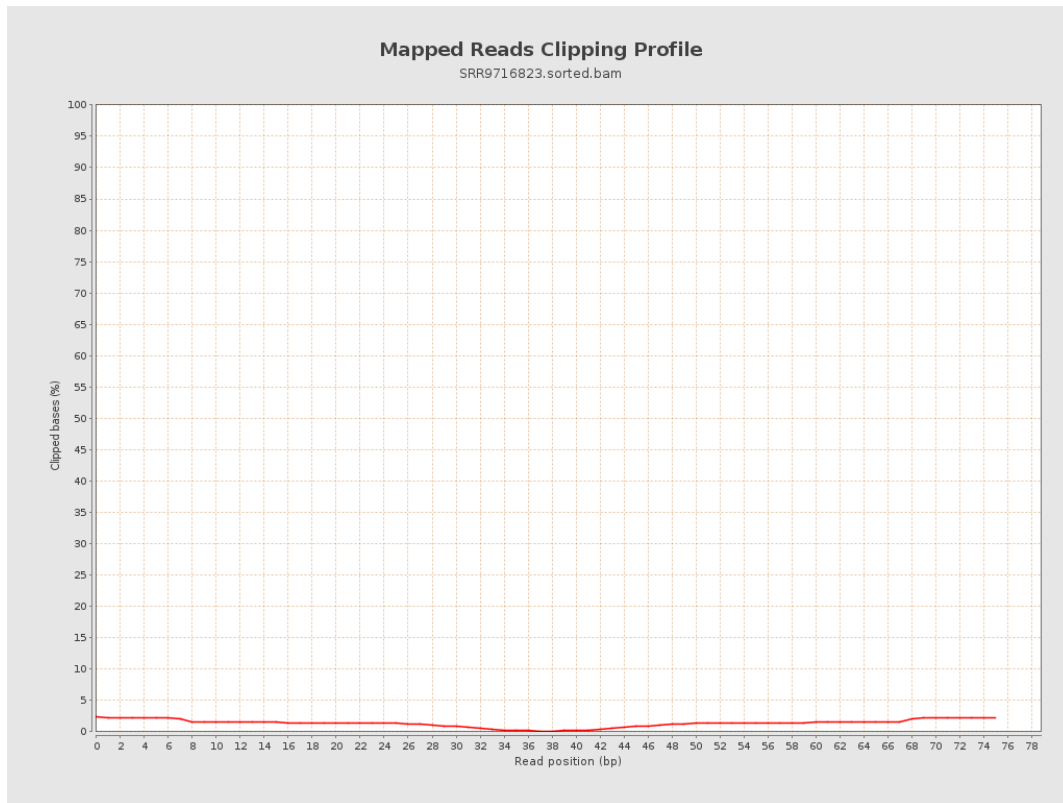
8. Results : Mapped Reads Nucleotide Content



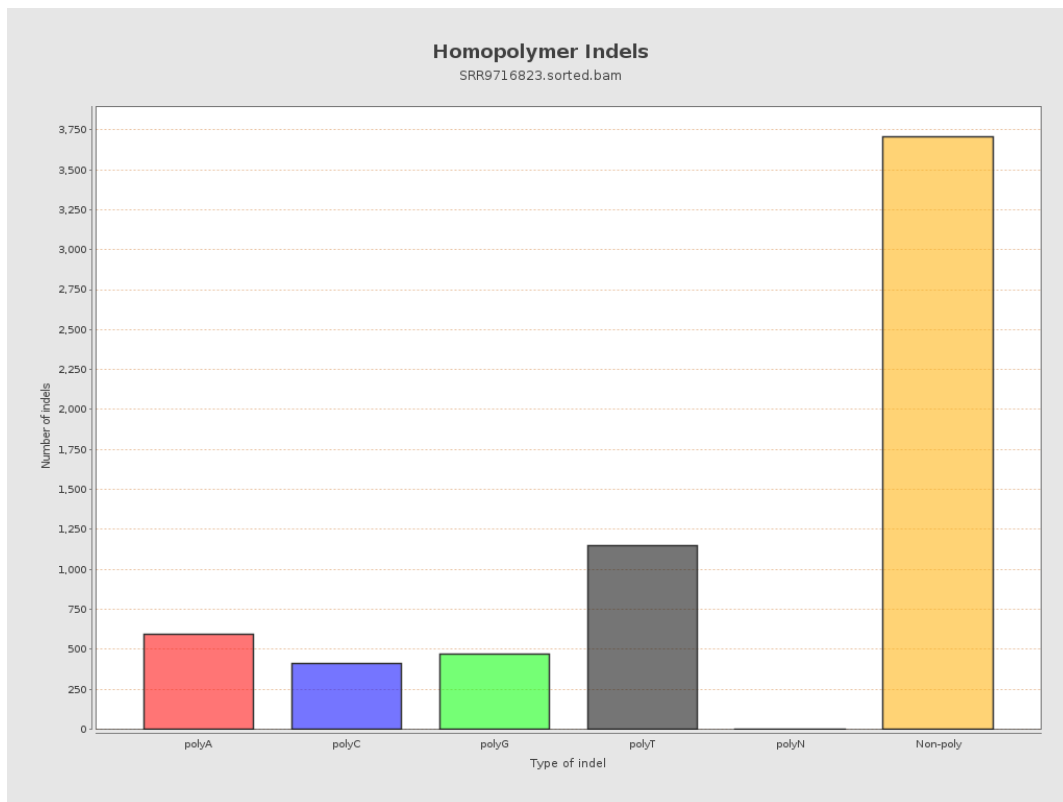
9. Results : Mapped Reads GC-content Distribution



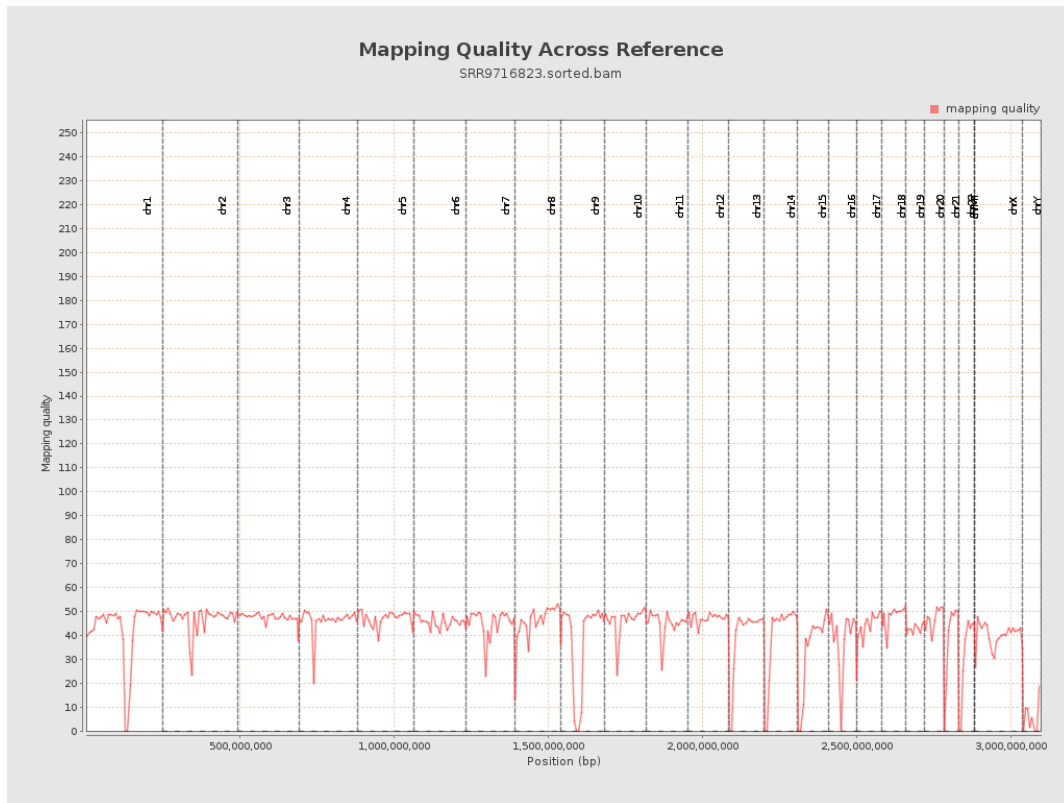
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

