

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/02 05:49:35

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR9716288.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_SRR9716288 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR9716288.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Mon Sep 02 05:49:34 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR9716288.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	724,812
Mapped reads	577,781 / 79.71%
Unmapped reads	147,031 / 20.29%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	1,762 / 0.24%
Read min/max/mean length	30 / 76 / 76.08
Duplicated reads (estimated)	9,852 / 1.36%
Duplication rate	1.16%
Clipped reads	578,224 / 79.78%

2.2. ACGT Content

Number/percentage of A's	7,559,136 / 23.7%
Number/percentage of C's	6,234,898 / 19.55%
Number/percentage of T's	9,523,065 / 29.86%
Number/percentage of G's	8,579,244 / 26.9%
Number/percentage of N's	900 / 0%
GC Percentage	46.44%

2.3. Coverage

Mean	0.0103

Standard Deviation	0.1187
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2.4. Mapping Quality

Mean Mapping Quality	39.03
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2.5. Mismatches and indels

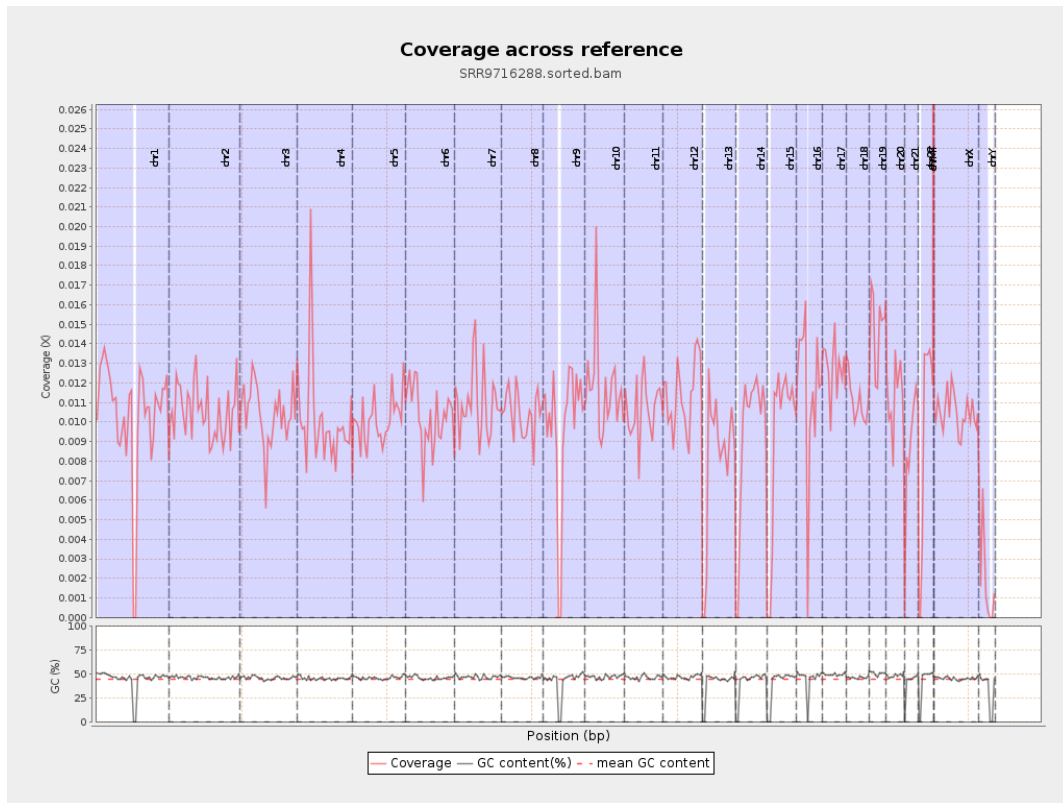
General error rate	0.55%
Mismatches	170,848
Insertions	2,544
Mapped reads with at least one insertion	0.44%
Deletions	5,581
Mapped reads with at least one deletion	0.96%
Homopolymer indels	36.74%

2.6. Chromosome stats

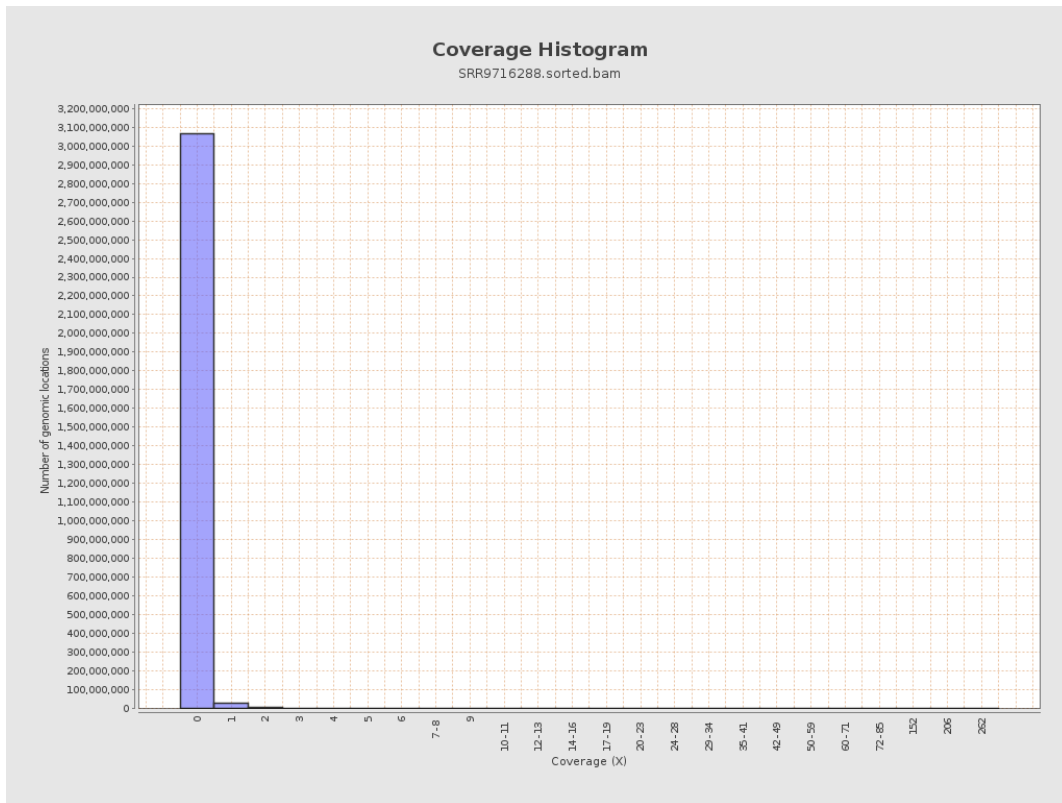
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	2577700	0.0103	0.1217
chr2	243199373	2562272	0.0105	0.1542
chr3	198022430	2061916	0.0104	0.1105
chr4	191154276	1904519	0.01	0.1153
chr5	180915260	1834105	0.0101	0.1063
chr6	171115067	1756485	0.0103	0.1102
chr7	159138663	1740889	0.0109	0.1275

chr8	146364022	1534363	0.0105	0.1141
chr9	141213431	1361830	0.0096	0.1087
chr10	135534747	1577615	0.0116	0.1396
chr11	135006516	1439167	0.0107	0.122
chr12	133851895	1516541	0.0113	0.1123
chr13	115169878	927737	0.0081	0.0942
chr14	107349540	1006238	0.0094	0.1039
chr15	102531392	946512	0.0092	0.1026
chr16	90354753	1052143	0.0116	0.1189
chr17	81195210	1032727	0.0127	0.1226
chr18	78077248	863408	0.0111	0.1319
chr19	59128983	870721	0.0147	0.1438
chr20	63025520	704947	0.0112	0.114
chr21	48129895	423281	0.0088	0.1082
chr22	51304566	468491	0.0091	0.1025
chrMT	16571	1476	0.0891	0.3043
chrX	155270560	1628828	0.0105	0.1144
chrY	59373566	112711	0.0019	0.0634

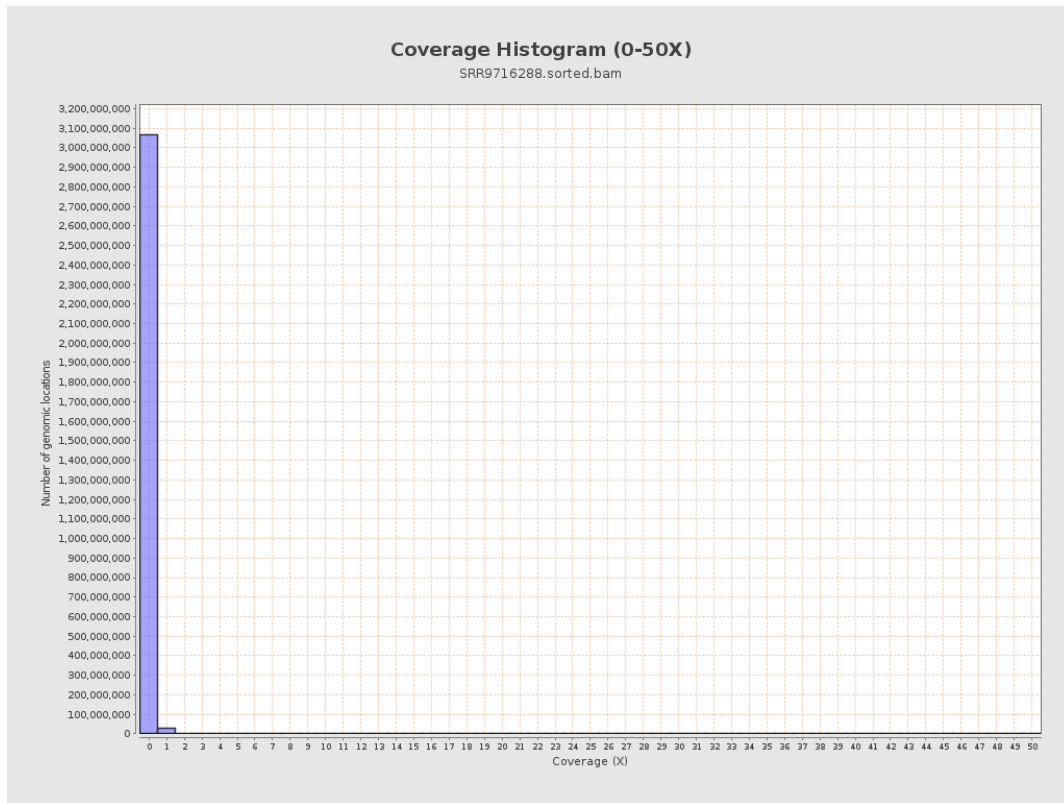
3. Results : Coverage across reference



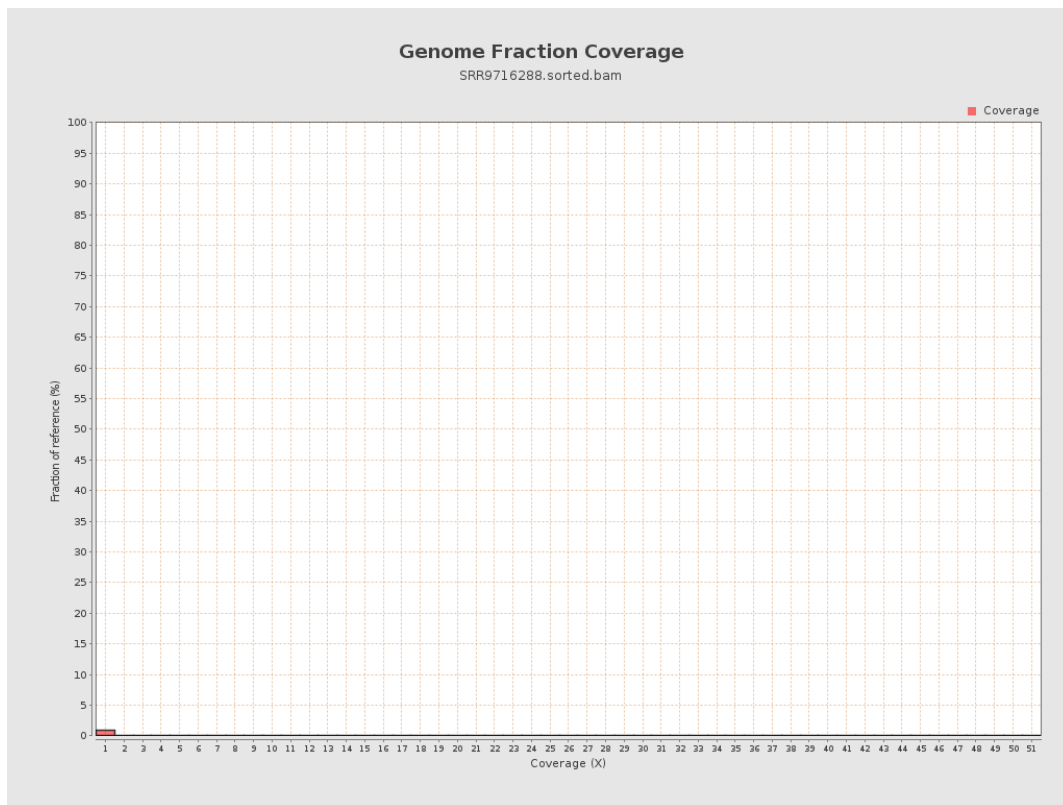
4. Results : Coverage Histogram



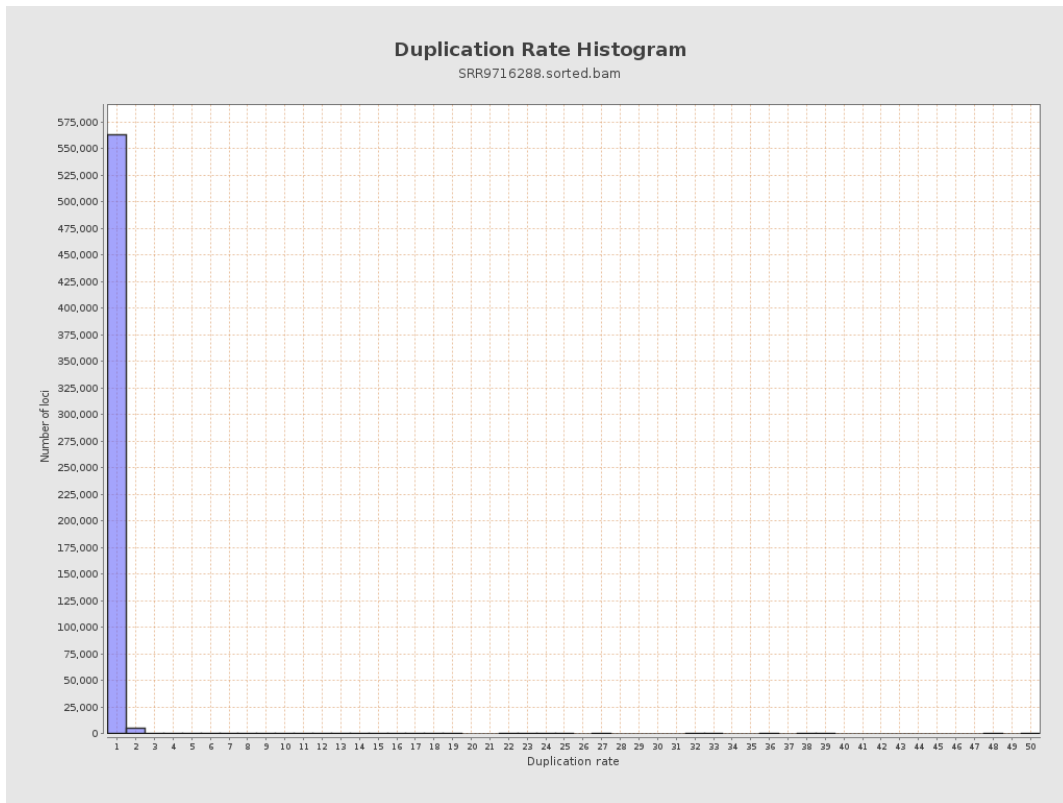
5. Results : Coverage Histogram (0-50X)



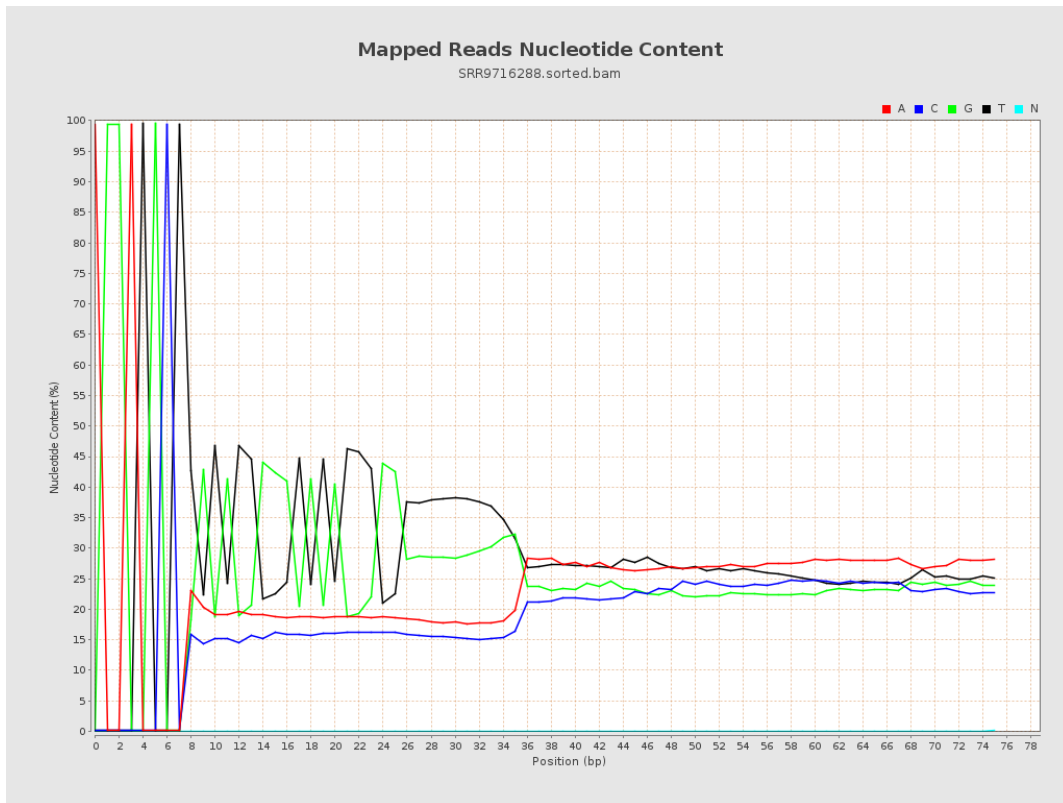
6. Results : Genome Fraction Coverage



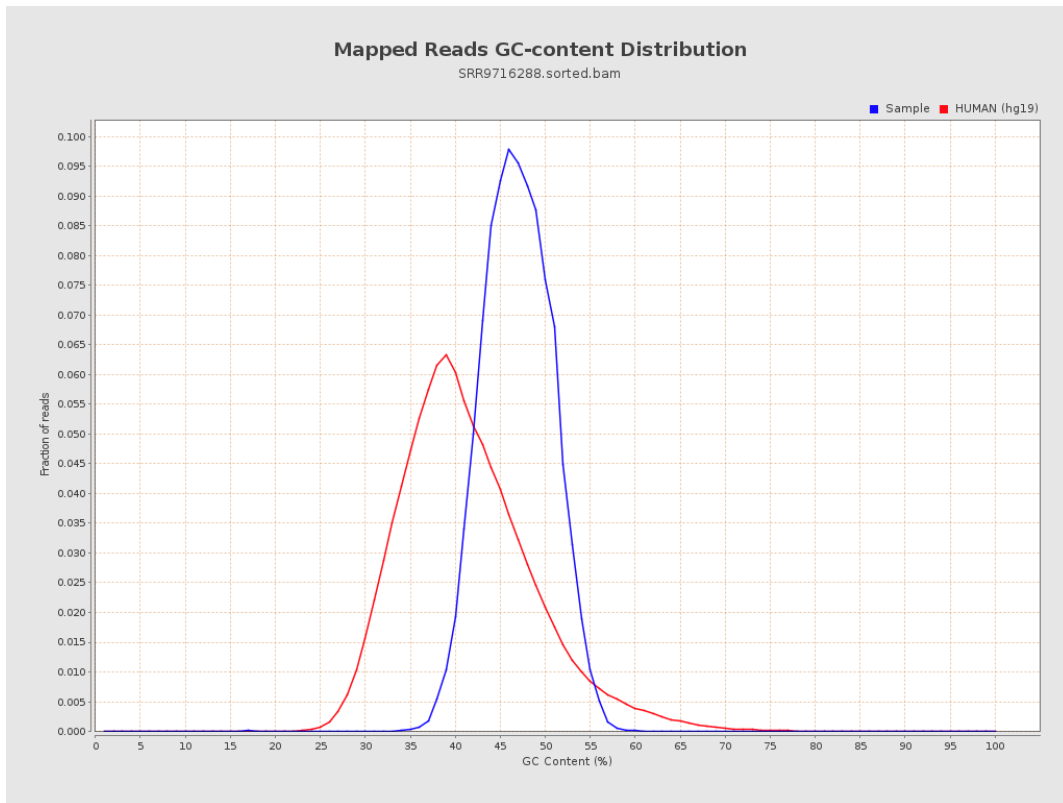
7. Results : Duplication Rate Histogram



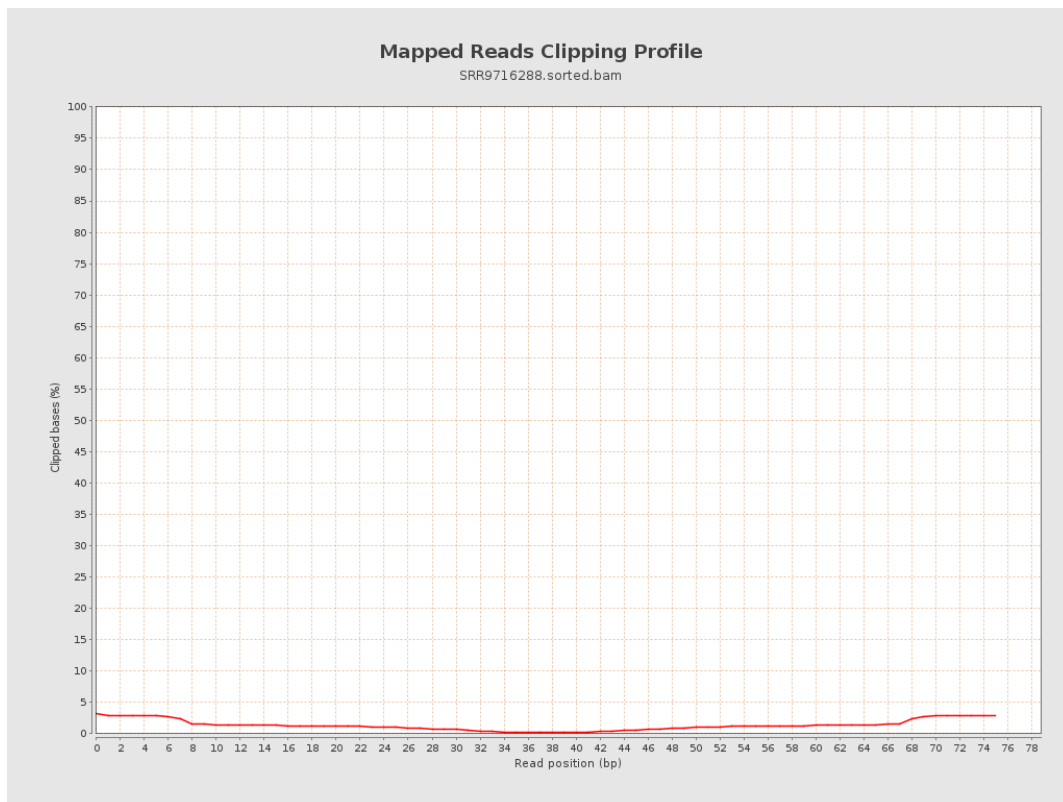
8. Results : Mapped Reads Nucleotide Content



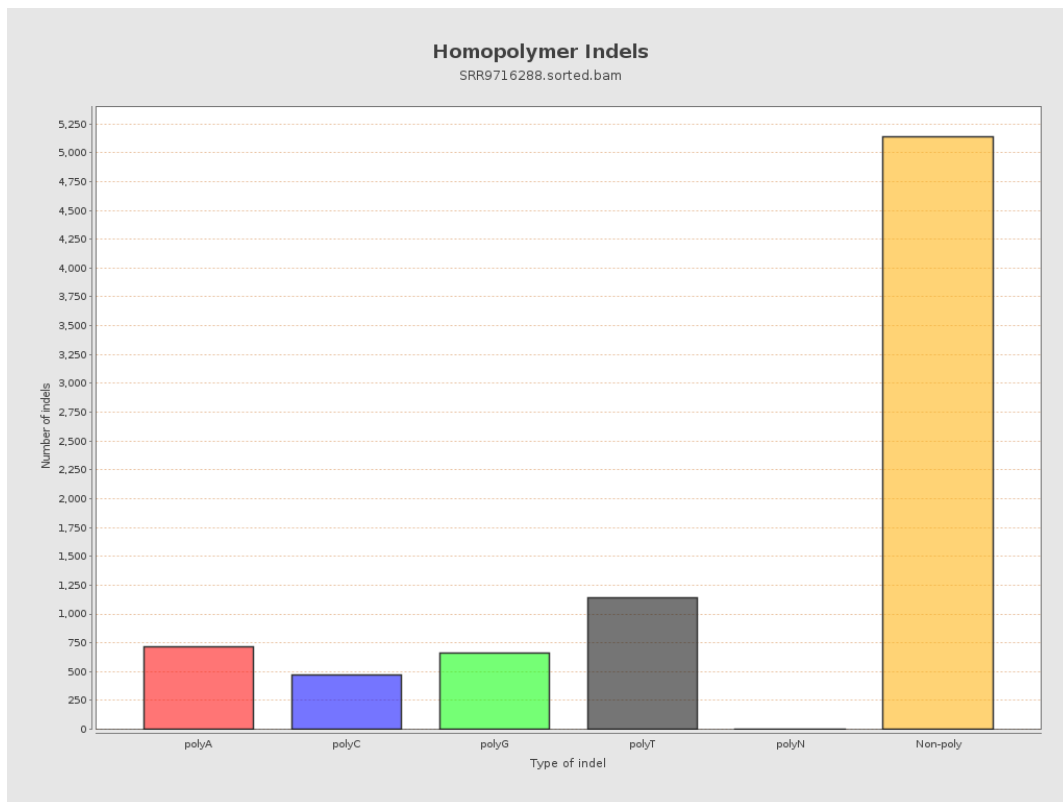
9. Results : Mapped Reads GC-content Distribution



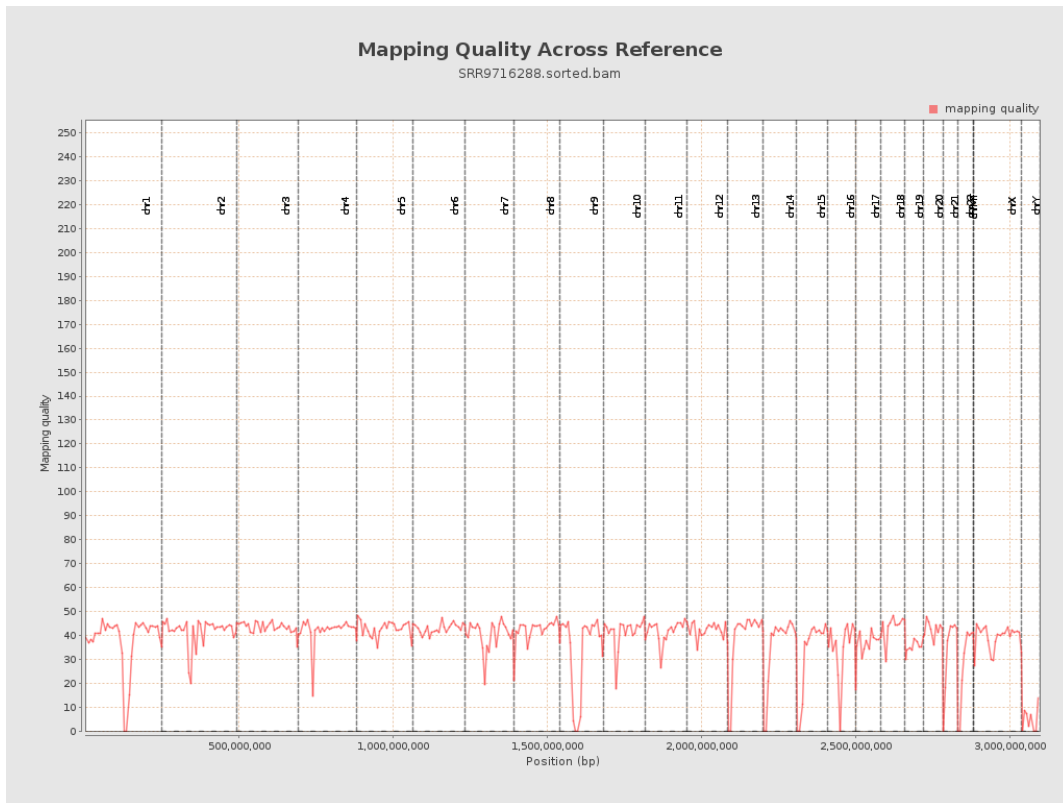
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

