

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/13 19:26:49

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6003991.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_SRR6003991 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6003991.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Fri Sep 13 19:26:48 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6003991.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,315,451
Mapped reads	2,764,987 / 83.4%
Unmapped reads	550,464 / 16.6%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	17,890 / 0.54%
Read min/max/mean length	30 / 76 / 76.19
Duplicated reads (estimated)	130,488 / 3.94%
Duplication rate	3.32%
Clipped reads	1,200,497 / 36.21%

2.2. ACGT Content

Number/percentage of A's	52,405,027 / 28.25%
Number/percentage of C's	33,199,206 / 17.9%
Number/percentage of T's	60,047,906 / 32.37%
Number/percentage of G's	39,804,220 / 21.46%
Number/percentage of N's	20,123 / 0.01%
GC Percentage	39.36%

2.3. Coverage

Mean	0.0599

Standard Deviation	0.6063
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2.4. Mapping Quality

Mean Mapping Quality	45.3
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2.5. Mismatches and indels

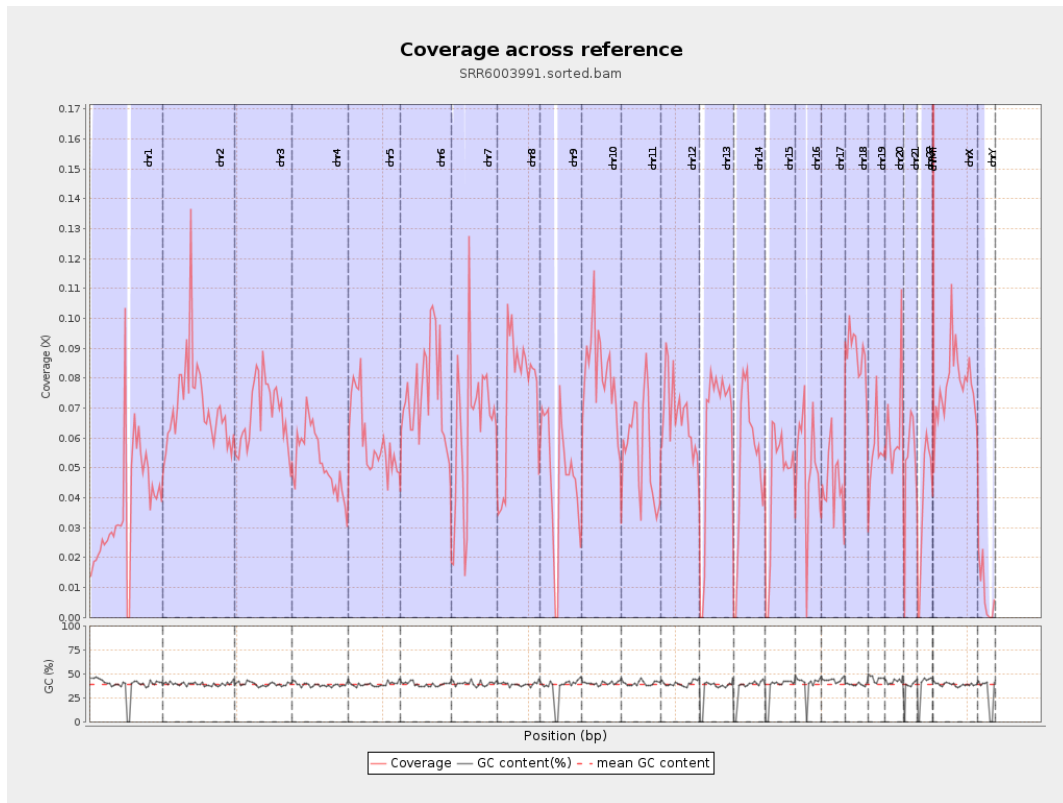
General error rate	0.97%
Mismatches	1,770,070
Insertions	16,925
Mapped reads with at least one insertion	0.61%
Deletions	49,811
Mapped reads with at least one deletion	1.78%
Homopolymer indels	46.8%

2.6. Chromosome stats

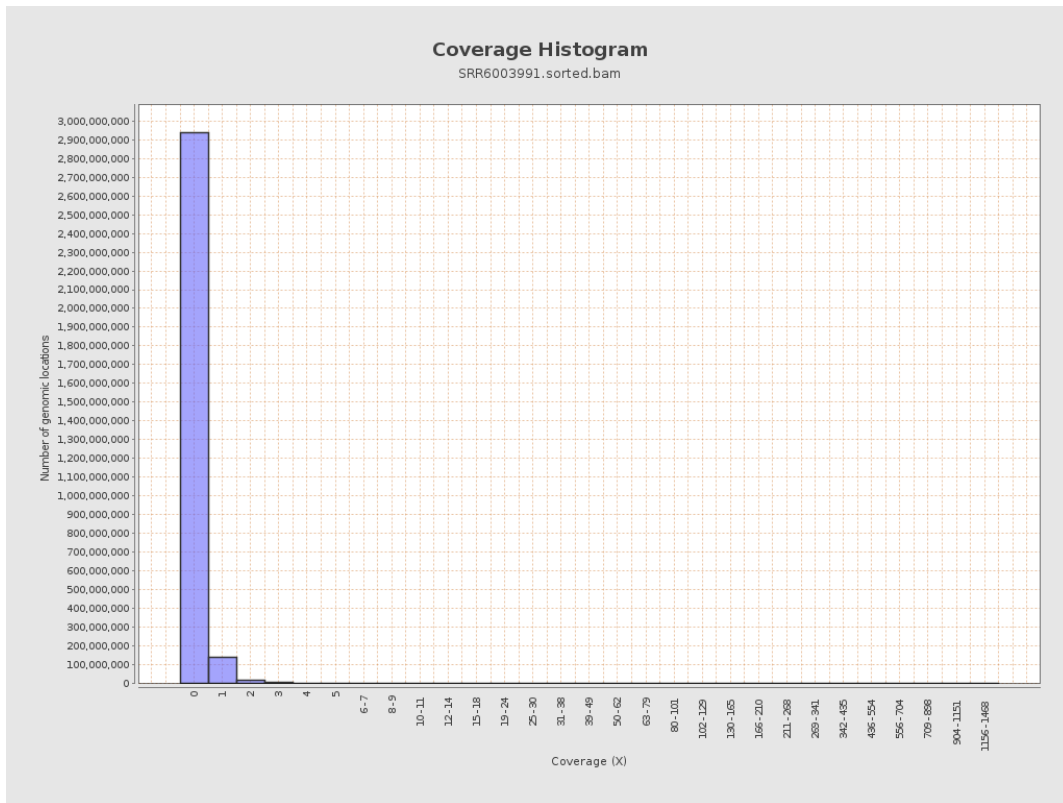
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	9152434	0.0367	1.1389
chr2	243199373	17161937	0.0706	0.6818
chr3	198022430	13448911	0.0679	0.2982
chr4	191154276	9963627	0.0521	0.2775
chr5	180915260	10697797	0.0591	0.2863
chr6	171115067	12753363	0.0745	0.3795
chr7	159138663	10105464	0.0635	0.8794

chr8	146364022	10961463	0.0749	0.9626
chr9	141213431	6749687	0.0478	0.5694
chr10	135534747	11017900	0.0813	0.5426
chr11	135006516	7496297	0.0555	0.4762
chr12	133851895	8962413	0.067	0.3134
chr13	115169878	7226724	0.0627	0.2834
chr14	107349540	5709984	0.0532	0.3133
chr15	102531392	4624173	0.0451	0.2447
chr16	90354753	4534713	0.0502	0.3542
chr17	81195210	3665794	0.0451	0.288
chr18	78077248	6901744	0.0884	1.2689
chr19	59128983	3305519	0.0559	0.7552
chr20	63025520	4041042	0.0641	0.3088
chr21	48129895	2478028	0.0515	0.3005
chr22	51304566	2008479	0.0391	0.2218
chrMT	16571	31899	1.925	1.7914
chrX	155270560	12080536	0.0778	0.3854
chrY	59373566	479460	0.0081	0.163

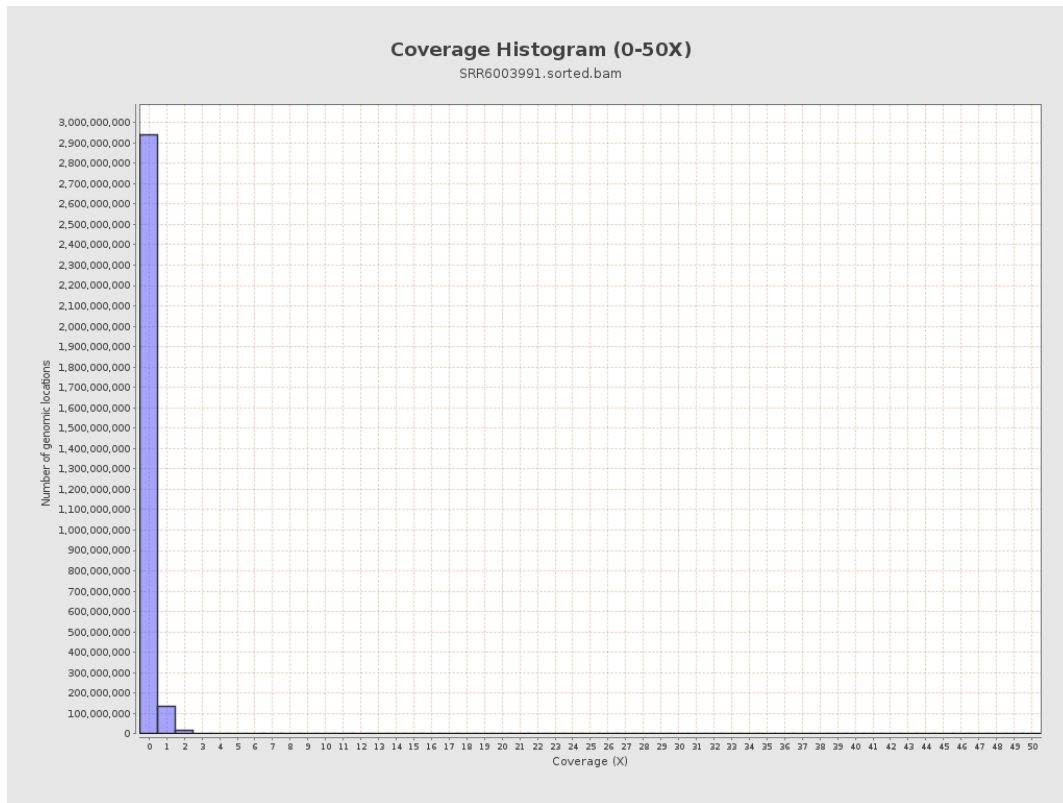
3. Results : Coverage across reference



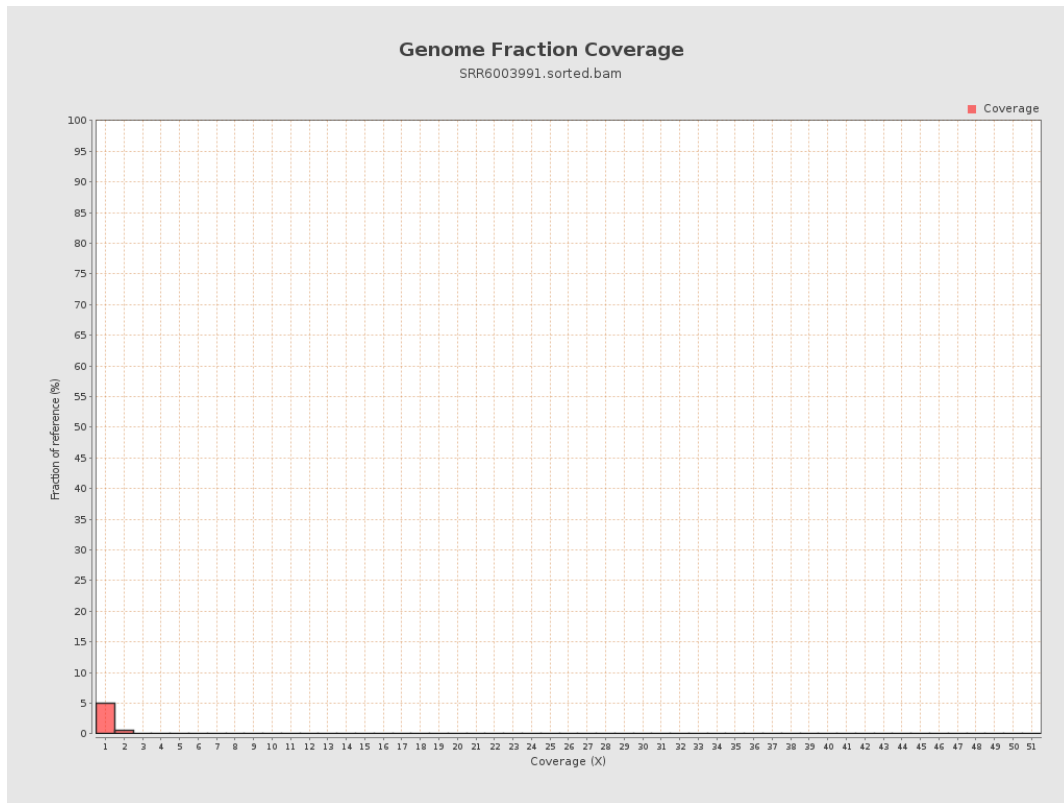
4. Results : Coverage Histogram



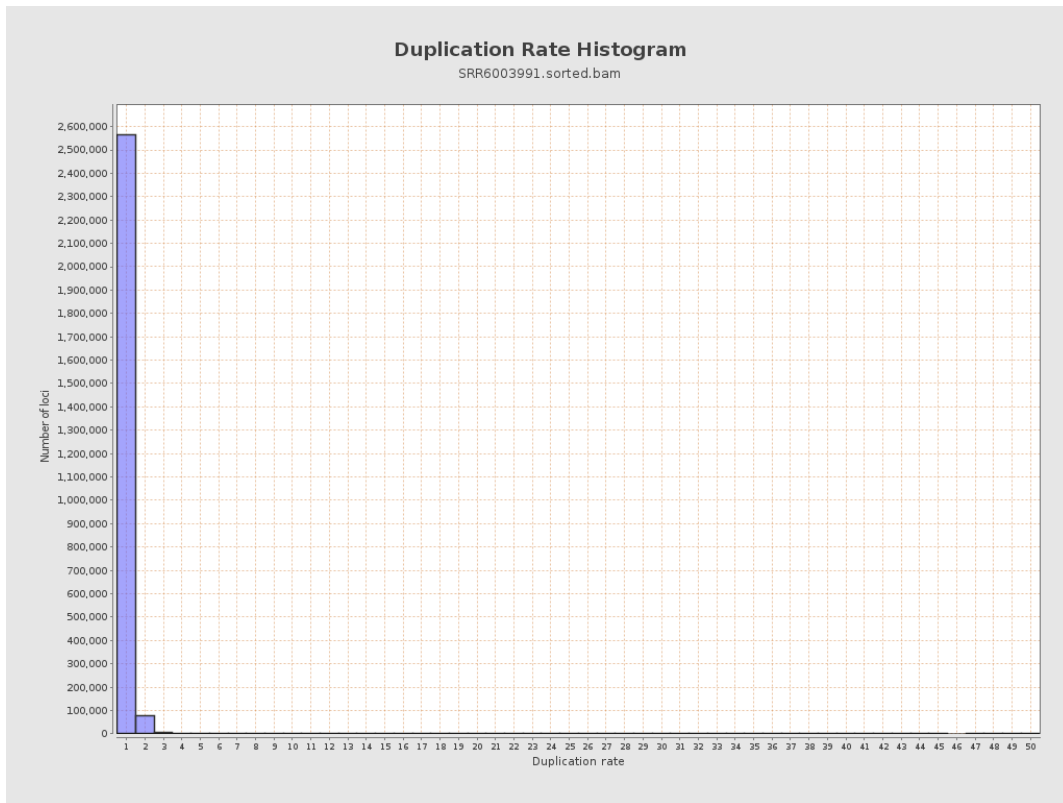
5. Results : Coverage Histogram (0-50X)



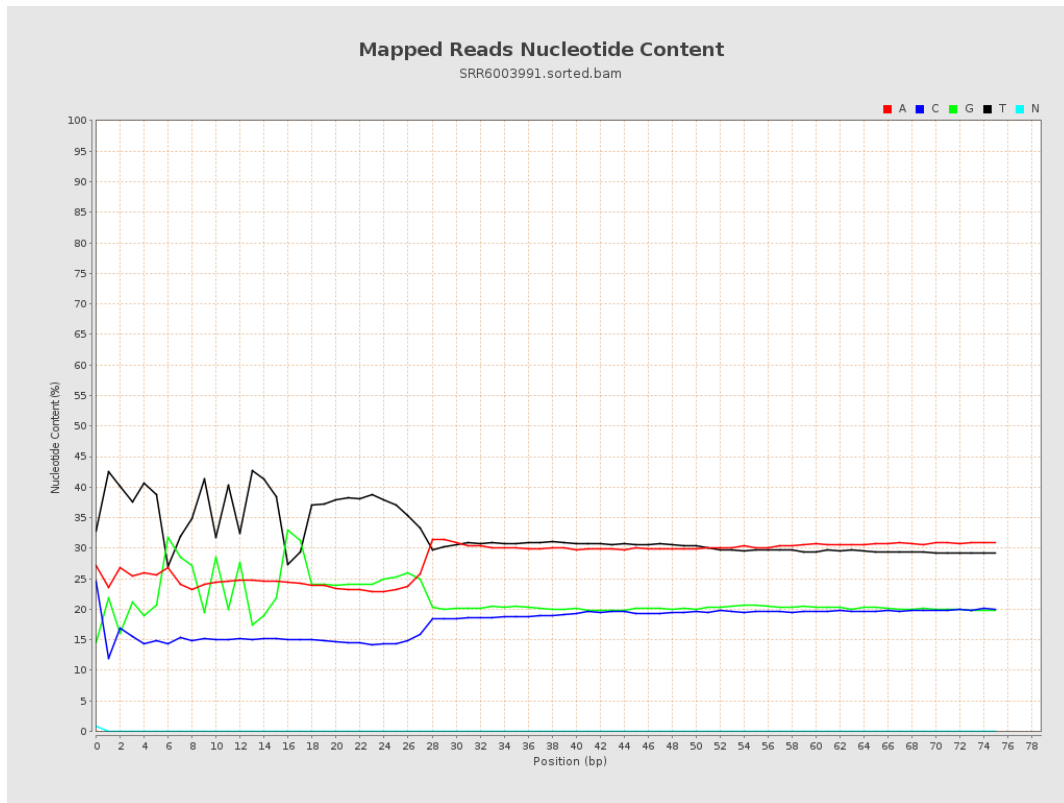
6. Results : Genome Fraction Coverage



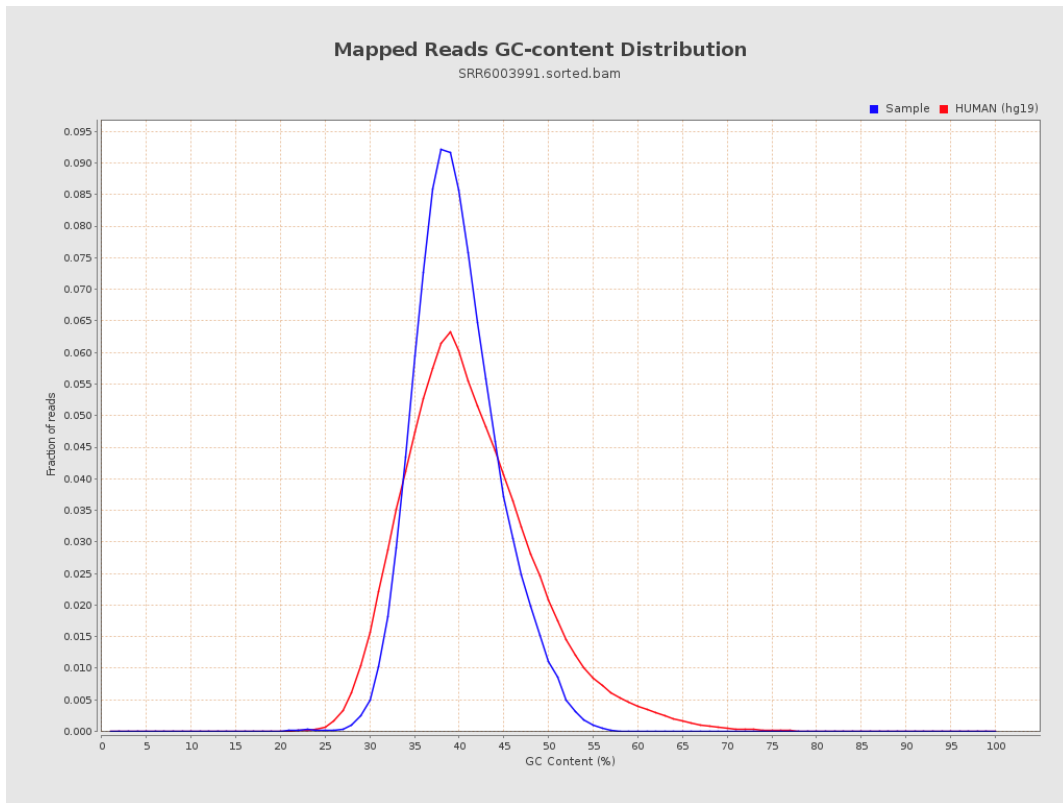
7. Results : Duplication Rate Histogram



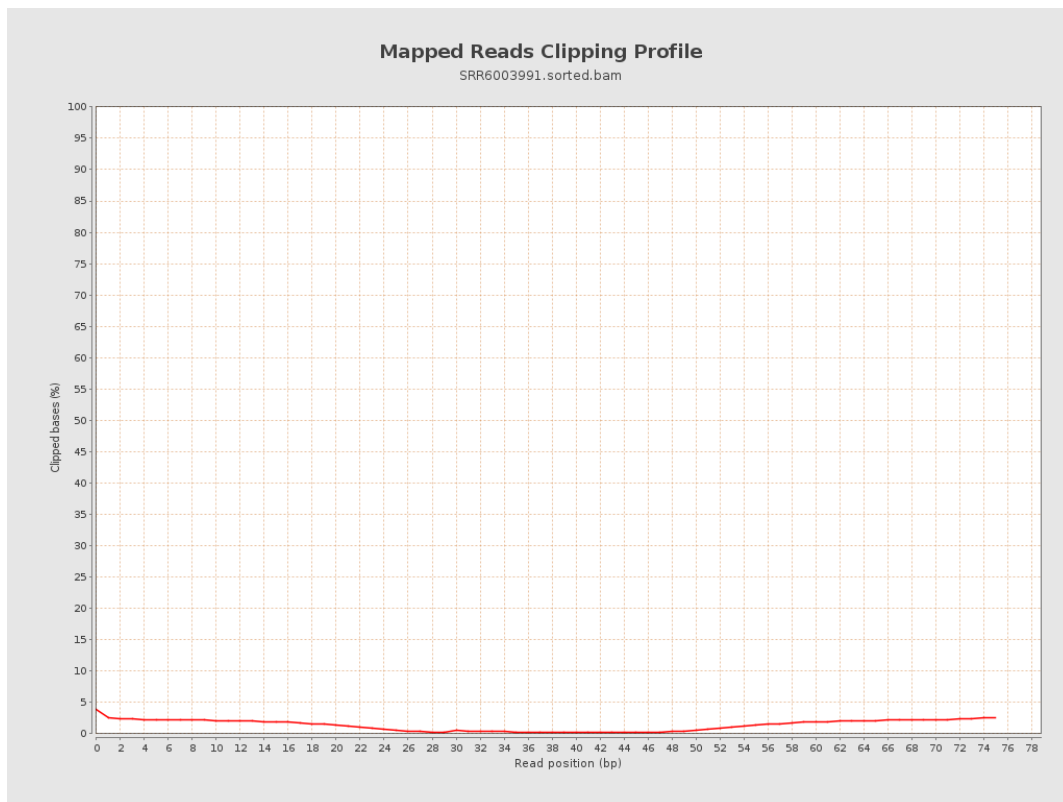
8. Results : Mapped Reads Nucleotide Content



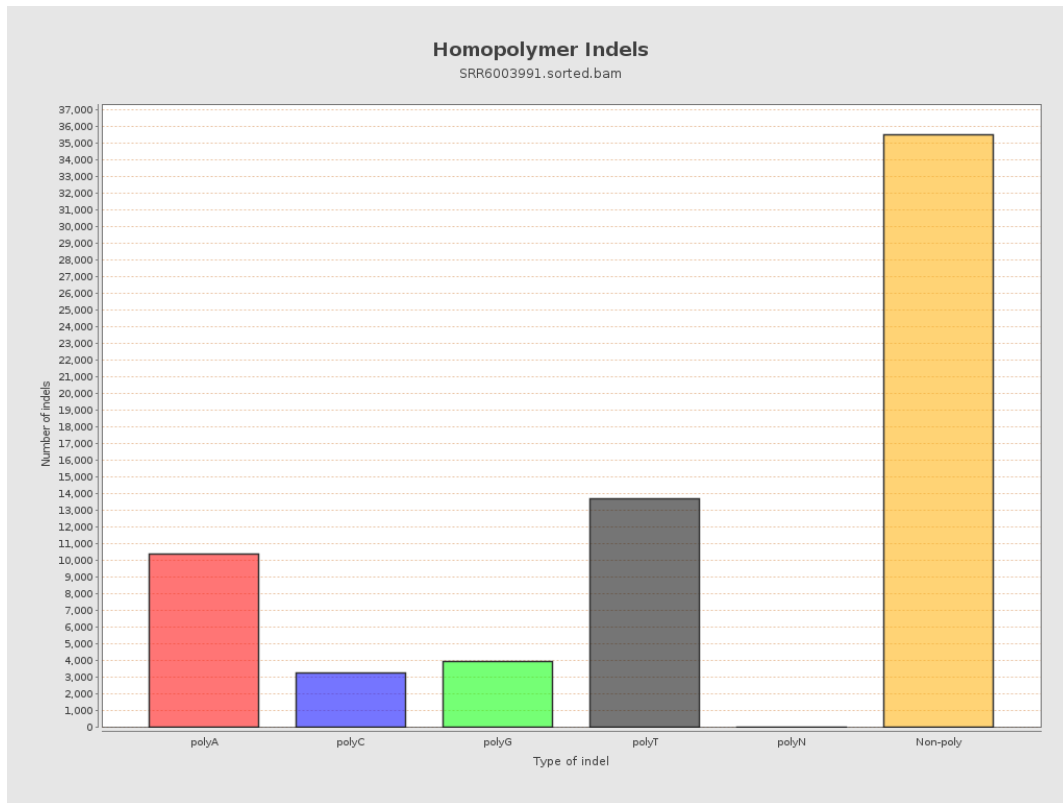
9. Results : Mapped Reads GC-content Distribution



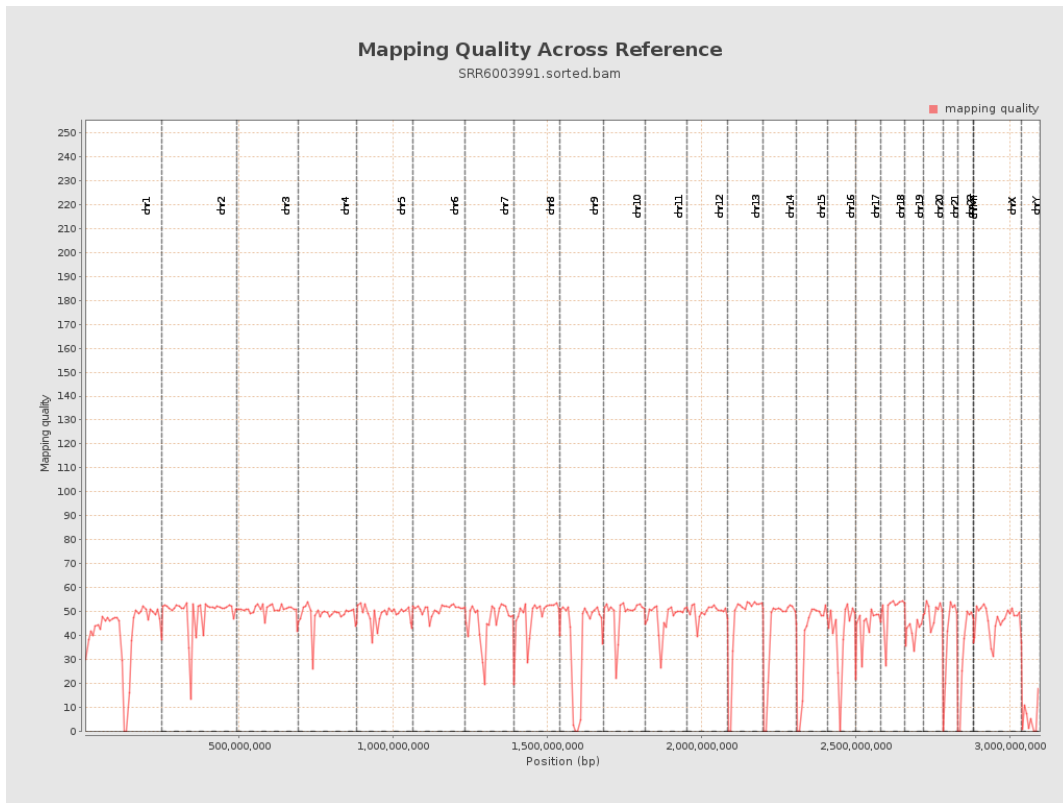
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

