

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/15 09:24:50

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6068089.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_SRR6068089 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6068089.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sun Sep 15 09:24:49 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6068089.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	4,373,237
Mapped reads	4,115,675 / 94.11%
Unmapped reads	257,562 / 5.89%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	26,000 / 0.59%
Read min/max/mean length	30 / 76 / 76.2
Duplicated reads (estimated)	614,429 / 14.05%
Duplication rate	12.63%
Clipped reads	2,139,420 / 48.92%

2.2. ACGT Content

Number/percentage of A's	70,629,833 / 26.61%
Number/percentage of C's	46,898,428 / 17.67%
Number/percentage of T's	87,417,827 / 32.93%
Number/percentage of G's	60,453,355 / 22.78%
Number/percentage of N's	30,159 / 0.01%
GC Percentage	40.44%

2.3. Coverage

Mean	0.0858

Standard Deviation	0.923
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2.4. Mapping Quality

Mean Mapping Quality	46.25
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2.5. Mismatches and indels

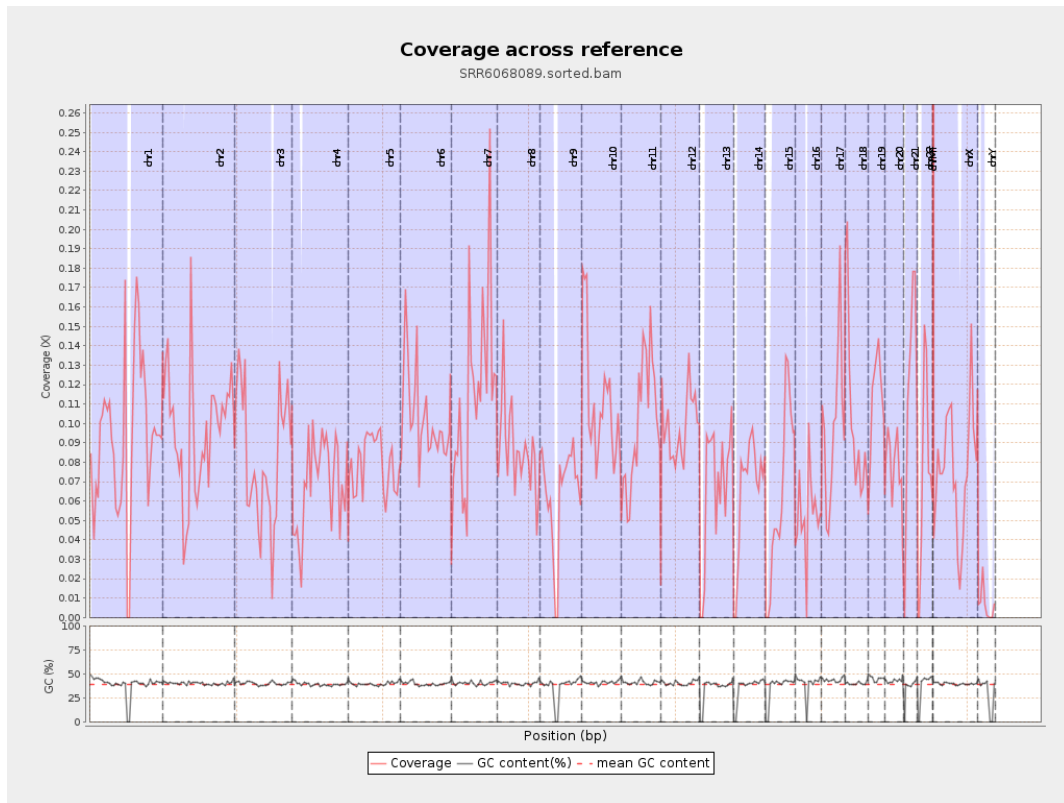
General error rate	0.54%
Mismatches	1,388,663
Insertions	17,154
Mapped reads with at least one insertion	0.41%
Deletions	59,003
Mapped reads with at least one deletion	1.42%
Homopolymer indels	45.81%

2.6. Chromosome stats

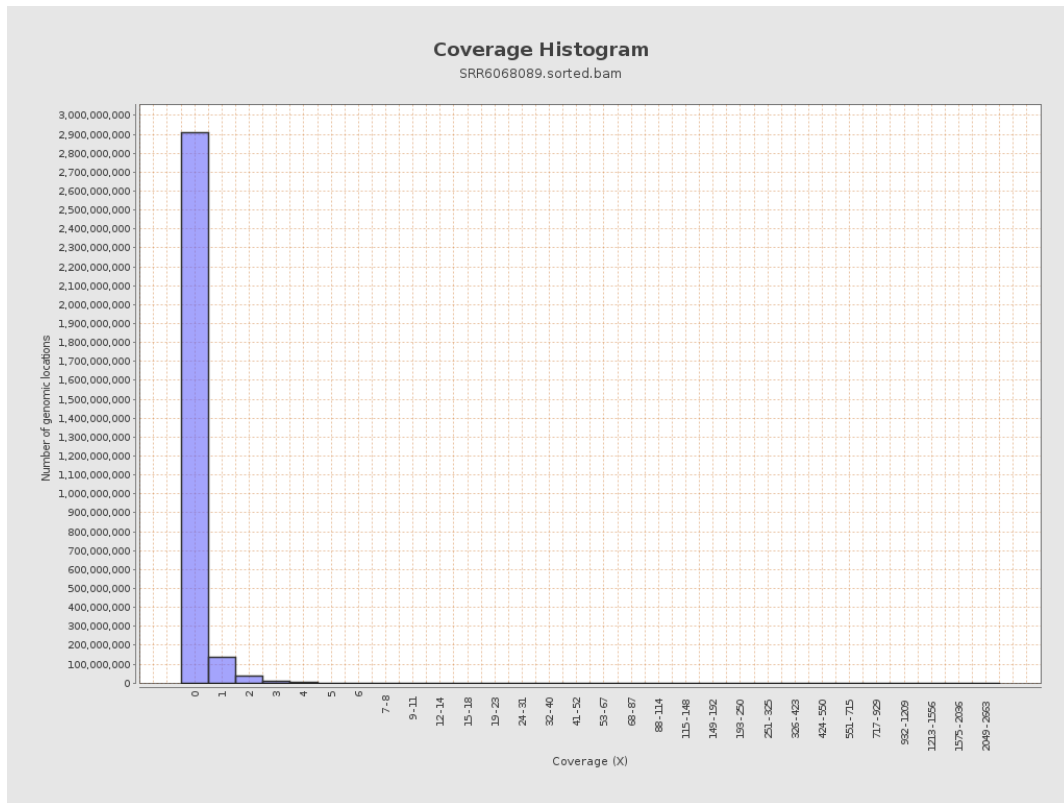
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	22930827	0.092	2.0487
chr2	243199373	23508563	0.0967	1.3622
chr3	198022430	16285317	0.0822	0.3883
chr4	191154276	13347717	0.0698	0.3757
chr5	180915260	14325448	0.0792	0.3821
chr6	171115067	17718132	0.1035	0.6035
chr7	159138663	18156945	0.1141	1.4179

chr8	146364022	12553099	0.0858	0.9734
chr9	141213431	9003181	0.0638	0.5967
chr10	135534747	15088789	0.1113	0.6119
chr11	135006516	13461623	0.0997	0.6826
chr12	133851895	13312724	0.0995	0.4465
chr13	115169878	7763766	0.0674	0.4027
chr14	107349540	7100346	0.0661	0.3785
chr15	102531392	6328418	0.0617	0.3711
chr16	90354753	4829939	0.0535	0.3504
chr17	81195210	8187440	0.1008	0.4732
chr18	78077248	7921837	0.1015	1.1404
chr19	59128983	6705980	0.1134	1.2628
chr20	63025520	4944423	0.0785	0.3886
chr21	48129895	5544307	0.1152	0.4724
chr22	51304566	3801656	0.0741	0.3623
chrMT	16571	266445	16.079	10.0665
chrX	155270560	12033013	0.0775	0.4725
chrY	59373566	411984	0.0069	0.2311

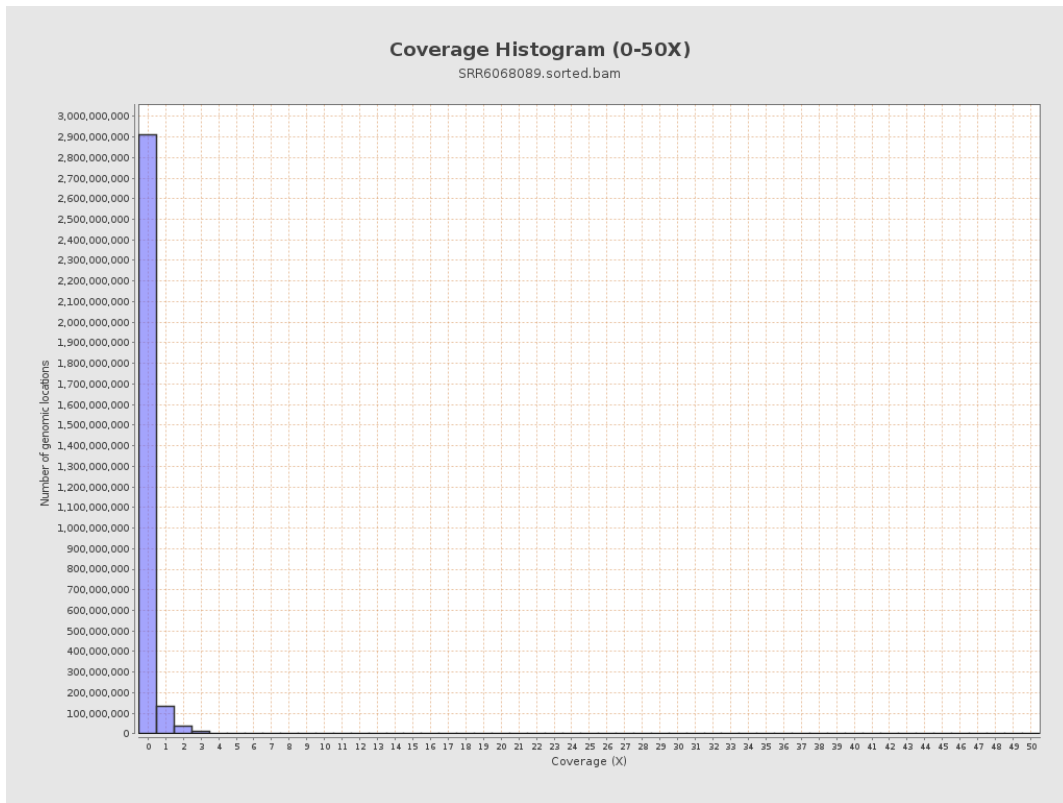
3. Results : Coverage across reference



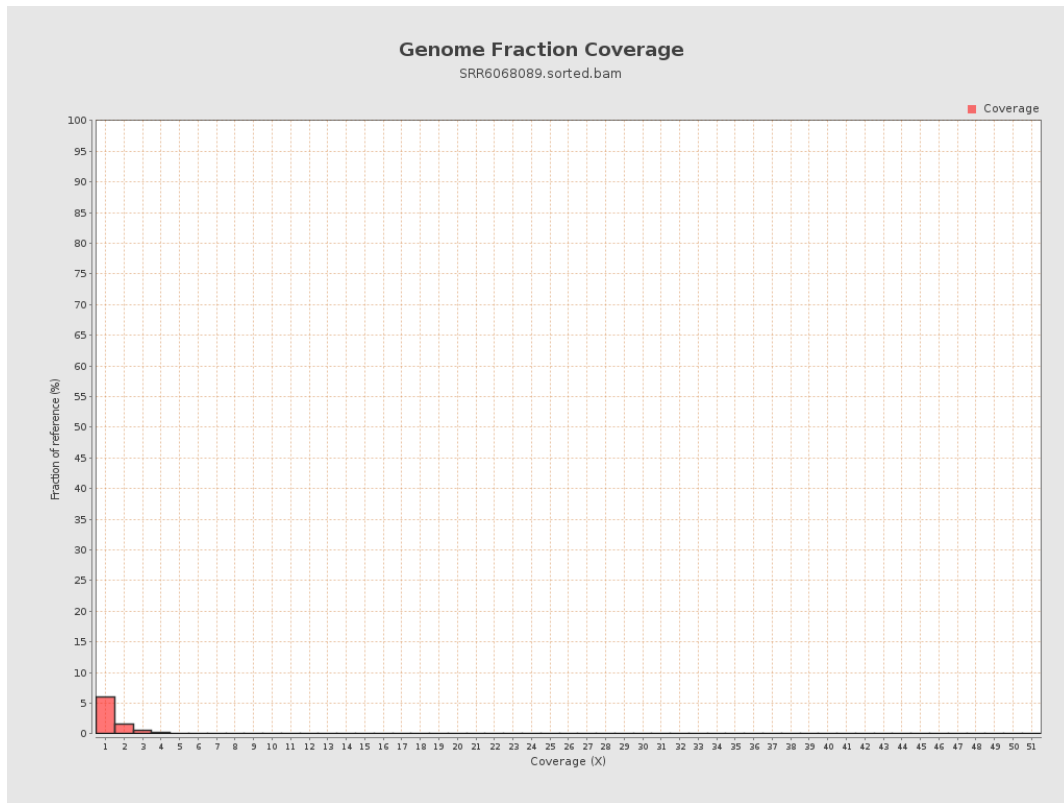
4. Results : Coverage Histogram



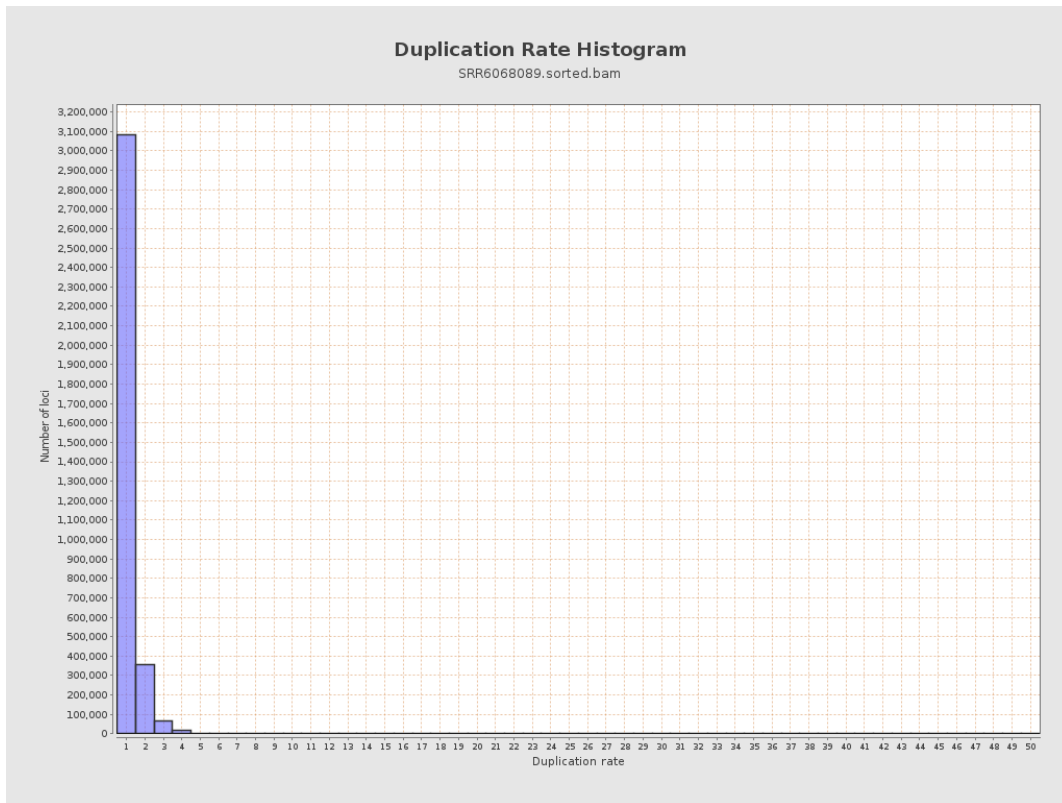
5. Results : Coverage Histogram (0-50X)



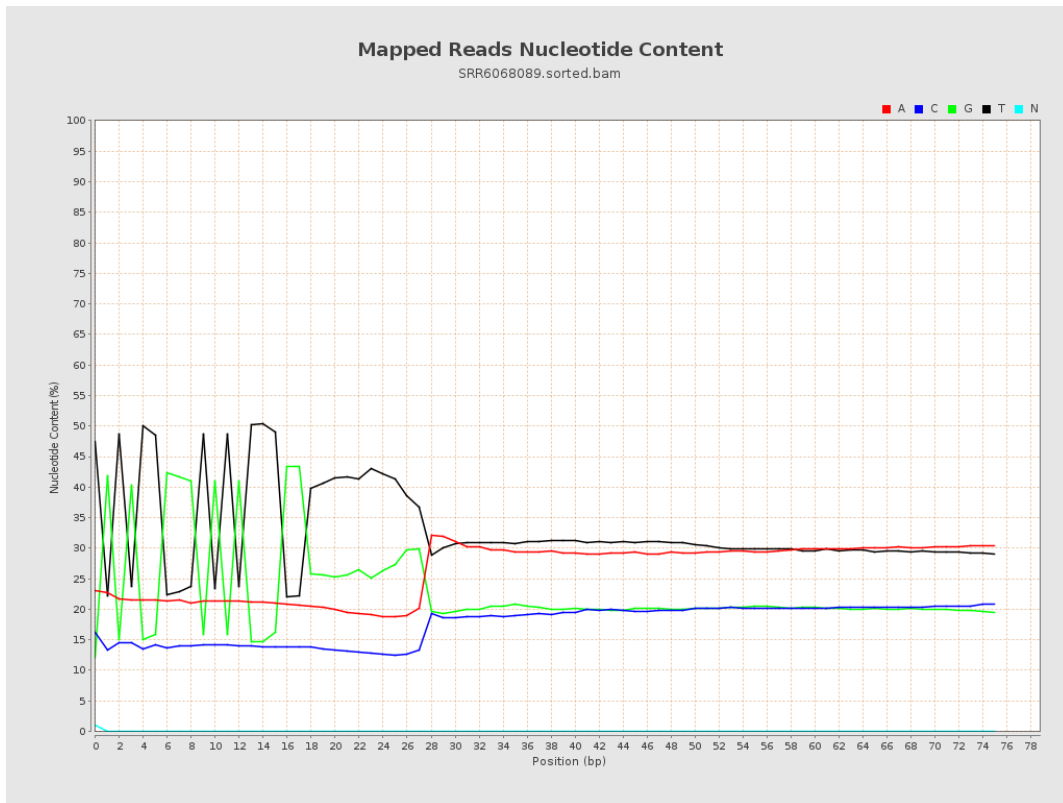
6. Results : Genome Fraction Coverage



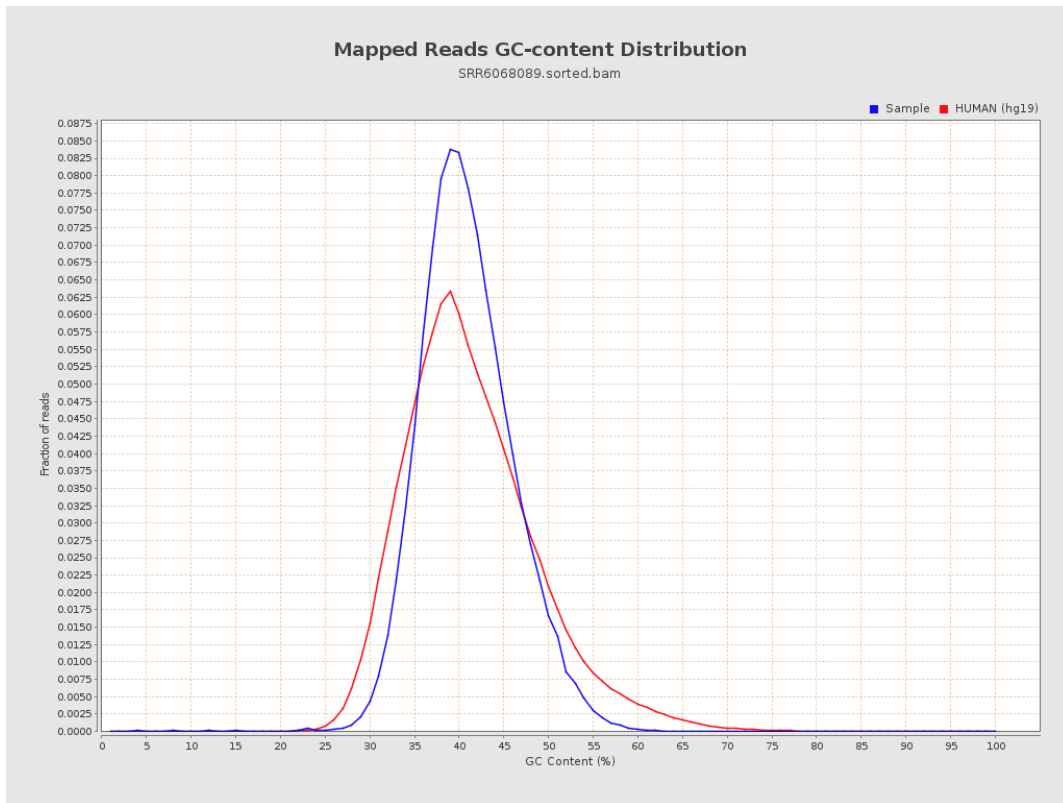
7. Results : Duplication Rate Histogram



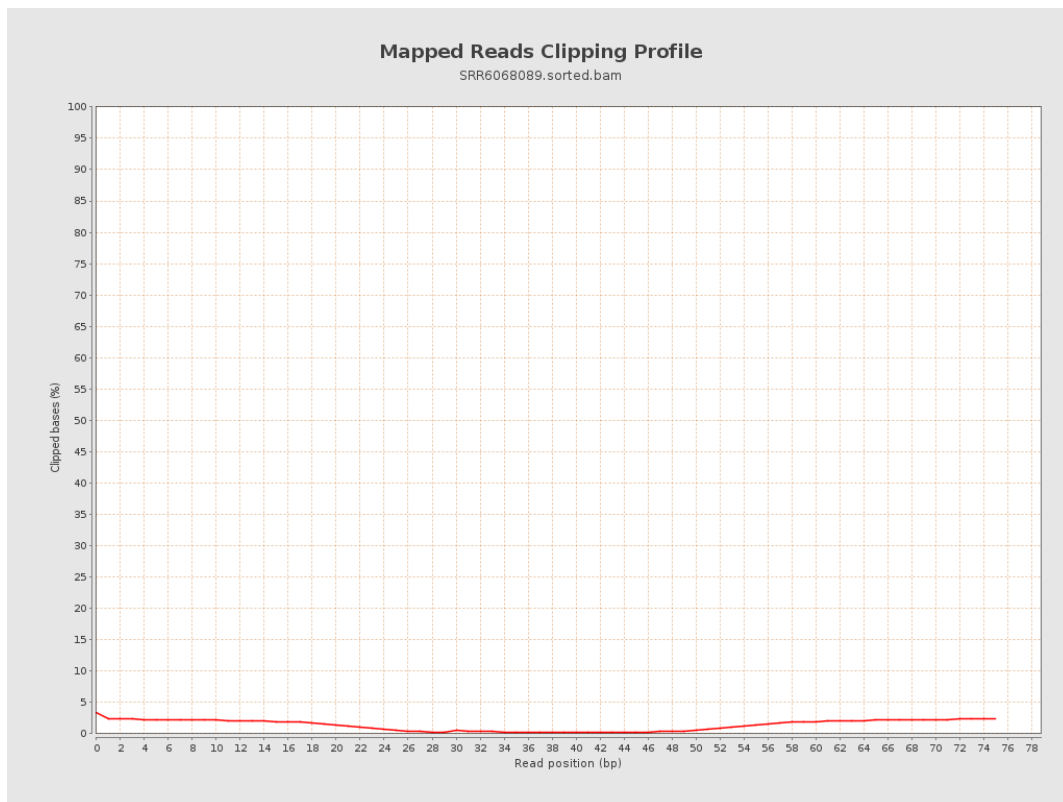
8. Results : Mapped Reads Nucleotide Content



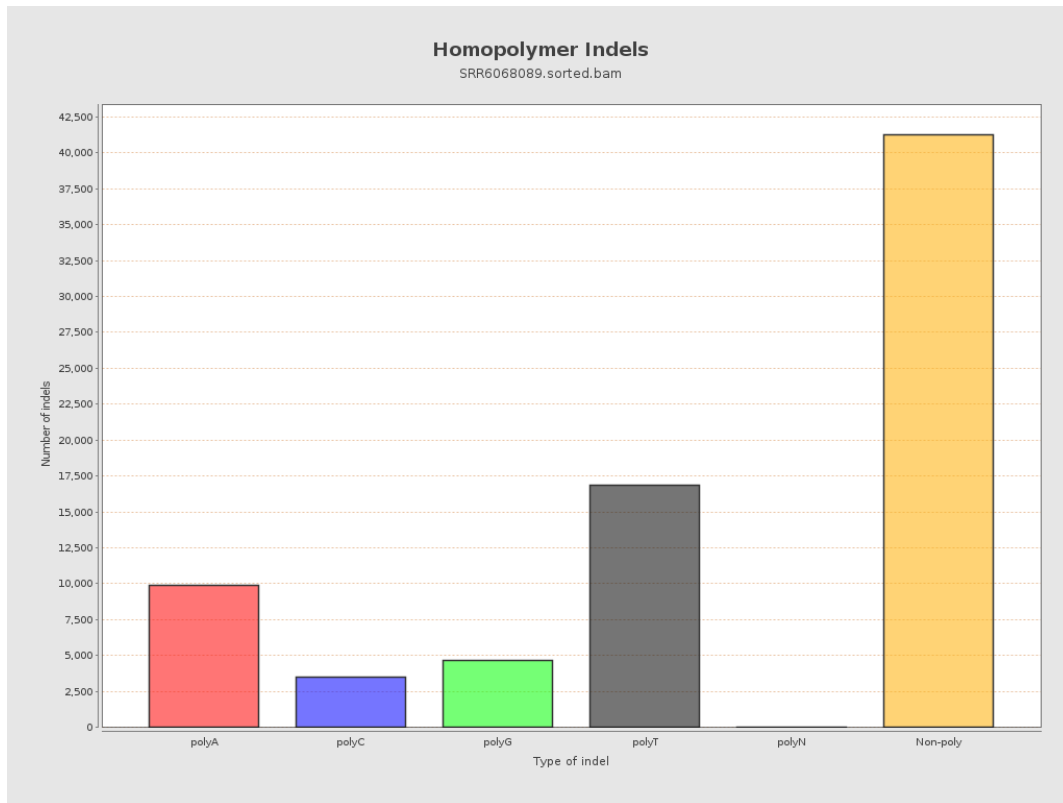
9. Results : Mapped Reads GC-content Distribution



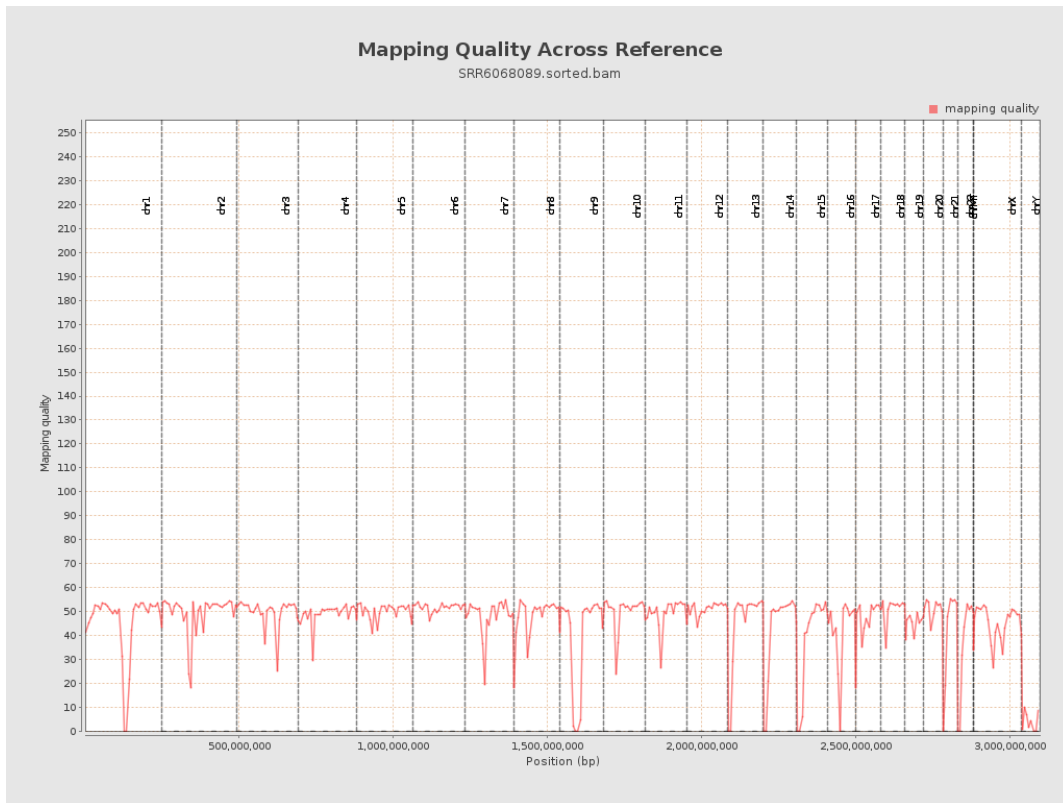
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

