

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/09/14 02:29:13

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR6004113.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_SRR6004113 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR6004113.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sat Sep 14 02:29:12 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR6004113.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	4,037,972
Mapped reads	3,579,835 / 88.65%
Unmapped reads	458,137 / 11.35%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	21,991 / 0.54%
Read min/max/mean length	30 / 76 / 76.19
Duplicated reads (estimated)	211,705 / 5.24%
Duplication rate	4.62%
Clipped reads	1,393,242 / 34.5%

2.2. ACGT Content

Number/percentage of A's	67,579,081 / 27.67%
Number/percentage of C's	46,111,348 / 18.88%
Number/percentage of T's	77,219,178 / 31.62%
Number/percentage of G's	53,257,954 / 21.81%
Number/percentage of N's	33,490 / 0.01%
GC Percentage	40.69%

2.3. Coverage

Mean	0.0789

Standard Deviation	0.6361
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2.4. Mapping Quality

Mean Mapping Quality	46.04
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2.5. Mismatches and indels

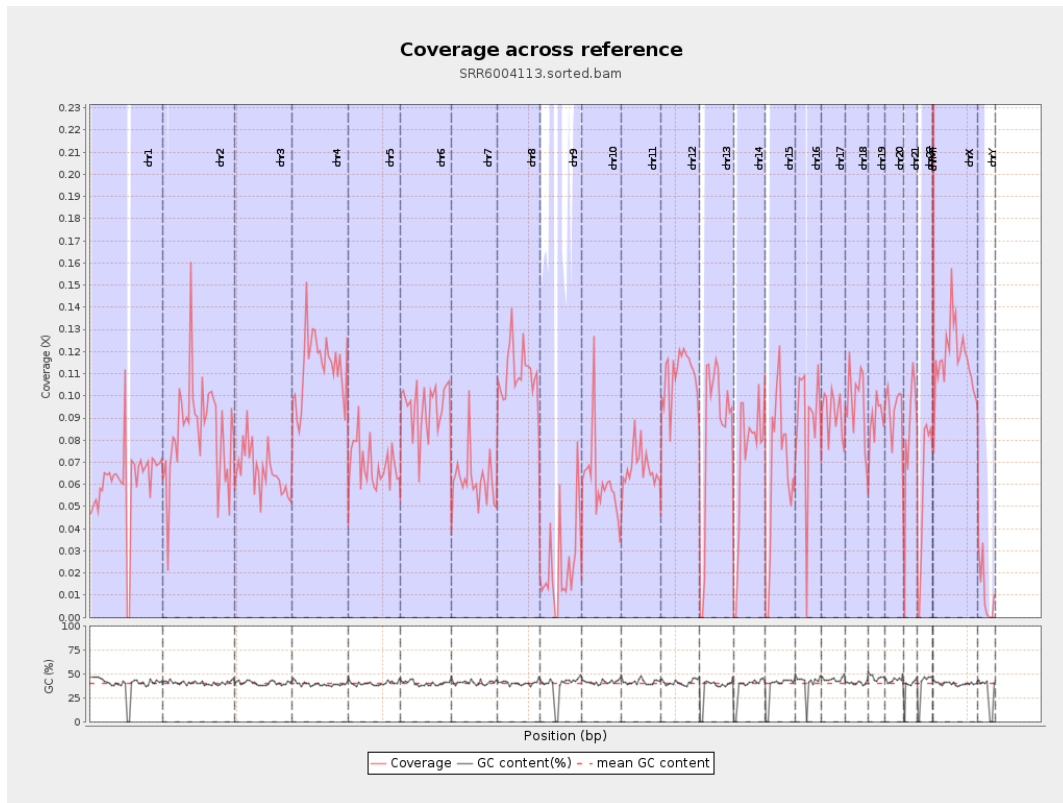
General error rate	0.95%
Mismatches	2,276,749
Insertions	19,773
Mapped reads with at least one insertion	0.55%
Deletions	63,829
Mapped reads with at least one deletion	1.76%
Homopolymer indels	46.37%

2.6. Chromosome stats

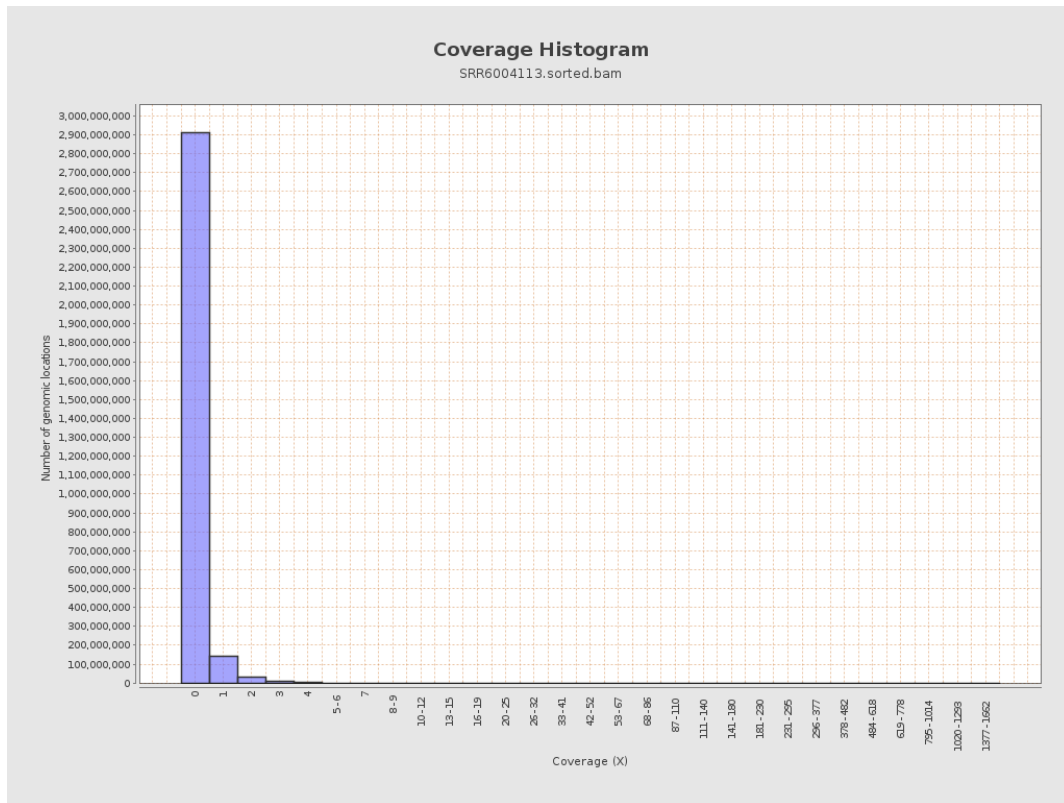
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	15161795	0.0608	1.1229
chr2	243199373	20382403	0.0838	0.7602
chr3	198022430	13058758	0.0659	0.3295
chr4	191154276	21718998	0.1136	0.4628
chr5	180915260	12432666	0.0687	0.3329
chr6	171115067	16159896	0.0944	0.47
chr7	159138663	9824867	0.0617	0.6359

chr8	146364022	16038294	0.1096	1.0949
chr9	141213431	3272656	0.0232	0.4644
chr10	135534747	8296365	0.0612	0.6245
chr11	135006516	9130450	0.0676	0.5042
chr12	133851895	14668783	0.1096	0.4222
chr13	115169878	9710973	0.0843	0.3652
chr14	107349540	7672363	0.0715	0.3639
chr15	102531392	6695418	0.0653	0.3239
chr16	90354753	7920798	0.0877	0.4893
chr17	81195210	7516638	0.0926	0.4343
chr18	78077248	7603940	0.0974	1.2327
chr19	59128983	5284339	0.0894	0.7581
chr20	63025520	5912676	0.0938	0.413
chr21	48129895	4026386	0.0837	0.4012
chr22	51304566	2997919	0.0584	0.3054
chrMT	16571	32550	1.9643	1.9516
chrX	155270560	18097073	0.1166	0.4918
chrY	59373566	691329	0.0116	0.2413

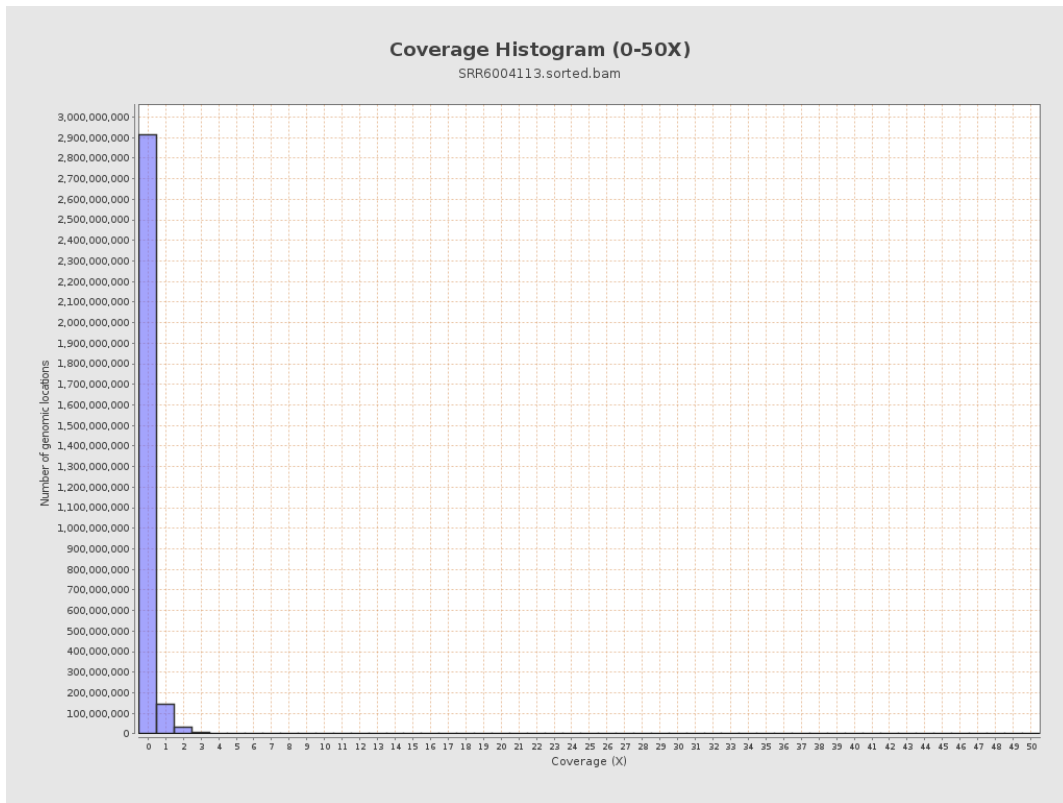
3. Results : Coverage across reference



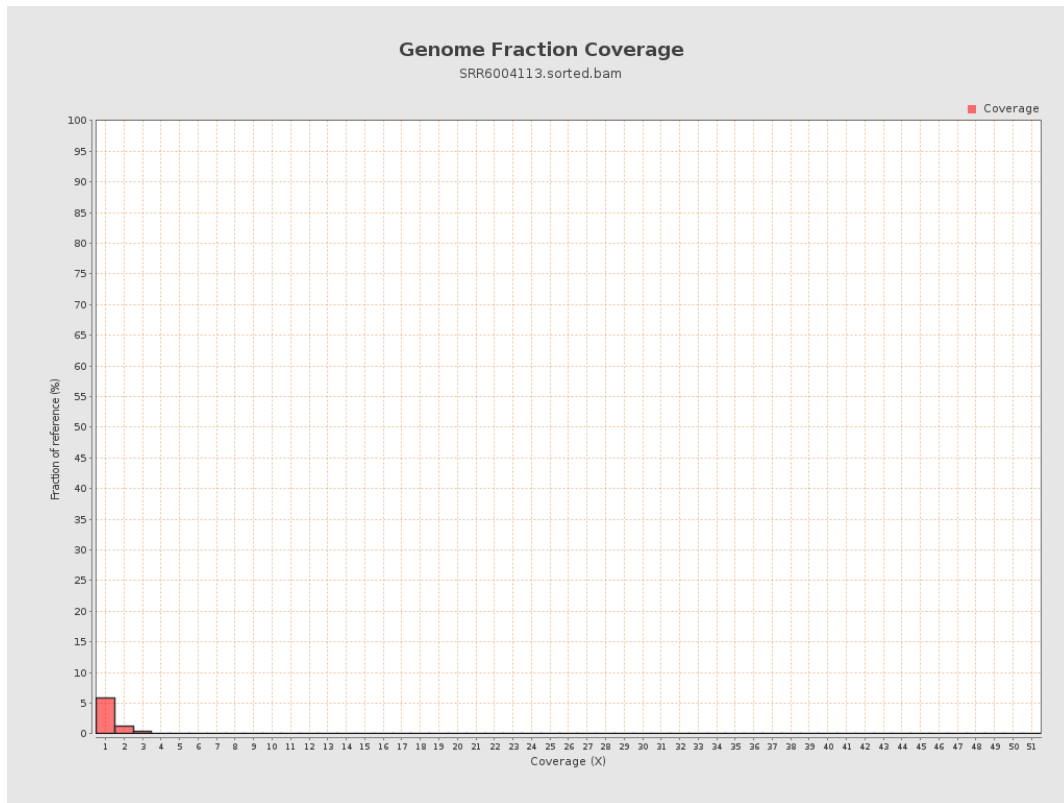
4. Results : Coverage Histogram



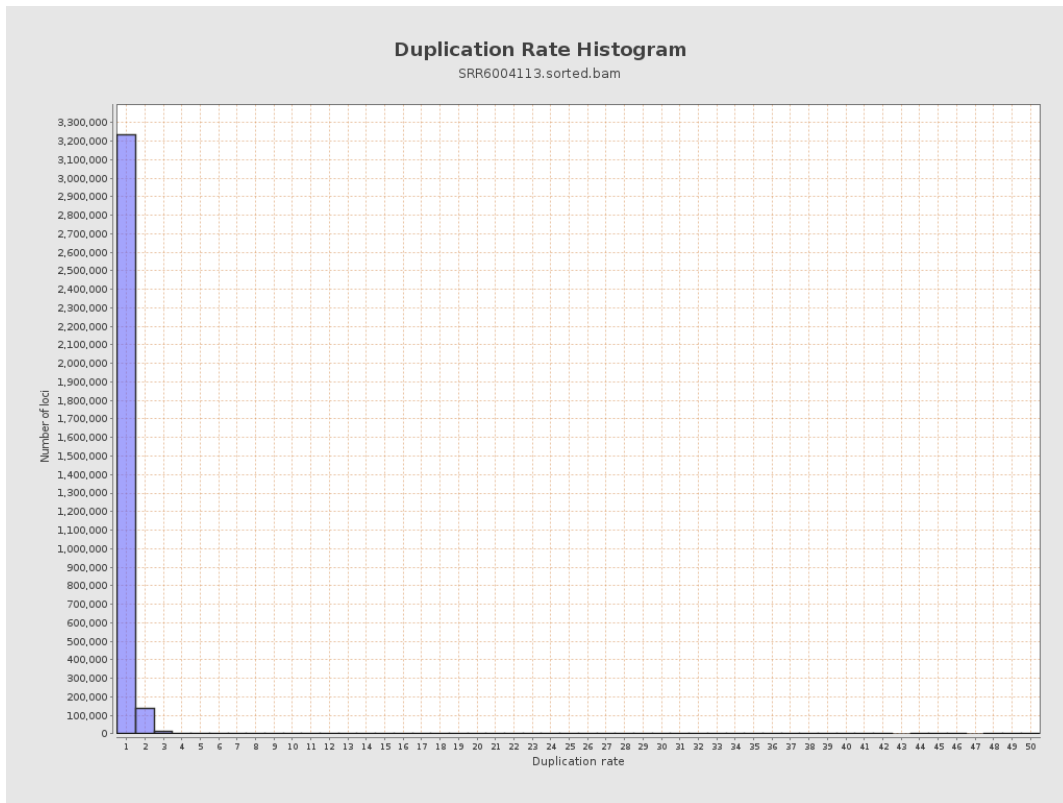
5. Results : Coverage Histogram (0-50X)



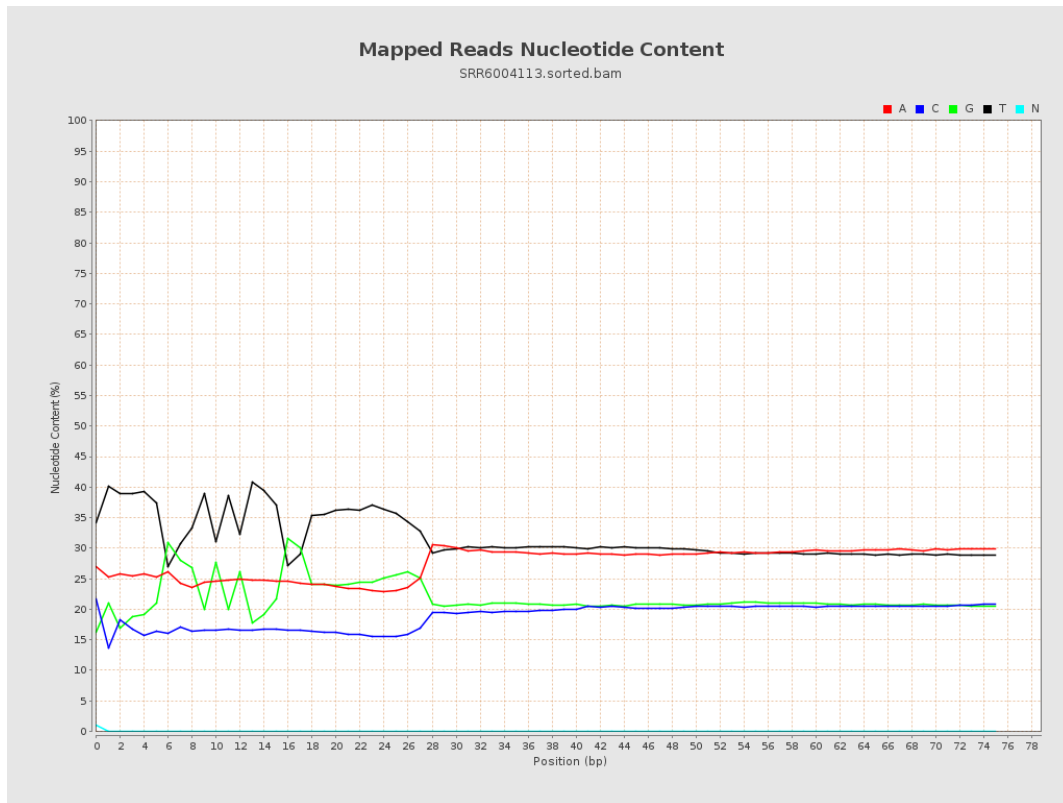
6. Results : Genome Fraction Coverage



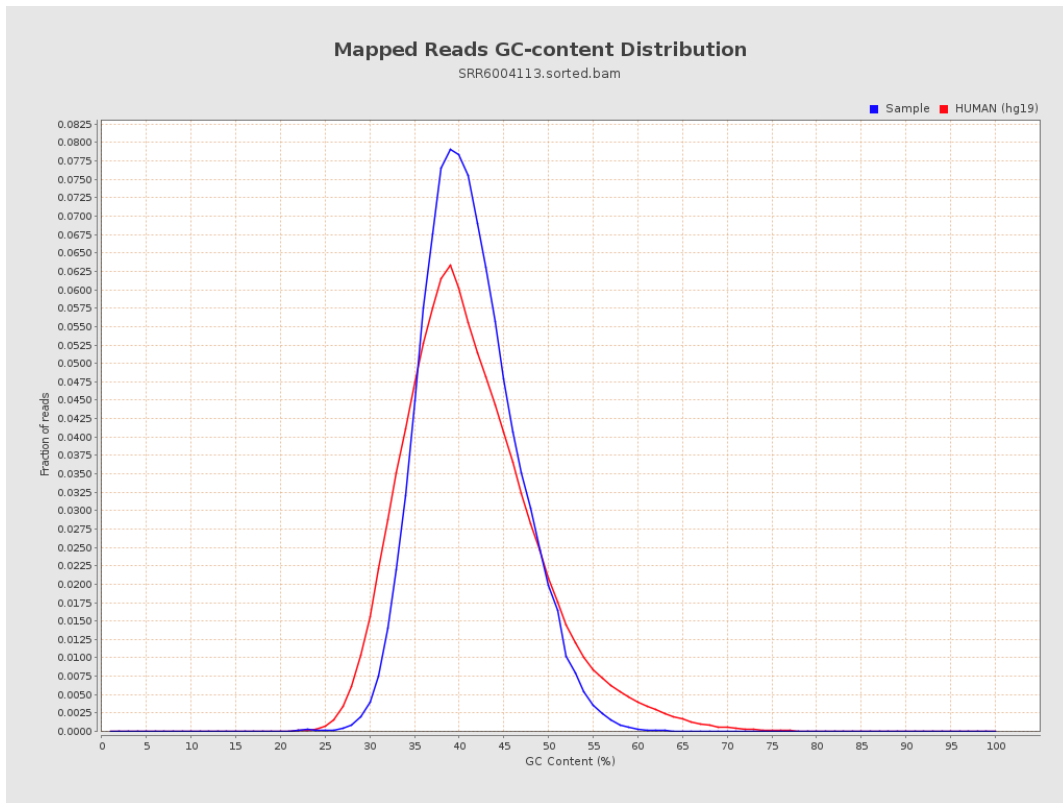
7. Results : Duplication Rate Histogram



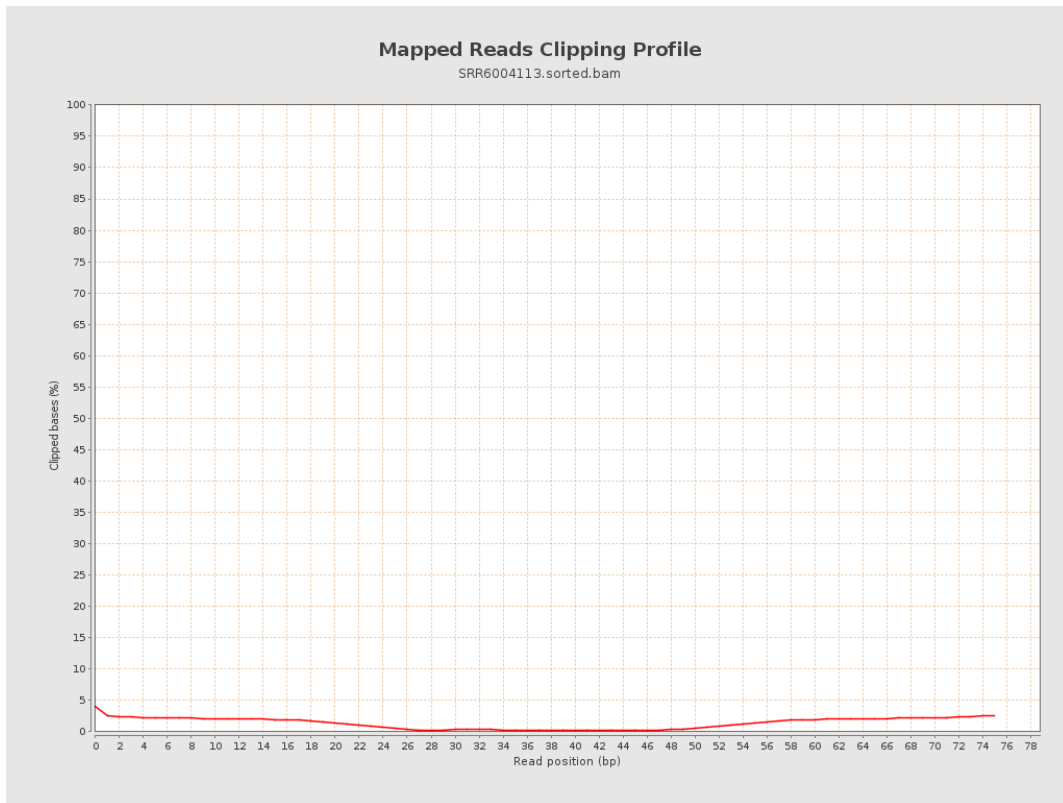
8. Results : Mapped Reads Nucleotide Content



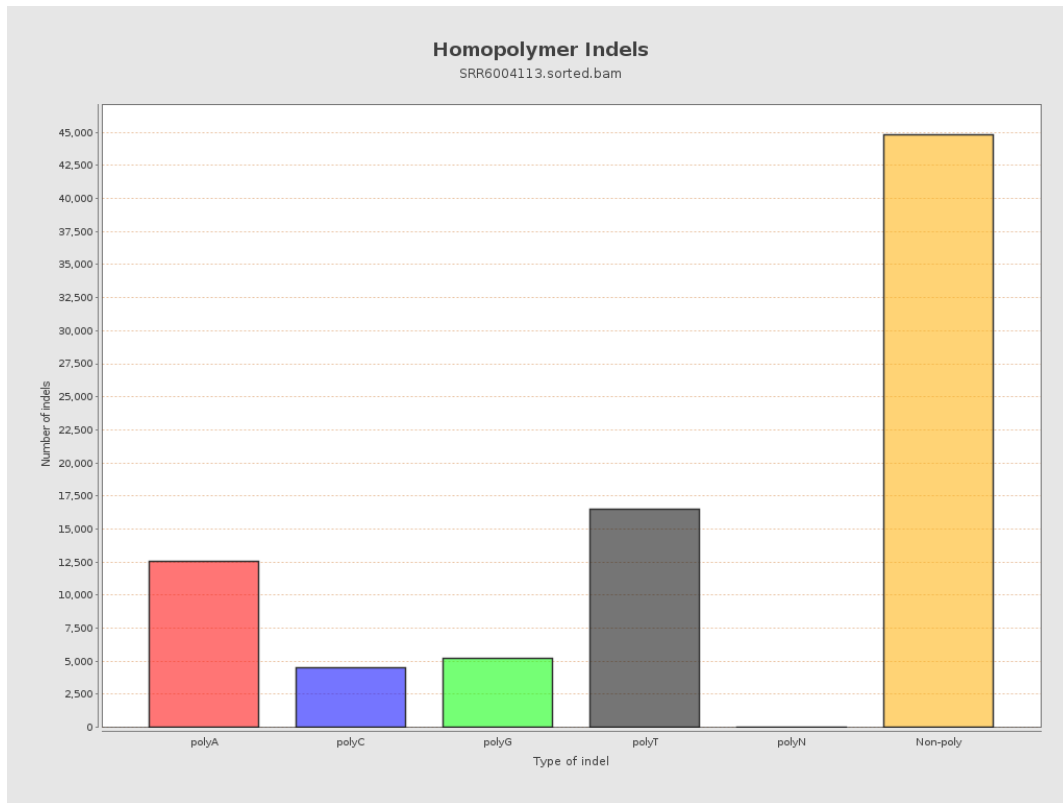
9. Results : Mapped Reads GC-content Distribution



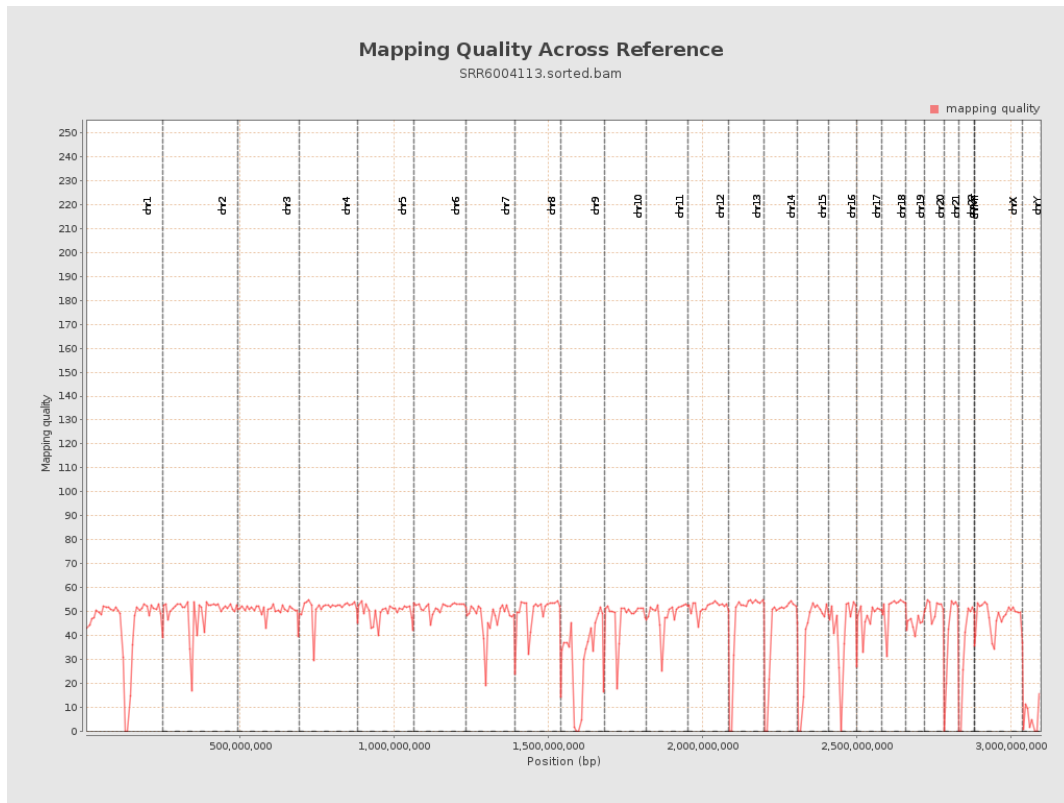
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

