

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2022/04/19 04:11:08

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR054567.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_SRR054567 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR054567.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Tue Apr 19 04:11:08 CST 2022
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR054567.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	14,459,313
Mapped reads	10,779,631 / 74.55%
Unmapped reads	3,679,682 / 25.45%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	306 / 0%
Read min/max/mean length	30 / 48 / 48
Duplicated reads (estimated)	4,175,370 / 28.88%
Duplication rate	25.81%
Clipped reads	1,017,731 / 7.04%

2.2. ACGT Content

Number/percentage of A's	158,846,405 / 31.24%
Number/percentage of C's	94,443,481 / 18.57%
Number/percentage of T's	157,819,809 / 31.04%
Number/percentage of G's	97,310,552 / 19.14%
Number/percentage of N's	80,759 / 0.02%
GC Percentage	37.71%

2.3. Coverage

Mean	0.1643

Standard Deviation	1.5951
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2.4. Mapping Quality

Mean Mapping Quality	45.86
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2.5. Mismatches and indels

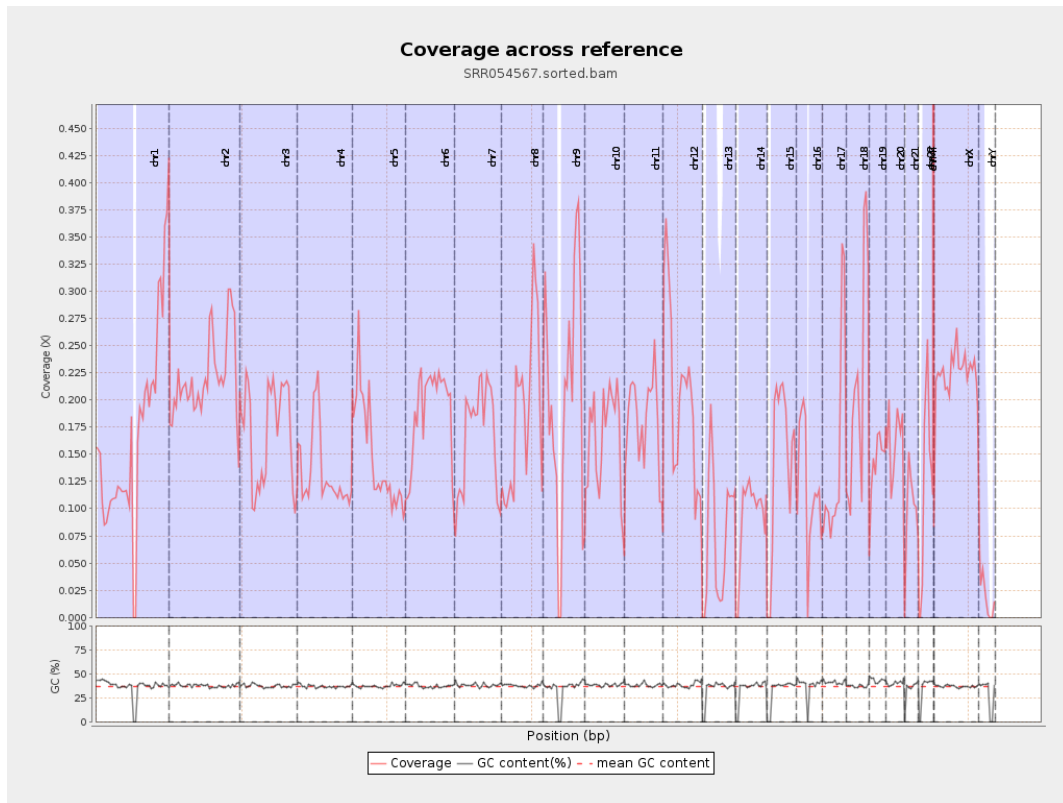
General error rate	0.55%
Mismatches	2,772,038
Insertions	24,409
Mapped reads with at least one insertion	0.23%
Deletions	80,968
Mapped reads with at least one deletion	0.75%
Homopolymer indels	49.05%

2.6. Chromosome stats

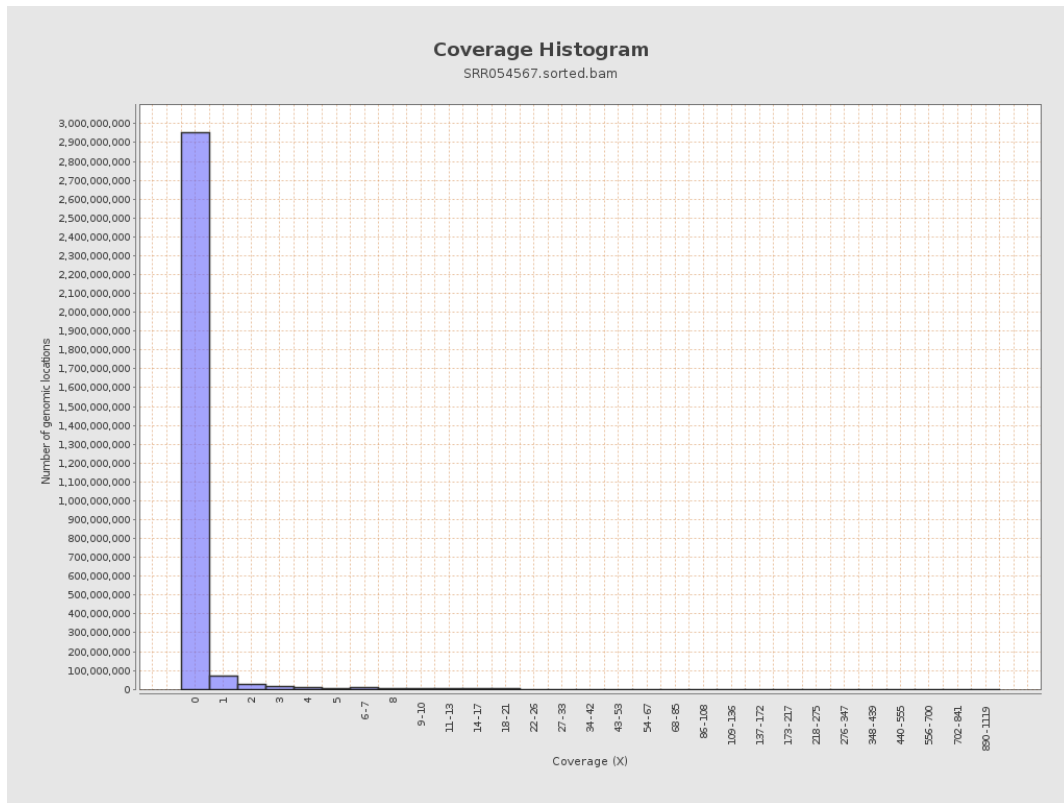
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	41789469	0.1677	1.6598
chr2	243199373	53781353	0.2211	1.9655
chr3	198022430	33798735	0.1707	1.5615
chr4	191154276	25486533	0.1333	1.4036
chr5	180915260	26607217	0.1471	1.4246
chr6	171115067	32148005	0.1879	1.7596
chr7	159138663	26699535	0.1678	1.6843

chr8	146364022	27857285	0.1903	1.756
chr9	141213431	29044561	0.2057	1.7735
chr10	135534747	22697340	0.1675	1.5577
chr11	135006516	24129161	0.1787	1.6433
chr12	133851895	27161628	0.2029	1.7782
chr13	115169878	8327466	0.0723	0.9917
chr14	107349540	9773770	0.091	1.272
chr15	102531392	14782408	0.1442	1.3964
chr16	90354753	10910977	0.1208	1.2439
chr17	81195210	12040161	0.1483	1.4245
chr18	78077248	16424835	0.2104	1.9104
chr19	59128983	8586552	0.1452	1.4331
chr20	63025520	10063616	0.1597	1.473
chr21	48129895	4780889	0.0993	1.1337
chr22	51304566	6052100	0.118	1.2246
chrMT	16571	40337	2.4342	6.4815
chrX	155270560	34482068	0.2221	1.8652
chrY	59373566	1155074	0.0195	0.4121

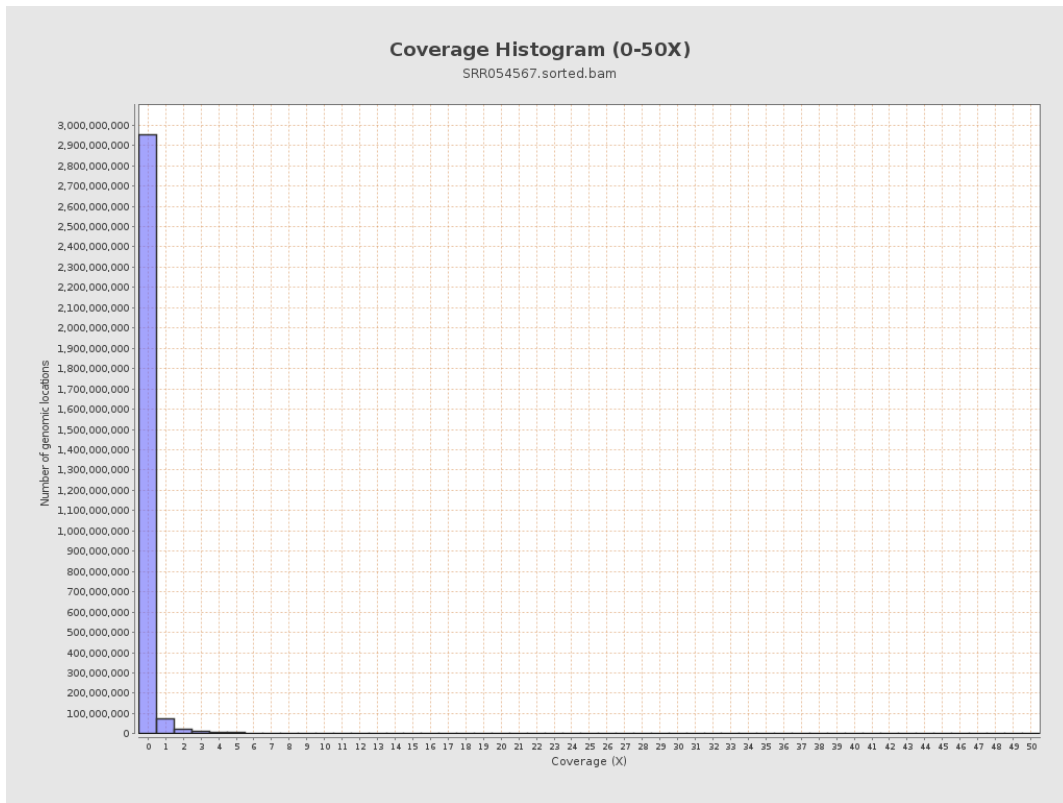
3. Results : Coverage across reference



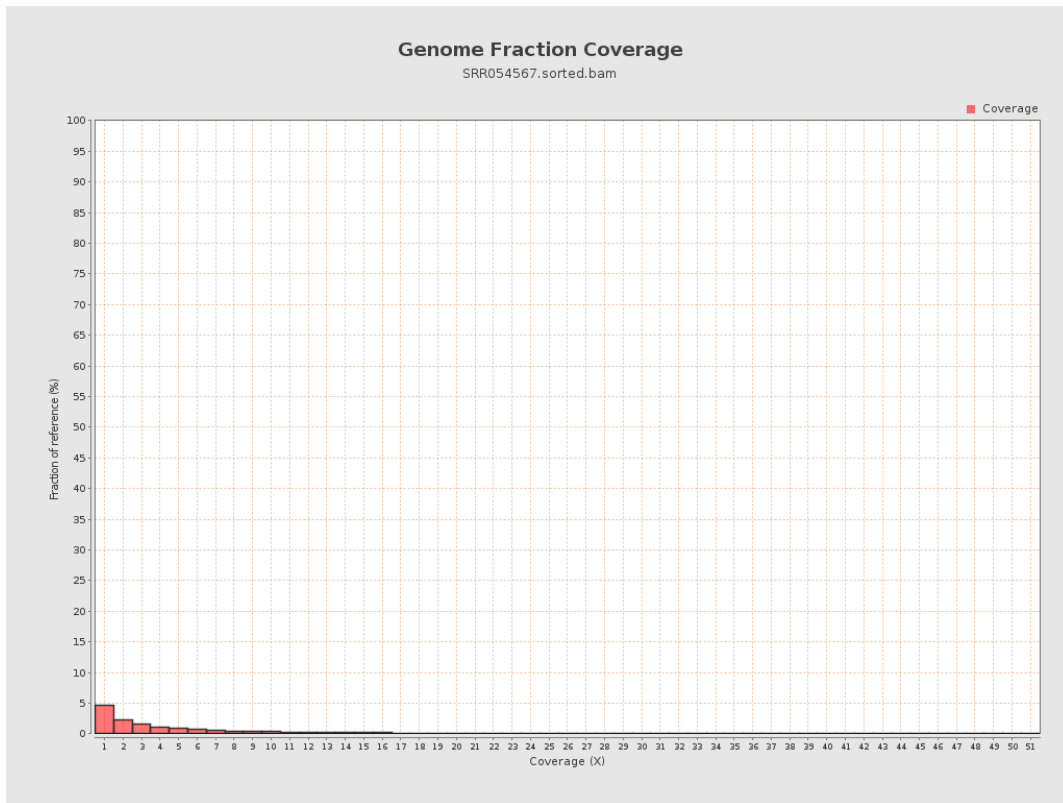
4. Results : Coverage Histogram



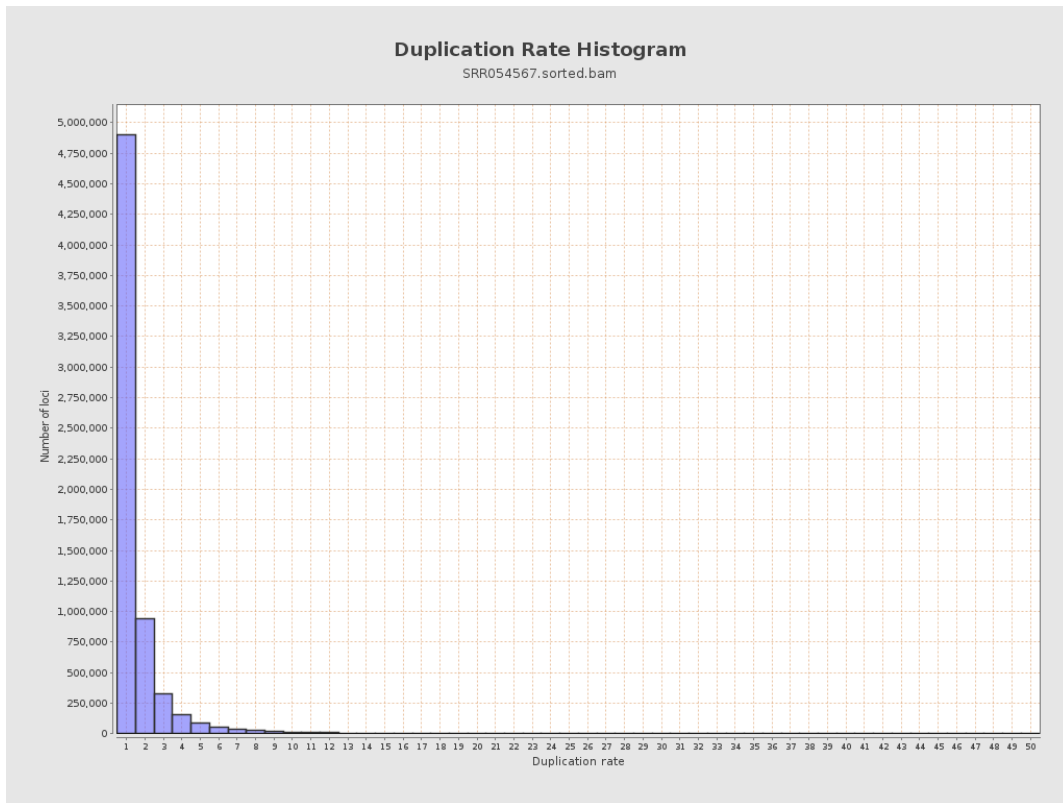
5. Results : Coverage Histogram (0-50X)



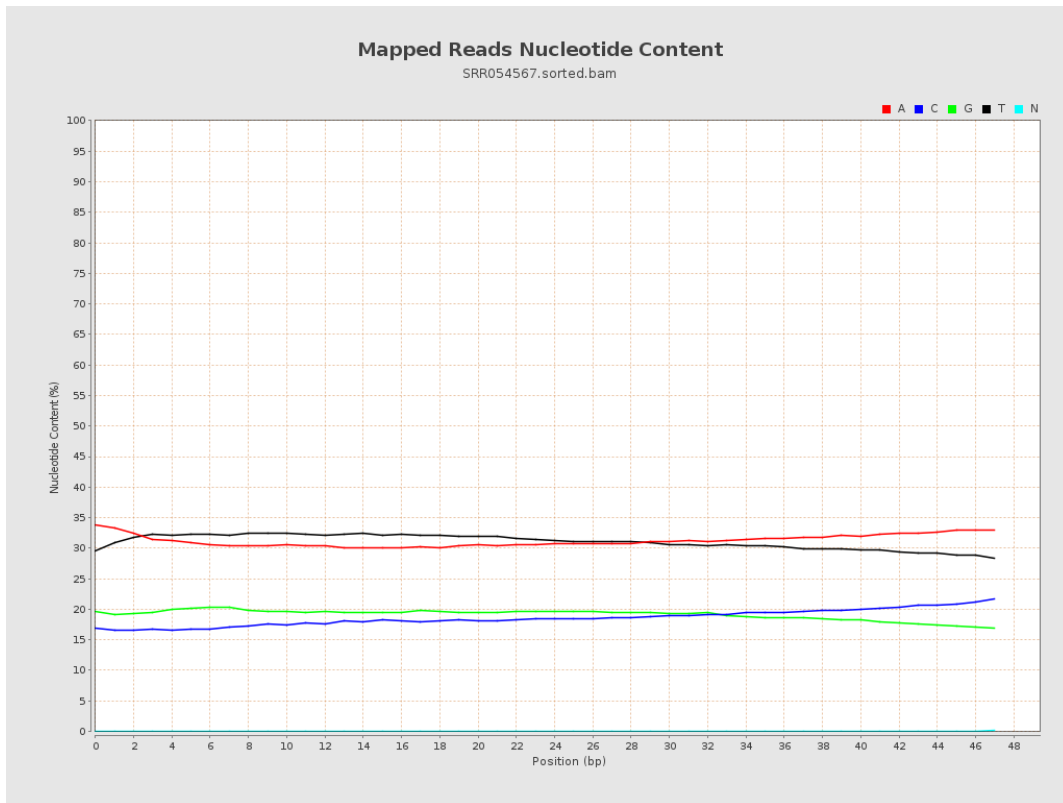
6. Results : Genome Fraction Coverage



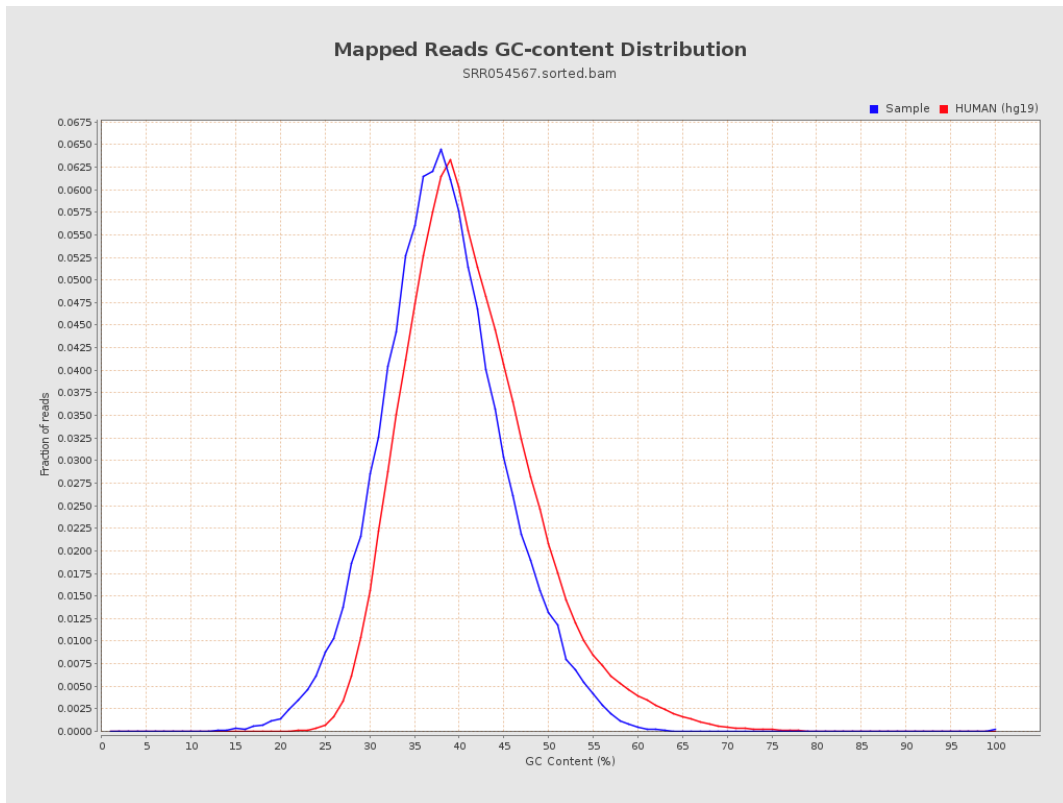
7. Results : Duplication Rate Histogram



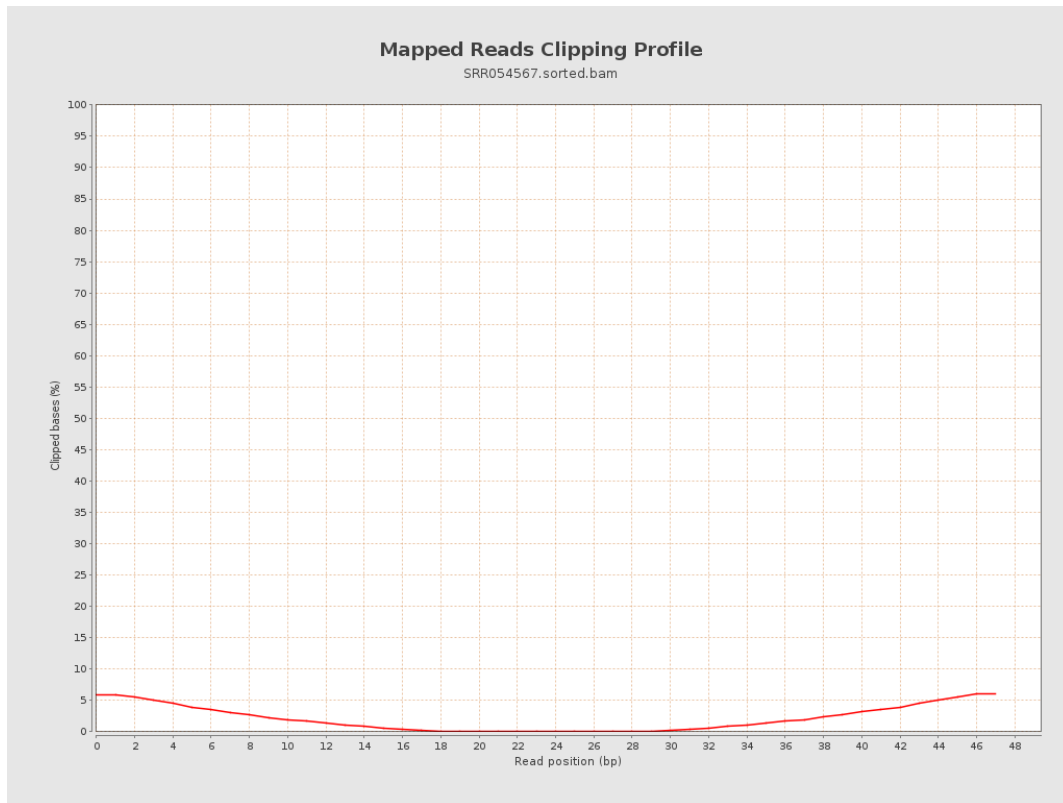
8. Results : Mapped Reads Nucleotide Content



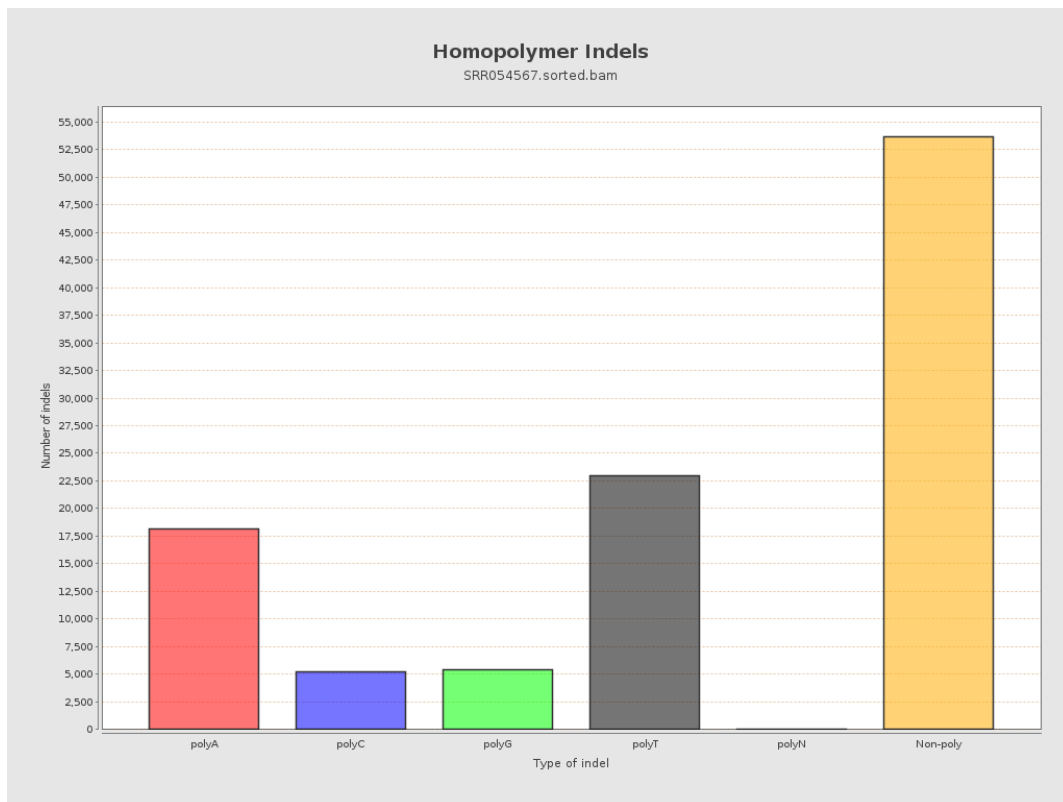
9. Results : Mapped Reads GC-content Distribution



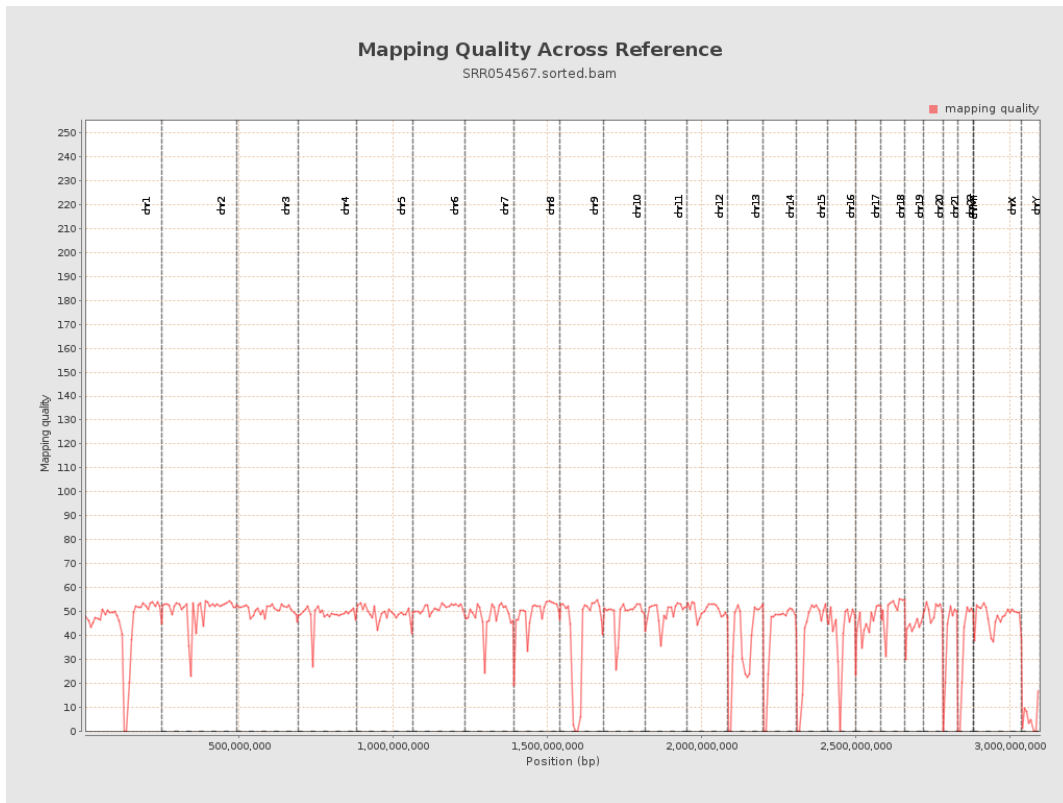
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

