

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/24 00:37:42

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR1549096.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_SRR1549096 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR1549096.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Sat Aug 24 00:37:41 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR1549096.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	7,586,928
Mapped reads	6,662,102 / 87.81%
Unmapped reads	924,826 / 12.19%
Mapped paired reads	0 / 0%
Secondary alignments	0
Read min/max/mean length	40 / 40 / 40
Duplicated reads (estimated)	215,320 / 2.84%
Duplication rate	2.21%
Clipped reads	311,831 / 4.11%

2.2. ACGT Content

Number/percentage of A's	78,392,371 / 29.62%
Number/percentage of C's	54,050,829 / 20.43%
Number/percentage of T's	79,468,534 / 30.03%
Number/percentage of G's	52,654,173 / 19.9%
Number/percentage of N's	64,060 / 0.02%
GC Percentage	40.32%

2.3. Coverage

Mean	0.0855
Standard Deviation	0.645

2.4. Mapping Quality

Mean Mapping Quality	44.25
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2.5. Mismatches and indels

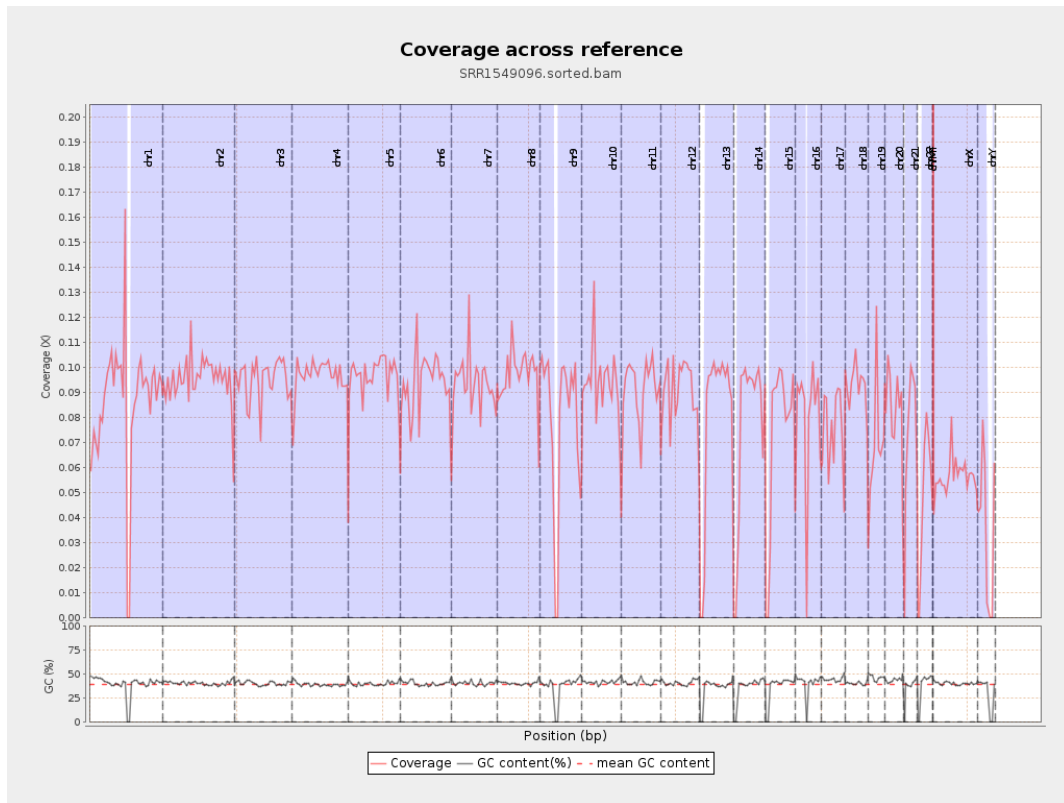
General error rate	0.39%
Mismatches	1,037,969
Insertions	5,807
Mapped reads with at least one insertion	0.09%
Deletions	19,239
Mapped reads with at least one deletion	0.29%
Homopolymer indels	45.78%

2.6. Chromosome stats

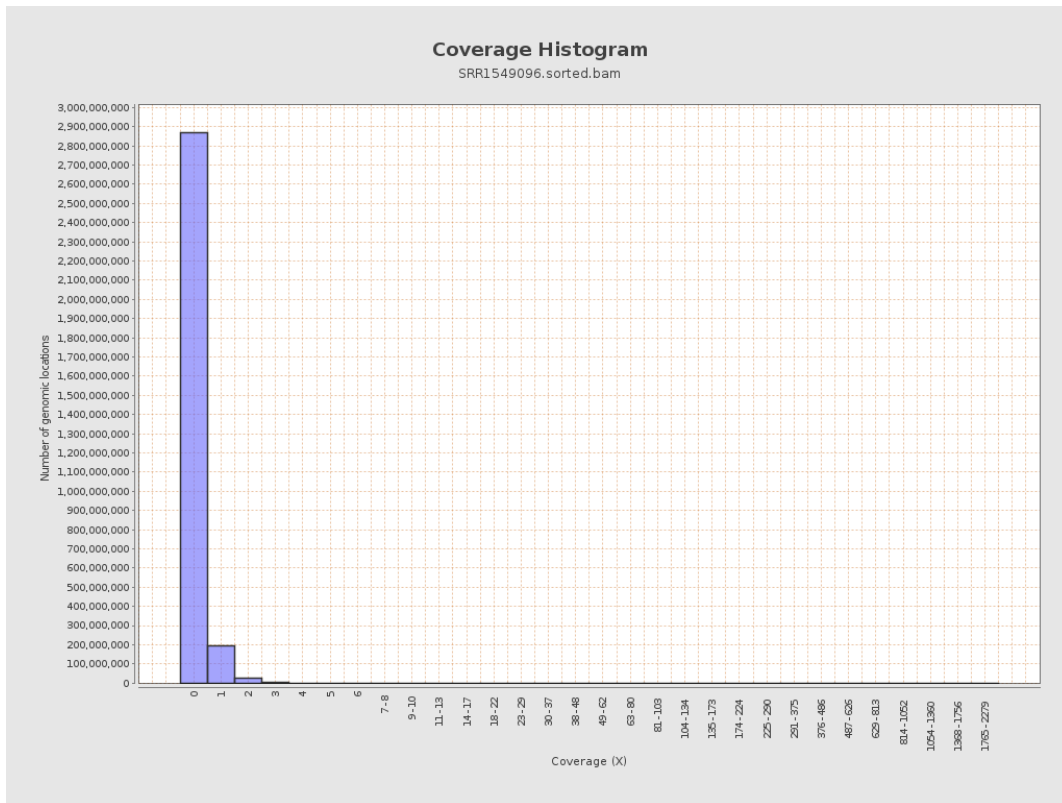
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	21480428	0.0862	1.5716
chr2	243199373	23196648	0.0954	0.4958
chr3	198022430	18832143	0.0951	0.3495
chr4	191154276	18506348	0.0968	0.3608
chr5	180915260	17384349	0.0961	0.3576
chr6	171115067	16143645	0.0943	0.4402
chr7	159138663	14757139	0.0927	0.6729
chr8	146364022	14182477	0.0969	0.6633

chr9	141213431	11171060	0.0791	0.4772
chr10	135534747	12755098	0.0941	0.564
chr11	135006516	12316427	0.0912	0.4964
chr12	133851895	12160443	0.0908	0.3557
chr13	115169878	9167040	0.0796	0.3123
chr14	107349540	8287008	0.0772	0.3621
chr15	102531392	7432771	0.0725	0.3002
chr16	90354753	7024877	0.0777	0.3823
chr17	81195210	6154507	0.0758	0.3661
chr18	78077248	7267418	0.0931	0.9228
chr19	59128983	4180868	0.0707	1.1519
chr20	63025520	5351330	0.0849	0.3405
chr21	48129895	3480352	0.0723	0.3481
chr22	51304566	2491929	0.0486	0.2833
chrMT	16571	14291	0.8624	1.4623
chrX	155270560	8748059	0.0563	0.3326
chrY	59373566	2167513	0.0365	0.3307

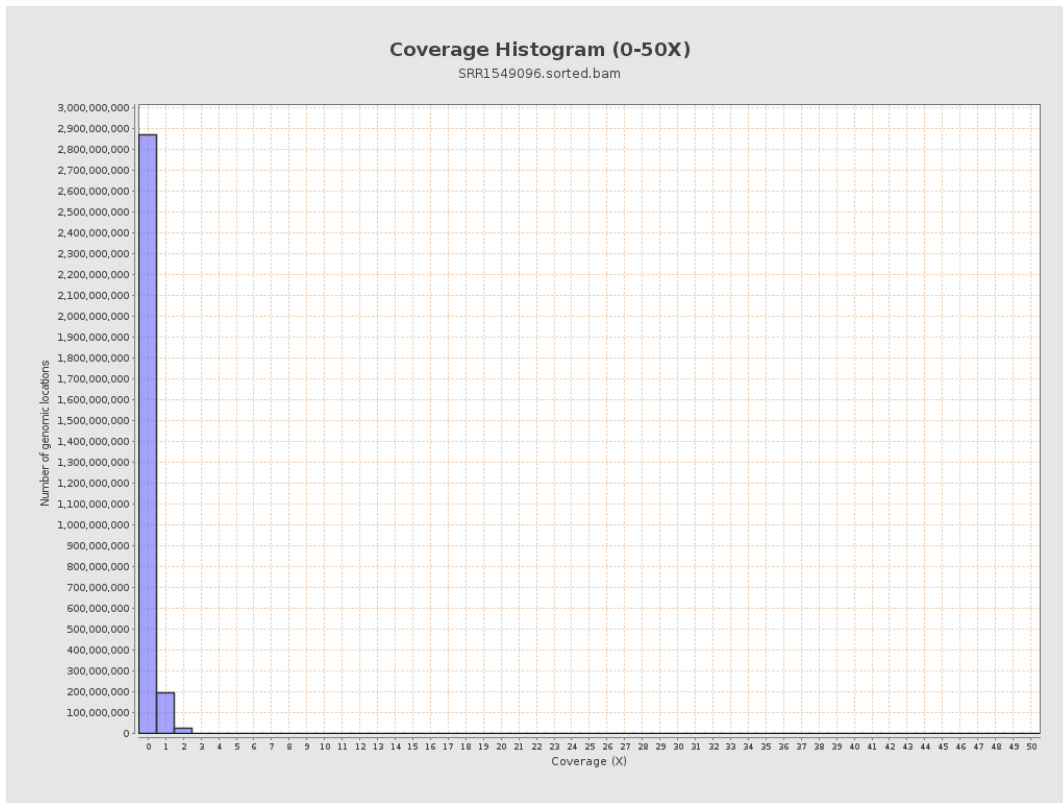
3. Results : Coverage across reference



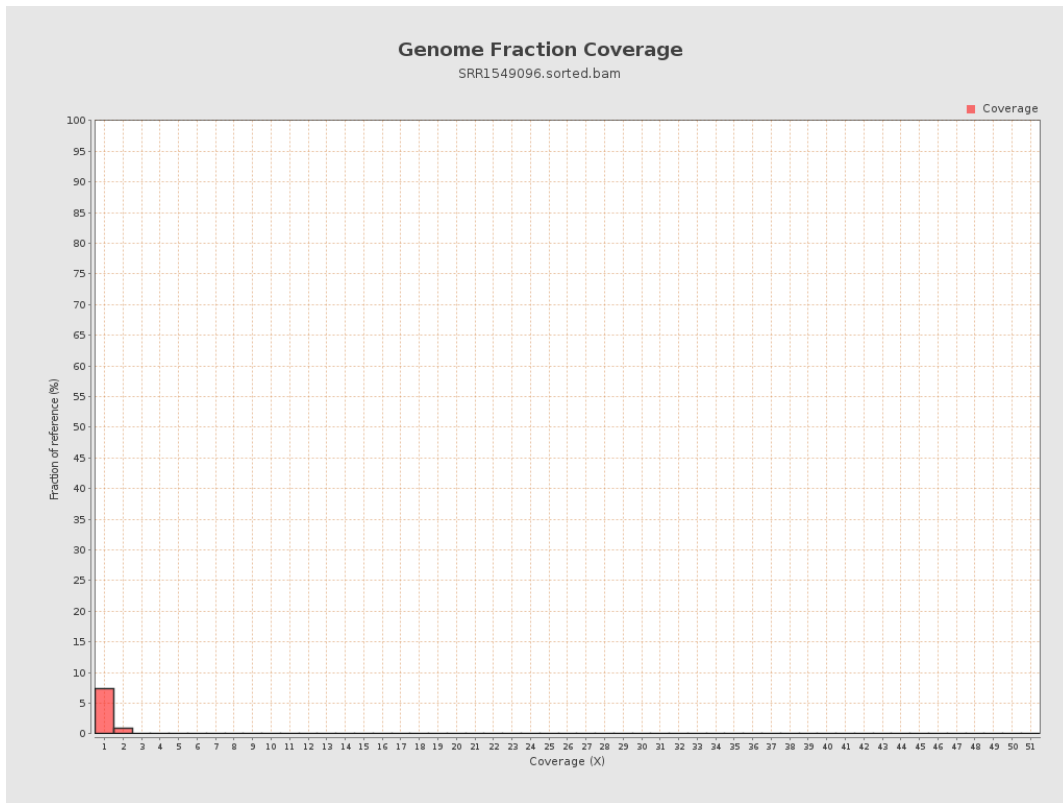
4. Results : Coverage Histogram



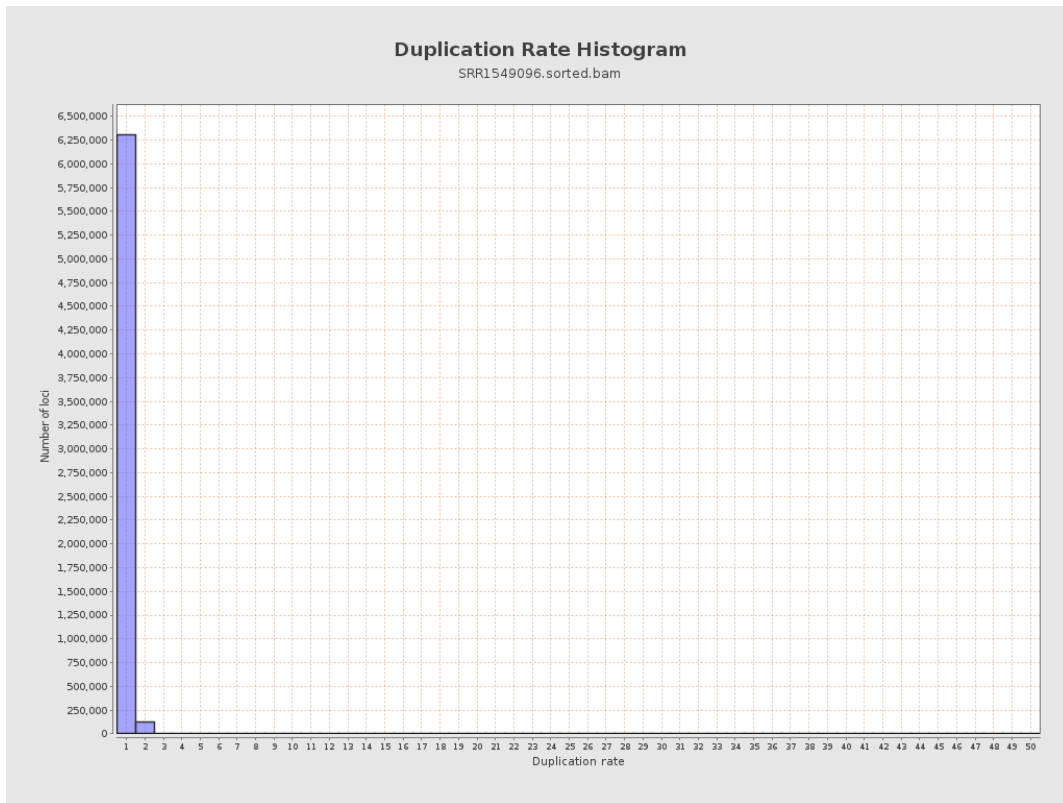
5. Results : Coverage Histogram (0-50X)



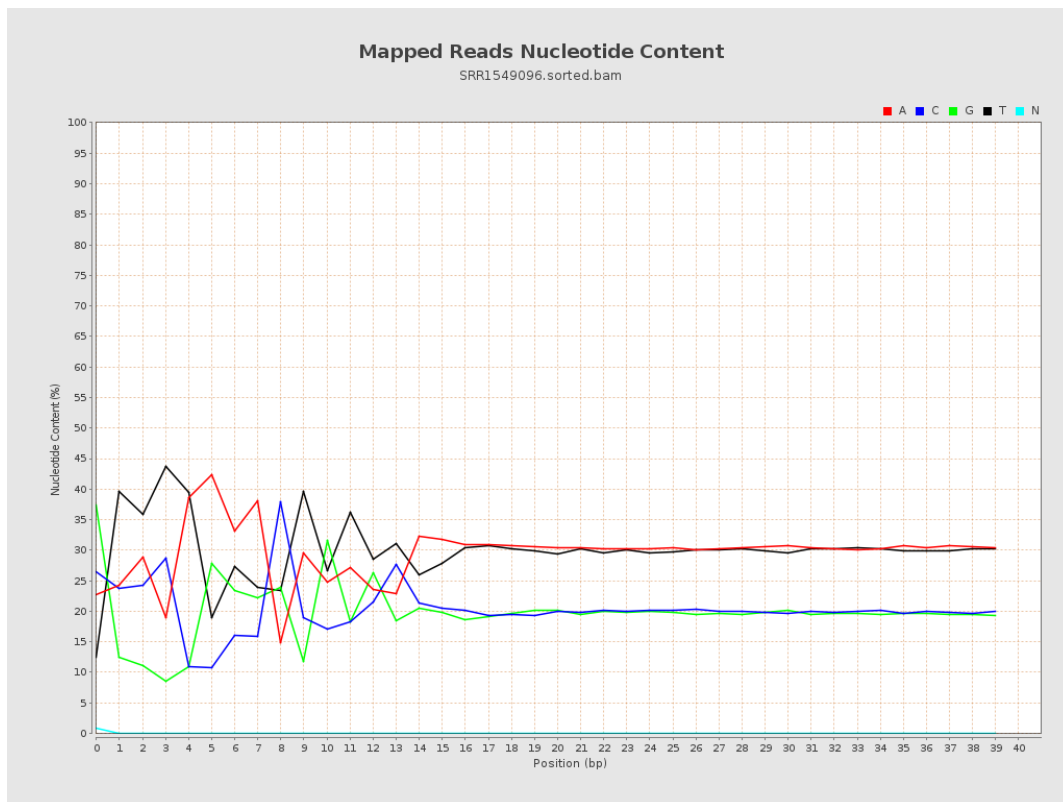
6. Results : Genome Fraction Coverage



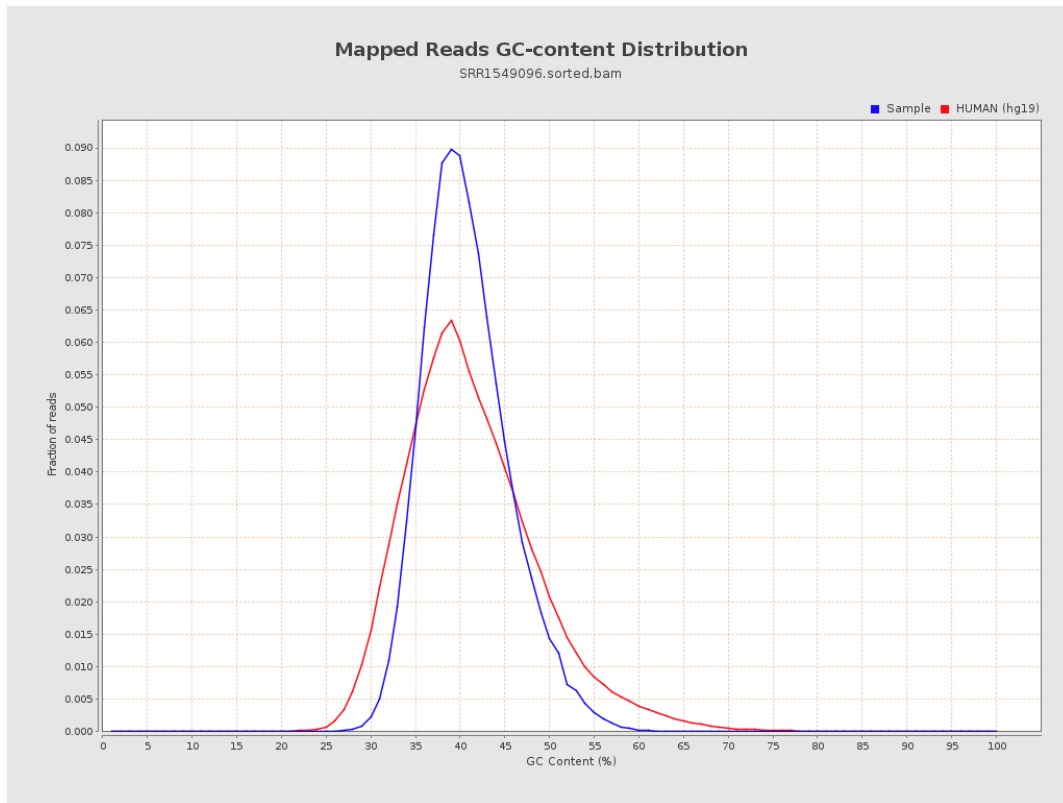
7. Results : Duplication Rate Histogram



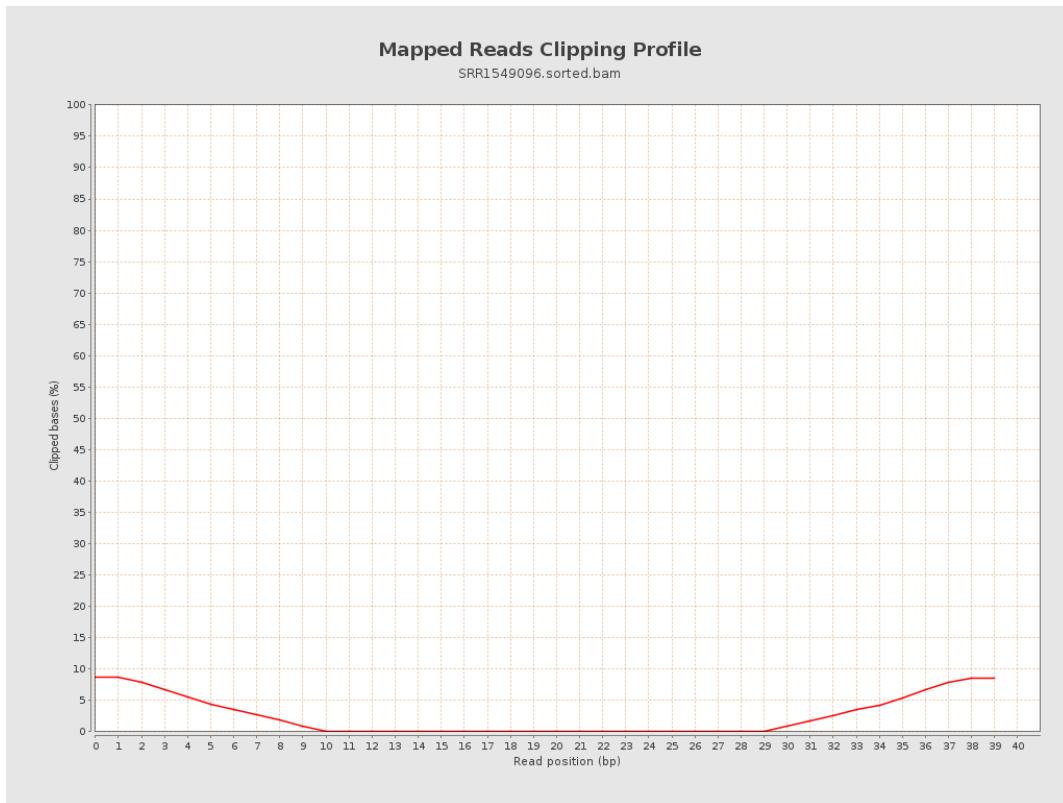
8. Results : Mapped Reads Nucleotide Content



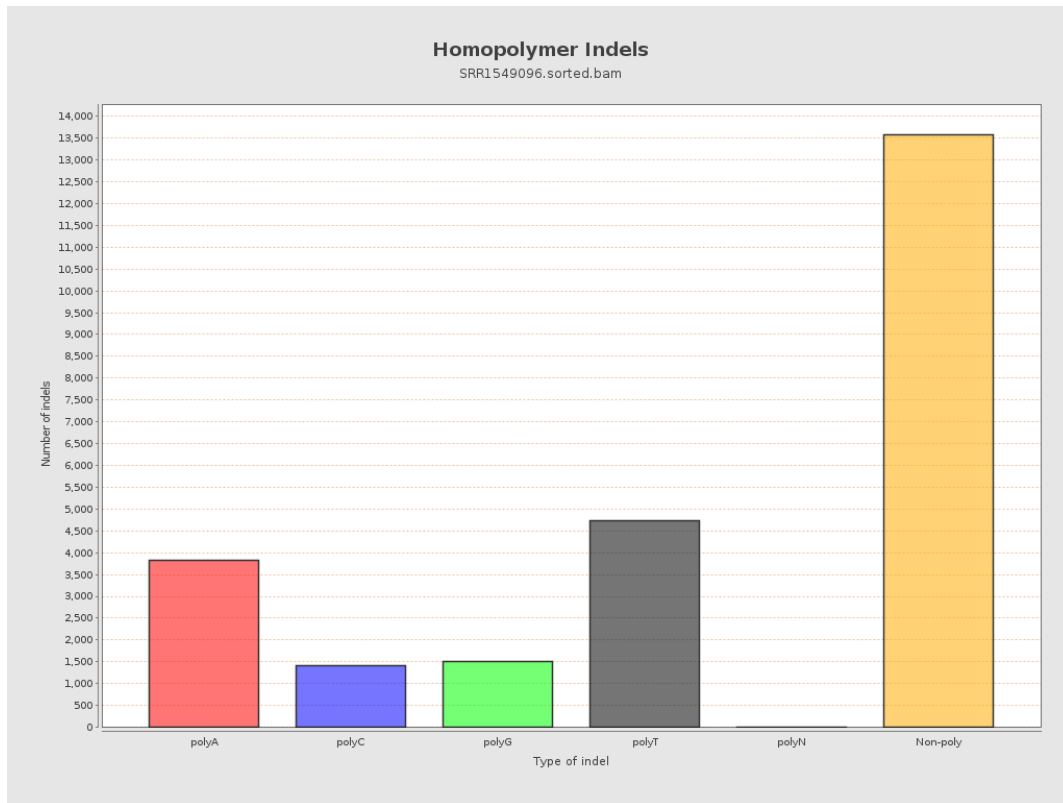
9. Results : Mapped Reads GC-content Distribution



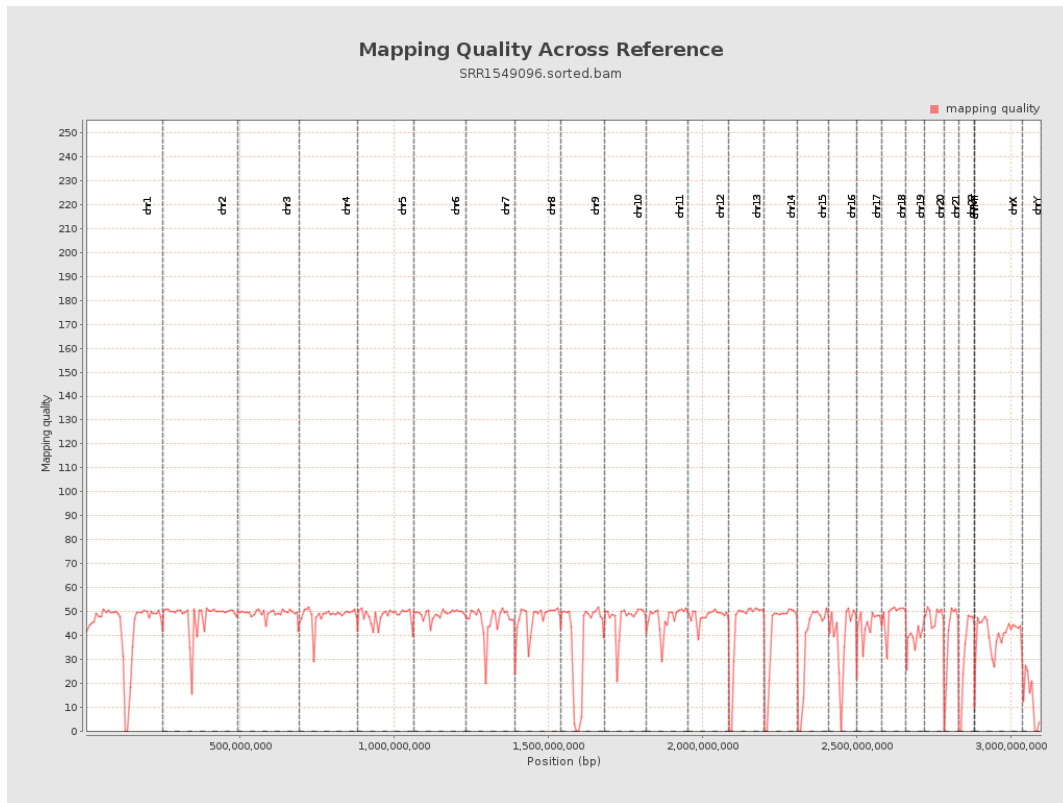
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

