

Qualimap Analysis Results

BAM QC analysis

Generated by Qualimap v.2.2.2-dev

2024/08/29 23:05:33

1. Input data & parameters

1.1. QualiMap command line

```
qualimap bamqc -bam SRR10525206.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_SRR10525206 /home/luna/Desktop/database/homo_bwa/hsa.fa SRR10525206.fastq.gz
Draw chromosome limits:	yes
Analyze overlapping paired-end reads:	no
Program:	bwa (0.7.17-r1188)
Analysis date:	Thu Aug 29 23:05:26 CST 2024
Size of a homopolymer:	3
Skip duplicate alignments:	no
Number of windows:	400
BAM file:	SRR10525206.sorted.bam

2. Summary

2.1. Globals

Reference size	3,095,693,983
Number of reads	3,339,911
Mapped reads	3,058,031 / 91.56%
Unmapped reads	281,880 / 8.44%
Mapped paired reads	0 / 0%
Secondary alignments	0
Supplementary alignments	9,476 / 0.28%
Read min/max/mean length	30 / 76 / 76.1
Duplicated reads (estimated)	155,056 / 4.64%
Duplication rate	3.68%
Clipped reads	3,058,017 / 91.56%

2.2. ACGT Content

Number/percentage of A's	43,673,169 / 24.67%
Number/percentage of C's	35,408,929 / 20%
Number/percentage of T's	57,263,762 / 32.34%
Number/percentage of G's	40,698,769 / 22.99%
Number/percentage of N's	4,270 / 0%
GC Percentage	42.99%

2.3. Coverage

Mean	0.0572

Standard Deviation	0.4907
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2.4. Mapping Quality

Mean Mapping Quality	44.53
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2.5. Mismatches and indels

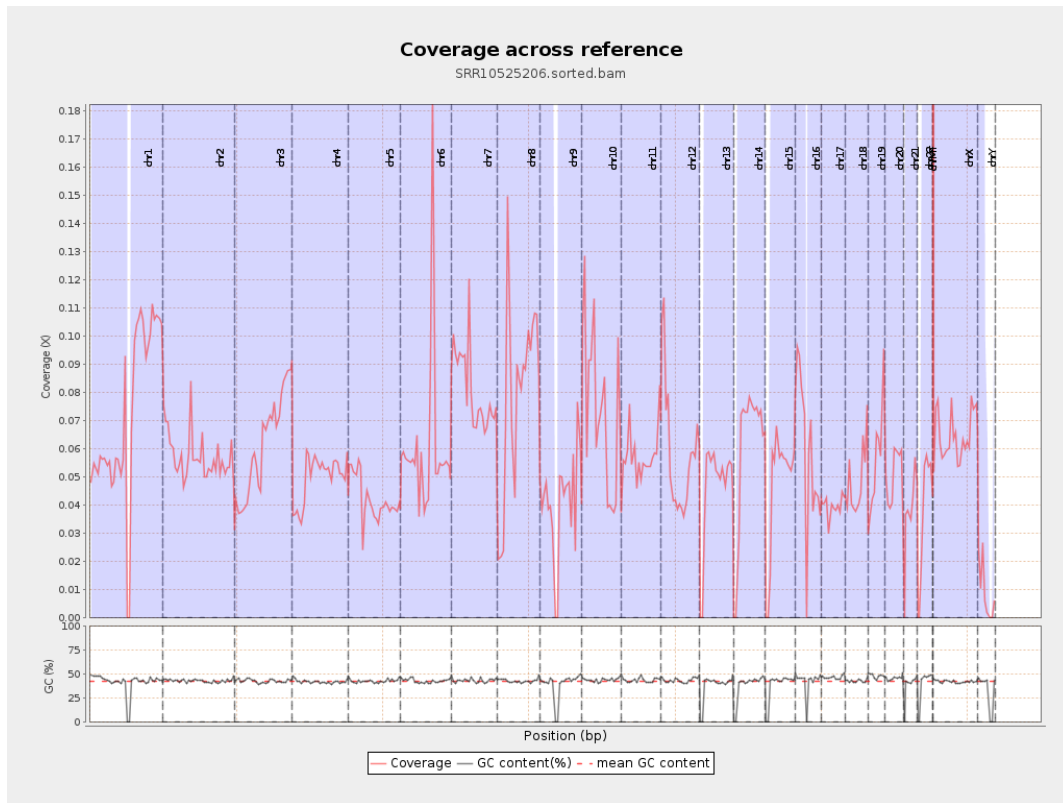
General error rate	0.53%
Mismatches	916,147
Insertions	11,365
Mapped reads with at least one insertion	0.37%
Deletions	36,640
Mapped reads with at least one deletion	1.19%
Homopolymer indels	43.26%

2.6. Chromosome stats

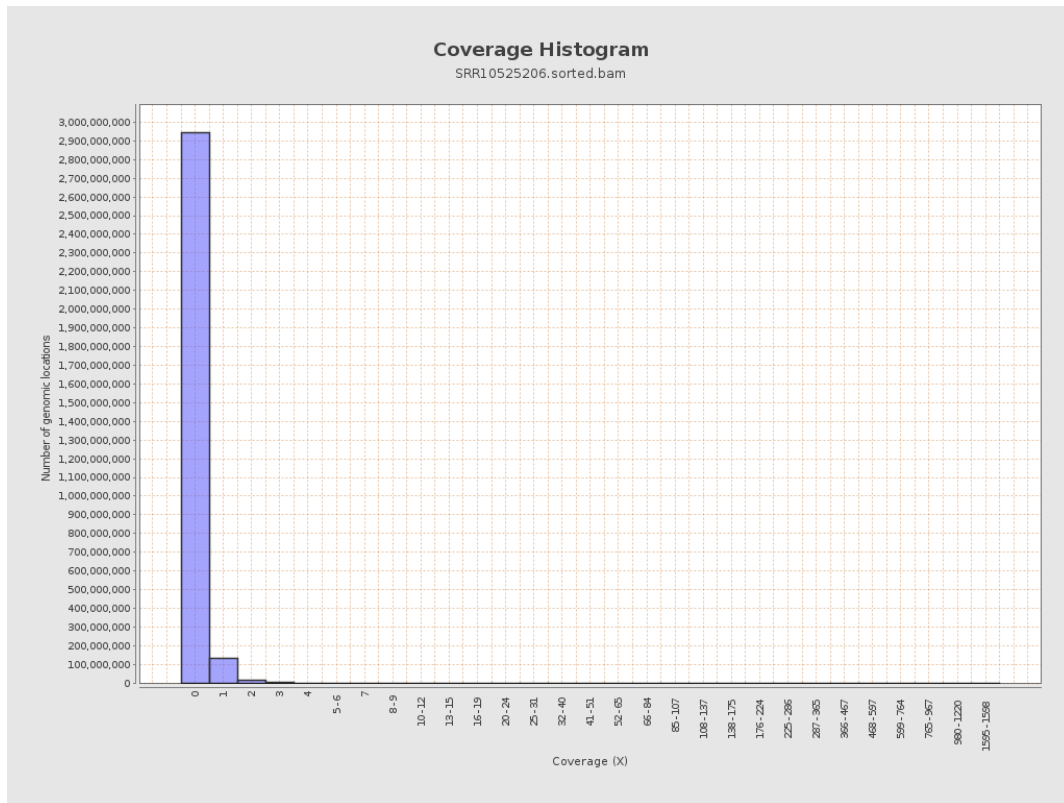
Name	Length	Mapped bases	Mean coverage	Standard deviation
chr1	249250621	18001734	0.0722	0.8854
chr2	243199373	13915943	0.0572	0.7066
chr3	198022430	12082456	0.061	0.2903
chr4	191154276	9539286	0.0499	0.2764
chr5	180915260	7581673	0.0419	0.2356
chr6	171115067	10278969	0.0601	0.338
chr7	159138663	12914141	0.0812	0.8545

chr8	146364022	11821505	0.0808	0.4684
chr9	141213431	5720896	0.0405	0.3373
chr10	135534747	9836018	0.0726	0.5171
chr11	135006516	7745082	0.0574	0.3865
chr12	133851895	7817942	0.0584	0.2823
chr13	115169878	5353930	0.0465	0.2553
chr14	107349540	6498416	0.0605	0.297
chr15	102531392	4714946	0.046	0.2656
chr16	90354753	5303092	0.0587	0.3283
chr17	81195210	3196255	0.0394	0.2433
chr18	78077248	3738228	0.0479	0.7341
chr19	59128983	3361484	0.0569	0.6214
chr20	63025520	3176351	0.0504	0.2658
chr21	48129895	1857401	0.0386	0.2484
chr22	51304566	1935010	0.0377	0.2221
chrMT	16571	161520	9.7471	5.7357
chrX	155270560	10039294	0.0647	0.3346
chrY	59373566	515326	0.0087	0.2107

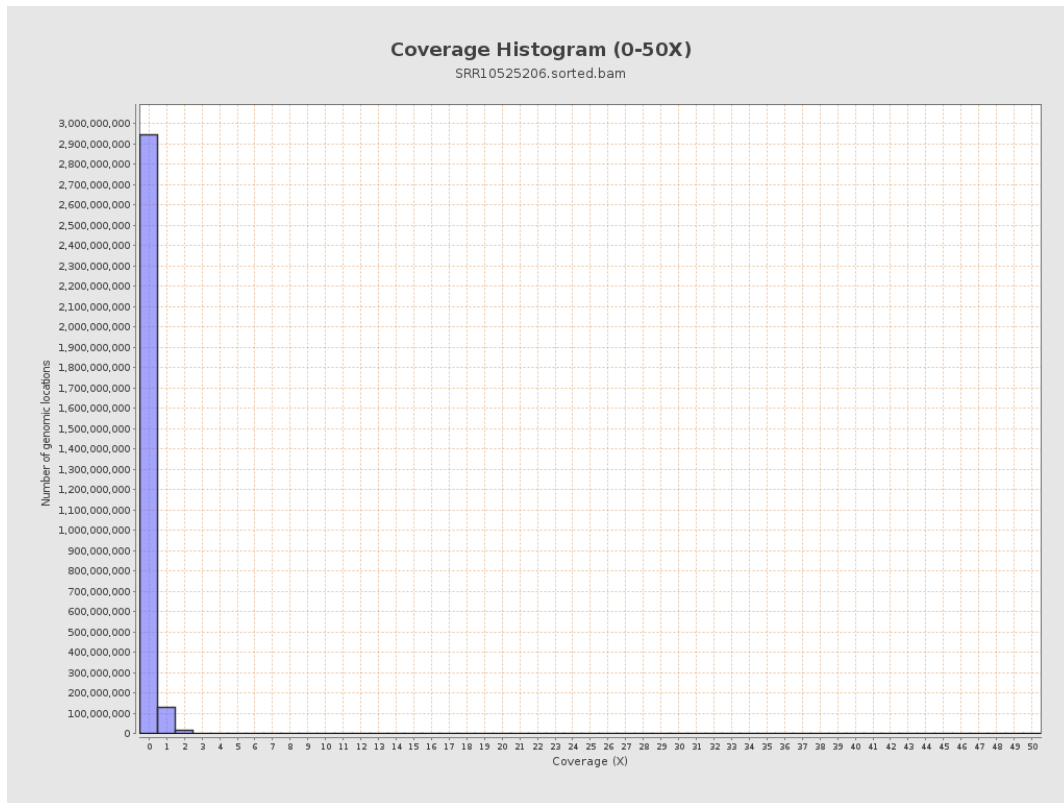
3. Results : Coverage across reference



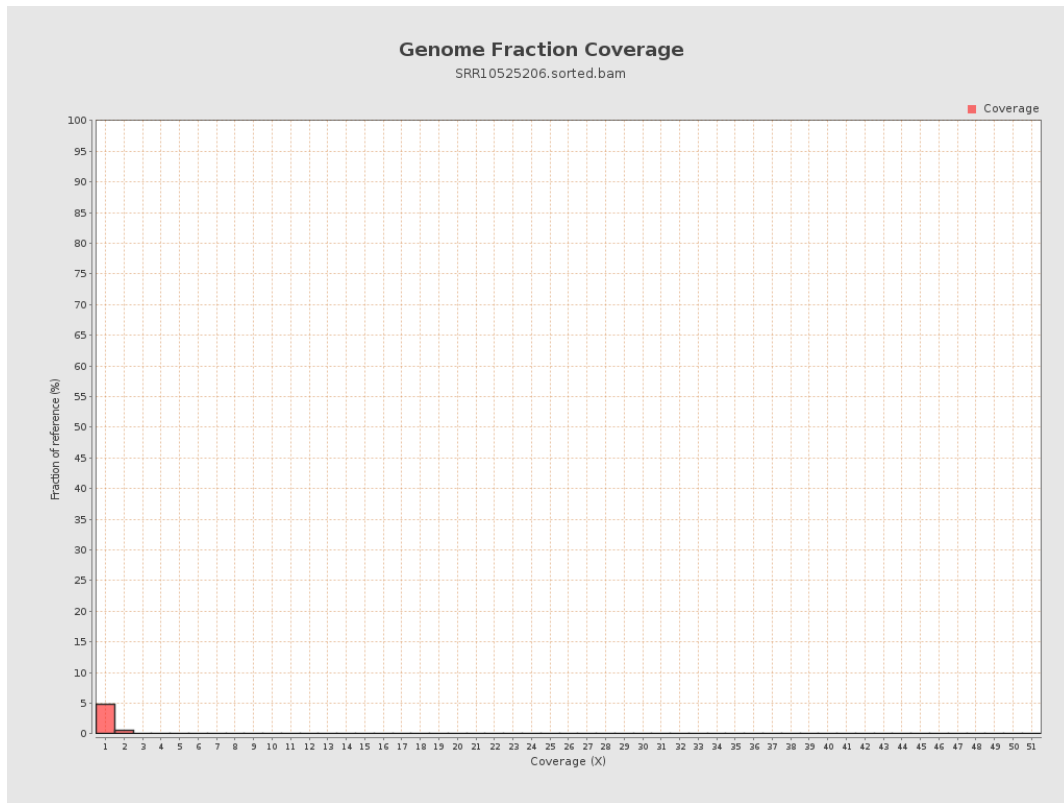
4. Results : Coverage Histogram



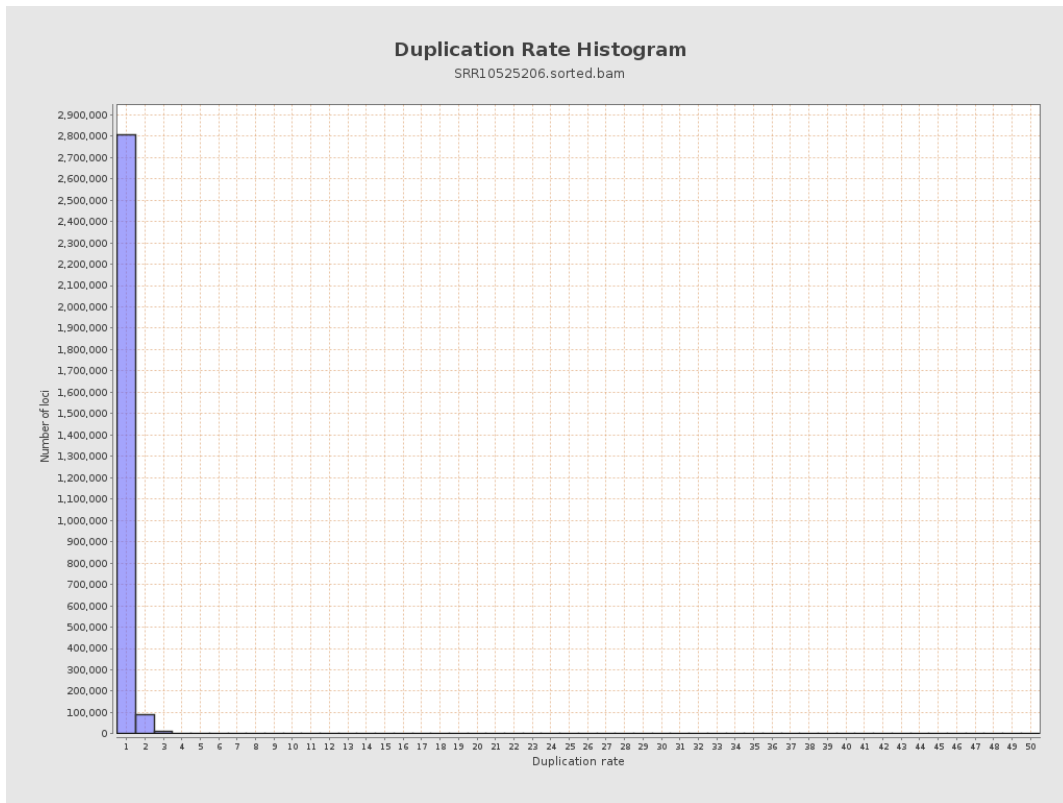
5. Results : Coverage Histogram (0-50X)



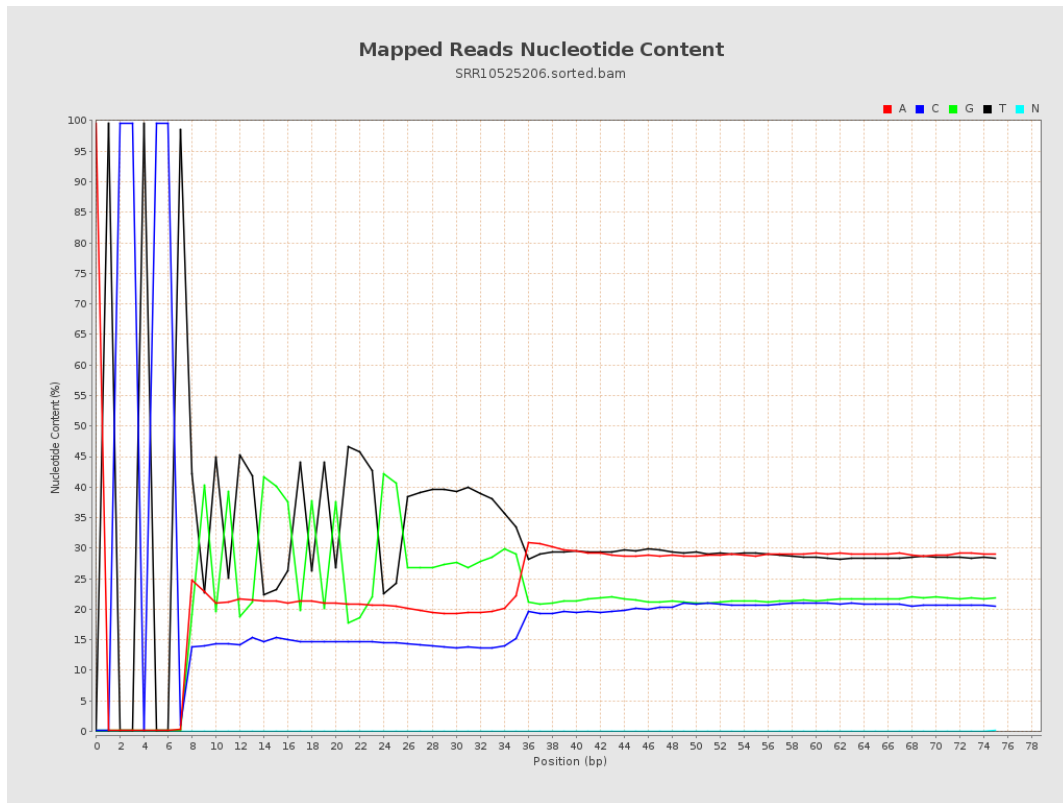
6. Results : Genome Fraction Coverage



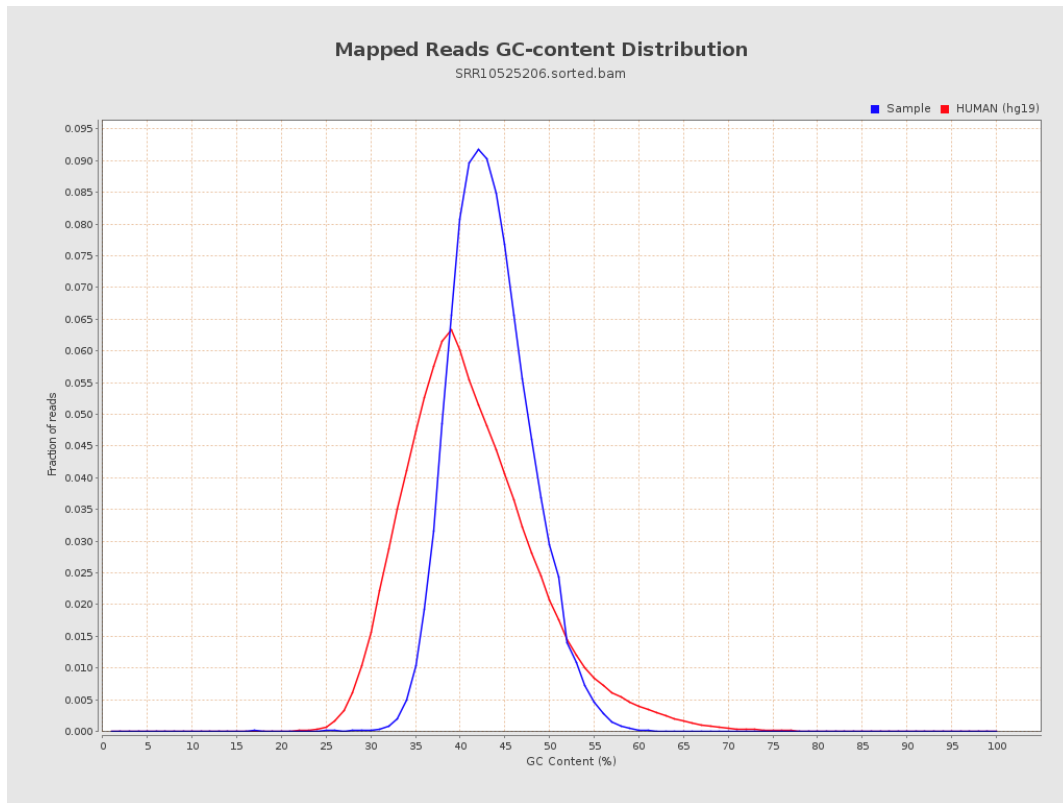
7. Results : Duplication Rate Histogram



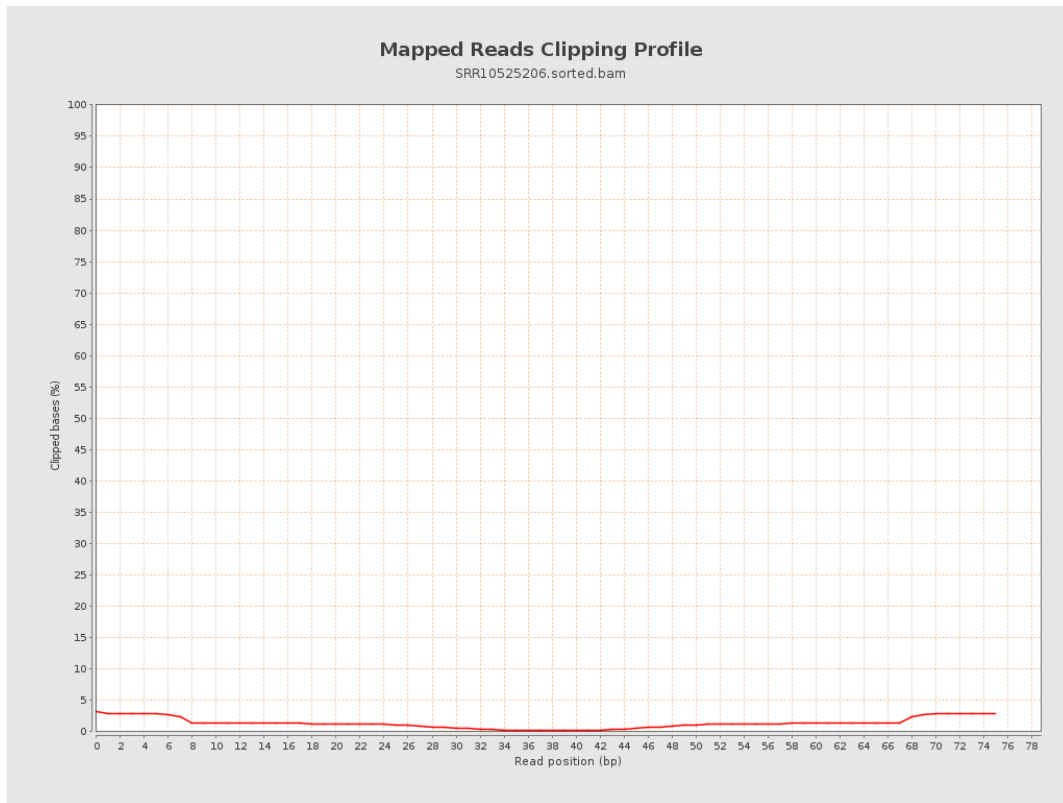
8. Results : Mapped Reads Nucleotide Content



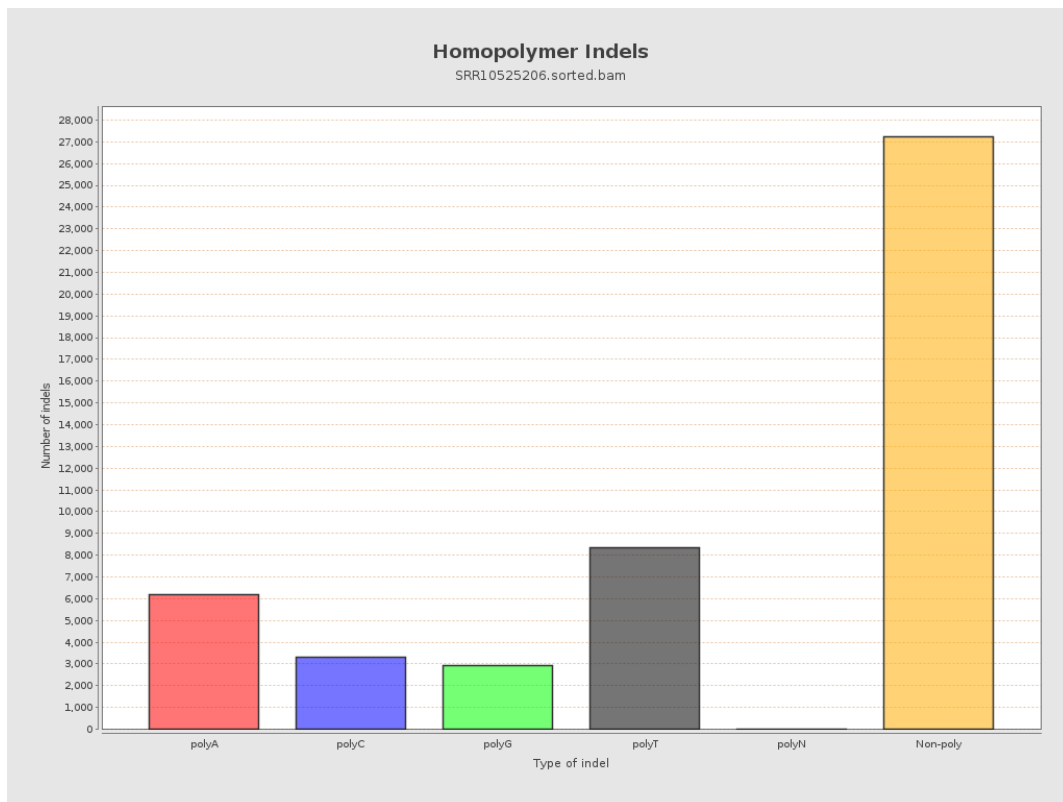
9. Results : Mapped Reads GC-content Distribution



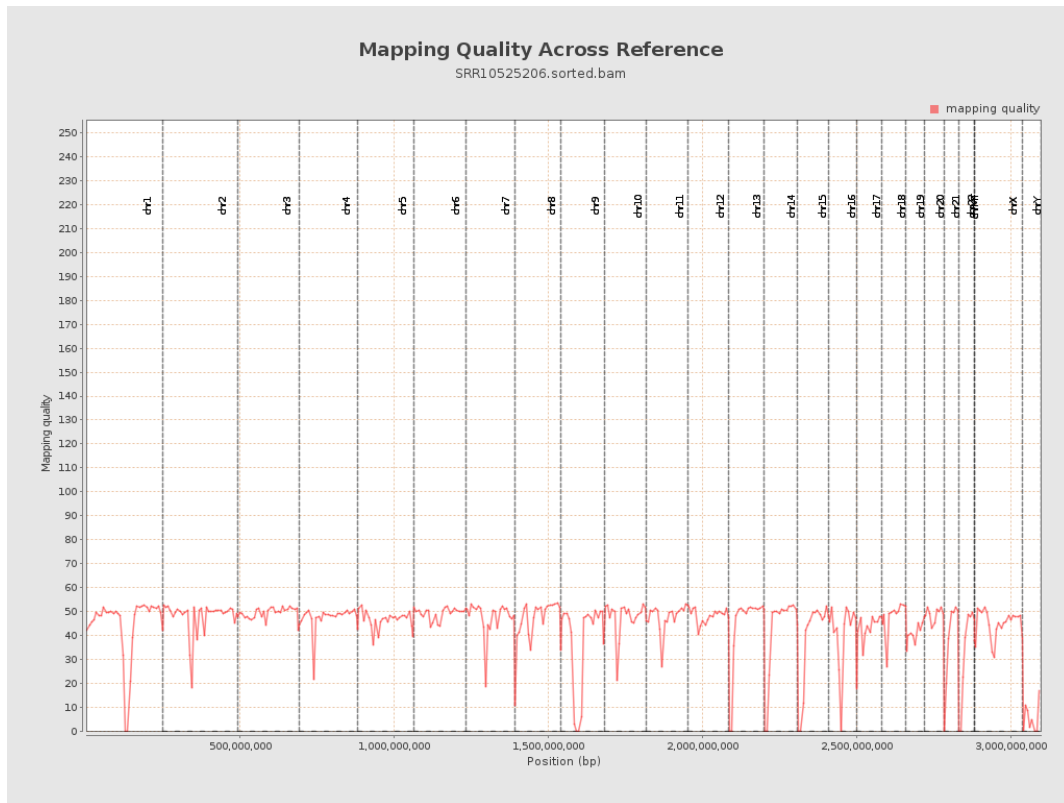
10. Results : Mapped Reads Clipping Profile



11. Results : Homopolymer Indels



12. Results : Mapping Quality Across Reference



13. Results : Mapping Quality Histogram

