Qualimap Analysis Results

BAM QC analysis Generated by Qualimap v.2.2.1 2023/05/29 21:27:13



1. Input data & parameters

1.1. QualiMap command line

qualimap bamqc -bam /proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/147 9 .nodup.bam -nw 400 -hm 3

1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/worki ng/Kate/Working/F2/MarkDuplicates/ 1479 .nodup.bam
Program:	bwa (0.7.17-r1188)
Analyze overlapping paired-end reads:	no
Command line:	bwa mem -M -t 8 -R @RG\tID:\unit\tPL:IIIumina\tLB:LibA\t SM:\unit\tPL:IIIumina\tLB:LibA\t SM:\unit\tPL:IIIumina\tLB:LibA\t SM:\unit\text{sample} /proj/uppstore2018210/Aalpina/data/r eference/GCA_900128785.1_MPIPZ. v5_genomic.fa /proj/uppstore2018210/Aalpina/data/r awdata/P26207/P26207_538/02- FASTQ/220906_A00187_0838_AHM G3KDSX3/P26207_538_S105_L004 _R1_001.fastq.gz /proj/uppstore2018210/Aalpina/data/r awdata/P26207/P26207_538/02- FASTQ/220906_A00187_0838_AHM G3KDSX3/P26207_538_S105_L004 _R2_001.fastq.gz
Size of a homopolymer:	3



Number of windows:	400
Analysis date:	Mon May 29 21:27:12 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no



2. Summary

2.1. Globals

Reference size	311,642,060
Number of reads	54,959,118
Mapped reads	50,737,668 / 92.32%
Unmapped reads	4,221,450 / 7.68%
Mapped paired reads	50,737,668 / 92.32%
Mapped reads, first in pair	25,478,874 / 46.36%
Mapped reads, second in pair	25,258,794 / 45.96%
Mapped reads, both in pair	49,298,726 / 89.7%
Mapped reads, singletons	1,438,942 / 2.62%
Read min/max/mean length	30 / 151 / 147.97
Duplicated reads (flagged)	7,591,122 / 13.81%
Clipped reads	12,985,543 / 23.63%

2.2. ACGT Content

Number/percentage of A's	2,146,854,799 / 30.99%		
Number/percentage of C's	1,315,334,359 / 18.99%		
Number/percentage of T's	2,146,606,454 / 30.99%		
Number/percentage of G's	1,318,137,457 / 19.03%		
Number/percentage of N's	47,870 / 0%		
GC Percentage	38.02%		

2.3. Coverage



Mean	22.2859
Standard Deviation	198.0774

2.4. Mapping Quality

Mean Mapping Quality	43.71

2.5. Insert size

Mean	253,154.22	
Standard Deviation	2,414,990.8	
P25/Median/P75	315 / 415 / 535	

2.6. Mismatches and indels

General error rate	2.65%
Mismatches	169,926,442
Insertions	5,044,234
Mapped reads with at least one insertion	8.87%
Deletions	4,907,092
Mapped reads with at least one deletion	8.58%
Homopolymer indels	56.83%

2.7. Chromosome stats

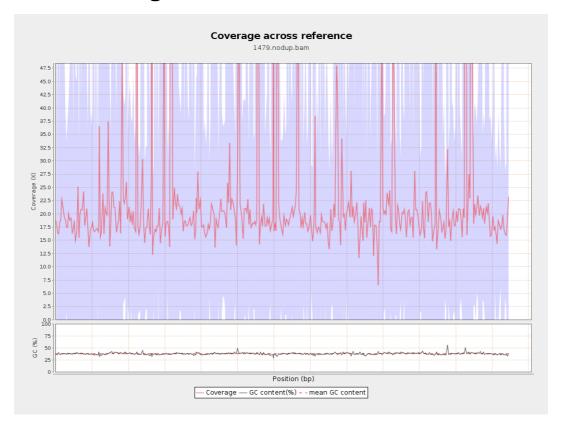
Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	547728033	18.4269	81.5777



LT669789.1	36598175	843509753	23.0479	210.1281
LT669790.1	30422129	802556746	26.3807	263.7921
LT669791.1	52758100	1158626484	21.9611	217.0047
LT669792.1	28376109	644603454	22.7164	209.3898
LT669793.1	33388210	681267430	20.4044	108.3817
LT669794.1	50579949	1084495802	21.4412	165.1775
LT669795.1	49795044	1182422143	23.7458	236.8942

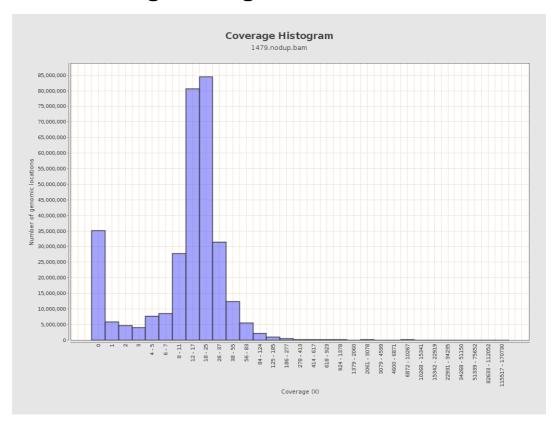


3. Results: Coverage across reference



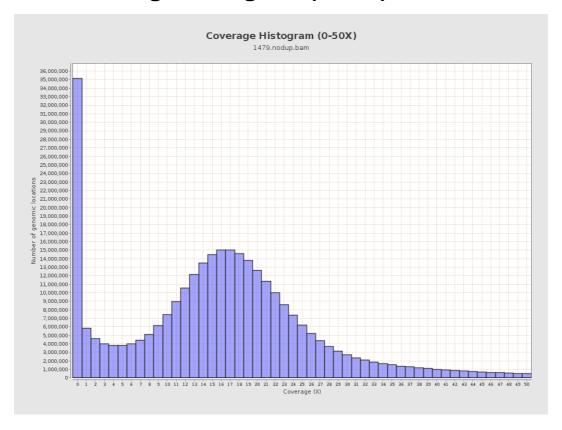


4. Results: Coverage Histogram



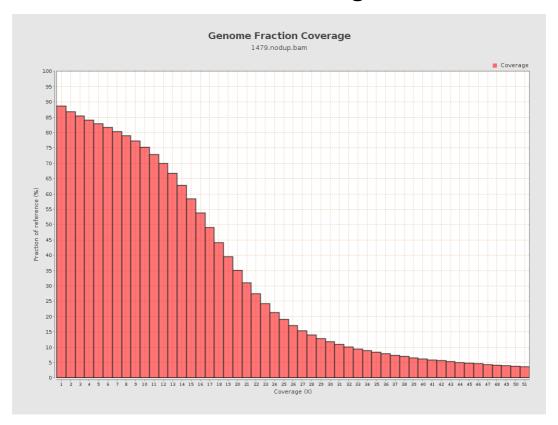


5. Results: Coverage Histogram (0-50X)



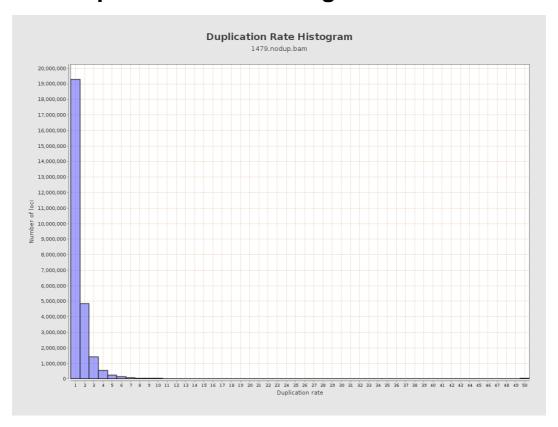


6. Results : Genome Fraction Coverage





7. Results : Duplication Rate Histogram



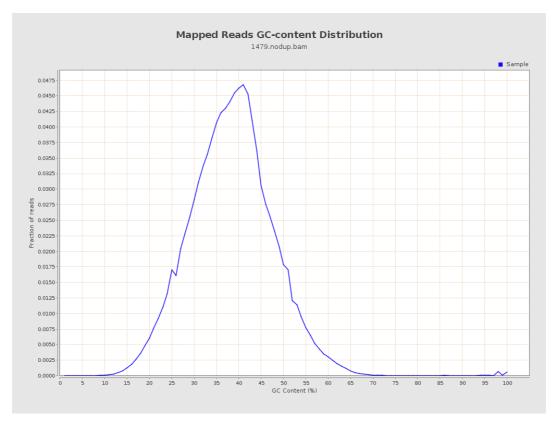


8. Results: Mapped Reads Nucleotide Content



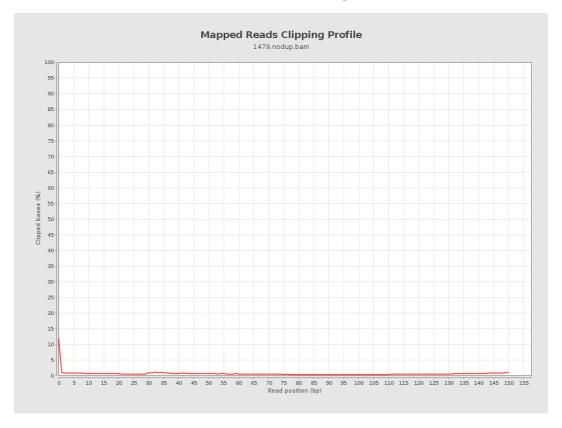


9. Results: Mapped Reads GC-content Distribution



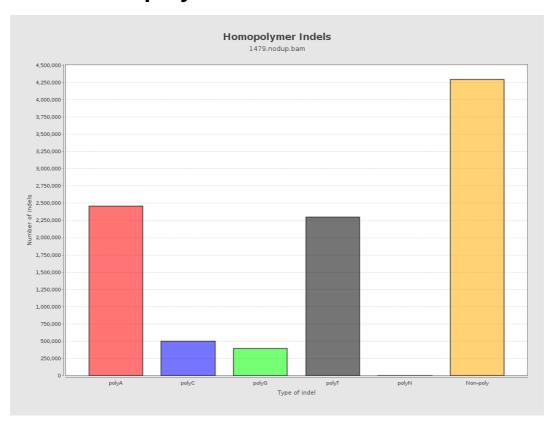


10. Results: Mapped Reads Clipping Profile





11. Results : Homopolymer Indels



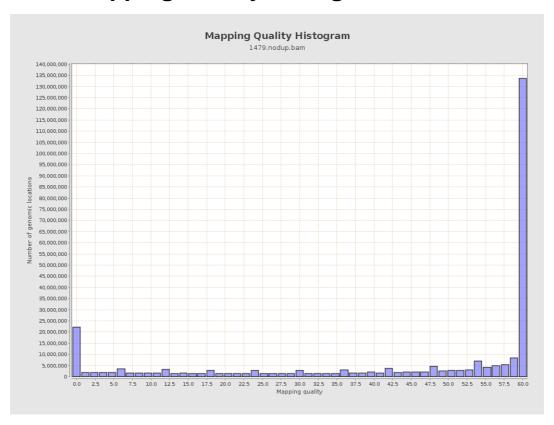


12. Results: Mapping Quality Across Reference



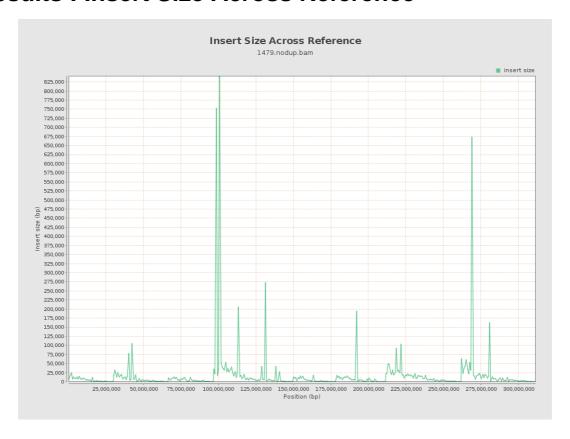


13. Results: Mapping Quality Histogram





14. Results: Insert Size Across Reference





15. Results: Insert Size Histogram

