Qualimap Analysis Results

BAM QC analysis Generated by Qualimap v.2.2.1 2023/05/29 21:30:46



1. Input data & parameters

1.1. QualiMap command line

qualimap bamqc -bam /proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/143 7 .nodup.bam -nw 400 -hm 3

1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/worki ng/Kate/Working/F2/MarkDuplicates/ 1437 .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:\tIllumina\tLB:\tibA\t SM:\unit\tPL:\tIllumina\tLB:\tibA\t SM:\unit\tPL:\tIllumina\tLB:\tibA\t SM:\unit\tPL:\tIllumina\tLB:\tibA\t SM:\unit\tPL:\tIllumina\tLB:\tibA\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_562/02- FASTQ/220906_A00187_0838_AHM G3KDSX3/P26207/P26207_562/02- FASTQ/220906_A00187_0838_AHM G3KDSX3/P26207_562_S129_L004 _R2_001.fastq.gz
Size of a homopolymer:	3



	CENTRO DE INVESTIGACION
Number of windows:	400
Analysis date:	Mon May 29 21:30:45 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no



2. Summary

2.1. Globals

Reference size	311,642,060
Number of reads	60,480,316
Mapped reads	57,548,201 / 95.15%
Unmapped reads	2,932,115 / 4.85%
Mapped paired reads	57,548,201 / 95.15%
Mapped reads, first in pair	28,846,816 / 47.7%
Mapped reads, second in pair	28,701,385 / 47.46%
Mapped reads, both in pair	56,588,023 / 93.56%
Mapped reads, singletons	960,178 / 1.59%
Read min/max/mean length	30 / 151 / 148.09
Duplicated reads (flagged)	7,822,395 / 12.93%
Clipped reads	13,302,839 / 22%

2.2. ACGT Content

Number/percentage of A's	2,467,842,252 / 30.95%
Number/percentage of C's	1,519,811,218 / 19.06%
Number/percentage of T's	2,468,934,635 / 30.97%
Number/percentage of G's	1,516,454,349 / 19.02%
Number/percentage of N's	59,902 / 0%
GC Percentage	38.08%

2.3. Coverage



Mean	25.6498
Standard Deviation	201.8073

2.4. Mapping Quality

Mean Mapping Quality	44 23
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2.5. Insert size

Mean	206,801.62
Standard Deviation	2,136,541.76
P25/Median/P75	306 / 403 / 522

2.6. Mismatches and indels

General error rate	2.44%
Mismatches	180,269,609
Insertions	5,245,617
Mapped reads with at least one insertion	8.22%
Deletions	5,428,362
Mapped reads with at least one deletion	8.41%
Homopolymer indels	56.22%

2.7. Chromosome stats

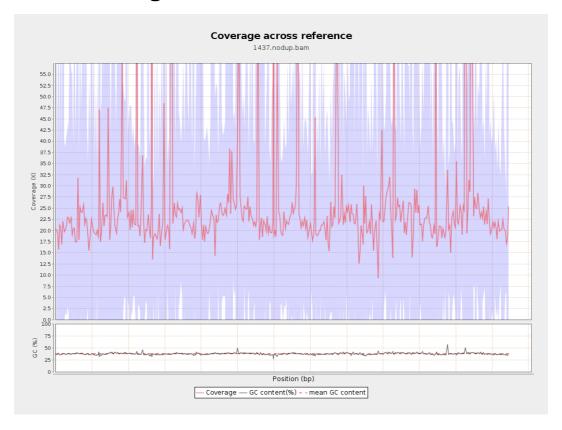
Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	619904581	20.8551	44.3469



LT669789.1	36598175	977636551	26.7127	216.5278
LT669790.1	30422129	807939028	26.5576	178.7853
LT669791.1	52758100	1326467386	25.1424	155.5692
LT669792.1	28376109	724101821	25.518	228.6529
LT669793.1	33388210	805202576	24.1164	142.7832
LT669794.1	50579949	1247290298	24.6598	177.3352
LT669795.1	49795044	1485008116	29.8224	314.0693

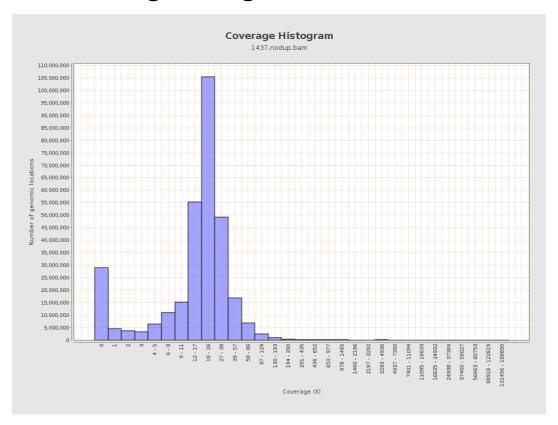


3. Results: Coverage across reference



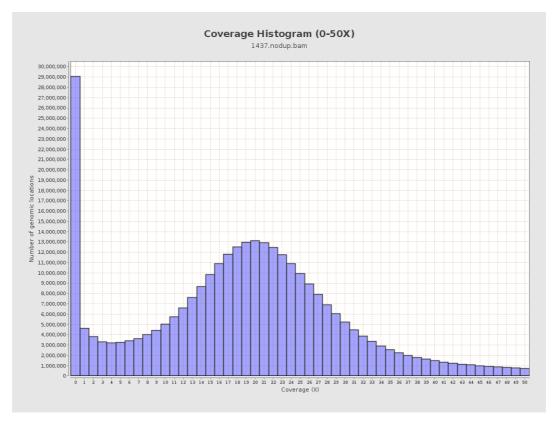


4. Results: Coverage Histogram



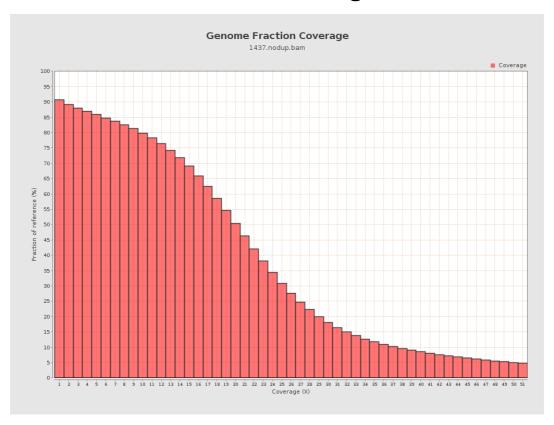


5. Results: Coverage Histogram (0-50X)



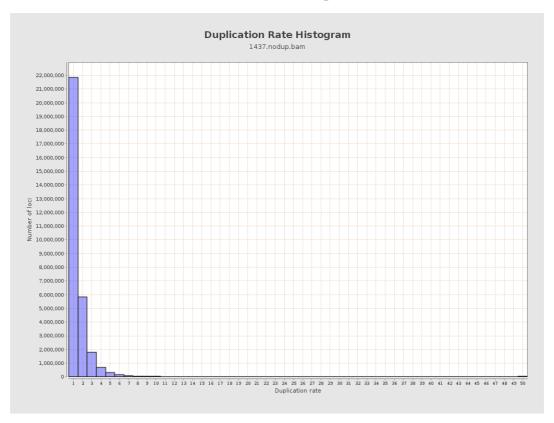


6. Results : Genome Fraction Coverage





7. Results : Duplication Rate Histogram



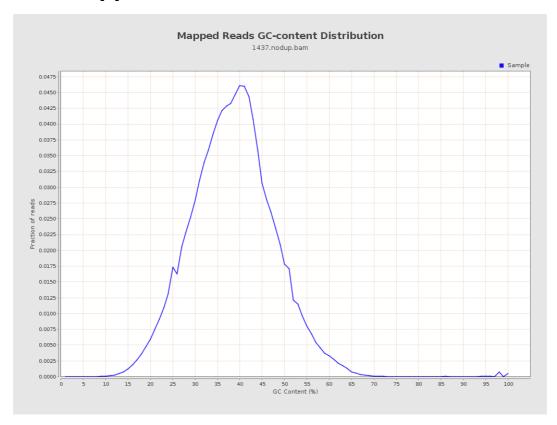


8. Results: Mapped Reads Nucleotide Content



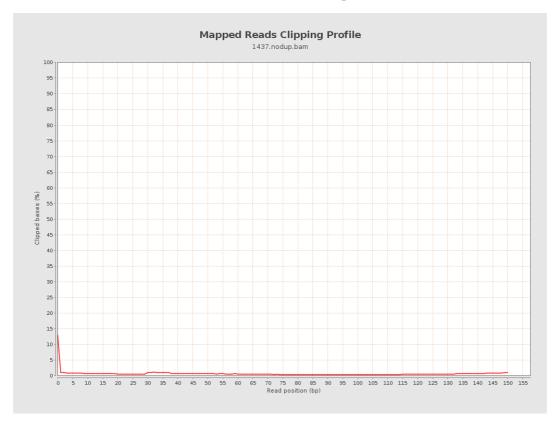


9. Results: Mapped Reads GC-content Distribution



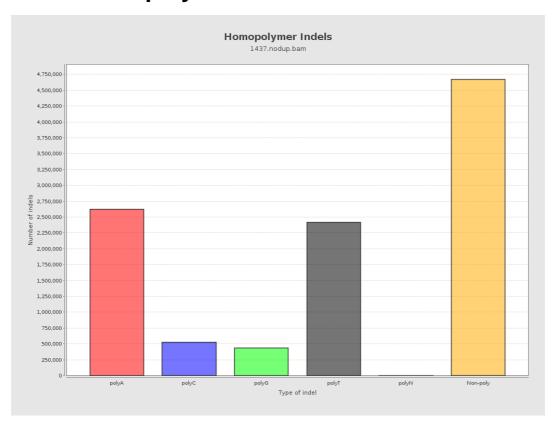


10. Results: Mapped Reads Clipping Profile





11. Results : Homopolymer Indels



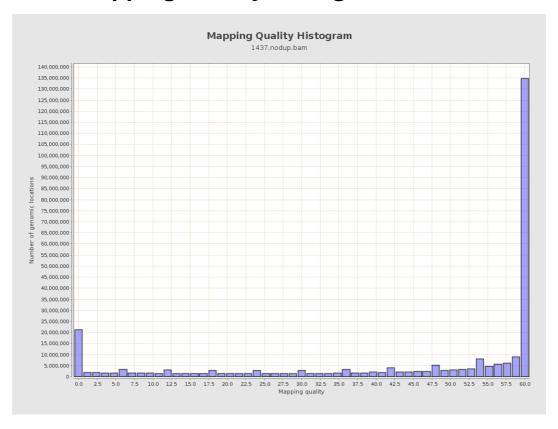


12. Results: Mapping Quality Across Reference



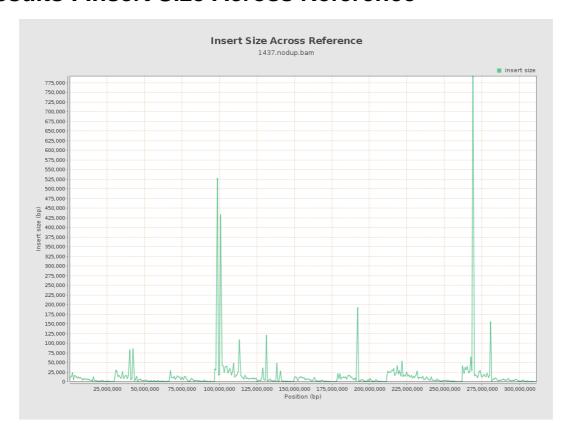


13. Results: Mapping Quality Histogram





14. Results: Insert Size Across Reference





15. Results: Insert Size Histogram

