Qualimap Analysis Results

BAM QC analysis Generated by Qualimap v.2.2.1 2023/05/29 21:26:29



1. Input data & parameters

1.1. QualiMap command line

qualimap bamqc -bam /proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/425 .nodup.bam -nw 400 -hm 3

1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/worki ng/Kate/Working/F2/MarkDuplicates/ 425 .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\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_212/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_212/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_212_S293_L003 _R2_001.fastq.gz
Size of a homopolymer:	3
Number of windows:	400



Analysis date:	Mon May 29 21:26:29 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no



2. Summary

2.1. Globals

Reference size	311,642,060
Number of reads	51,716,823
Mapped reads	47,602,356 / 92.04%
Unmapped reads	4,114,467 / 7.96%
Mapped paired reads	47,602,356 / 92.04%
Mapped reads, first in pair	23,883,938 / 46.18%
Mapped reads, second in pair	23,718,418 / 45.86%
Mapped reads, both in pair	46,270,928 / 89.47%
Mapped reads, singletons	1,331,428 / 2.57%
Read min/max/mean length	30 / 151 / 147.91
Duplicated reads (flagged)	6,928,934 / 13.4%
Clipped reads	12,139,974 / 23.47%

2.2. ACGT Content

Number/percentage of A's	1,998,048,267 / 30.85%		
Number/percentage of C's	1,239,349,156 / 19.13%		
Number/percentage of T's	2,000,295,524 / 30.88%		
Number/percentage of G's	1,239,855,137 / 19.14%		
Number/percentage of N's	23,556 / 0%		
GC Percentage	38.27%		

2.3. Coverage



Mean	20.8391
Standard Deviation	193.2373

2.4. Mapping Quality

Mean Mapping Quality	44.13

2.5. Insert size

Mean	252,660.79
Standard Deviation	2,410,354.54
P25/Median/P75	321 / 434 / 565

2.6. Mismatches and indels

General error rate	2.37%
Mismatches	140,970,353
Insertions	4,541,160
Mapped reads with at least one insertion	8.53%
Deletions	4,477,892
Mapped reads with at least one deletion	8.33%
Homopolymer indels	56.71%

2.7. Chromosome stats

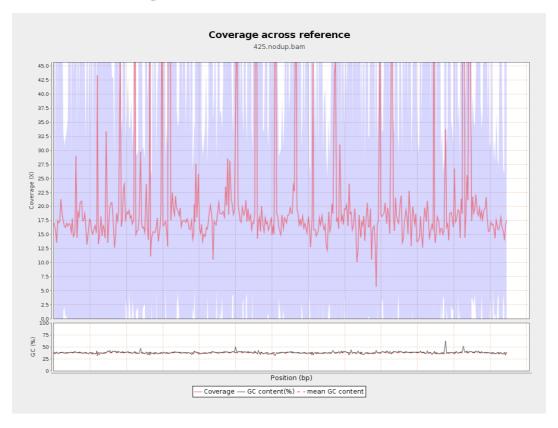
Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	510559662	17.1765	66.5634



LT669789.1	36598175	752169949	20.5521	187.3184
LT669790.1	30422129	739766562	24.3167	233.5768
LT669791.1	52758100	1083117487	20.5299	185.3139
LT669792.1	28376109	586706442	20.6761	188.4465
LT669793.1	33388210	634280578	18.9971	130.0103
LT669794.1	50579949	977712872	19.33	151.8771
LT669795.1	49795044	1210036625	24.3003	283.7482

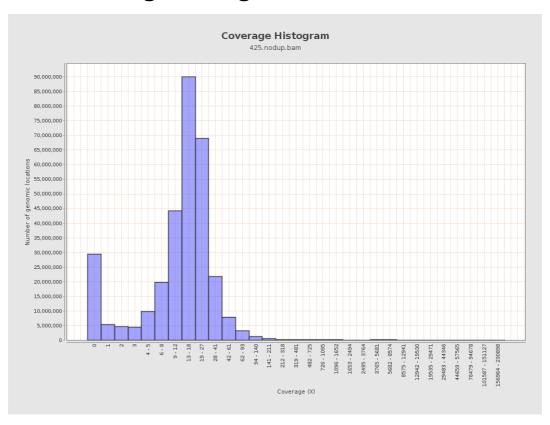


3. Results: Coverage across reference



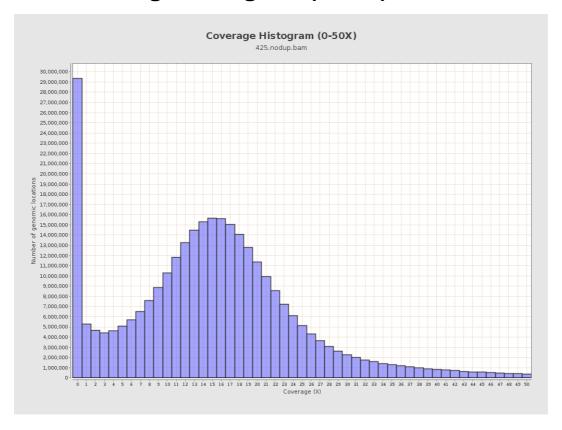


4. Results: Coverage Histogram



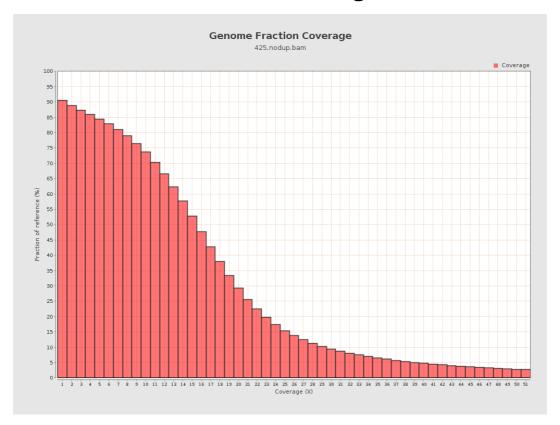


5. Results: Coverage Histogram (0-50X)



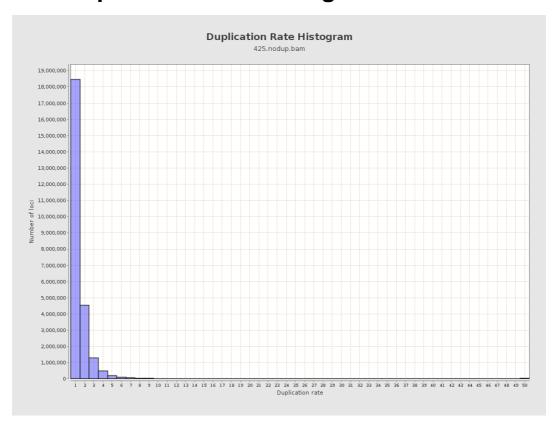


6. Results : Genome Fraction Coverage





7. Results : Duplication Rate Histogram



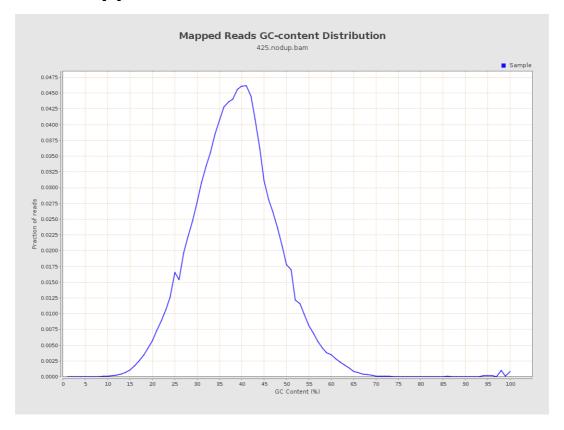


8. Results: Mapped Reads Nucleotide Content



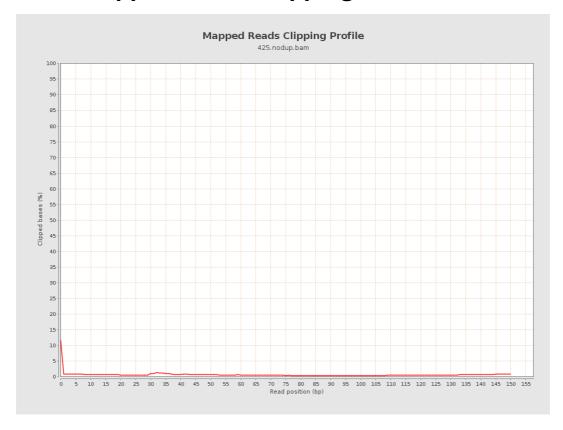


9. Results: Mapped Reads GC-content Distribution



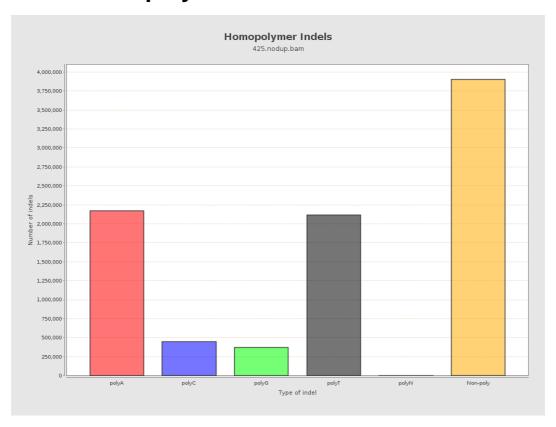


10. Results: Mapped Reads Clipping Profile



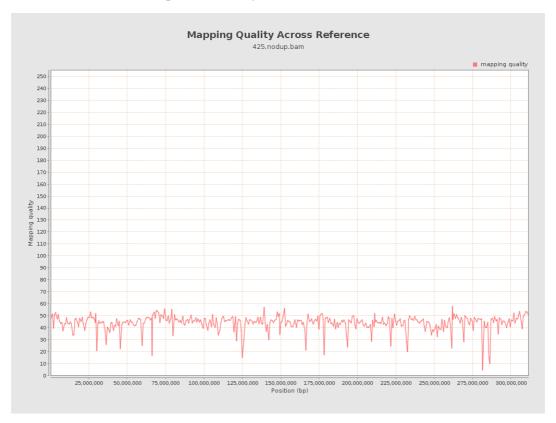


11. Results : Homopolymer Indels



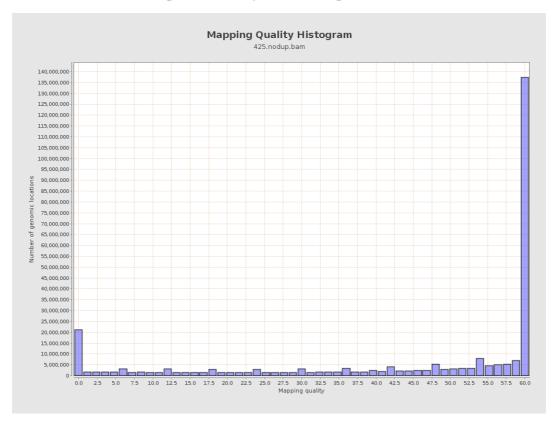


12. Results: Mapping Quality Across Reference



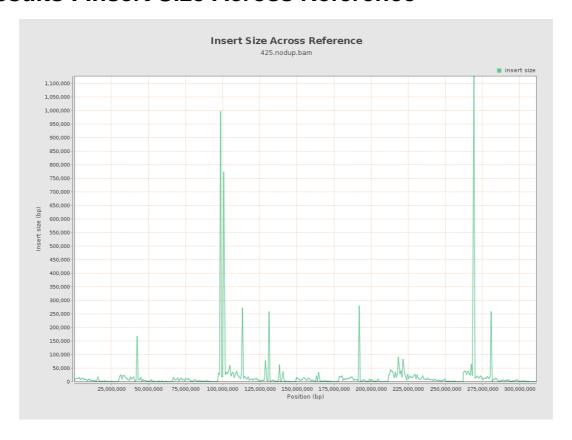


13. Results: Mapping Quality Histogram





14. Results: Insert Size Across Reference





15. Results: Insert Size Histogram

