# Qualimap Analysis Results

BAM QC analysis Generated by Qualimap v.2.2.1 2023/05/29 21:40:19



#### 1. Input data & parameters

#### 1.1. QualiMap command line

qualimap bamqc -bam /proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/145 4 .nodup.bam -nw 400 -hm 3

#### 1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/worki ng/Kate/Working/F2/MarkDuplicates/ 1454 .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_444/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_444_S419_L004 _R1_001.fastq.gz /proj/uppstore2018210/Aalpina/data/r awdata/P26207/P26207_444/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_444_S419_L004 _R2_001.fastq.gz
Size of a homopolymer:	3



Number of windows:	400
Analysis date:	Mon May 29 21:40:19 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no



## 2. Summary

#### 2.1. Globals

Reference size	311,642,060
Number of reads	94,146,788
Mapped reads	88,327,200 / 93.82%
Unmapped reads	5,819,588 / 6.18%
Mapped paired reads	88,327,200 / 93.82%
Mapped reads, first in pair	44,248,353 / 47%
Mapped reads, second in pair	44,078,847 / 46.82%
Mapped reads, both in pair	86,485,628 / 91.86%
Mapped reads, singletons	1,841,572 / 1.96%
Read min/max/mean length	30 / 151 / 148.06
Duplicated reads (flagged)	14,248,172 / 15.13%
Clipped reads	19,320,916 / 20.52%

#### 2.2. ACGT Content

Number/percentage of A's	3,792,223,898 / 30.93%		
Number/percentage of C's	2,336,571,810 / 19.06%		
Number/percentage of T's	3,799,298,434 / 30.99%		
Number/percentage of G's	2,331,938,780 / 19.02%		
Number/percentage of N's	42,864 / 0%		
GC Percentage	38.08%		

#### 2.3. Coverage



Mean	39.4436
Standard Deviation	296.4663

## 2.4. Mapping Quality

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Mean Mapping Quality	44.43
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#### 2.5. Insert size

Mean	243,640.46
Standard Deviation	2,361,032.65
P25/Median/P75	350 / 458 / 599

#### 2.6. Mismatches and indels

General error rate	2.25%
Mismatches	252,576,154
Insertions	8,390,170
Mapped reads with at least one insertion	8.53%
Deletions	8,488,907
Mapped reads with at least one deletion	8.53%
Homopolymer indels	56.52%

#### 2.7. Chromosome stats

Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	1019331451	34.2928	94.8139



LT669789.1	36598175	1468678425	40.1298	342.6311
LT669790.1	30422129	1320879655	43.4184	320.0131
LT669791.1	52758100	2011739546	38.1314	293.9105
LT669792.1	28376109	1116264149	39.3382	287.9653
LT669793.1	33388210	1234238168	36.9663	204.7884
LT669794.1	50579949	1894084991	37.4473	284.0384
LT669795.1	49795044	2227073593	44.7248	383.3356

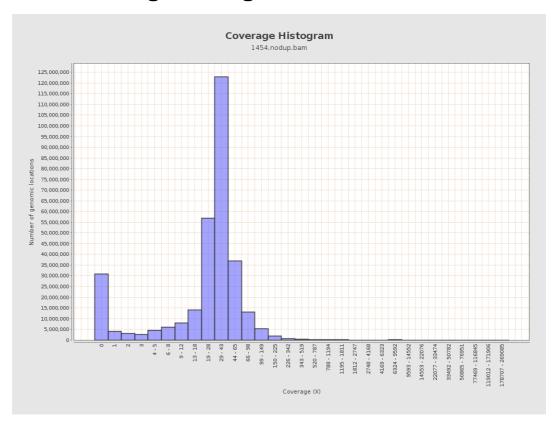


## 3. Results : Coverage across reference



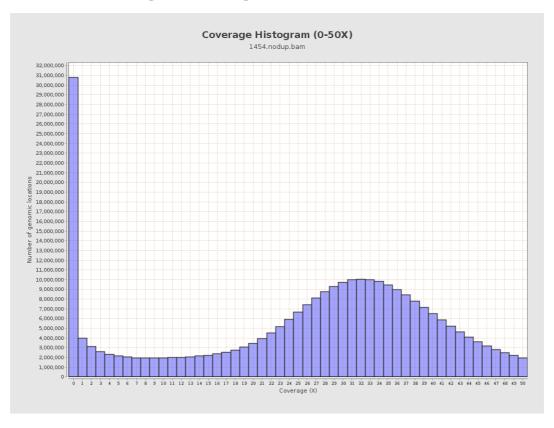


## 4. Results: Coverage Histogram



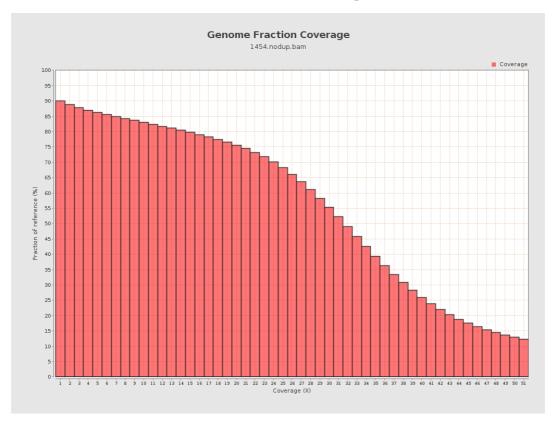


## 5. Results: Coverage Histogram (0-50X)



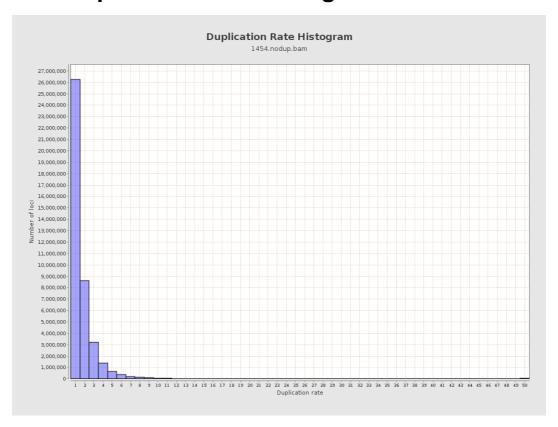


## 6. Results : Genome Fraction Coverage



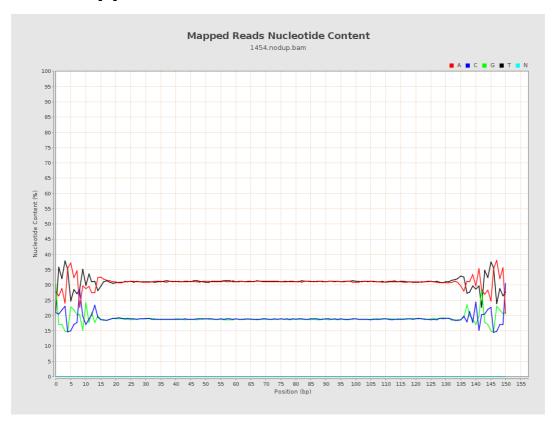


## 7. Results: Duplication Rate Histogram



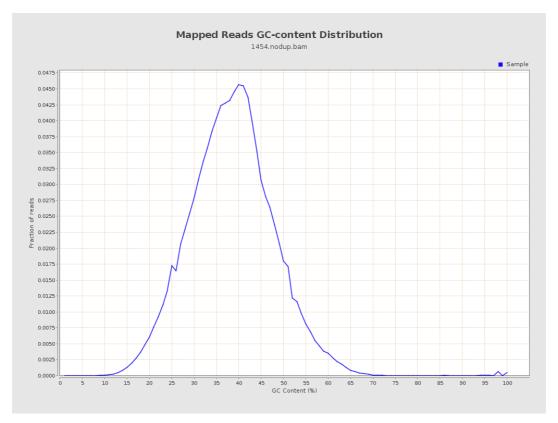


## 8. Results: Mapped Reads Nucleotide Content



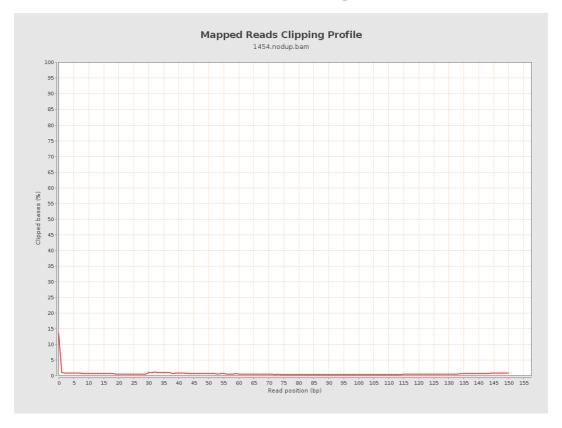


## 9. Results: Mapped Reads GC-content Distribution



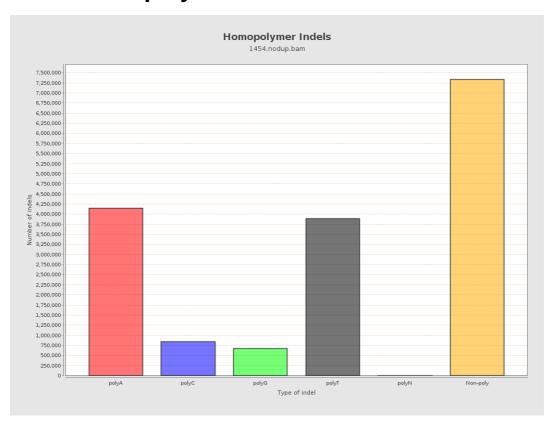


## 10. Results: Mapped Reads Clipping Profile





## 11. Results: Homopolymer Indels



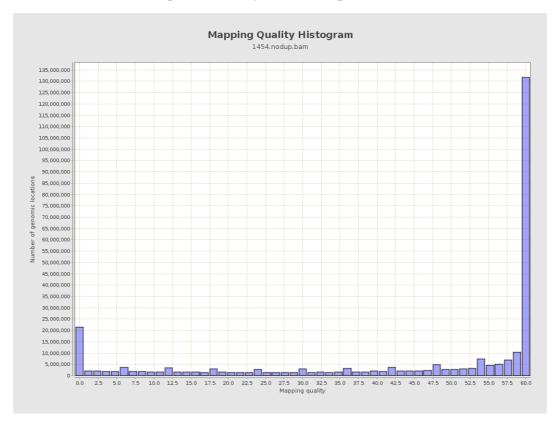


# 12. Results: Mapping Quality Across Reference



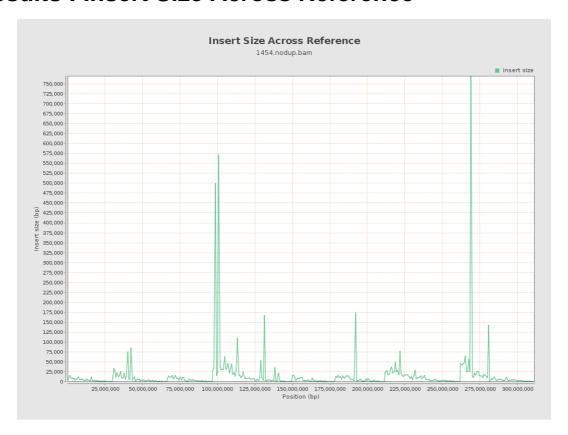


## 13. Results: Mapping Quality Histogram





#### 14. Results: Insert Size Across Reference





## 15. Results: Insert Size Histogram

