# Qualimap Analysis Results

BAM QC analysis Generated by Qualimap v.2.2.1 2023/05/29 21:32:46



#### 1. Input data & parameters

#### 1.1. QualiMap command line

qualimap bamqc -bam /proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/596 .nodup.bam -nw 400 -hm 3

#### 1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/worki ng/Kate/Working/F2/MarkDuplicates/ 596 .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:Illumina\tLB:LibA\t SM:\unit\tPL:Illumina\tLB:LibA\t SM:\unit\tPL:Illumina\tLB:LibA\t SM:\unit\sample /proj/uppstore2018210/Aalpina/data/r eference/GCA_900128785.1_MPIPZ. v5_genomic.fa /proj/uppstore2018210/Aalpina/data/r awdata/P26207/P26207_488/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_488_S463_L004 _R1_001.fastq.gz /proj/uppstore2018210/Aalpina/data/r awdata/P26207/P26207_488/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_488_S463_L004 _R2_001.fastq.gz
Size of a homopolymer:	3
Number of windows:	400



Analysis date:	Mon May 29 21:32:46 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no



## 2. Summary

#### 2.1. Globals

Reference size	311,642,060
Number of reads	72,885,411
Mapped reads	66,501,996 / 91.24%
Unmapped reads	6,383,415 / 8.76%
Mapped paired reads	66,501,996 / 91.24%
Mapped reads, first in pair	33,299,449 / 45.69%
Mapped reads, second in pair	33,202,547 / 45.55%
Mapped reads, both in pair	64,428,785 / 88.4%
Mapped reads, singletons	2,073,211 / 2.84%
Read min/max/mean length	30 / 151 / 148.21
Duplicated reads (flagged)	12,027,472 / 16.5%
Clipped reads	15,066,650 / 20.67%

#### 2.2. ACGT Content

Number/percentage of A's	2,828,646,702 / 30.87%		
Number/percentage of C's	1,752,371,306 / 19.12%		
Number/percentage of T's	2,833,341,166 / 30.92%		
Number/percentage of G's	1,749,320,192 / 19.09%		
Number/percentage of N's	30,417 / 0%		
GC Percentage	38.21%		

#### 2.3. Coverage



Mean	29.4796
Standard Deviation	279.2221

## 2.4. Mapping Quality

Mean Mapping Quality	44.27

#### 2.5. Insert size

Mean	255,038.07	
Standard Deviation	2,431,669.4	
P25/Median/P75	331 / 431 / 564	

#### 2.6. Mismatches and indels

General error rate	2.3%
Mismatches	192,707,050
Insertions	6,598,749
Mapped reads with at least one insertion	8.84%
Deletions	6,289,082
Mapped reads with at least one deletion	8.4%
Homopolymer indels	57.37%

#### 2.7. Chromosome stats

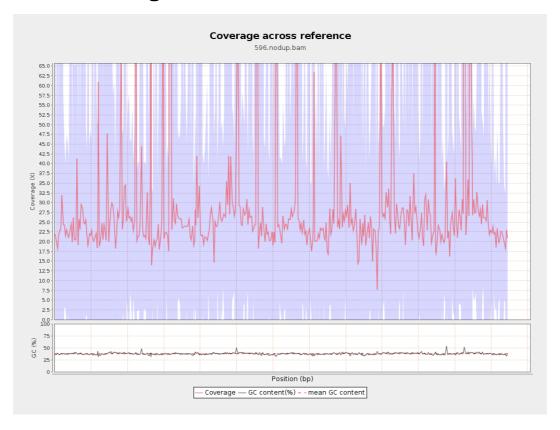
Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	691448746	23.262	107.049



LT669789.1	36598175	1085818259	29.6686	306.578
LT669790.1	30422129	1058118711	34.7812	391.7069
LT669791.1	52758100	1547489569	29.3318	298.8683
LT669792.1	28376109	822617355	28.9898	292.8588
LT669793.1	33388210	898854321	26.9213	185.2418
LT669794.1	50579949	1415795228	27.9912	223.563
LT669795.1	49795044	1666931554	33.4759	317.6964

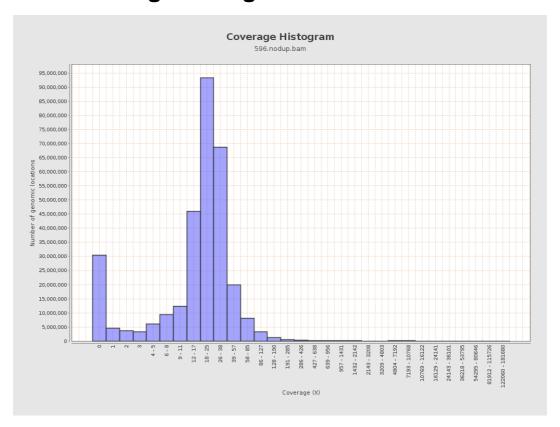


## 3. Results: Coverage across reference



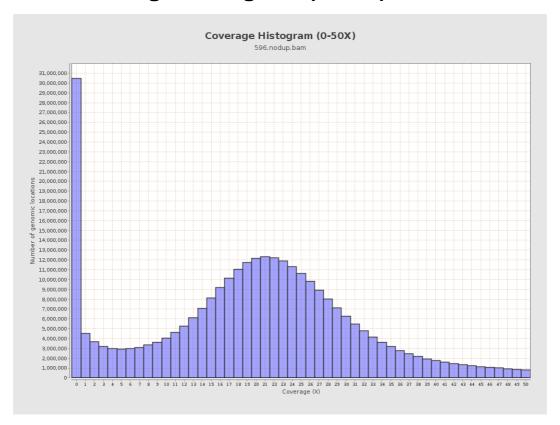


## 4. Results : Coverage Histogram



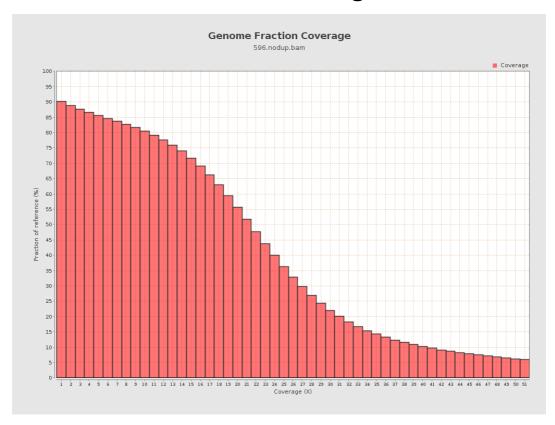


## 5. Results: Coverage Histogram (0-50X)



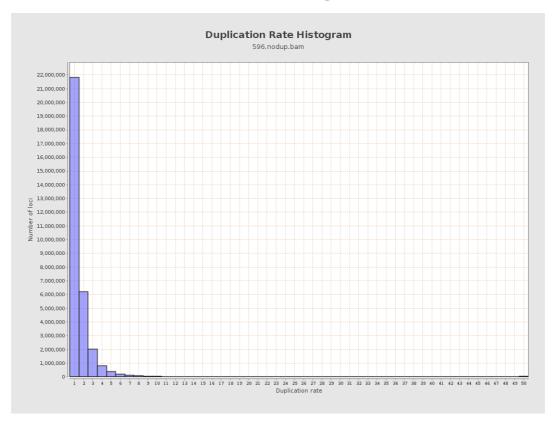


## 6. Results : Genome Fraction Coverage



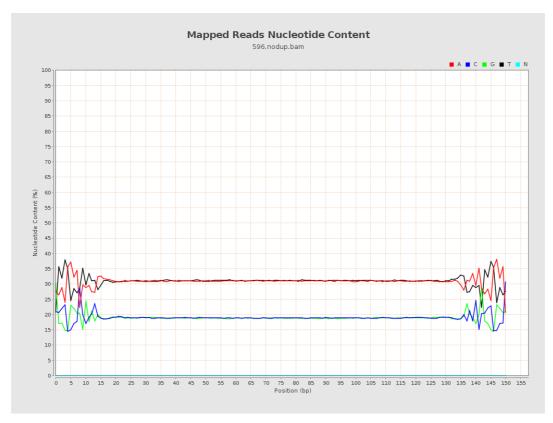


## 7. Results: Duplication Rate Histogram



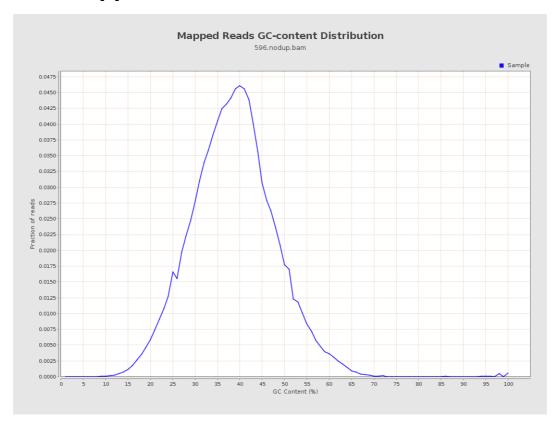


## 8. Results: Mapped Reads Nucleotide Content



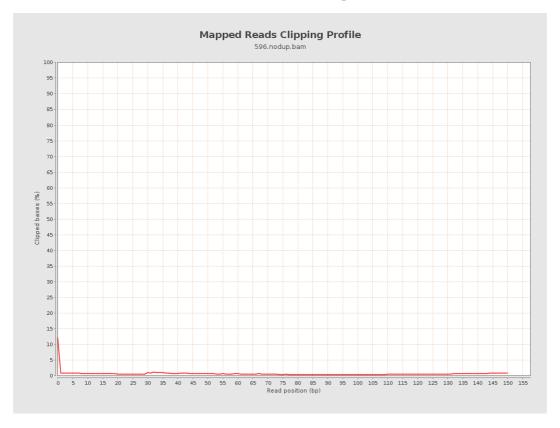


## 9. Results: Mapped Reads GC-content Distribution



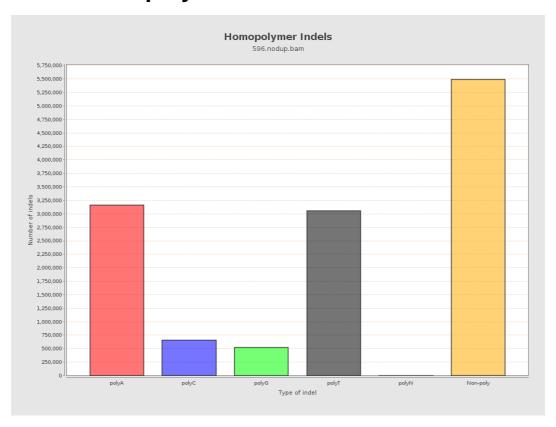


## 10. Results: Mapped Reads Clipping Profile



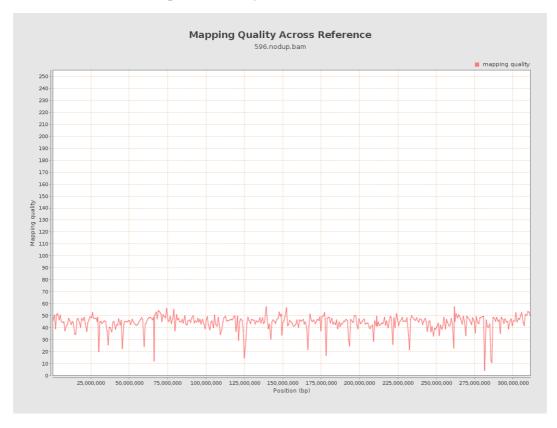


# 11. Results : Homopolymer Indels



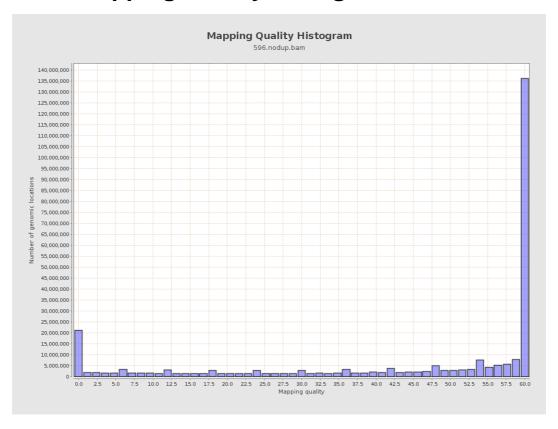


# 12. Results: Mapping Quality Across Reference



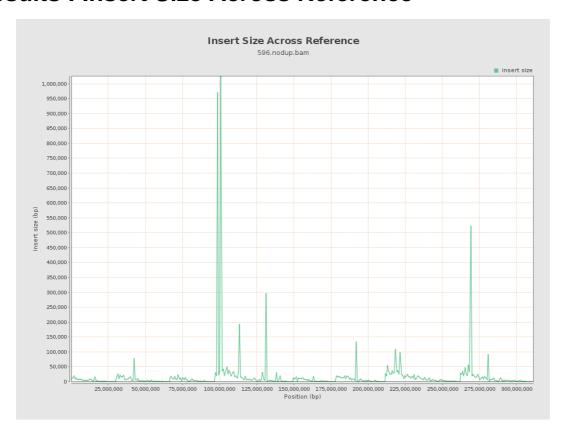


## 13. Results: Mapping Quality Histogram





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





## 15. Results: Insert Size Histogram

