# Qualimap Analysis Results

BAM QC analysis Generated by Qualimap v.2.2.1 2023/05/29 21:33:41



#### 1. Input data & parameters

#### 1.1. QualiMap command line

qualimap bamqc -bam /proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/146 7 .nodup.bam -nw 400 -hm 3

#### 1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/worki ng/Kate/Working/F2/MarkDuplicates/ 1467 .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_109/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_109_S199_L002 _R1_001.fastq.gz /proj/uppstore2018210/Aalpina/data/r awdata/P26207/P26207_109/02- FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_109_S199_L002 _R2_001.fastq.gz
Size of a homopolymer:	3



Number of windows:	400
Analysis date:	Mon May 29 21:33:40 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no



## 2. Summary

#### 2.1. Globals

Reference size	311,642,060
Number of reads	75,555,748
Mapped reads	68,847,751 / 91.12%
Unmapped reads	6,707,997 / 8.88%
Mapped paired reads	68,847,751 / 91.12%
Mapped reads, first in pair	34,482,653 / 45.64%
Mapped reads, second in pair	34,365,098 / 45.48%
Mapped reads, both in pair	66,826,331 / 88.45%
Mapped reads, singletons	2,021,420 / 2.68%
Read min/max/mean length	30 / 151 / 148.09
Duplicated reads (flagged)	12,913,762 / 17.09%
Clipped reads	16,093,506 / 21.3%

#### 2.2. ACGT Content

Number/percentage of A's	2,932,011,393 / 30.97%
Number/percentage of C's	1,801,513,394 / 19.03%
Number/percentage of T's	2,933,654,686 / 30.99%
Number/percentage of G's	1,799,897,832 / 19.01%
Number/percentage of N's	39,441 / 0%
GC Percentage	38.04%

#### 2.3. Coverage



Mean	30.4598
Standard Deviation	257.3336

## 2.4. Mapping Quality

Mean Mapping Quality	44.07

#### 2.5. Insert size

Mean	254,205.88	
Standard Deviation	2,417,531.12	
P25/Median/P75	317 / 412 / 531	

#### 2.6. Mismatches and indels

General error rate	2.38%
Mismatches	205,698,622
Insertions	6,926,700
Mapped reads with at least one insertion	8.98%
Deletions	6,772,329
Mapped reads with at least one deletion	8.71%
Homopolymer indels	57.28%

#### 2.7. Chromosome stats

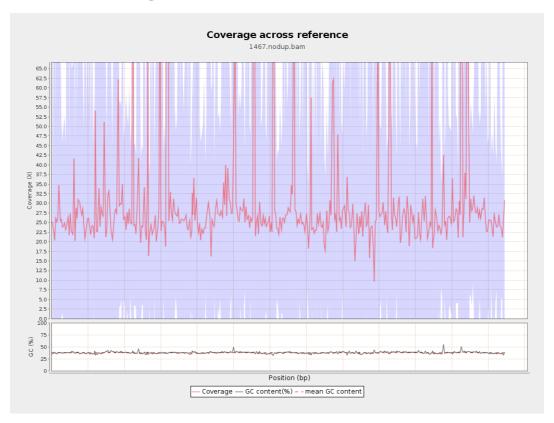
Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	760157466	25.5736	100.9821



LT669789.1	36598175	1134534772	30.9998	279.8638
LT669790.1	30422129	1106038466	36.3564	370.129
LT669791.1	52758100	1586011229	30.0619	274.4363
LT669792.1	28376109	867083139	30.5568	268.5849
LT669793.1	33388210	927881161	27.7907	167.4511
LT669794.1	50579949	1436949396	28.4095	219.9632
LT669795.1	49795044	1673901559	33.6158	279.6136

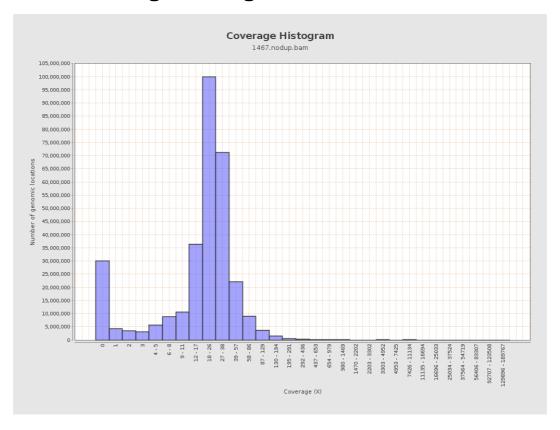


## 3. Results: Coverage across reference



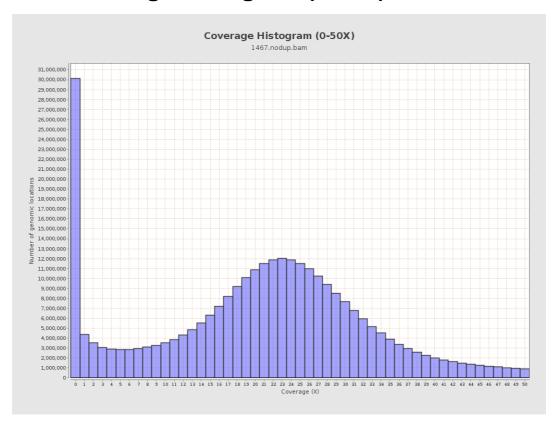


## 4. Results: Coverage Histogram



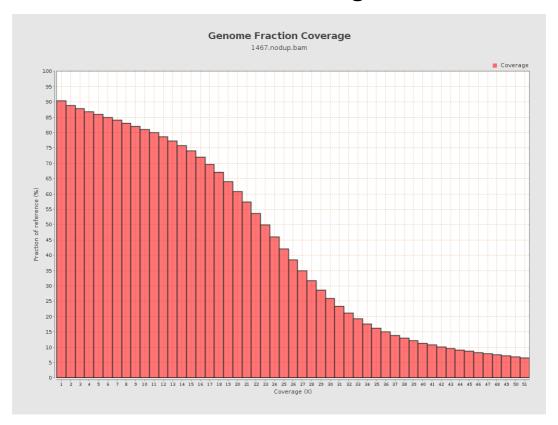


## 5. Results: Coverage Histogram (0-50X)



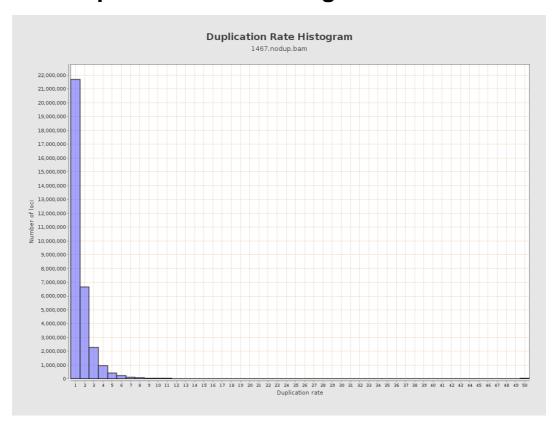


## 6. Results : Genome Fraction Coverage





## 7. Results: Duplication Rate Histogram



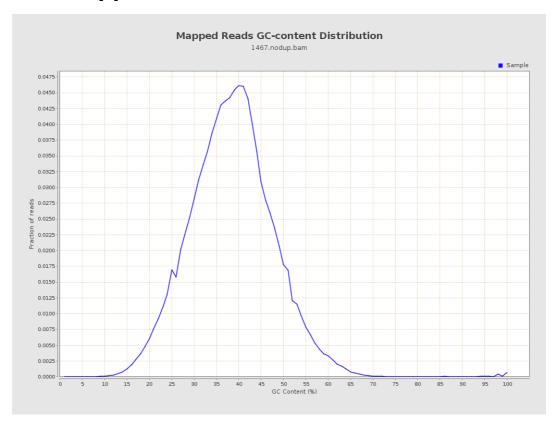


## 8. Results: Mapped Reads Nucleotide Content



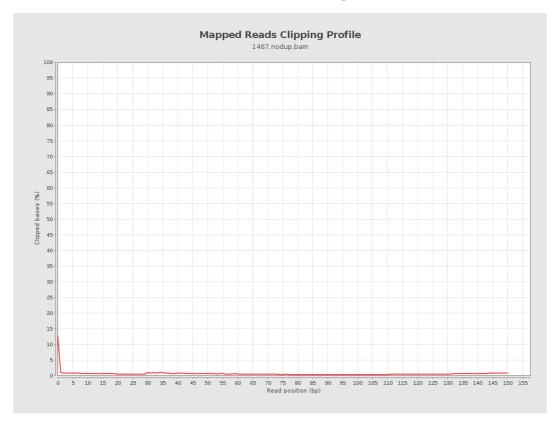


## 9. Results: Mapped Reads GC-content Distribution



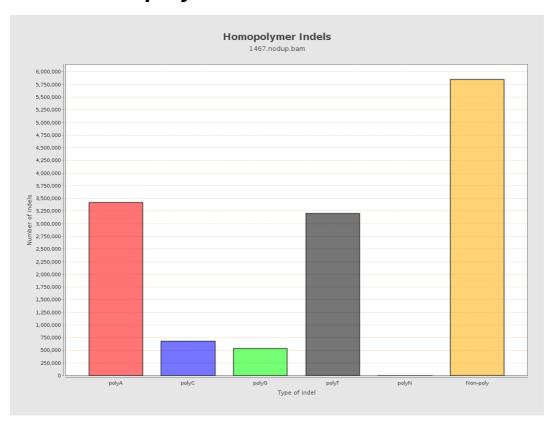


## 10. Results: Mapped Reads Clipping Profile



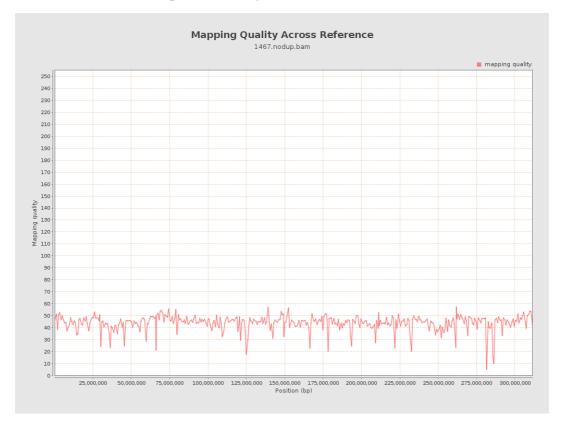


# 11. Results : Homopolymer Indels



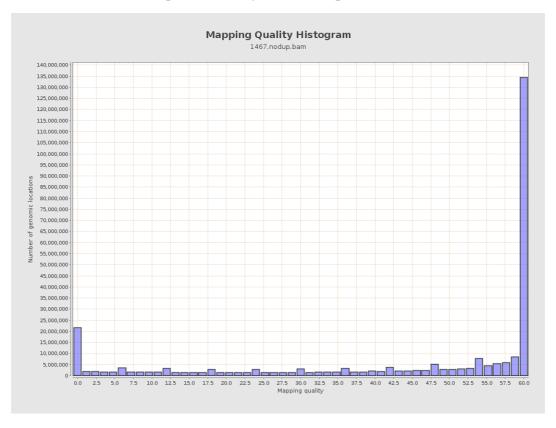


# 12. Results: Mapping Quality Across Reference



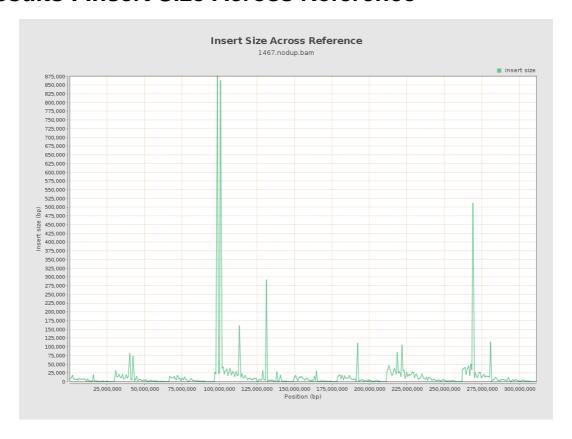


## 13. Results: Mapping Quality Histogram





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





## 15. Results: Insert Size Histogram

