

# Qualimap Analysis Results

*BAM QC analysis*

*Generated by Qualimap v.2.2.1*

*2023/05/29 21:28:48*

# 1. Input data & parameters

## 1.1. QualiMap command line

```
qualimap bamqc -bam
/proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/531
.nodup.bam -nw 400 -hm 3
```

## 1.2. Alignment

BAM file:	/proj/uppstore2018210/Aalpina/working/Kate/Working/F2/MarkDuplicates/531.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\tSM:\$sample /proj/uppstore2018210/Aalpina/data/reference/GCA_900128785.1_MPIPZ.v5_genomic.fa /proj/uppstore2018210/Aalpina/data/awdata/P26207/P26207_218/02-FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_218_S299_L003_R1_001.fastq.gz /proj/uppstore2018210/Aalpina/data/awdata/P26207/P26207_218/02-FASTQ/220902_A00621_0737_BHM GCVDSX3/P26207_218_S299_L003_R2_001.fastq.gz
Size of a homopolymer:	3
Number of windows:	400

Analysis date:	Mon May 29 21:28:48 CEST 2023
Draw chromosome limits:	no
Skip duplicate alignments:	no

## 2. Summary

### 2.1. Globals

Reference size	311,642,060
Number of reads	60,366,934
Mapped reads	55,809,140 / 92.45%
Unmapped reads	4,557,794 / 7.55%
Mapped paired reads	55,809,140 / 92.45%
Mapped reads, first in pair	27,971,014 / 46.33%
Mapped reads, second in pair	27,838,126 / 46.11%
Mapped reads, both in pair	54,326,256 / 89.99%
Mapped reads, singletons	1,482,884 / 2.46%
Read min/max/mean length	30 / 151 / 148.17
Duplicated reads (flagged)	8,335,015 / 13.81%
Clipped reads	12,872,610 / 21.32%

### 2.2. ACGT Content

Number/percentage of A's	2,377,461,461 / 30.9%
Number/percentage of C's	1,469,553,013 / 19.1%
Number/percentage of T's	2,378,355,131 / 30.92%
Number/percentage of G's	1,467,714,723 / 19.08%
Number/percentage of N's	28,732 / 0%
GC Percentage	38.18%

### 2.3. Coverage

Mean	24.7478
Standard Deviation	216.5484

## 2.4. Mapping Quality

Mean Mapping Quality	44.61
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## 2.5. Insert size

Mean	236,222.79
Standard Deviation	2,327,801.04
P25/Median/P75	317 / 416 / 542

## 2.6. Mismatches and indels

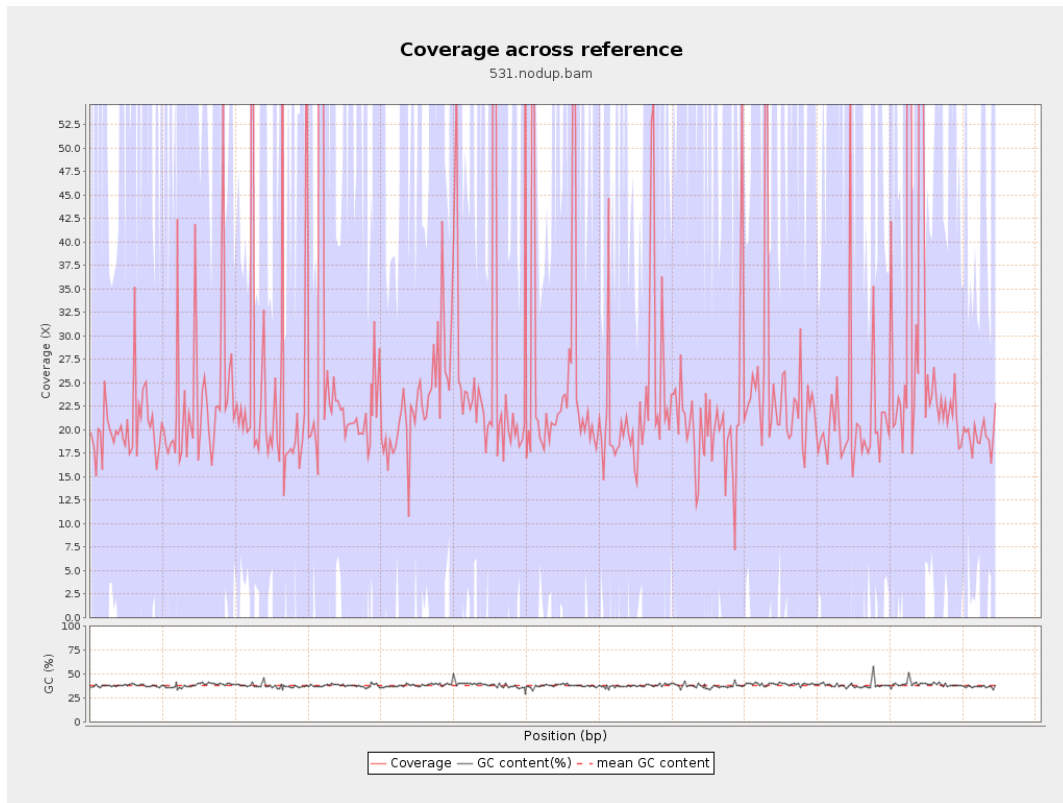
General error rate	2.3%
Mismatches	162,284,219
Insertions	5,319,019
Mapped reads with at least one insertion	8.54%
Deletions	5,173,388
Mapped reads with at least one deletion	8.25%
Homopolymer indels	57.14%

## 2.7. Chromosome stats

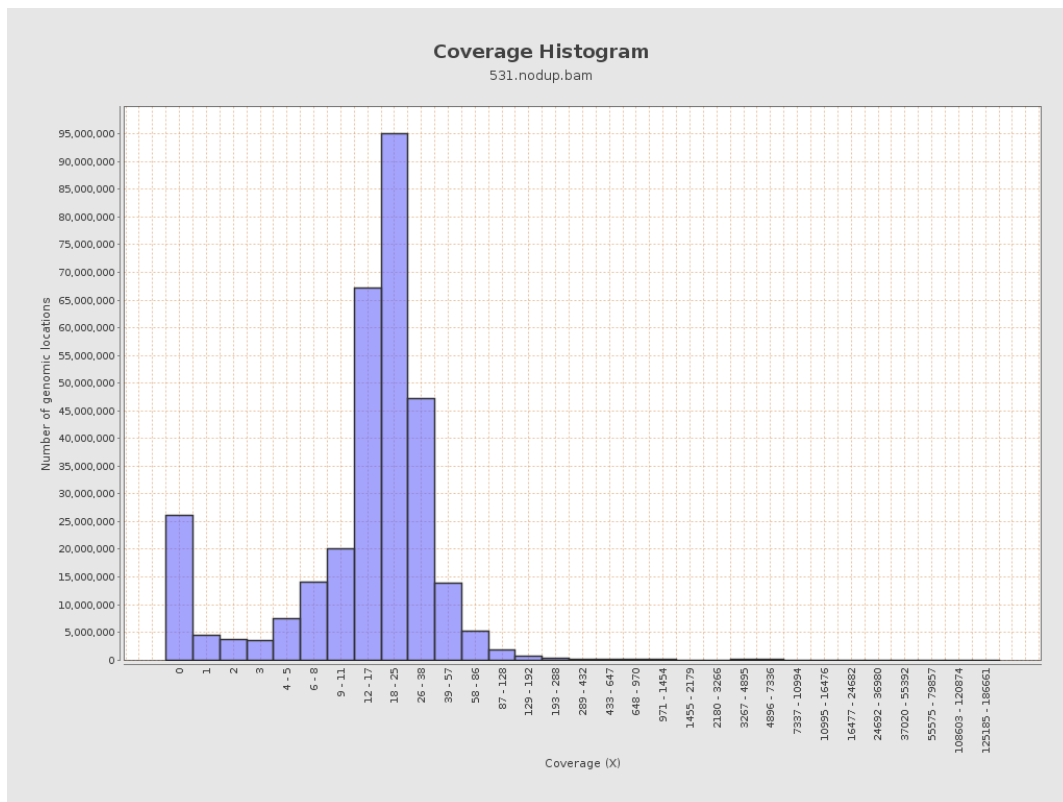
Name	Length	Mapped bases	Mean coverage	Standard deviation
LT669788.1	29724344	594574884	20.003	75.3365

LT669789.1	36598175	901763053	24.6396	215.3242
LT669790.1	30422129	876589121	28.8142	280.0244
LT669791.1	52758100	1298222226	24.6071	213.9739
LT669792.1	28376109	700117807	24.6728	231.1224
LT669793.1	33388210	745738655	22.3354	143.0172
LT669794.1	50579949	1164958199	23.032	171.1817
LT669795.1	49795044	1430498308	28.7277	292.6331

### 3. Results : Coverage across reference

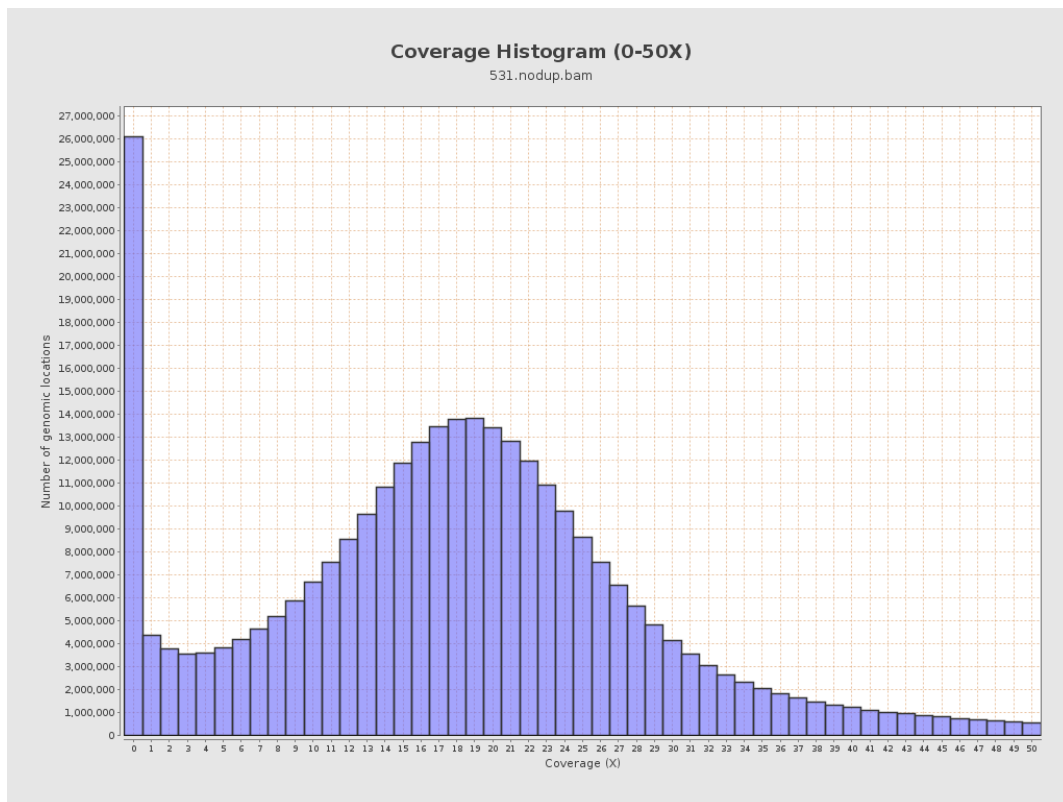


## 4. Results : Coverage Histogram

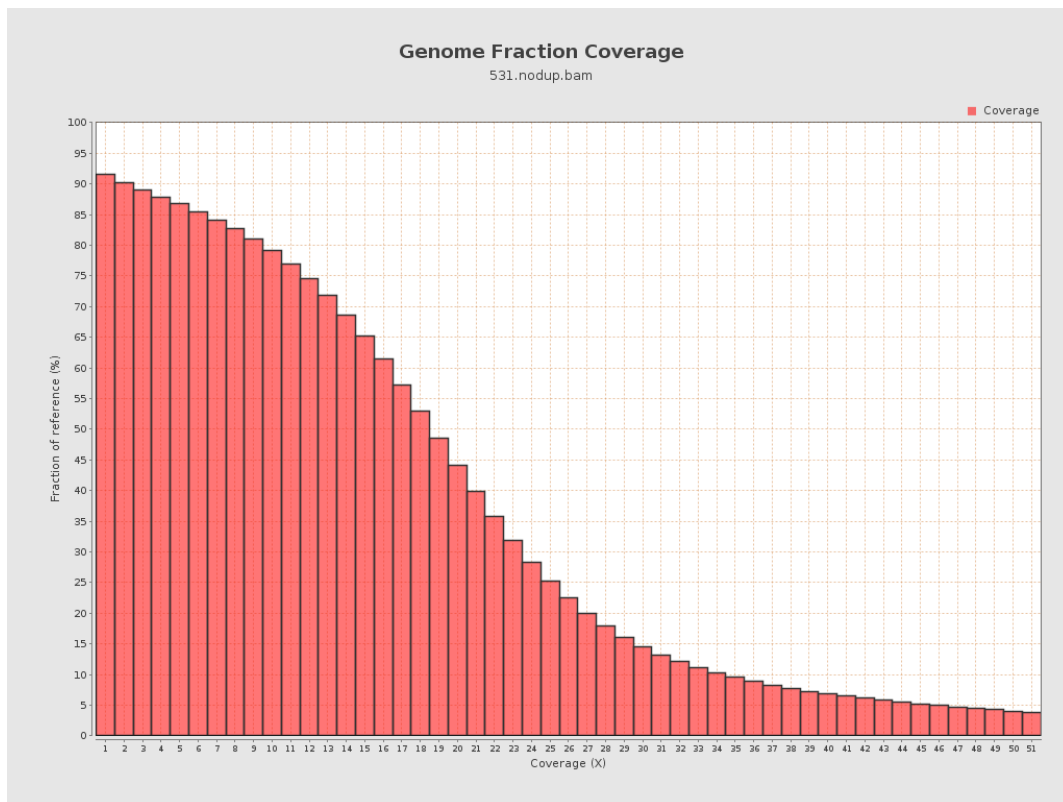




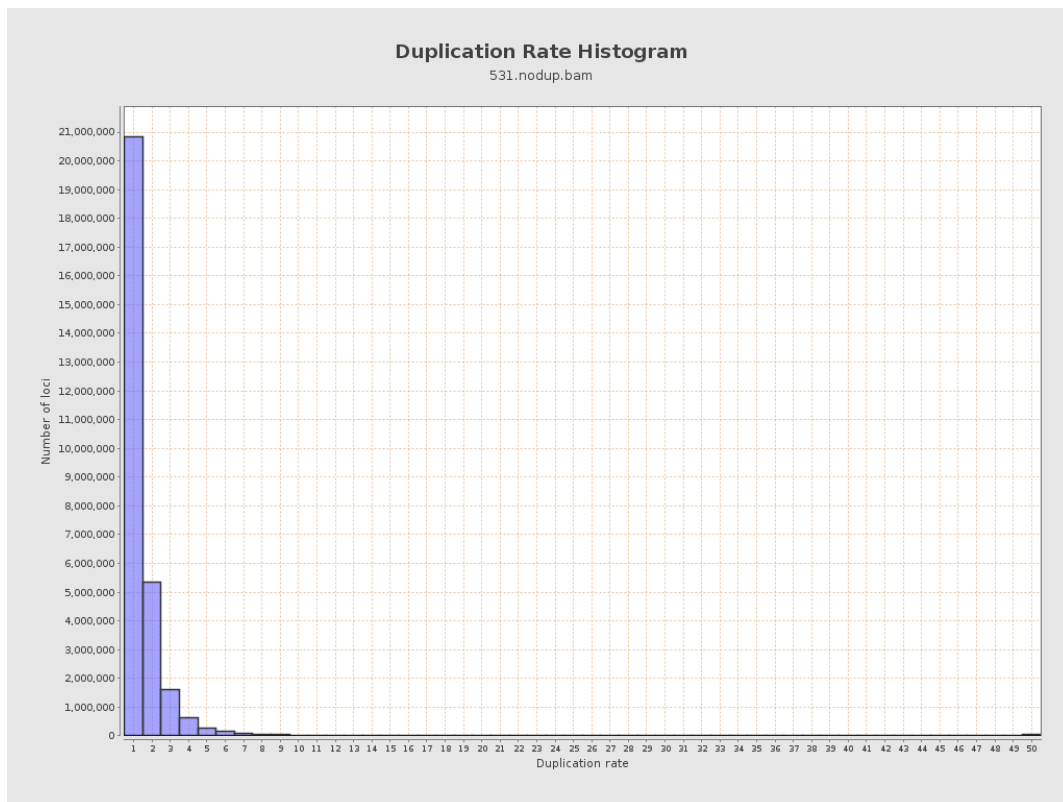
## 5. Results : Coverage Histogram (0-50X)



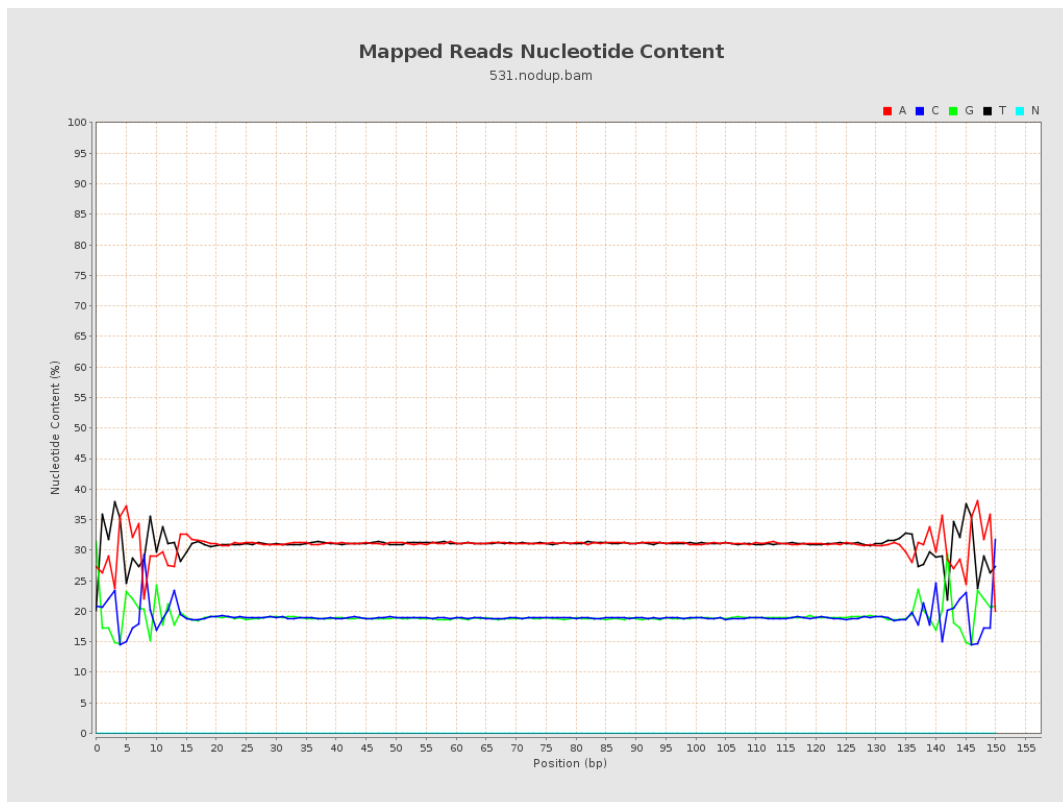
## 6. Results : Genome Fraction Coverage



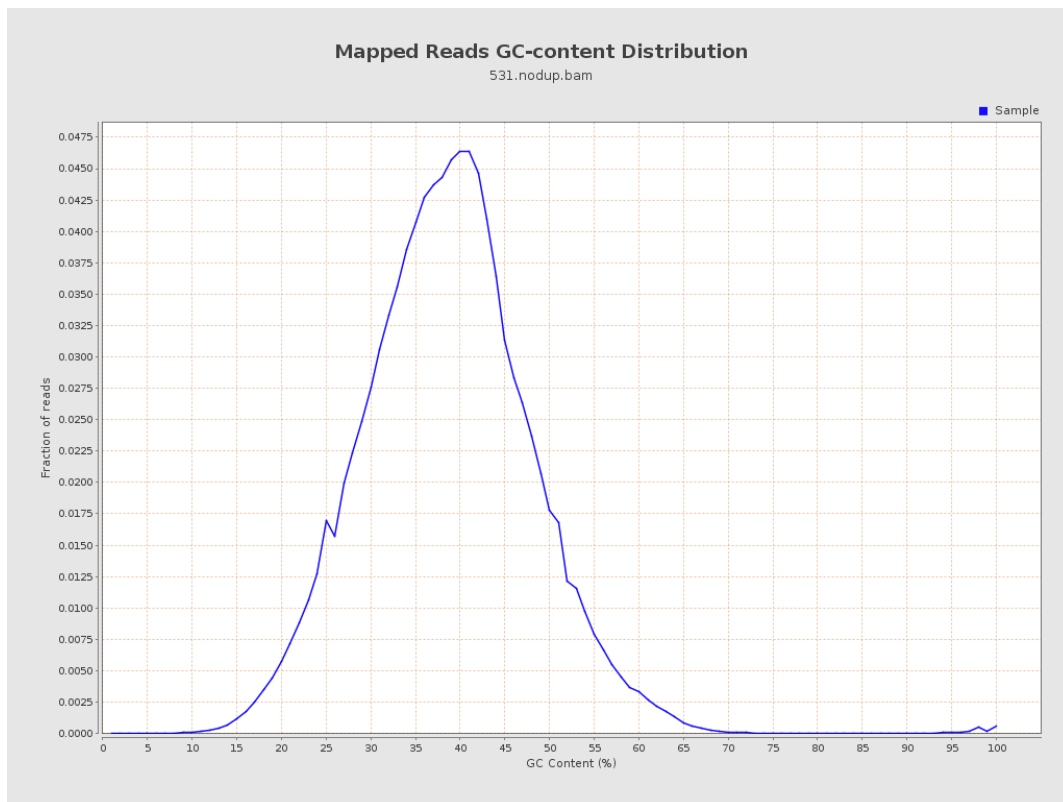
## 7. Results : Duplication Rate Histogram



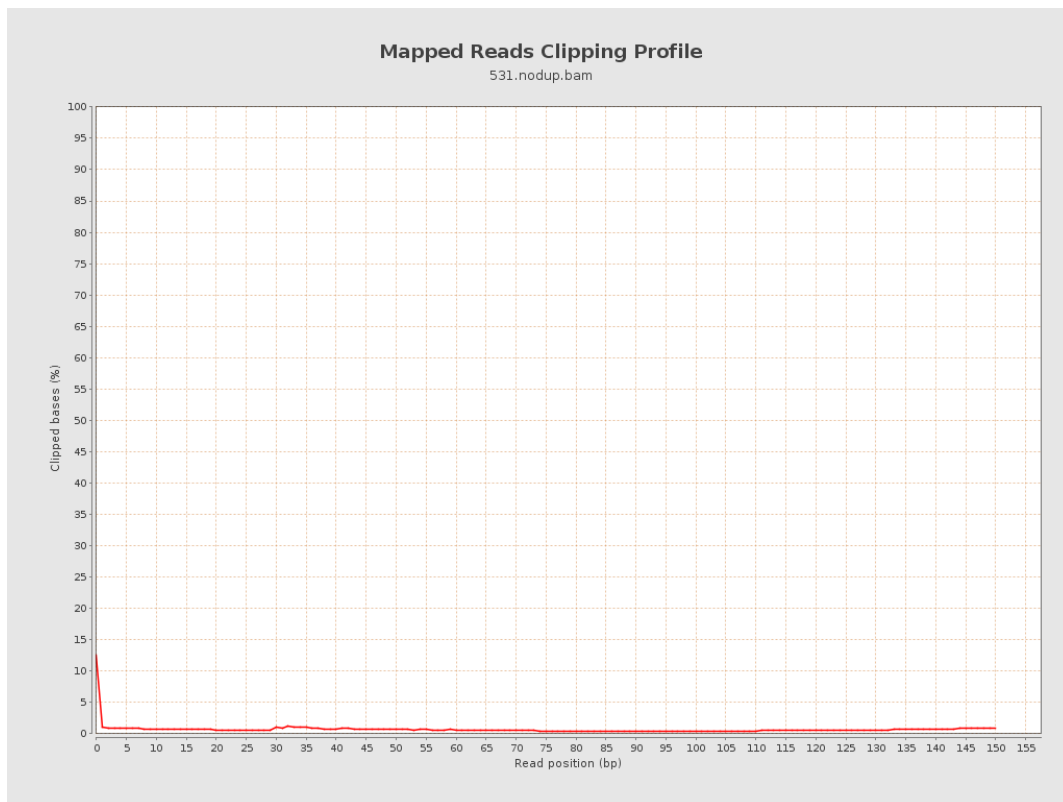
## 8. Results : Mapped Reads Nucleotide Content



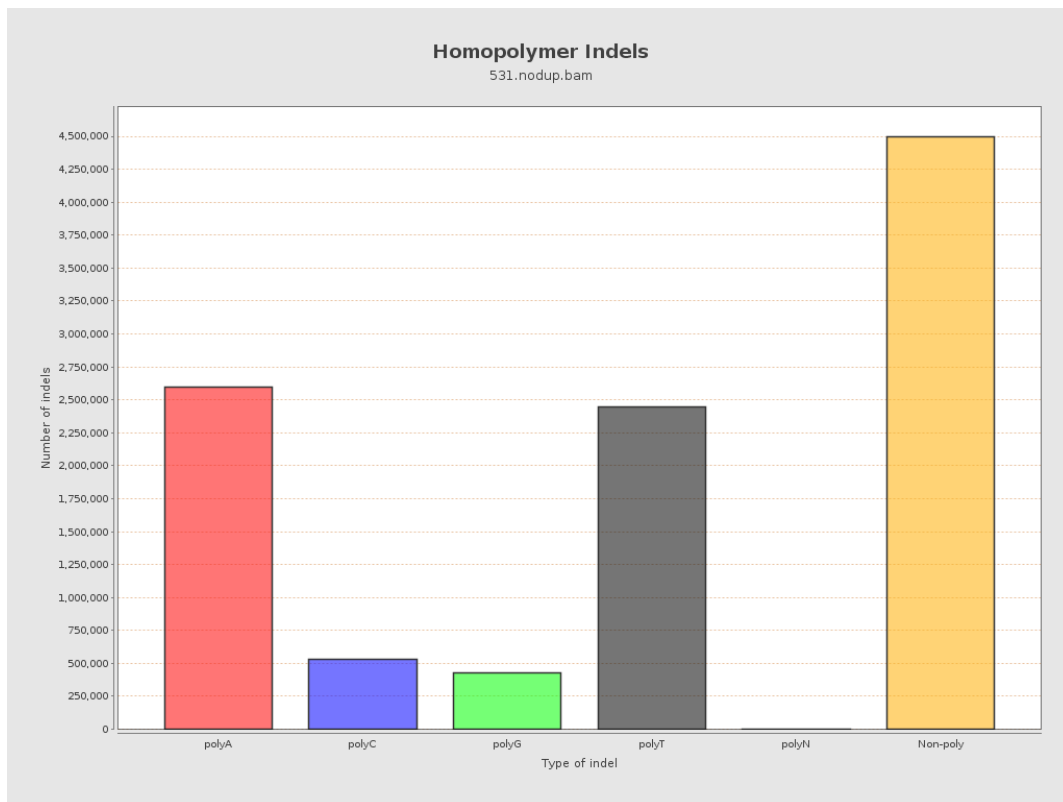
## 9. Results : Mapped Reads GC-content Distribution



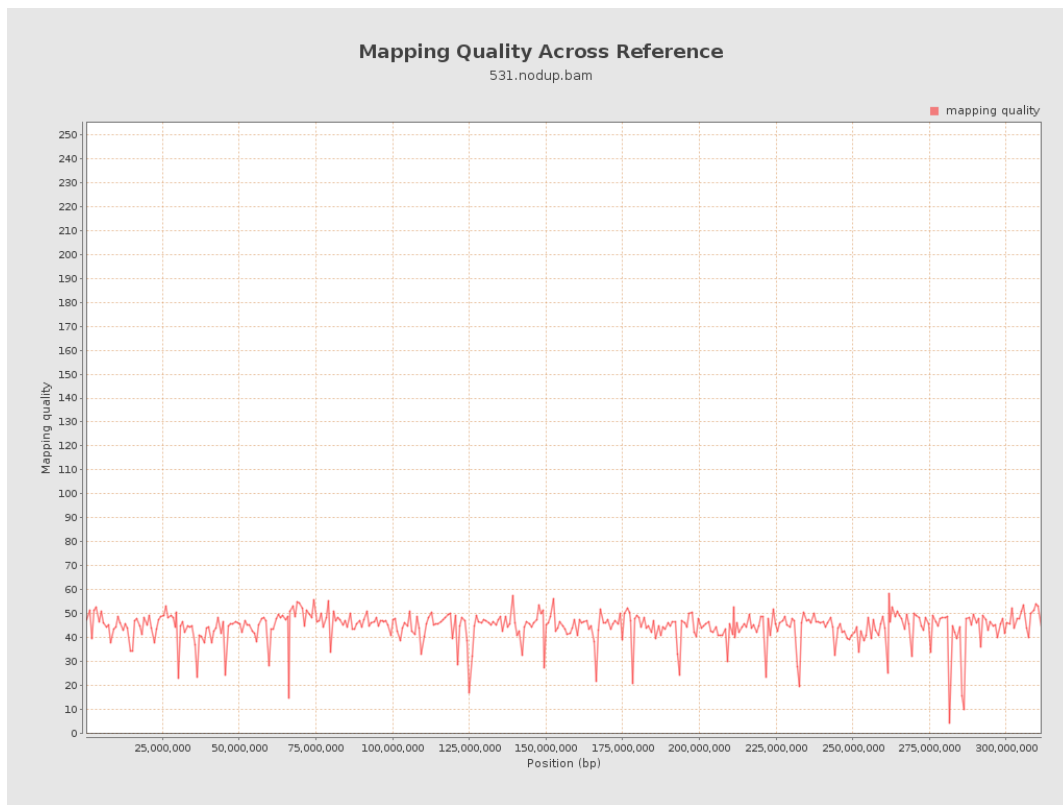
## 10. Results : Mapped Reads Clipping Profile



## 11. Results : Homopolymer Indels

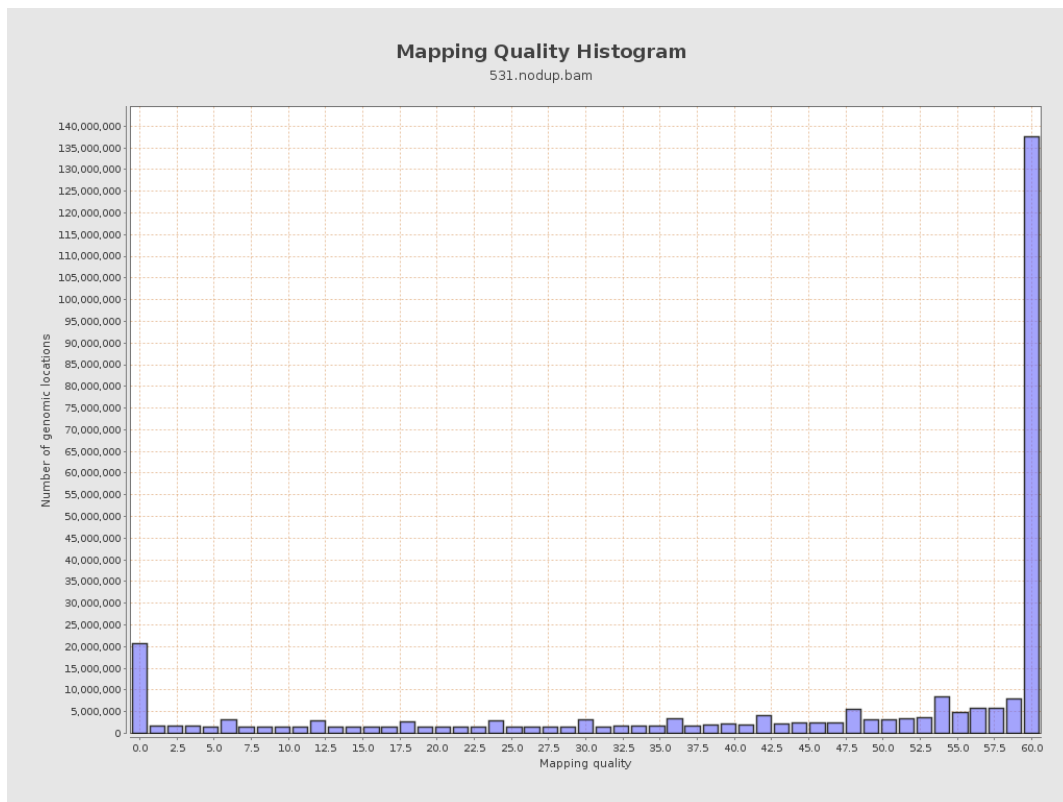


## 12. Results : Mapping Quality Across Reference

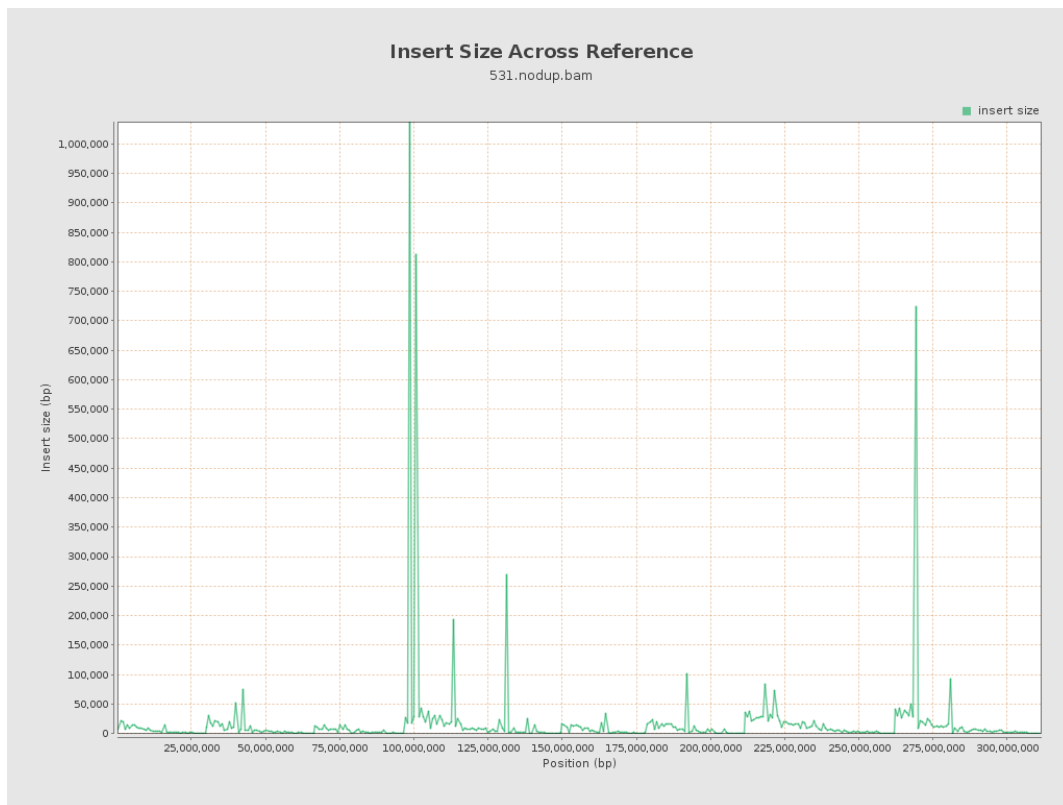




## 13. Results : Mapping Quality Histogram



## 14. Results : Insert Size Across Reference



## 15. Results : Insert Size Histogram

