

# SnpEff: Variant analysis

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## Summary

Genome	pppncal39
Date	2012-09-03 21:28
SnpEff version	SnpEff 3.0c (build 2012-07-19), by Pablo Cingolani
Command line arguments	SnpEff pppncal39 -i /VCF /Volumes/USB_DISK/VCF4/GR_VCF -r /Volumes/USB_DISK/VCF4/GR_mask -v
Warnings	0
Number of lines (input file)	633,812
Number of variants (before filter)	639,062
Filter	
Number of variants filtered out	0
Number of not variants (i.e. reference equals alternative)	0
Number of variants processed (i.e. after filter and non-variants)	639,062
Number of known variants (i.e. non-empty ID)	0 (0%)
Number of effects	1,028,718
Genome total length	227,252,108
Genome effective length	227,156,785
Change rate	1 change every 355 bases

## Change rate details

Chromosome	Length	Changes	Change rate
1	45,877,626	101,258	462
2	28,807,724	113,949	238
3	22,028,550	69,178	366
4	36,528,727	92,257	336
5	18,502,877	49,814	370
6	28,902,262	81,503	354
7	22,795,183	56,588	402
8	21,829,783	48,275	482
Total	227,156,785	639,062	355

## Number changes by type

Type	Total	Ratio	Ratio
SNP	541,876	84.8%	84.8%
INS	0	0%	0%
DEL	20,535	3.2%	3.2%
MIXED	0	0%	0%
INDEL	0	0%	0%
Total	639,062	100.0%	100.0%

## Number of effects by impact

Type (alphabetical order)	Count	Percent
HIGH	2,007	0.20%
LOW	18,502	1.80%
MODERATE	35,028	3.41%
MODIFIER	952,881	92.59%

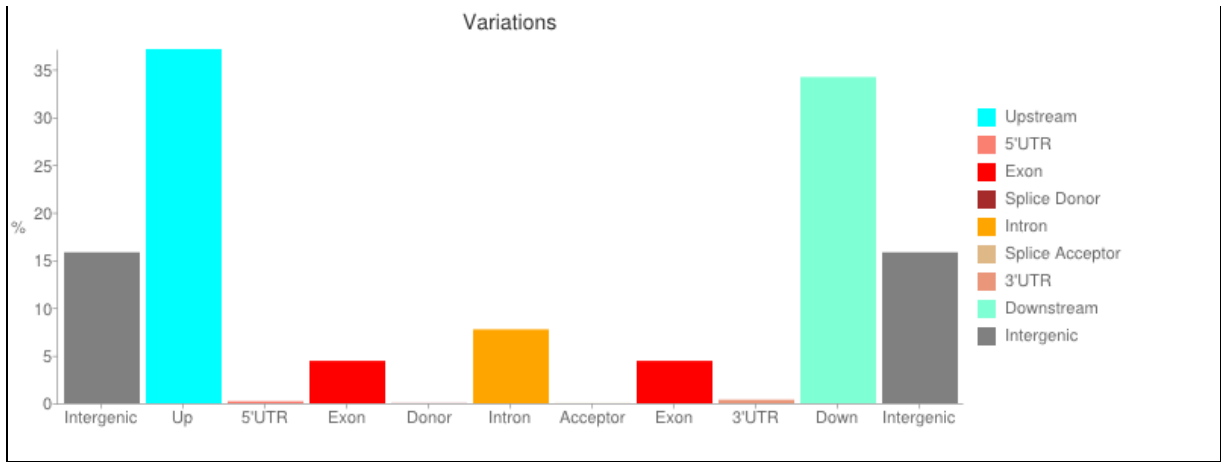
## Number of effects by functional class

Type (alphabetical order)	Count	Percent
MISSENSE	25,120	2.45%
NONSENSE	819	0.08%
SILENT	926,942	90.47%

Missense / Silent ratio: 1.4481

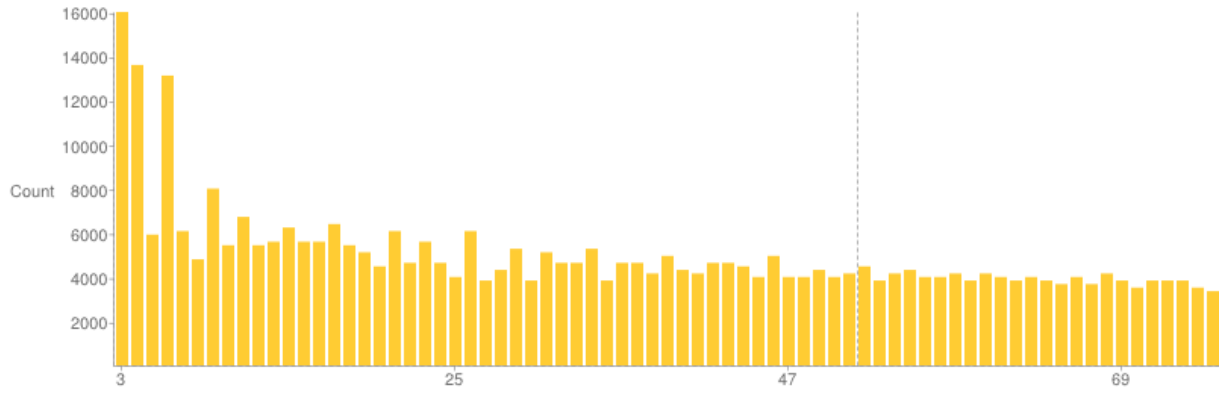
## Number of effects by type and region

Type (alphabetical order)	Count	Percent	Region (alphabetical order)	Count	Percent
CODON_CHANGE_PLUS_CODON_DELETION	98	0.01%	DOWNSTREAM	35,095	3.41%
CODON_CHANGE_PLUS_CODON_INSERTION	128	0.01%	EXON	45,048	4.40%
CODON_DELETION	143	0.01%	INTERGENIC	162,880	15.83%
CODON_INSERTION	98	0.01%	INTRON	79,877	7.76%
DOWNSTREAM	35,095	3.41%	INTRON	79,877	7.76%
FRAME_SHIFT	1,174	0.11%	SPLICE_SITE_ACCEPTOR	191	0.01%
INTERGENIC	162,880	15.83%	SPLICE_SITE_DONOR	191	0.01%
INTRON	79,877	7.76%	UPSTREAM	1,028,718	100.00%
NON_SYNONYMOUS_CODING	25,939	2.53%	UTR_3_PRIME	1,981	0.19%
NON_SYNONYMOUS_START	6	0.00%	UTR_5_PRIME	2,198	0.21%
SPLICE_SITE_ACCEPTOR	191	0.01%			
SPLICE_SITE_DONOR	191	0.01%			
START_GAINED	298	0.03%			
START_LOST	49	0.00%			
STOP_GAINED	638	0.06%			
STOP_LOST	75	0.00%			
SYNONYMOUS_CODING	17,742	1.72%			
SYNONYMOUS_STOP	28	0.00%			
UPSTREAM	1,028,718	100.00%			
UTR_3_PRIME	1,981	0.19%			
UTR_5_PRIME	2,198	0.21%			



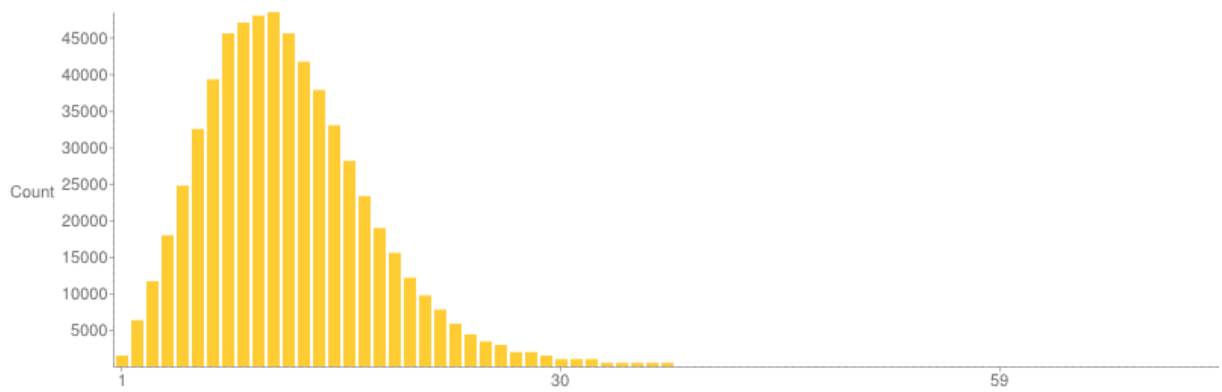
### Quality:

Min	3
Max	255
Mean	71.268
Median	61
Standard deviation	53.159



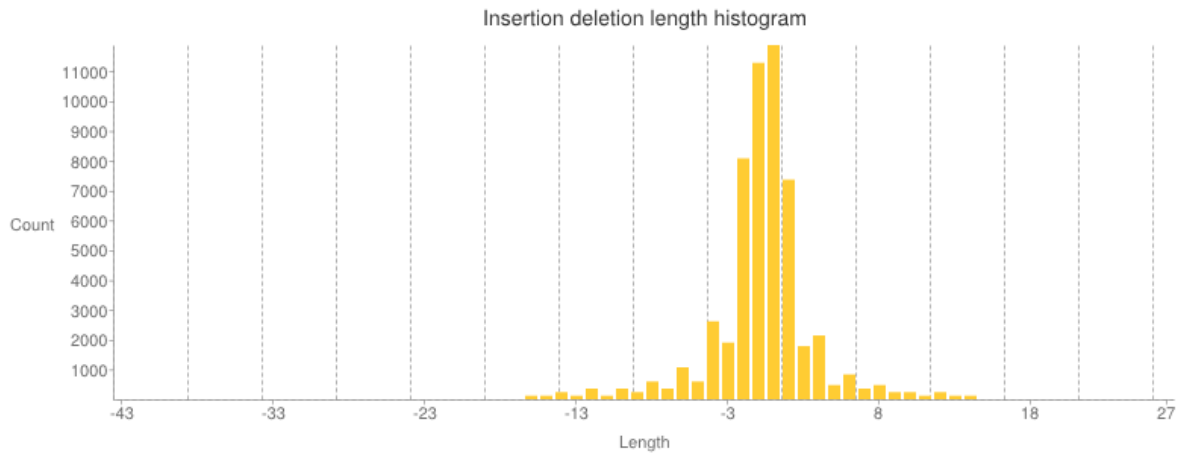
### Coverage:

Min	1
Max	1,763
Mean	12.855
Median	11
Standard deviation	11.535



### Insertions and deletions length:

Min	-43
Max	27
Mean	-0.459
Median	-1
Standard deviation	5.022



### Base changes (SNPs)

	A	C	G	T
A	0	374726	205356	205356
C	374726	0	205356	205356
G	205356	205356	0	374726
T	205356	205356	374726	0

### Ts/Tv (transitions / transversions)

**Note:** Only SNPs are used for this statistic.

**Note:** This Ts/Tv ratio is a 'raw' ratio. Some people prefer to use a ratio of rates, not observed events. In that case, you need to multiply by 2.0 (since there are twice as many possible transitions than transversions, E[Ts/Tv] ratio is twice the ratio of events).

Transitions	374726
Transversions	205356
Ts/Tv ratio	1.825

#### All variants:

```

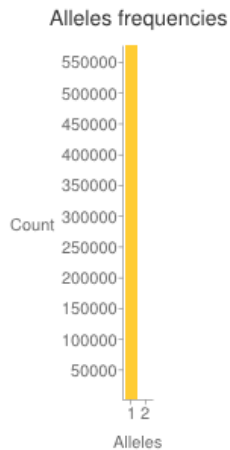
Sample      : /Users/macpro/GenBioPeach/Sequences_DD_F8_GB/BAMS/combined_GB.bam      Total
Transitions : 374726  374726
Transversions : 205356  205356
Ts/Tv      : 1.825  1.825
  
```

#### Only known variants (i.e. the ones having a non-empty ID field):

No results available (empty input?)

### Frequency of alleles

Note: Number of times an allele appears once (singleton), twice (doubletons), etc.



#### All variants:

Min	1
Max	2
Mean	1.023
Median	1

Standard deviation	0.00
Values	12
Count	8771601481

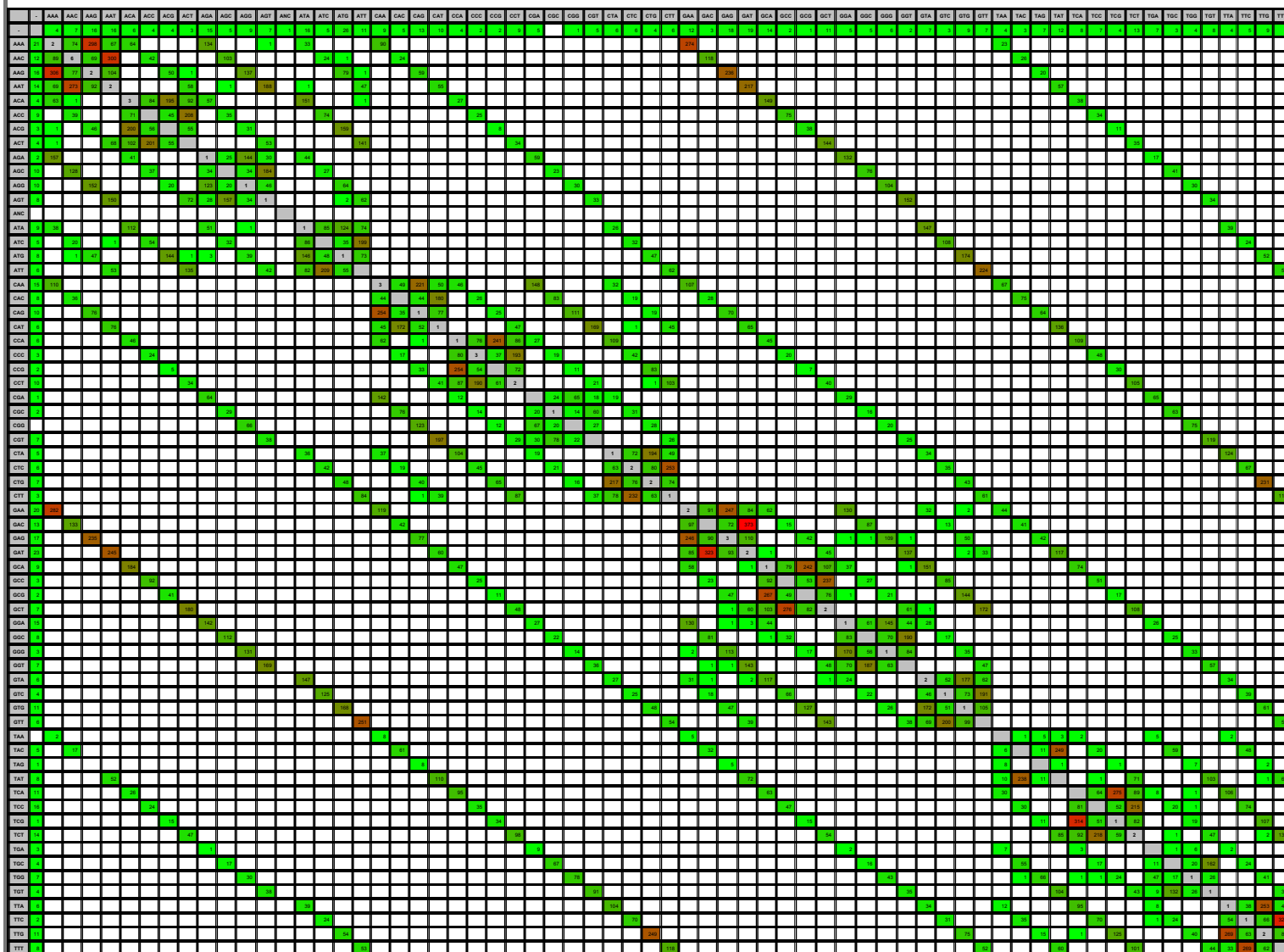
Only known variants (i.e. the ones having a non-empty ID field):

Min	2147483347
Max	2147483348
Mean	0
Median	0
Standard deviation	0
Values	
Count	

### Codon changes

How to read this table:

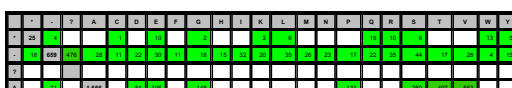
- Rows are reference codons and columns are changed codons. E.g. Row 'AAA' column 'TAA' indicates how many 'AAA' codons have been replaced by 'TAA' codons.
- Red background colors indicate that more changes happened (heat-map).
- Diagonals are indicated using grey background color
- WARNING: This table may include different translation codon tables (e.g. mamalian DNA and mitochondrial DNA).



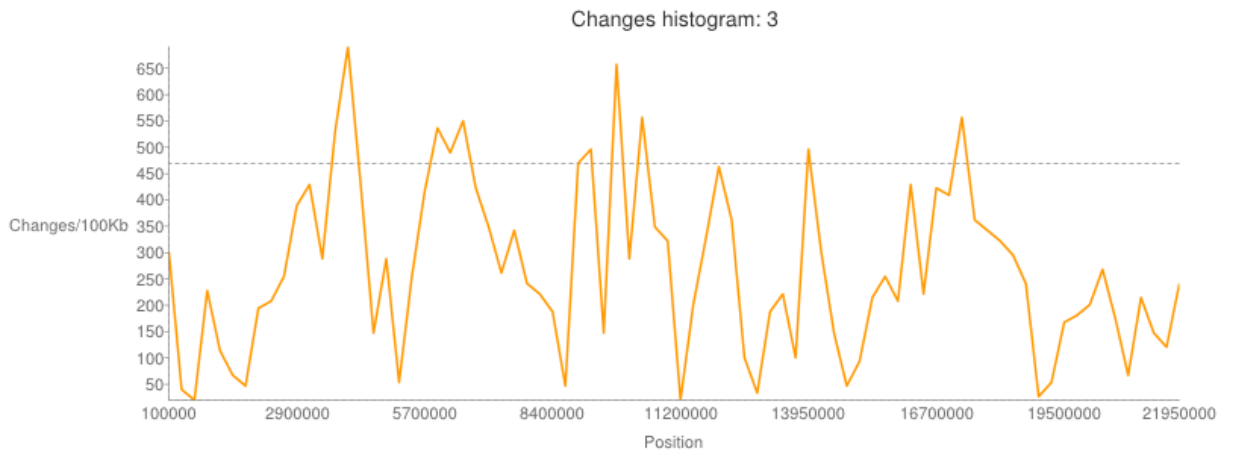
### Amino acid changes

How to read this table:

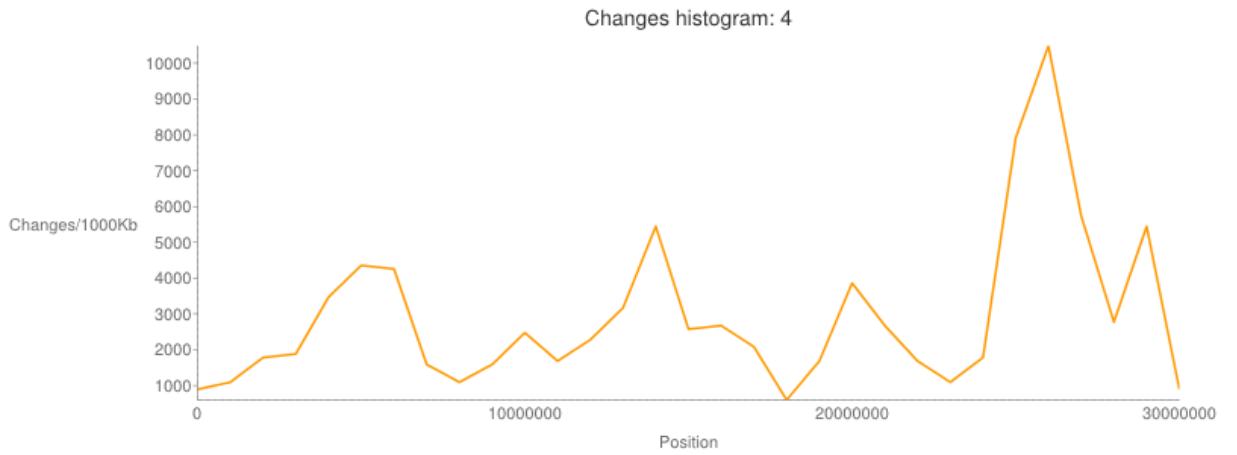
- Rows are reference amino acids and columns are changed amino acids. E.g. Row 'A' column 'E' indicates how many 'A' amino acids have been replaced by 'E' amino acids.
- Red background colors indicate that more changes happened (heat-map).
- Diagonals are indicated using grey background color
- WARNING: This table may include different translation codon tables (e.g. mamalian DNA and mitochondrial DNA).



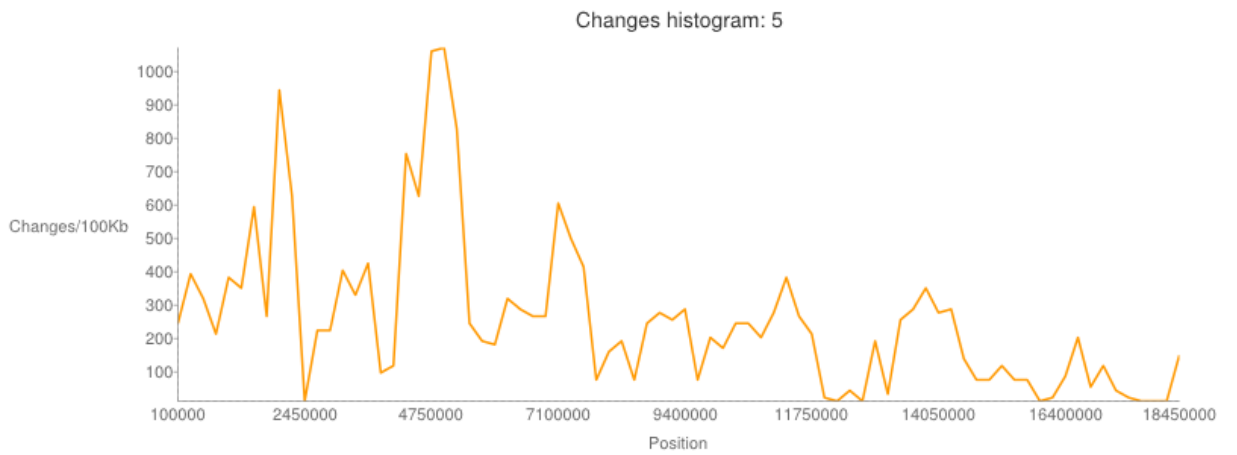




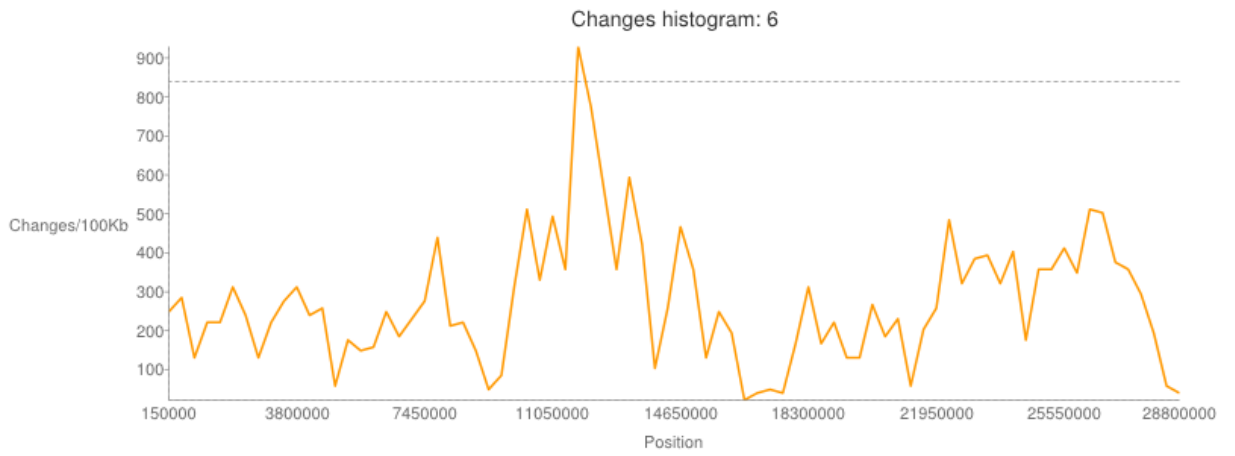
Chromosome: 3



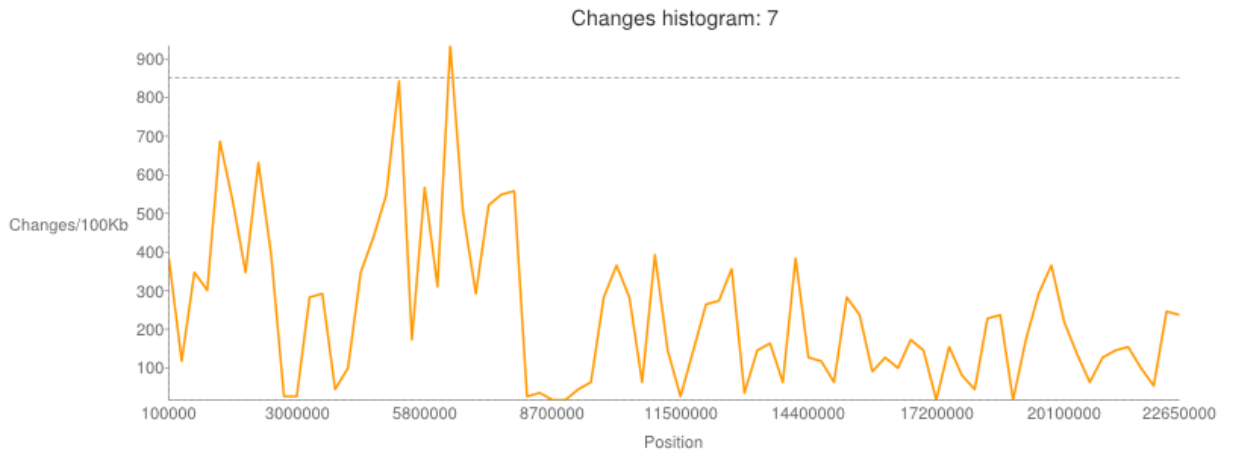
Chromosome: 4



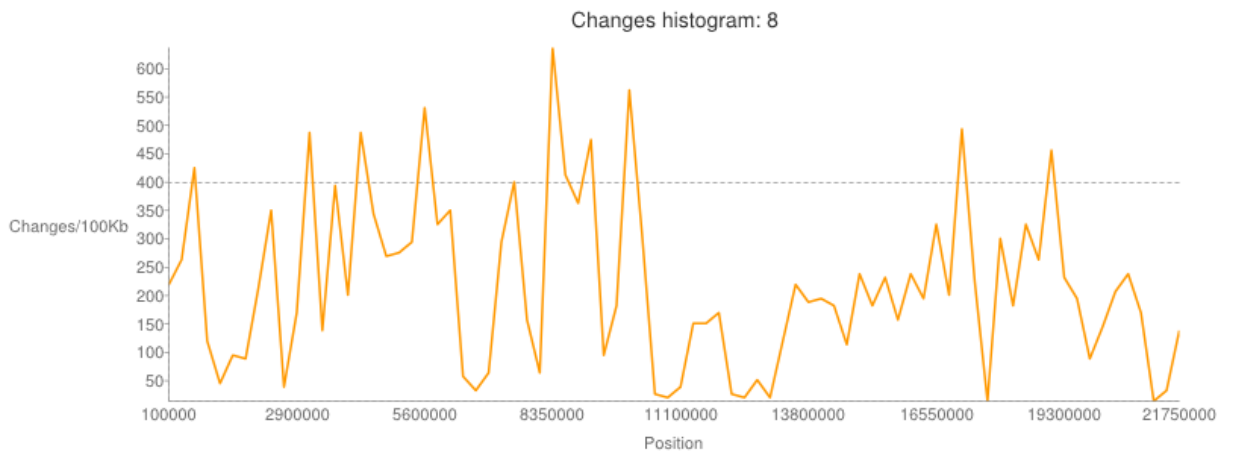
Chromosome: 5



Chromosome: 6



Chromosome: 7



Chromosome: 8

**Details by gene**

[Here](#) you can find a tab-separated table.