

SnpEff: Variant analysis

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Summary

Genome	pppnc139
Date	2012-09-03 21:34
SnpEff version	SnpEff 3.0c (build 2012-07-19), by Pablo Cispedal
Command line arguments	SnpEff pppnc139 -i ref /Volumes/USB_DISK/VCF4/00_vcf -r /Volumes/USB_DISK/VCF4/00_mask -v
Warnings	0
Number of lines (input file)	395,433
Number of variants (before filter)	399,649
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)	399,649
Number of known variants (i.e. non-empty ID)	0 (0%)
Number of effects	630,488
Genome total length	227,252,108
Genome effective length	227,128,408
Change rate	1 change every 568 bases

Change rate details

Chromosome	Length	Changes	Change rate
1	45,877,626	89,337	524
2	28,807,724	87,058	307
3	22,028,950	21,520	1,023
4	30,528,727	60,371	504
5	18,502,877	33,854	545
6	28,902,262	32,718	881
7	22,795,183	58,020	392
8	21,829,783	17,208	1,288
Total	227,128,408	399,649	568

Number changes by type

Type	Total	Ratio	Ratio
SNP	326,646	1,000	172,227
SNP	0	0	0
INS	10,148	4,734	15,382
DEL	21,858	2,048	14,958
MIXED	0	0	0
Interval	0	0	0
Total	399,649	16,179	184,479

Number of effects by impact

Type (alphabetical order)	Count	Percent
HIGH	1,687	0.266%
LOW	10,404	1.65%
MODERATE	15,797	2.505%
MODIFIER	462,599	73.579%

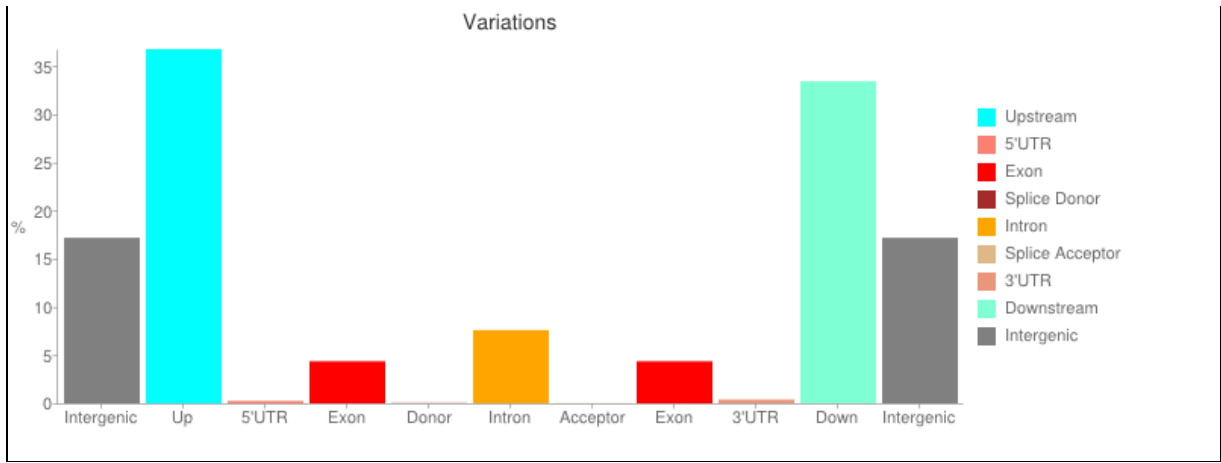
Number of effects by functional class

Type (alphabetical order)	Count	Percent
MISSENSE	15,813	2.508%
NONSENSE	491	0.078%
SILENT	19,628	3.11%

Missense / Silent ratio: 1.5262

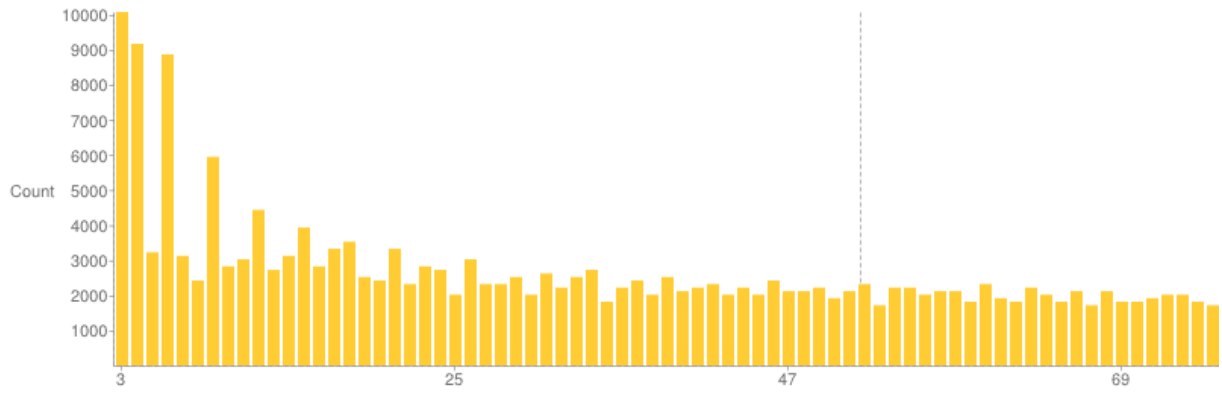
Number of effects by type and region

Type (alphabetical order)	Count	Percent	Region (alphabetical order)	Count	Percent
CODON_CHANGE_PLUS_CODON_DELETION	84	0.013%	DOWNSTREAM	250,787	39.78%
CODON_CHANGE_PLUS_CODON_INSERTION	79	0.012%	EXON	21,408	3.403%
CODON_DELETION	82	0.013%	INTERGENIC	108,303	17.176%
CODON_INSERTION	35	0.005%	INTRON	47,807	7.582%
DOWNSTREAM	250,787	39.78%	INTRON	47,807	7.582%
FRAME_SHIFT	941	0.148%	SPLICE_SITE_ACCEPTOR	131	0.021%
INTERGENIC	108,303	17.176%	SPLICE_SITE_DONOR	134	0.021%
INTRON	47,807	7.582%	UPSTREAM	19,628	3.11%
NON_SYNONYMOUS_CODING	15,813	2.508%	UTR_3_PRIME	424	0.067%
NON_SYNONYMOUS_START	2	0%	UTR_5_PRIME	1,054	0.166%
SPLICE_SITE_ACCEPTOR	131	0.021%			
SPLICE_SITE_DONOR	134	0.021%			
START_GAINED	196	0.031%			
START_LOST	35	0.005%			
STOP_GAINED	498	0.078%			
STOP_LOST	45	0.007%			
SYNONYMOUS_CODING	10,817	1.65%			
SYNONYMOUS_STOP	19	0.003%			
UPSTREAM	19,628	3.11%			
UTR_3_PRIME	424	0.067%			
UTR_5_PRIME	1,054	0.166%			



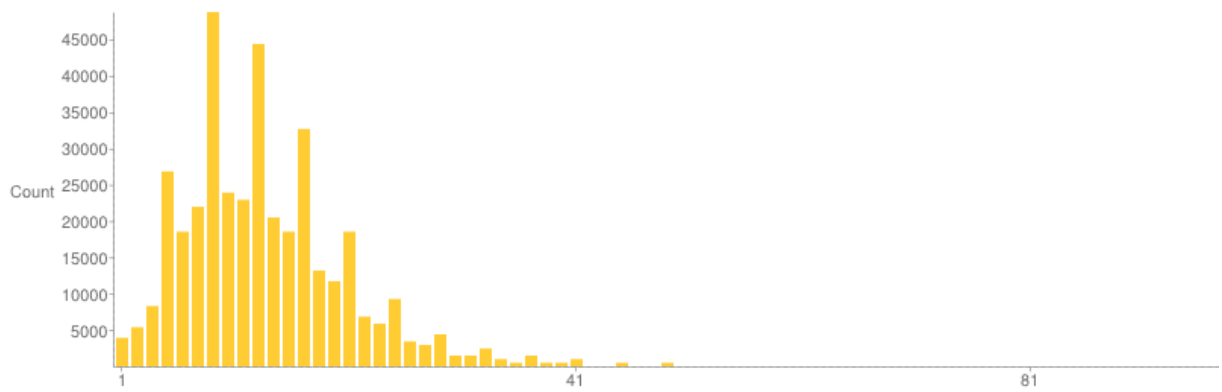
Quality:

Min	3
Max	227
Mean	46.614
Median	75
Standard deviation	63.136



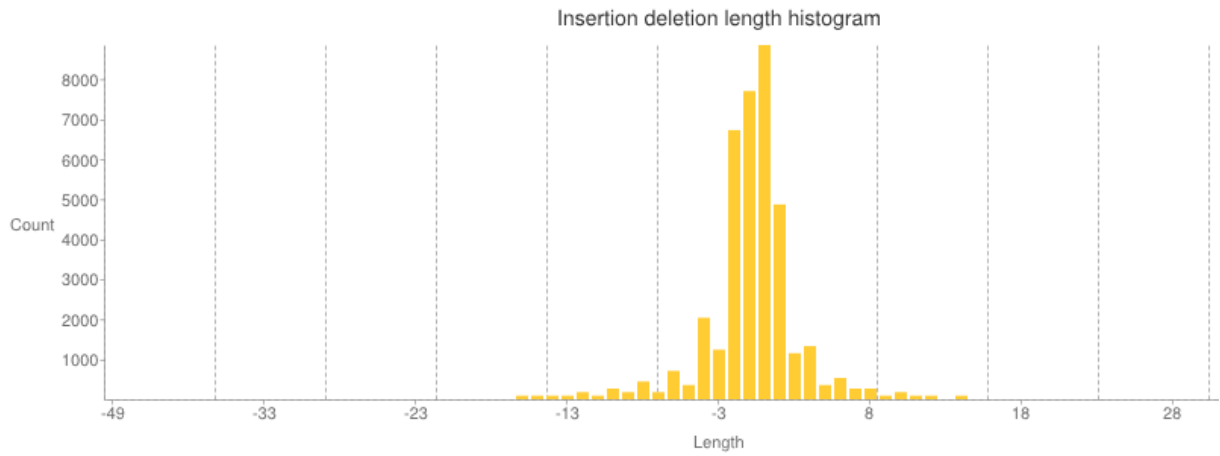
Coverage:

Min	1
Max	2,887
Mean	15,504
Median	13
Standard deviation	22,705



Insertions and deletions length:

Min	-49
Max	42
Mean	-0.528
Median	-1
Standard deviation	5.131



Base changes (SNPs)

	A	C	G	T
A	0	1752	2272	2128
C	1752	0	1752	2272
G	2272	1752	0	1752
T	2128	2272	1752	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	227722
Transversions	130402
Ts/Tv ratio	1.746

All variants:

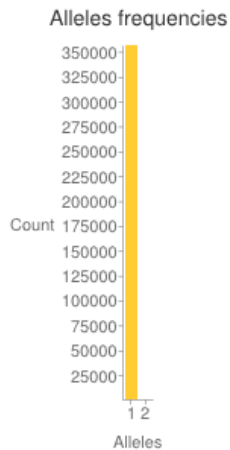
Sample : pp Total
 Transitions : 227723 227723
 Transversions : 130402 130402
 Ts/Tv : 1.746 1.746

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	1.022
Median	1

Standard deviation	0.045
Values	12
Count	39681732

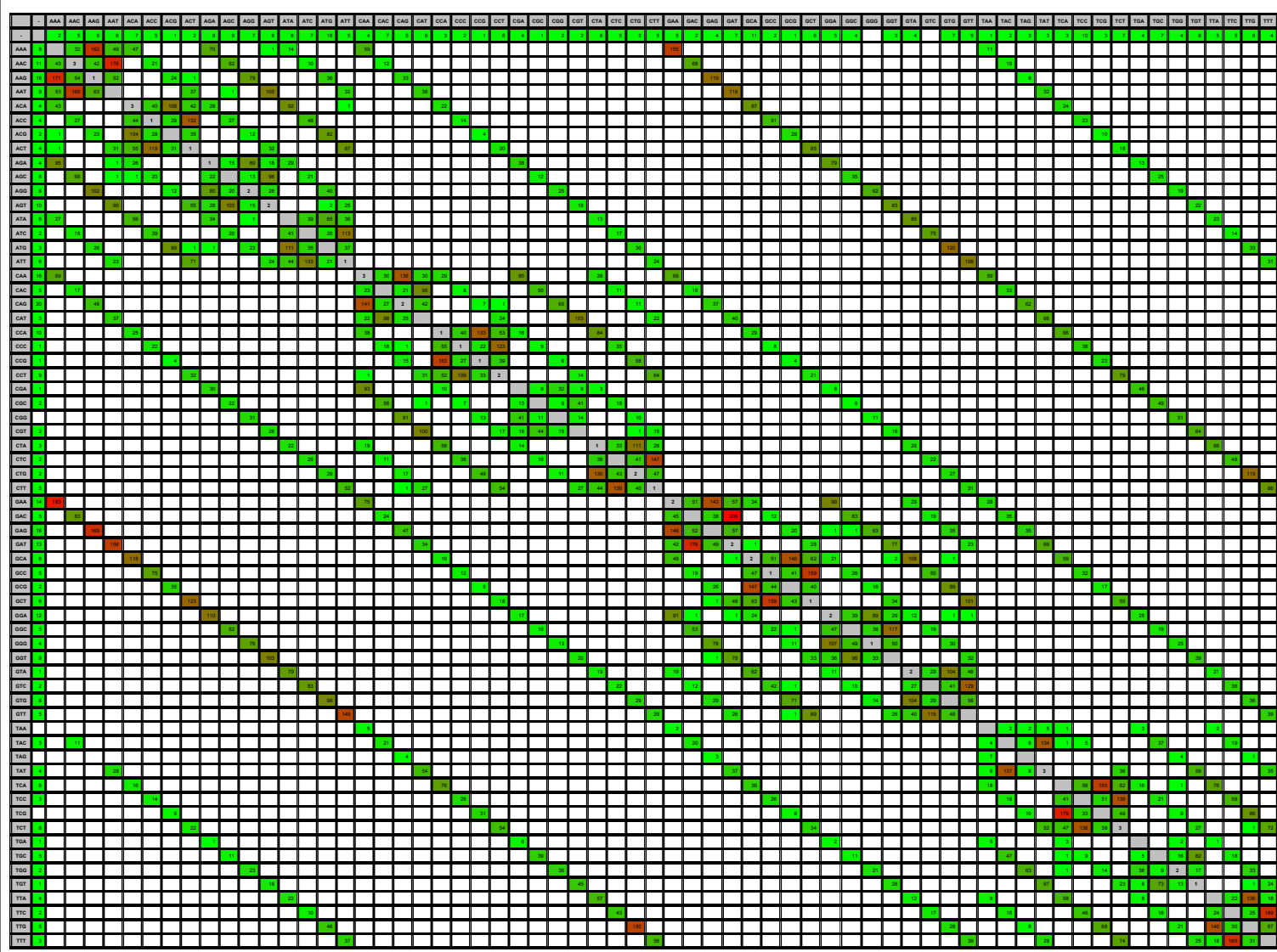
Only known variants (i.e. the ones having a non-empty ID field):

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

Codon changes

How to read this table:

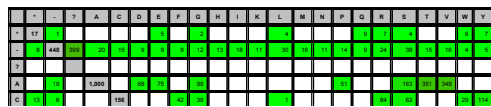
- Rows are reference codons and columns are changed codons. E.g. Row 'AAA' column 'TAA' indicates how many 'AAA' codons have been replaced by 'TAA' codons.
- Red background colors indicate that more changes happened (heat-map).
- Diagonals are indicated using grey background color
- WARNING: This table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



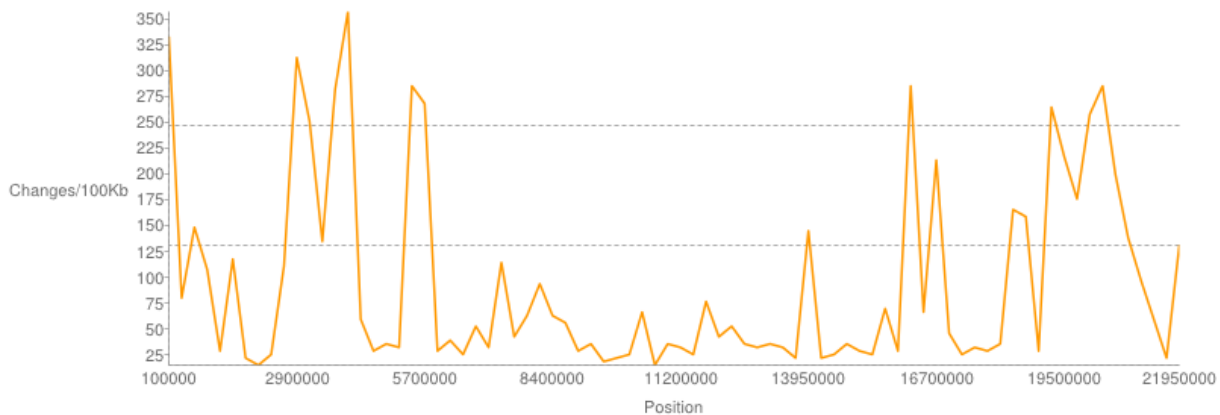
Amino acid changes

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).

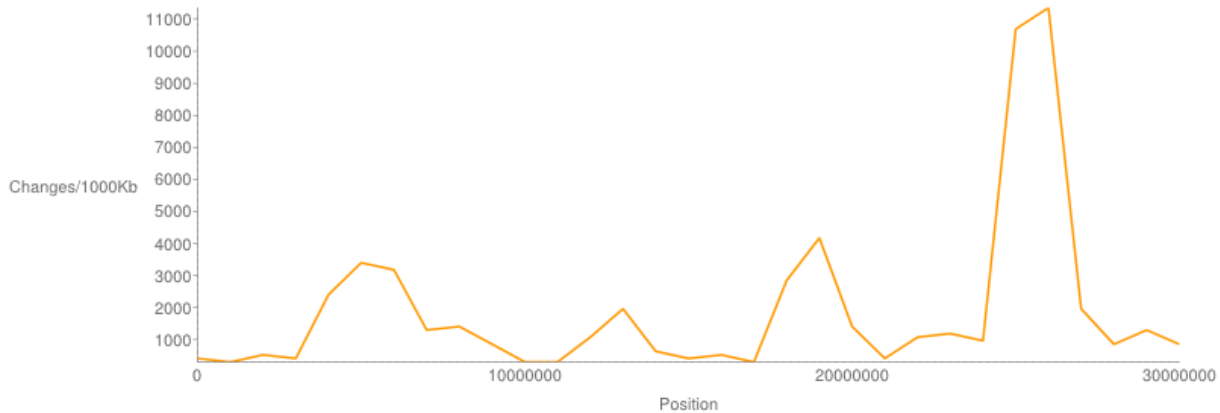


Changes histogram: 3



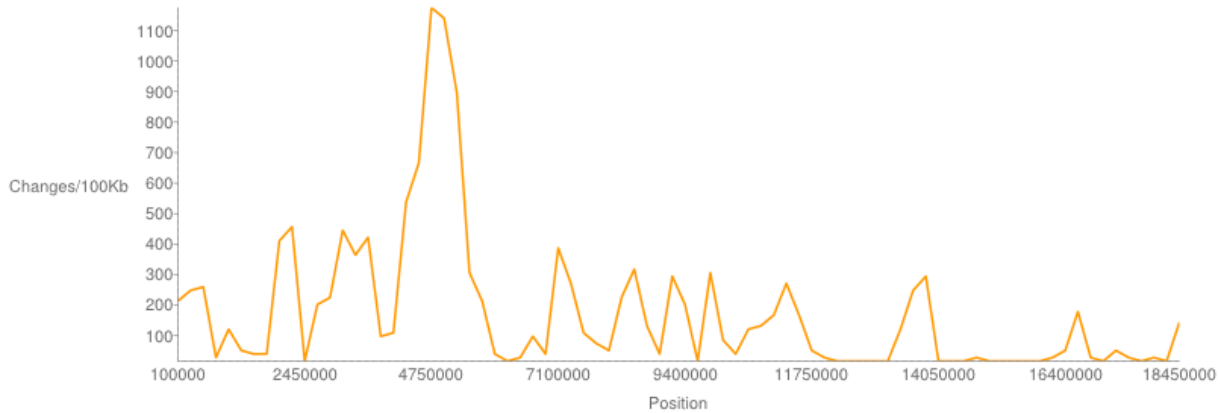
Chromosome: 3

Changes histogram: 4

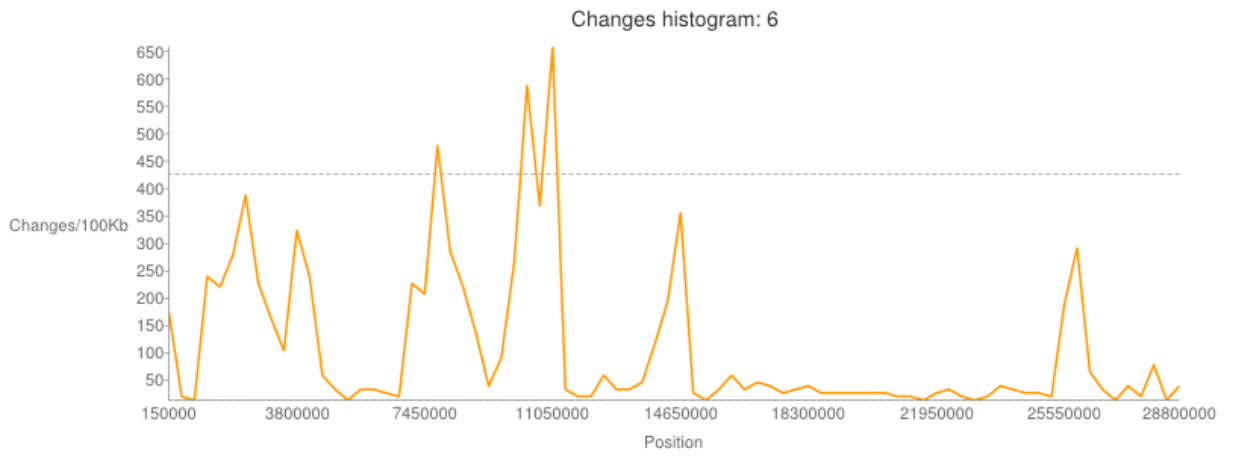


Chromosome: 4

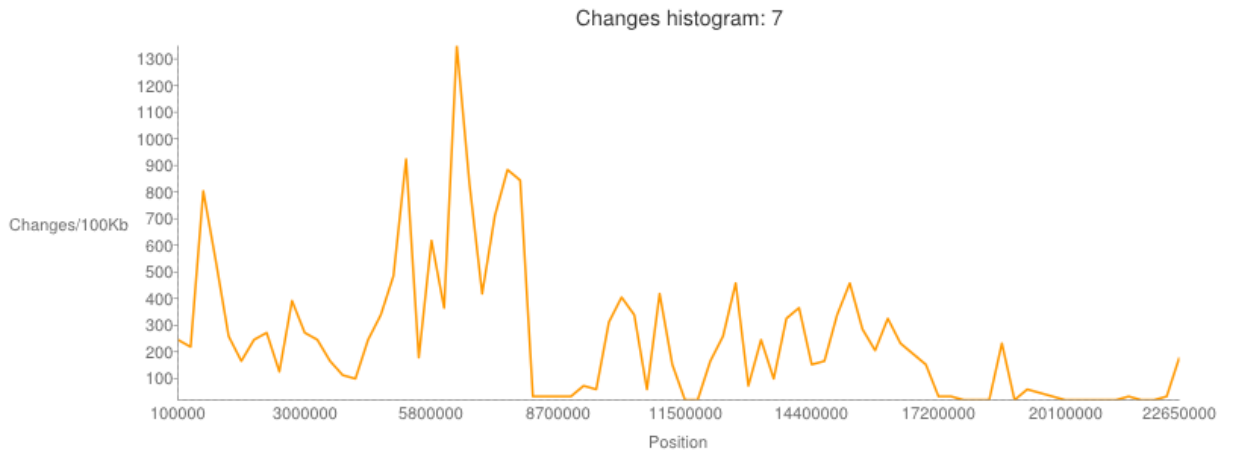
Changes histogram: 5



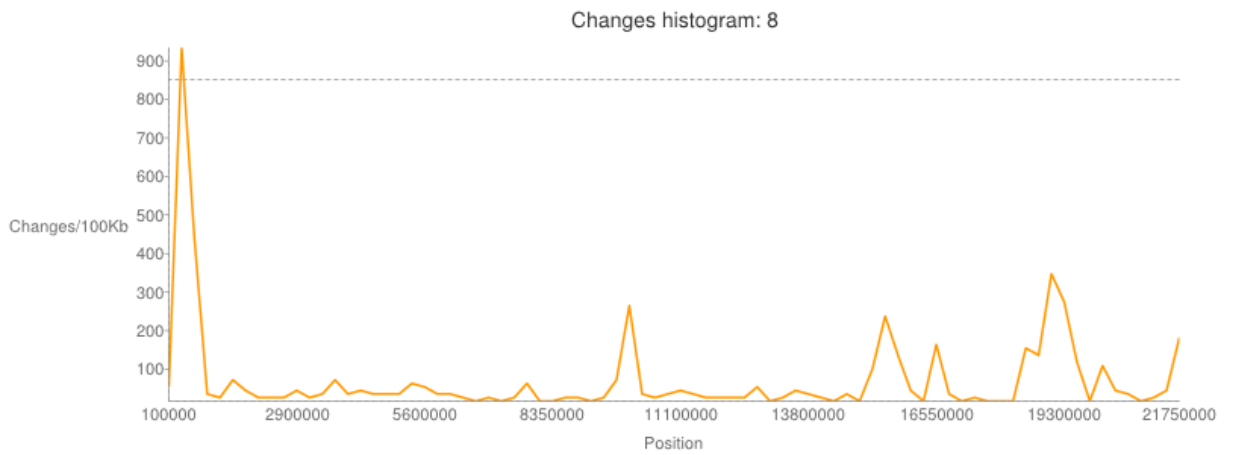
Chromosome: 5



Chromosome: 6



Chromosome: 7



Chromosome: 8

Details by gene

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