Exploring chromosome 12

Going to look closer at chromosome 12

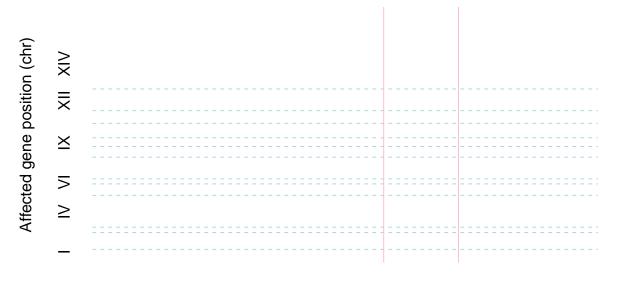
```
library("data.table")
source("myfunctions.R")
genepos <- fread("results/gene pos.gz")</pre>
if (!exists("find.effects TF")){
  find.effects_TF <- fread("results/findeffects_TF_newparams.gz")</pre>
# Add chomosome and start position to each gene
causal.pos.A <- merge(find.effects_TF, genepos, by.x="geneA", by.y="gene", all.x=T)
colnames(causal.pos.A) <- c("geneA", "geneB", "eqtl.A", "eqtl.B", "A->B", "B->A", "strand.A", "start.A"
causal.pos.B <- merge(causal.pos.A, genepos, by.x="geneB", by.y="gene", all.x=T)</pre>
# Keep olnly columns with genes, start positions and chromosomes
causal.pos.B.2 <- unique(causal.pos.B[,.(geneA, geneB, start.A, chr.A, chr.start, chr.id)])</pre>
colnames(causal.pos.B.2) <- c("geneA", "geneB", "start.A", "chr.A", "start.B", "chr.B")</pre>
## transform chromosome ids into numbers
# remove "chr" part of the chromosome name
causal.pos.B.2$chr.A <-gsub('chr', '', causal.pos.B.2$chr.A)</pre>
causal.pos.B.2$chr.B <-gsub('chr', '', causal.pos.B.2$chr.B)</pre>
colnames(causal.pos.B.2) <- c("geneA", "geneB", "start.A", "chr.A", "start.B", "chr.B")</pre>
# convert roman chromosome numbers to numbers
causal.pos.B.2$chr.A <- as.numeric(as.roman(causal.pos.B.2$chr.A))</pre>
causal.pos.B.2$chr.B <- as.numeric(as.roman(causal.pos.B.2$chr.B))</pre>
# order values
causal.pos.B.2.order <- causal.pos.B.2[order(chr.A, start.A, chr.B, start.B)]</pre>
# organize coordinates so that they are ordered by chromosome
# vector of chromosomes
vchr <- 1:16
# how much space will be separating chromosomes
separator <- 1e5
coordinates_plot <- sort_by_chr(vchr = vchr, causal.pos.B.2.order, separator = separator)</pre>
if (!exists("find.effects")){
  find.effects <- fread("results/findeffects_all_newparams.gz")</pre>
#Create a function to generate a continuous color palette
```

XII

Causal gene position (chr)

Chromosome 3 seems to mostly be affected by the genes in the vertical bands so I'm going to look closer at chromosome 3

```
# the first gene of chromosome 12 that affects chromosome 3
lower_limit <- coordinates_plot_cor[chr.A==12 & chr.B==3][order(start.A)][1]$start.A
# the last gene on chromosome 12 that affects chromosome 3 and that's in the band (there's an extra gen
top_limit <- coordinates_plot_cor[chr.A==12 & chr.B==3][order(start.A)][nrow(coordinates_plot_cor[chr.A==12 & chr.B==3]]]
# plot chromosome 12 - the pink lines are the limits of the group of vertical bands
plot_sorted_coordinates(coordinates_plot_cor[chr.A==12], separator = separator, col = coordinates_plot_abline(v = c(lower_limit-1500, top_limit+1500), col="pink")</pre>
```



XII

Causal gene position (chr)

there are 30 genes between the two pink bands

There are

 $nrow (unique (coordinates_plot_cor[chr.A==12 \& chr.B==3][order(start.A)][,.(geneA,start.A)]) [between (start.A,lower_limit,top_limit)]) \#genes that between the two pink bands$

 $\label{eq:cordinates_plot_corf} $$\operatorname{plot_sorted_coordinates_plot_cor[chr.A==12]}, $$ \operatorname{separator} = \operatorname{separator}, $\operatorname{col} = \operatorname{coordinates_plot_cor$col}, $\operatorname{xlim=c}(7422300, \operatorname{top_limit}+1000))$ $$\#$ there are abline($\operatorname{v=c}(7422391-1500, 7425907+1500), \operatorname{col="pink"}) $\operatorname{nrow}(\operatorname{unique}(\operatorname{coordinates_plot_cor[chr.A==12 \& \operatorname{chr.B==3}][\operatorname{order}(\operatorname{start.A})][,.(\operatorname{geneA}, \operatorname{start.A})])$ $$\operatorname{between}(\operatorname{start.A}, 7422391-1500, 7425907+1500)])$ $$$

 $plot_sorted_coordinates(coordinates_plot_cor[chr.A==12], separator = separator, col = coordinates_plot_cor$col, xlim=c(7422391-1500, 7425907+1500))$