## main exons

September 8, 2021

## 1 eQTL boxplot: Enrichment and Overlap of PGC2+CLOZUK

This is script ported from python to fix unknown plotting error.

```
[1]: suppressPackageStartupMessages({
    library(tidyverse)
    library(ggpubr)
})
```

#### 1.1 Functions

```
[2]: feature = "exons"
```

#### 1.1.1 Cached functions

```
[3]: get_de_df <- function(){
         de_file = paste0("../../differential_expression/_m/", feature,
                           "/diffExpr_szVctl_full.txt")
         return(data.table::fread(de_file))
     memDE <- memoise::memoise(get_de_df)</pre>
     get_eqtl_df <- function(){</pre>
         eGenes_file = paste0('../../eqtl/caudate/summary_table/_m/',
                              'Brainseq_LIBD_caudate_4features.signifpairs.txt.gz')
         eGenes = data.table::fread(eGenes_file) %>%
             filter(Type == feature_map(feature)) %>%
             arrange(pval_nominal)
         return(eGenes)
     memEQTL <- memoise::memoise(get_eqtl_df)</pre>
     get_pheno_df <- function(){</pre>
         phenotype_file = paste0('/ceph/projects/v4_phase3_paper/inputs/',
                                   'phenotypes/_m/merged_phenotypes.csv')
         return(data.table::fread(phenotype_file))
     memPHENO <- memoise::memoise(get_pheno_df)</pre>
```

```
get_residualized_df <- function(){</pre>
    expr_file = paste0("../../differential_expression/_m/", feature,
                        "/residualized_expression.tsv")
    return(data.table::fread(expr_file) %>% column_to_rownames("V1"))
}
memRES <- memoise::memoise(get_residualized_df)</pre>
get genotypes <- function(){</pre>
    traw_file = paste0("/ceph/projects/brainseq/genotype/download/topmed/
"filter_maf_01/a_transpose/_m/LIBD_Brain_TopMed.traw")
    traw = data.table::fread(traw_file) %>% rename_with(~ gsub('\\_.*', '', .x))
    return(traw)
memSNPs <- memoise::memoise(get_genotypes)</pre>
get_gwas_snps <- function(){</pre>
    gwas_snp_file = paste0('/ceph/projects/v4_phase3_paper/inputs/sz_gwas/',
                            'pgc2_clozuk/map_phase3/_m/libd_hg38_pgc2sz_snps.tsv')
    gwas df = data.table::fread(gwas snp file) %>% arrange(P)
    return(gwas df)
memGWAS <- memoise::memoise(get_gwas_snps)</pre>
get_integration_df <- function(){</pre>
    return(inner_join(memGWAS(), memEQTL(),
                       by=c("our_snp_id"="variant_id"),
                       suffix=c("_PGC2", "_eQTL")) %>%
            inner_join(memDE(), by=c("gene_id"="V1")) %>%
            mutate(agree_direction=sign(OR -1) * sign(slope) * sign(t) *_
\rightarrowifelse(pgc2 a1 same as our counted, 1, -1)))
memMERGE <- memoise::memoise(get integration df)</pre>
get_geno_annot <- function(){</pre>
    return(memSNPs() %>% select(CHR, SNP, POS, COUNTED, ALT))
memANNOT <- memoise::memoise(get_geno_annot)</pre>
get_snps_df <- function(){</pre>
    return(memSNPs() %>% select("SNP", starts_with("Br")))
memGENOTYPES <- memoise::memoise(get_snps_df)</pre>
get_snp_df <- function(variant_id, gene_id){</pre>
    zz = memANNOT() %>% filter(SNP == variant id)
```

```
xx = memGENOTYPES() %>% filter(SNP == variant_id) %>%
        column_to_rownames("SNP") %>% t %>% as.data.frame %>%
        rownames to column("BrNum") %>% mutate(COUNTED=zz$COUNTED, ALT=zz$ALT)_
 →%>%
        rename("SNP"=all_of(variant_id))
    yy = memRES()[gene id, ] %>% t %>% as.data.frame %>%
        rownames_to_column("RNum") %>% inner_join(memPHENO(), by="RNum")
    ## Annotated SNPs
    letters = c()
    for(ii in seq_along(xx$COUNTED)){
        a0 = xx$COUNTED[ii]; a1 = xx$ALT[ii]; number = xx$SNP[ii]
        letters <- append(letters, letter_snp(number, a0, a1))</pre>
    }
    xx = xx %>% mutate(LETTER=letters, ID=paste(SNP, LETTER, sep="\n"))
    df = inner_join(xx, yy, by="BrNum") %>% mutate_if(is.character, as.factor)
    return(df)
memDF <- memoise::memoise(get_snp_df)</pre>
```

#### 1.1.2 Simple functions

```
[4]: feature_map <- function(feature){
         return(list("genes"="Gene", "transcripts"= "Transcript",
                      "exons"= "Exon", "junctions"= "Junction")[[feature]])
     }
     letter_snp <- function(number, a0, a1){</pre>
         if(is.na(number)){ return(NA) }
         if( length(a0) == 1 & length(a1) == 1){
             seps = ""; collapse=""
         } else {
             seps = " "; collapse=NULL
         }
         return(paste(paste0(rep(a0, number), collapse = collapse),
                       paste0(rep(a1, (2-number)), collapse = collapse), sep=seps))
     }
     save_ggplots <- function(fn, p, w, h){</pre>
         for(ext in c('.pdf', '.png', '.svg')){
             ggsave(paste0(fn, ext), plot=p, width=w, height=h)
         }
     }
     get_biomart_df <- function(){</pre>
         biomart = data.table::fread("../ h/biomart.csv")
     memMART <- memoise::memoise(get_biomart_df)</pre>
```

```
get_gene_symbol <- function(gene_id){</pre>
    ensemblID = gsub("\\..*", "", gene_id)
    geneid = memMART() %>% filter(ensembl_gene_id == gsub("\\..*", "", gene_id))
    if(dim(geneid)[1] == 0){
        return("")
    } else {
        return(geneid$external_gene_name)
    }
}
plot_simple_eqtl <- function(fn, gene_id, variant_id, eqtl_annot){</pre>
    bxp = memDF(variant_id, gene_id) %>%
        ggboxplot(x="ID", y=gene_id, fill="red", add="jitter", xlab="",
                  ylab="Residualized Expression", outlier.shape=NA,
                  add.params=list(alpha=0.5), alpha=0.4,
                  ggtheme=theme_pubr(base_size=20, border=TRUE)) +
        font("xy.title", face="bold") +
        ggtitle(paste(get_gene_symbol(gene_id), gene_id, eqtl_annot, sep='\n'))_u
\hookrightarrow+
        theme(plot.title = element text(hjust = 0.5, face="bold"))
    print(bxp)
    save_ggplots(fn, bxp, 7, 7)
}
```

#### 1.1.3 GWAS plots

```
[5]: get risk allele <- function(OR, A1, A2){
         ra = ifelse(OR > 1, A1, A2)
         return(ra)
     }
     get_df <- function(){</pre>
         return(memEQTL() %>% inner_join(memGWAS(), by="variant_id"))
     }
     get_gwas_ordered_snp_df <- function(variant_id, gene_id,__
      →pgc2_a1_same_as_our_counted, OR){
         df = memDF(variant_id, gene_id)
         if(!pgc2 a1 same as our counted){ # Fix bug with matching alleles!
             if(OR < 1){ df = df %>% mutate(SNP = 2-SNP, ID=paste(SNP, LETTER,,)
      →sep="\n")) }
         } else {
             if(OR > 1){ df = df %>% mutate(SNP = 2-SNP, ID=paste(SNP, LETTER, __
      \rightarrowsep="\n")) }
         }
         return(df)
```

```
}
plot_gwas_eqtl_pheno <- function(fn, gene_id, variant_id,__</pre>
 →pgc2_a1_same_as_our_counted, OR, title){
    bxp = get_gwas_ordered_snp_df(variant_id, gene_id,__
 \rightarrowpgc2_a1_same_as_our_counted, OR) %>%
        mutate_if(is.character, as.factor) %>% filter(Dx %in% c("CTL", "SZ"),_
 →Age > 17) %>%
        ggboxplot(x="ID", y=gene_id, fill="Dx", color="Dx", add="jitter", u
⇒xlab=variant id,
                  ylab="Residualized Expression", outlier.shape=NA,
                  add.params=list(alpha=0.5), alpha=0.4, legend="bottom",
                  ggtheme=theme_pubr(base_size=20, border=TRUE)) +
        font("xy.title", face="bold") + ggtitle(title) +
        theme(plot.title = element_text(hjust = 0.5, face="bold"))
    print(bxp)
    save_ggplots(fn, bxp, 7, 9)
}
```

#### 1.1.4 Exon specific

## 1.2 Integration analysis

```
[7]: dir.create(feature)
```

#### 1.2.1 Enrichment

Integrate DEG with PGC2+CLOZUK SNPs

1. 3650186 2. 53

```
[9]: table(dft$agree_direction)
```

```
[10]: table = matrix(c(sum((dft$P<5e-8) & (dft$adj.P.Val < 0.05)),
                        sum((dft$P>=5e-8) & (dft$adj.P.Val < 0.05)),</pre>
                        sum((dft\$P<5e-8) & (dft\$adj.P.Val >= 0.05)),
                        sum((dft$P>=5e-8) & (dft$adj.P.Val >= 0.05))),
                      nrow=2)
      print(table)
      fisher.test(table)
             [,1]
                     [,2]
     [1,]
            3962
                    67556
     [2,] 256601 3322067
             Fisher's Exact Test for Count Data
     data: table
     p-value < 2.2e-16
     alternative hypothesis: true odds ratio is not equal to 1
     95 percent confidence interval:
      0.734971 0.784211
     sample estimates:
     odds ratio
      0.7592562
[11]: dft2 = dft %>% filter(P <= 5e-8, `adj.P.Val` < 0.05) %>%
          mutate(eqtl_gwas_dir=sign(OR -1) * sign(slope) *__
       →ifelse(pgc2_a1_same_as_our_counted, 1, -1),
                 de_dir=sign(t), eqtl_slope=sign(OR<sub>□</sub>
       \rightarrow-1)*sign(slope)*ifelse(pgc2 a1 same as our counted, 1, -1)) %>%
          #rowwise() %>% mutate(risk_allele=get_risk_allele(our_snp_id)) %>%
          select(gene_id, Symbol, our_snp_id, rsid, A1, A2, OR, P, pval_nominal, adj.
       →P.Val, logFC,
                 t, eqtl_slope, de_dir, eqtl_gwas_dir, agree_direction,_
       →pgc2_a1_same_as_our_counted) %>%
          rename("variant_id"="our_snp_id") %>% mutate_all(list(~na_if(.,""))) %>%
          mutate(Symbol = coalesce(Symbol,gene id))
      dft2 %>% data.table::fwrite(paste0(feature, "/integration_by_symbol.txt"),__
       \rightarrowsep='\t')
      dim(dft2)
     1. 3964 2. 17
[12]: df = dft2 %>% group_by(Symbol) %>% slice(1) %>% arrange(P)
      table(df$agree_direction)
```

No

3615 1820221 1826350

Yes

No Yes 11 14

[13]: df

	$gene\_id$	Symbol	variant_id	rsid	A1	A2	OR
_	<chr></chr>	<chr></chr>	<chr></chr>	<chr></chr>	<chr></chr>	<chr></chr>	<dbl></dbl>
A grouped_df: $25 \times 17$	e378566	ZSCAN26	chr6:28744470:A:G	rs1233578	A	G	1.2228
	e378375	ZNF391	chr6:27837477:A:C	rs34706883	A	$\mathbf{C}$	1.2622
	e379384	HCG4	chr6:29298706:G:A	rs3117439	G	A	1.2636
	e380557	FLOT1	chr6:31204374:T:C	rs3132510	${ m T}$	$\mathbf{C}$	1.2068
	e386363	BRD2	chr6:32669525:G:A	$\mathrm{rs}2395231$	$\mathbf{G}$	A	1.2197
	e380196	PPP1R10	chr6:30894646:C:T	rs1264319	$\mathbf{C}$	${ m T}$	1.1415
	e598887	CNNM2	chr10:102852578:T:A	rs11191419	${ m T}$	A	1.0875
	e378174	HCG11	chr6:26466161:G:A	rs1977199	G	A	0.9143
	e385196	ATF6B	chr6:31758911:C:A	rs707939	$\mathbf{C}$	A	0.9263
	e805625	ZFYVE21	chr14:103847845:G:A	rs10083370	$\mathbf{G}$	A	1.0780
	e193333	NGEF	chr2:232930111:C:T	rs2944591	$\mathbf{C}$	${ m T}$	0.9246
	e805477	e805477	chr14:103778246:C:T	rs12890837	$\mathbf{C}$	${ m T}$	0.9286
	e4306	PLCH2	chr1:2455662:C:T	rs4648845	$\mathbf{C}$	${ m T}$	0.9264
	e361367	LINC01470	chr5:152797561:A:G	rs111294930	A	G	1.0858
	e251235	PCCB	chr3:136435986:C:T	rs66691851	$\mathbf{C}$	${ m T}$	1.0680
	e387950	IP6K3	chr6:33773939:A:G	rs4711350	A	G	0.9238
	e378369	ZNF204P	chr6:27487359:G:A	rs6456793	G	A	1.0680
	e251474	SLC35G2	chr3:136789166:C:A	$\mathrm{rs}1280622$	$\mathbf{C}$	A	0.9404
	e880446	HIRIP3	chr16:30004701:C:T	rs12928610	$\mathbf{C}$	${ m T}$	1.0571
	e353080	REEP2	chr5:138439892:C:G	rs982085	$\mathbf{C}$	G	1.0571
	e226781	PPM1M	chr3:52304836:T:C	rs9311474	${ m T}$	$\mathbf{C}$	1.0558
	e1136648	ZC3H7B	chr22:41357599:G:A	rs11090045	G	A	0.9410
	e678063	B3GAT1	chr11:134426490:C:T	rs893949	$\mathbf{C}$	${ m T}$	1.0548
	e1037535	ZNF14	chr19:19633270:T:C	$\mathrm{rs}11878202$	${ m T}$	$\mathbf{C}$	1.0619
	e897871	SLC7A6	chr16:68251944:A:G	rs1975802	A	G	0.9329

#### 1.2.2 Plot with PGC2 risk allele

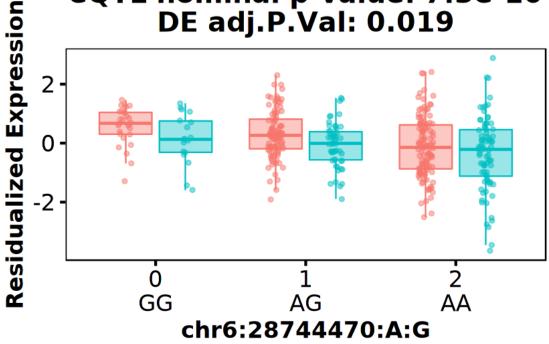
```
[14]: for(num in seq_along(df$gene_id)){
    variant_id = df$variant_id[num]
    gene_id = df$gene_id[num]
    exon_id = exon_annotation(gene_id)
    gene_name = df$Symbol[num]
    gencode_id = filter(memDE(), V1 == gene_id)$gencodeID
    pgc2_a1_same_as_our_counted = df$pgc2_a1_same_as_our_counted[num]
    OR = df$OR[num]; A1 = df$A1[num]; A2 = df$A2[num]
    fn = pasteO(feature, "/eqtl_gwas_", gsub("\\.", "_", gene_name))
    de_annot = paste('DE adj.P.Val:', signif(df$adj.P.Val[num], 2))
    eqtl_annot = paste("eQTL nominal p-value:", signif(df$pval_nominal[num], 2))
    gwas_annot = paste("SZ GWAS pvalue:", signif(df$P[num], 2))
```

Warning message in data.table::fread(expr\_file):

"Detected 393 column names but the data has 394 columns (i.e. invalid file). Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct, or fix the file write command that created the file to create a valid file." Warning message in data.table::fread(expr\_file):

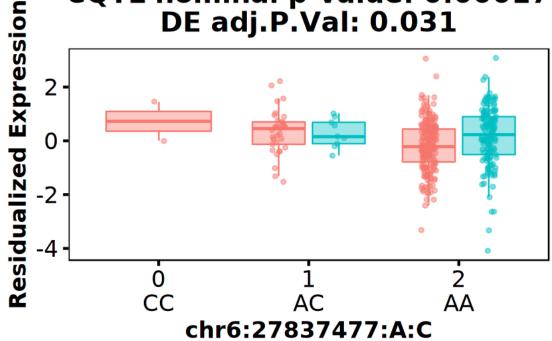
"Detected 393 column names but the data has 394 columns (i.e. invalid file). Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct, or fix the file write command that created the file to create a valid file." Warning message in data.table::fread(expr\_file):

ZSCAN26
ENSE00001651520.2
SZ GWAS pvalue: 4.3e-44
SZ risk allele: A
eQTL nominal p-value: 7.3e-10
DE adj.P.Val: 0.019



Warning message in data.table::fread(expr\_file):

ZNF391
ENSE00001733042.2
SZ GWAS pvalue: 6.5e-41
SZ risk allele: A
eQTL nominal p-value: 0.00017
DE adj.P.Val: 0.031



Warning message in data.table::fread(expr\_file):

HCG4
ENSE00001735745.1
SZ GWAS pvalue: 2.5e-39
SZ risk allele: G
eQTL nominal p-value: 6.9e-05
DE adj.P.Val: 0.019

chr6:29298706:G:A

Warning message in data.table::fread(expr\_file):
"Detected 393 column names but the data has 394 columns (i.e. invalid file).
Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct,

ENSE00003656841.1
SZ GWAS pvalue: 2.3e-31
SZ risk allele: T
eQTL nominal p-value: 2e-04
DE adj.P.Val: 0.043

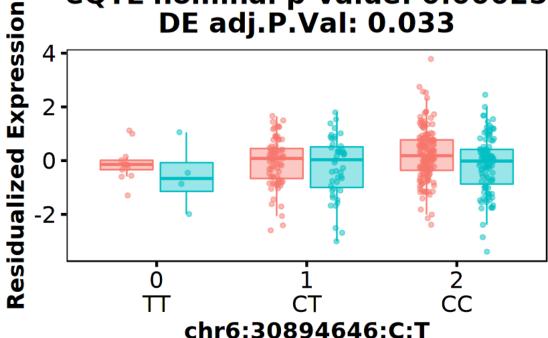
Warning message in data.table::fread(expr\_file):
"Detected 393 column names but the data has 394 columns (i.e. invalid file).
Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct,

BRD2 ENSE00003691002.1 SZ GWAS pvalue: 6.8e-30 SZ risk allele: G eQTL nominal p-value: 3e-06 Residualized Expression **DE adj.P.Val: 0.046** 4 2 0 -2 2 0 GA AA GG chr6:32669525:G:A

Warning message in data.table::fread(expr\_file):
"Detected 393 column names but the data has 394 columns (i.e. invalid file).
Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct,

# PPP1R10 ENSE00001919521.1 SZ GWAS pvalue: 2e-19 SZ risk allele: C

eQTL nominal p-value: 0.00025



Dx 😑 CTL 😑 SZ

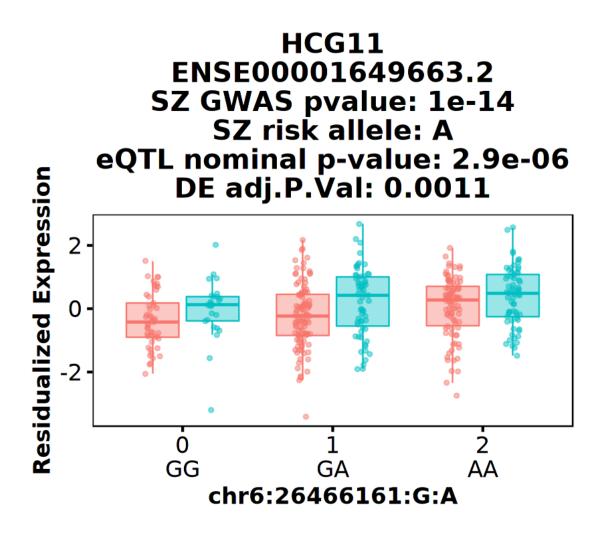
Warning message in data.table::fread(expr\_file):

CNNM2 ENSE00001288096.4 SZ GWAS pvalue: 2.1e-16 SZ risk allele: T eQTL nominal p-value: 4.3e-10 Residualized Expression DE adj.P.Val: 0.0018 2 0 -2 -4 -6 0

TA chr10:102852578:T:A

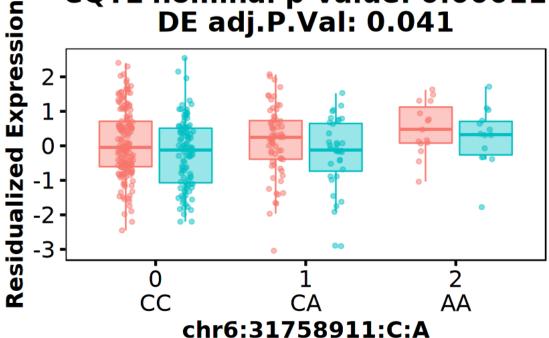
Warning message in data.table::fread(expr\_file):

AA



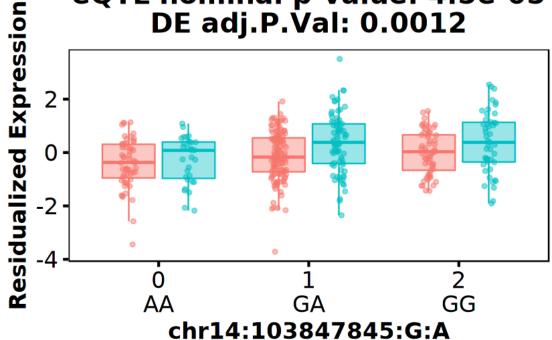
Warning message in data.table::fread(expr\_file):
"Detected 393 column names but the data has 394 columns (i.e. invalid file).
Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct,

ATF6B ENSE00001919167.1 SZ GWAS pvalue: 1.5e-14 SZ risk allele: A eQTL nominal p-value: 0.00011 DE adj.P.Val: 0.041



Warning message in data.table::fread(expr\_file):

ZFYVE21
ENSE00003418764.1
SZ GWAS pvalue: 3.4e-14
SZ risk allele: G
eQTL nominal p-value: 4.5e-05
DE adj.P.Val: 0.0012

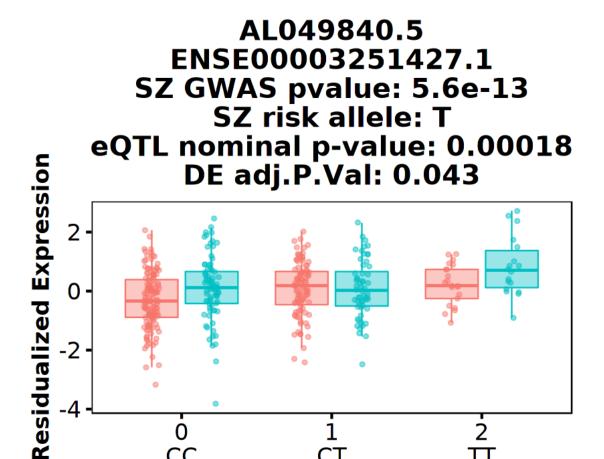


Warning message in data.table::fread(expr\_file):

NGEF
ENSE00001584949.1
SZ GWAS pvalue: 1.1e-13
SZ risk allele: T
eQTL nominal p-value: 4.2e-05
DE adj.P.Val: 0.039

chr2:232930111:C:T

Warning message in data.table::fread(expr\_file):

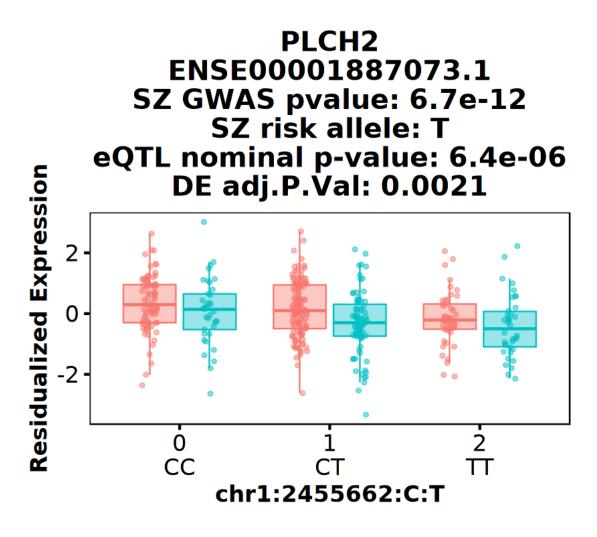


chr14:103778246:C:T

Warning message in data.table::fread(expr\_file): "Detected 393 column names but the data has 394 columns (i.e. invalid file). Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct,

CC

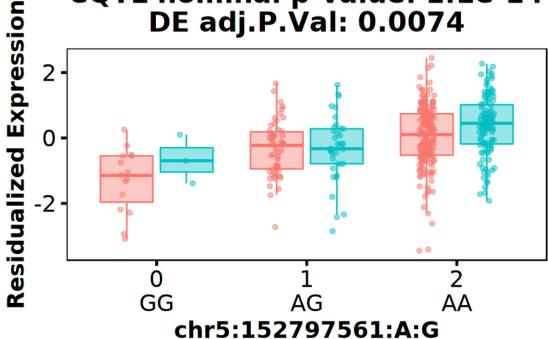
20



Warning message in data.table::fread(expr\_file):
"Detected 393 column names but the data has 394 columns (i.e. invalid file).

Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct, or fix the file write command that created the file to create a valid file."

# LINC01470 ENSE00002139369.1 SZ GWAS pvalue: 9e-12 SZ risk allele: A eQTL nominal p-value: 1.1e-14 DE adj.P.Val: 0.0074



Dx 😑 CTL 😑 SZ

Warning message in data.table::fread(expr\_file):

**PCCB** ENSE00001857296.1 SZ GWAS pvalue: 1.4e-11 SZ risk allele: C eQTL nominal p-value: 0.00037 Residualized Expression DE adj.P.Val: 0.046 2 0

chr3:136435986:C:T

Warning message in data.table::fread(expr\_file):

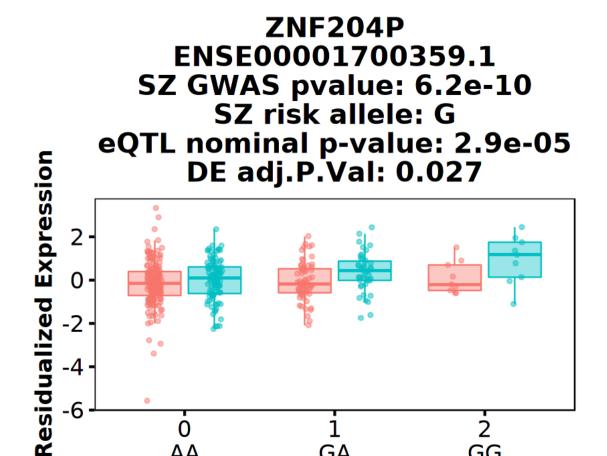
0

-2

IP6K3 ENSE00001061200.1 SZ GWAS pvalue: 2.3e-10 SZ risk allele: G eQTL nominal p-value: 5.9e-10 Residualized Expression **DE adj.P.Val: 0.025** 2.5 0.0 -2.5 0 2 AA AG GG chr6:33773939:A:G

Warning message in data.table::fread(expr\_file):
"Detected 393 column names but the data has 394 columns (i.e. invalid file).
Added 1 extra default column name for the first column which is guessed to be

row names or an index. Use setnames() afterwards if this guess is not correct, or fix the file write command that created the file to create a valid file."



GA

chr6:27487359:G:A

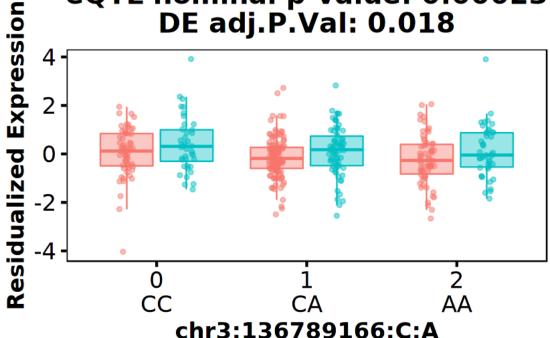
GG

Warning message in data.table::fread(expr\_file): "Detected 393 column names but the data has 394 columns (i.e. invalid file). Added 1 extra default column name for the first column which is guessed to be row names or an index. Use setnames() afterwards if this guess is not correct,

or fix the file write command that created the file to create a valid file."

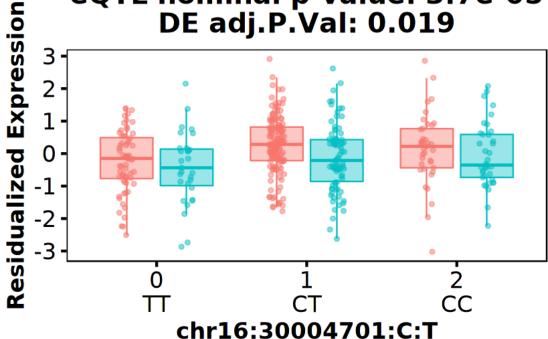
AA

SLC35G2 ENSE00001924650.1 SZ GWAS pvalue: 6.4e-10 SZ risk allele: A eQTL nominal p-value: 0.00023 DE adj.P.Val: 0.018



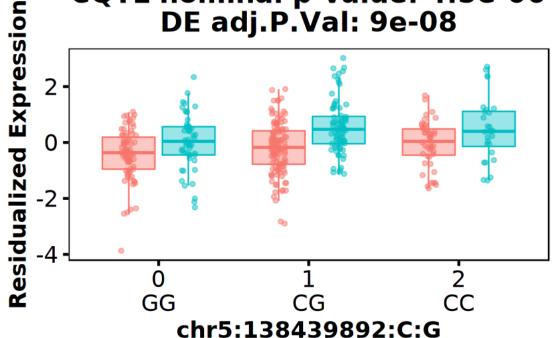
Warning message in data.table::fread(expr\_file):

HIRIP3
ENSE00002628577.2
SZ GWAS pvalue: 7.3e-09
SZ risk allele: C
eQTL nominal p-value: 3.7e-05
DE adj.P.Val: 0.019



Warning message in data.table::fread(expr\_file):

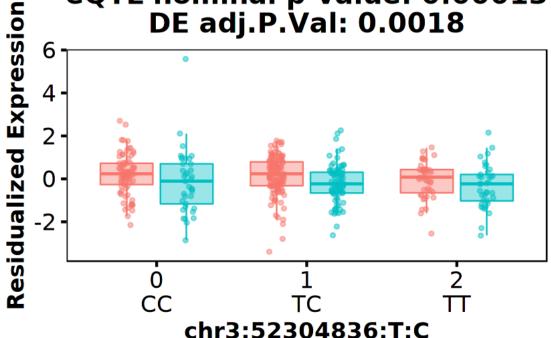
# REEP2 ENSE00001320042.5 SZ GWAS pvalue: 1e-08 SZ risk allele: C eQTL nominal p-value: 4.5e-06



Dx 🖶 CTL 😑 SZ

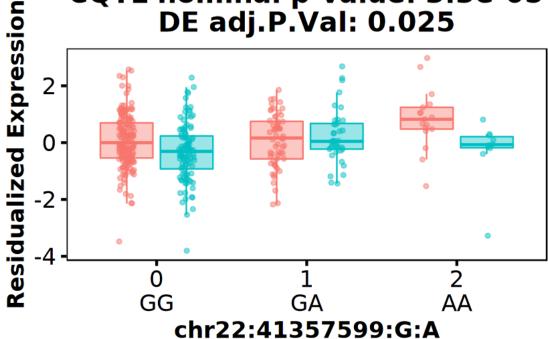
Warning message in data.table::fread(expr\_file):

PPM1M ENSE00001949438.1 SZ GWAS pvalue: 1.2e-08 SZ risk allele: T eQTL nominal p-value: 0.00013



Warning message in data.table::fread(expr\_file):

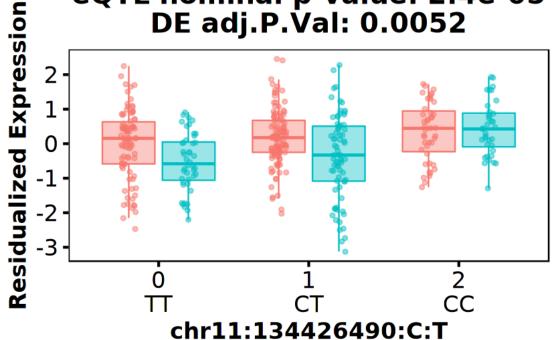
ZC3H7B
ENSE00000655656.1
SZ GWAS pvalue: 1.8e-08
SZ risk allele: A
eQTL nominal p-value: 3.3e-05
DE adj.P.Val: 0.025



Warning message in data.table::fread(expr\_file):

**B3GAT1** ENSE00002197011.1 SZ GWAS pvalue: 3e-08 SZ risk allele: C

eQTL nominal p-value: 2.4e-05 DE adj.P.Val: 0.0052

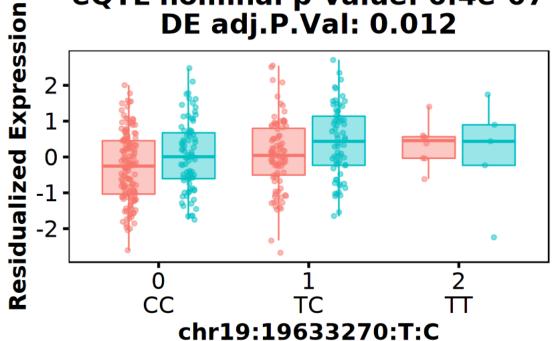


Dx 😑 CTL 😑 SZ

**ZNF14** ENSE00001407032.2 SZ GWAS pvalue: 3.1e-08 SZ risk allele: T

eQTL nominal p-value: 6.4e-07 DE adj.P.Val: 0.012

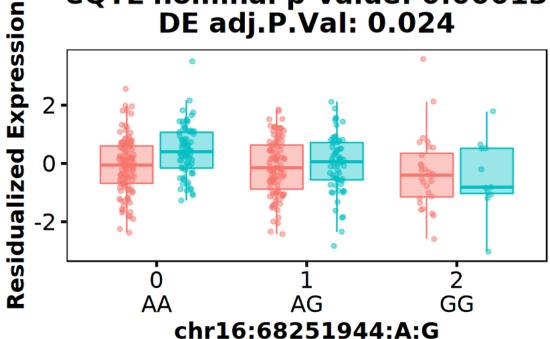




Dx 😑 CTL 😑 SZ

# SLC7A6 ENSE00001616168.2 SZ GWAS pvalue: 3.6e-08 SZ risk allele: G

eQTL nominal p-value: 0.00013 DE adj.P.Val: 0.024



Dx 🖶 CTL 🖶 SZ

#### 1.3 Session Info

[15]: Sys.time() proc.time() options(width = 120) sessioninfo::session\_info() [1] "2021-09-08 11:31:59 EDT"

system elapsed user 10734.485 5689.653 2323.851

Session info setting value version R version 4.0.3 (2020-10-10)

os Arch Linux

system x86\_64, linux-gnu

ui X11 language (EN)

collate en\_US.UTF-8
ctype en\_US.UTF-8
tz America/New\_York

date 2021-09-08

## Packages

package	*	version	date	lib	sourc	ce
abind		1.4-5	2016-07-21	[1]	CRAN	(R 4.0.2)
assertthat		0.2.1	2019-03-21	[1]	CRAN	(R 4.0.2)
backports		1.2.1	2020-12-09	[1]	CRAN	(R 4.0.2)
base64enc		0.1-3	2015-07-28	[1]	CRAN	(R 4.0.2)
broom		0.7.9	2021-07-27	[1]	CRAN	(R 4.0.3)
cachem		1.0.6	2021-08-19	[1]	CRAN	(R 4.0.3)
Cairo		1.5-12.2	2020-07-07	[1]	CRAN	(R 4.0.2)
car		3.0-11	2021-06-27	[1]	CRAN	(R 4.0.3)
carData		3.0-4	2020-05-22	[1]	CRAN	(R 4.0.2)
cellranger		1.1.0	2016-07-27	[1]	CRAN	(R 4.0.2)
cli		3.0.1	2021-07-17	[1]	CRAN	(R 4.0.3)
colorspace		2.0-2	2021-06-24	[1]	CRAN	(R 4.0.3)
crayon		1.4.1	2021-02-08	[1]	CRAN	(R 4.0.3)
curl		4.3.2	2021-06-23	[1]	CRAN	(R 4.0.3)
data.table		1.14.0	2021-02-21	[1]	CRAN	(R 4.0.3)
DBI		1.1.1	2021-01-15	[1]	CRAN	(R 4.0.2)
dbplyr		2.1.1	2021-04-06	[1]	CRAN	(R 4.0.3)
digest		0.6.27	2020-10-24	[1]	CRAN	(R 4.0.2)
dplyr	*	1.0.7	2021-06-18	[1]	CRAN	(R 4.0.3)
ellipsis		0.3.2	2021-04-29	[1]	CRAN	(R 4.0.3)
evaluate		0.14	2019-05-28	[1]	CRAN	(R 4.0.2)
fansi		0.5.0	2021-05-25	[1]	CRAN	(R 4.0.3)
farver		2.1.0	2021-02-28	[1]	CRAN	(R 4.0.3)
fastmap		1.1.0	2021-01-25	[1]	CRAN	(R 4.0.2)
forcats	*	0.5.1	2021-01-27	[1]	CRAN	(R 4.0.2)
foreign		0.8-80	2020-05-24	[2]	CRAN	(R 4.0.3)
fs		1.5.0	2020-07-31	[1]	CRAN	(R 4.0.2)
generics		0.1.0	2020-10-31	[1]	CRAN	(R 4.0.2)
ggplot2	*	3.3.5	2021-06-25	[1]	CRAN	(R 4.0.3)
ggpubr	*	0.4.0	2020-06-27	[1]	CRAN	(R 4.0.2)
ggsignif		0.6.2	2021-06-14	[1]	CRAN	(R 4.0.3)
glue		1.4.2	2020-08-27	[1]	CRAN	(R 4.0.2)
gtable		0.3.0	2019-03-25	[1]	CRAN	(R 4.0.2)
haven		2.4.3	2021-08-04	[1]	CRAN	(R 4.0.3)
hms		1.1.0	2021-05-17	[1]	CRAN	(R 4.0.3)
htmltools		0.5.2	2021-08-25	[1]	CRAN	(R 4.0.3)

```
1.4.2
                        2020-07-20 [1] CRAN (R 4.0.2)
httr
IRdisplay
              1.0
                        2021-01-20 [1] CRAN (R 4.0.2)
IRkernel
              1.2
                        2021-05-11 [1] CRAN (R 4.0.3)
jsonlite
              1.7.2
                        2020-12-09 [1] CRAN (R 4.0.2)
labeling
              0.4.2
                        2020-10-20 [1] CRAN (R 4.0.2)
lifecycle
                        2021-02-15 [1] CRAN (R 4.0.3)
              1.0.0
lubridate
              1.7.10
                        2021-02-26 [1] CRAN (R 4.0.3)
magrittr
              2.0.1
                        2020-11-17 [1] CRAN (R 4.0.2)
              2.0.0
                        2021-01-26 [1] CRAN (R 4.0.2)
memoise
modelr
              0.1.8
                        2020-05-19 [1] CRAN (R 4.0.2)
              0.5.0
                        2018-06-12 [1] CRAN (R 4.0.2)
munsell
              4.2.4
                        2021-06-16 [1] CRAN (R 4.0.3)
openxlsx
              0.3 - 5
                        2021-02-10 [1] CRAN (R 4.0.3)
pbdZMQ
pillar
              1.6.2
                        2021-07-29 [1] CRAN (R 4.0.3)
pkgconfig
              2.0.3
                        2019-09-22 [1] CRAN (R 4.0.2)
            * 0.3.4
                        2020-04-17 [1] CRAN (R 4.0.2)
purrr
R.methodsS3
              1.8.1
                        2020-08-26 [1] CRAN (R 4.0.3)
R.oo
              1.24.0
                        2020-08-26 [1] CRAN (R 4.0.3)
              2.10.1
                        2020-08-26 [1] CRAN (R 4.0.3)
R.utils
R6
              2.5.1
                        2021-08-19 [1] CRAN (R 4.0.3)
Rcpp
              1.0.7
                        2021-07-07 [1] CRAN (R 4.0.3)
                        2021-08-10 [1] CRAN (R 4.0.3)
readr
            * 2.0.1
readxl
              1.3.1
                        2019-03-13 [1] CRAN (R 4.0.2)
              1.1.3
                        2021-01-21 [1] CRAN (R 4.0.2)
repr
              2.0.1
                        2021-08-05 [1] CRAN (R 4.0.3)
reprex
              0.5.27
rio
                        2021-06-21 [1] CRAN (R 4.0.3)
              0.4.11
                        2021-04-30 [1] CRAN (R 4.0.3)
rlang
rstatix
              0.7.0
                        2021-02-13 [1] CRAN (R 4.0.3)
              0.13
                        2020-11-12 [1] CRAN (R 4.0.2)
rstudioapi
              1.0.1
                        2021-07-26 [1] CRAN (R 4.0.3)
rvest
                        2020-05-11 [1] CRAN (R 4.0.2)
scales
              1.1.1
sessioninfo
              1.1.1
                        2018-11-05 [1] CRAN (R 4.0.2)
stringi
              1.7.4
                        2021-08-25 [1] CRAN (R 4.0.3)
            * 1.4.0
                        2019-02-10 [1] CRAN (R 4.0.2)
stringr
                        2021-02-20 [1] CRAN (R 4.0.3)
svglite
              2.0.0
systemfonts
              1.0.2
                        2021-05-11 [1] CRAN (R 4.0.3)
tibble
            * 3.1.4
                        2021-08-25 [1] CRAN (R 4.0.3)
tidyr
            * 1.1.3
                        2021-03-03 [1] CRAN (R 4.0.3)
tidyselect
                        2021-04-30 [1] CRAN (R 4.0.3)
              1.1.1
tidyverse
            * 1.3.1
                        2021-04-15 [1] CRAN (R 4.0.3)
tzdb
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                        2021-07-20 [1] CRAN (R 4.0.3)
utf8
              1.2.2
                        2021-07-24 [1] CRAN (R 4.0.3)
uuid
              0.1-4
                        2020-02-26 [1] CRAN (R 4.0.2)
              0.3.8
                        2021-04-29 [1] CRAN (R 4.0.3)
vctrs
withr
              2.4.2
                        2021-04-18 [1] CRAN (R 4.0.3)
xml2
              1.3.2
                        2020-04-23 [1] CRAN (R 4.0.2)
              2.2.0
                        2021-05-31 [1] CRAN (R 4.0.3)
zip
```

- [1] /home/jbenja13/R/x86\_64-pc-linux-gnu-library/4.0
  [2] /usr/lib/R/library