main exons

March 8, 2022

1 eQTL boxplot

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 Basic loading functions

```
[3]: get residualized df <- function(){
        expr_file = paste0("/ceph/projects/v4_phase3_paper/analysis/eqtl_analysis/
     ⇒all/",
                          feature,"/expression_gct/covariates/
     feature, "_residualized_expression.csv")
        return(data.table::fread(expr_file) %>% column_to_rownames("gene_id"))
    memRES <- memoise::memoise(get_residualized_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_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>
```

1.1.2 eQTL and helpful functions

```
[4]: feature_map <- function(feature){
         return(list("genes"="Gene", "transcripts"= "Transcript",
                      "exons"= "Exon", "junctions"= "Junction")[[feature]])
     }
     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_caudate_eqtls <- function(){</pre>
         mashr_file = "../../summary_table/_m/BrainSeq_caudate_eQTL.txt.gz"
         return(data.table::fread(mashr_file) %>%
                filter(Type == feature_map(feature)) %>%
                 select(gene_id, variant_id, AA, EA))
     }
     memCAUDATE <- memoise::memoise(get_caudate_eqtls)</pre>
     get_eqtl_df <- function(){</pre>
         eGenes_file = paste0("../../_m/", feature, "/lfsr_allpairs_ancestry.txt.gz")
         eGenes = data.table::fread(eGenes_file)
         return(eGenes)
     memEQTL <- memoise::memoise(get_eqtl_df)</pre>
```

1.1.3 Basic eQTL plotting functions

```
[5]: get_geno_annot <- function(){
    return(memSNPs() %>% select(CHR, SNP, POS, COUNTED, ALT))
}

get_snps_df <- function(){
    return(memSNPs() %>% select("SNP", starts_with("Br")))
}

letter_snp <- function(number, a0, a1){
    if(is.na(number)){ return(NA) }
    if( length(a0) == 1 & length(a1) == 1){
        seps = ""; collapse=""
    } else {
        seps = " "; collapse=NULL</pre>
```

```
return(paste(paste0(rep(a0, number), collapse = collapse),
                       pasteO(rep(a1, (2-number)), collapse = collapse), sep=seps))
     }
     get_snp_df <- function(variant_id, gene_id){</pre>
         zz = get_geno_annot() %>% filter(SNP == variant_id)
         xx = get_snps_df() %>% 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("BrNum") %>% inner_join(memPHENO(), by="BrNum")
         ## 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>
     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)
         }
     }
[6]: plot_simple_eqtl <- function(fn, gene_id, variant_id, eqtl_annot, prefix,_
      \rightarrow y0=NULL, y1=NULL){
         if(is.null(y0)){ y0 = quantile(memDF(variant_id, gene_id)[[gene_id]],__
      \rightarrowprobs=c(0.01))[[1]] - 0.2}
         if(is.null(y1)){ y1 = quantile(memDF(variant_id, gene_id)[[gene_id]],__
      \rightarrowprobs=c(0.99))[[1]] + 0.2}
         bxp = memDF(variant_id, gene_id) %>%
             ggboxplot(x="ID", y=gene id, fill="Race", color="Race", add="jitter",
                        xlab=variant_id, ylab="Residualized Expression", outlier.
```

add.params=list(alpha=0.5), alpha=0.4, legend="bottom",

⇒shape=NA,

1.1.4 GWAS plots

```
[7]: get_gwas_snps <- function(){
         gwas_snp_file = paste0('/ceph/projects/v4_phase3_paper/inputs/sz_gwas/pgc3/
      \hookrightarrow ',
                                  'map_phase3/_m/libd_hg38_pgc2sz_snps_p5e_minus8.tsv')
         gwas_df = data.table::fread(gwas_snp_file) %>% arrange(P)
         return(gwas_df)
     memGWAS <- memoise::memoise(get_gwas_snps)</pre>
     get_gwas_snp <- function(variant){</pre>
         return(memGWAS() %>% filter(our_snp_id == variant))
     }
     get risk allele <- function(variant){</pre>
         gwas_snp = get_gwas_snp(variant)
         if(gwas snp$OR > 1){
             ra = gwas_snp$A1
         }else{
             ra = gwas_snp$A2
         }
         return(ra)
     }
     get_eqtl_gwas_df <- function(){</pre>
         return(memCAUDATE() %>% inner_join(memGWAS(),__
      ⇔by=c("variant_id"="our_snp_id")))
     get_gwas_ordered_snp_df <- function(variant_id, gene_id,__
      →pgc3_a1_same_as_our_counted, OR){
         df = memDF(variant id, gene id)
         if(!pgc3_a1_same_as_our_counted){ # Fix bug with matching alleles!
              if(OR < 1) { df = df %>% mutate(SNP = 2-SNP, ID=paste(SNP, LETTER, I
      \rightarrow sep="\langle n"\rangle) }
         } else {
```

```
if(OR > 1){ df = df %>% mutate(SNP = 2-SNP, ID=paste(SNP, LETTER, __
 return(df)
}
plot_gwas_eqtl <- function(fn, gene_id, variant_id, eqtl_annot,_
→pgc3_a1_same_as_our_counted,
                           OR, title){
    dt = get_gwas_ordered_snp_df(variant_id, gene_id,__
→pgc3_a1_same_as_our_counted, OR)
    bxp = dt %>% mutate_if(is.character, as.factor) %>%
        ggboxplot(x="ID", y=gene_id, fill="Race", color="Race", add="jitter",
                  xlab=variant_id, ylab="Residualized Expression", outlier.
 \rightarrowshape=NA,
                  add.params=list(alpha=0.5), alpha=0.4, legend="bottom", __
\rightarrow#ylim=c(y0,y1),
                  palette="npg", 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, 8)
}
```

1.2 Plot eQTL

1.2.1 DRD2 plot

```
[9]: drd2_short = "ENST00000346454.7"; drd2_long = "ENST00000362072.7" exons_df = data.table::fread("../../../DE_gwas_eqtl_integration/_h/exons.

→csv") %>%
```

```
filter(transcript_id %in% c(drd2_short, drd2_long)) %>%
          select(gene_id, gene_name, exon_id, brainseq_exon_id) %>%
          distinct(brainseq_exon_id, .keep_all=TRUE)
      all_drd2_exons = exons_df$brainseq_exon_id
      all_drd2_exons
     1. 'e667152' 2. 'e667153' 3. 'e667154' 4. 'e667155' 5. 'e667156' 6. 'e667157' 7. 'e667158' 8. 'e667159'
     9. 'e667164'
[10]: drd2 df0 = memCAUDATE() %>% filter(gene id %in% all drd2 exons) %>%
          arrange(AA, EA) %>% group_by(gene_id) %>% slice(1) %>% arrange(AA, EA)
      drd2 df0
                          gene_id variant_id
                                                        AA
                                                                    EA
                                   \langle chr \rangle
                                                        < dbl >
                                                                    <dbl>
                          <chr>
     A grouped df: 2 \times 4
                          e667159
                                   chr11:113528769:T:C 0.06756843
                                                                   0.01091546
                          e667157 chr11:113371155:C:T 0.11607231
                                                                   0.01183844
[11]: drd2_df = memEQTL() %>% filter(gene_id %in% all_drd2_exons) %>%
          arrange(EA, AA) %>% group_by(gene_id) %>% slice(1) %>% arrange(AA, EA)
      drd2_df
                         effect
                                                      gene id variant id
                                                                                    AA
                         <chr>
                                                      <chr>
                                                                <chr>
                                                                                    <dbl>
                         e667159_chr11:113524125:G:C
                                                      e667159
                                                               chr11:113524125:G:C
                                                                                    0.07281645
     A grouped df: 4 \times 5
                         e667158 chr11:112955463:T:G
                                                      e667158
                                                               chr11:112955463:T:G 0.07634220
                         e667157 chr11:113358943:T:C
                                                      e667157
                                                               chr11:113358943:T:C
                                                                                    0.13383811
                         e667153 chr11:113446345:G:A e667153 chr11:113446345:G:A 0.47320254
[12]: for(x in seq_along(drd2_df$gene_id)){
          anno = get_drd2_exon_annotation(drd2_df$gene_id[x])
          en = gsub(" ", "_", anno)
          ids = exons_df %>% filter(brainseq_exon_id == drd2_df$gene_id[x]) %>%_
       ⇔select("exon_id")
          fn = paste("drd2_eqtl", en, sep="_")
          eqtl_annot = paste(paste("eQTL (AA) lfsr:", signif(drd2_df$AA[x], 2)),
                               paste("eQTL (EA) lfsr:", signif(drd2_df$EA[x], 2)),__
       \hookrightarrowsep='\n')
          prefix = paste(anno, ids$exon_id, sep='\n')
          plot_simple_eqtl(fn, drd2_df$gene_id[x], drd2_df$variant_id[x], eqtl_annot,_
```

EA

<dbl>

0.0045421

0.0859227

0.0039085

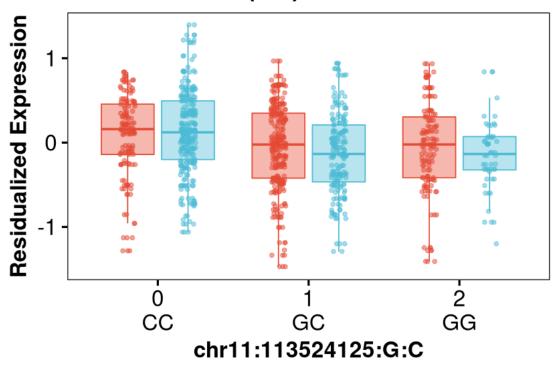
0.1383174

→prefix)

#print(eqtl_annot)

DRD2 exon 1L ENSE00002282098.1

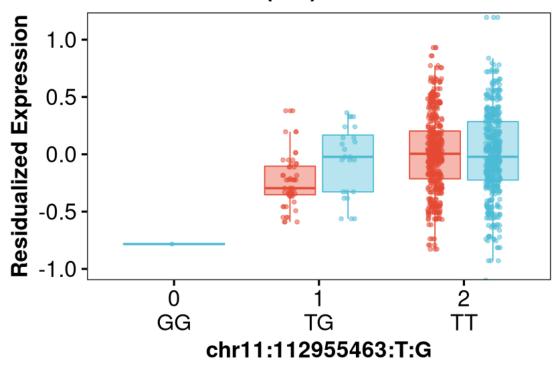
eQTL (AA) Ifsr: 0.073 eQTL (EA) Ifsr: 0.0045



Race 🖶 AA 😑 EA

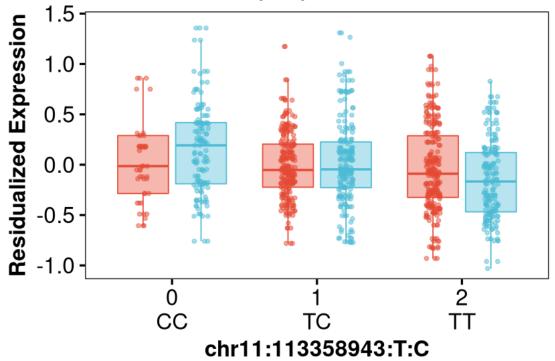
DRD2 exon 8
ENSE00002210277.1

eQTL (AA) Ifsr: 0.076 eQTL (EA) Ifsr: 0.086



Race = AA = EA

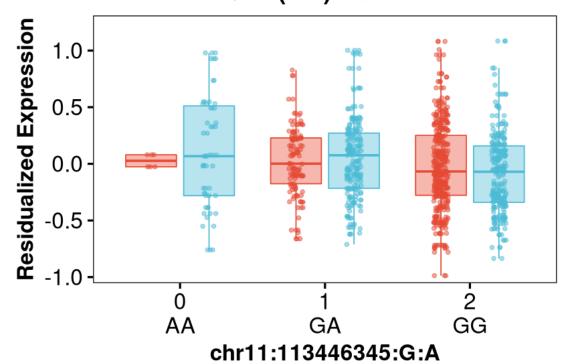
DRD2 exon 7 ENSE00001247012.2 eQTL (AA) Ifsr: 0.13 eQTL (EA) Ifsr: 0.0039



Race = AA = EA

DRD2 exon 2 ENSE00003665261.1 eQTL (AA) Ifsr: 0.47

eQTL (EA) lfsr: 0.47



Race 🖶 AA 🖶 EA

1.2.2 GWAS association

```
[13]: eGenes_gwas = get_eqtl_gwas_df() %>% filter(gene_id %in% all_drd2_exons) %>% arrange(EA, AA, P) %>% group_by(gene_id) %>% slice(1) %>% arrange(EA, AA, P) eGenes_gwas
```

Warning message in cbind(parts\$left, ellip_h, parts\$right, deparse.level = OL):
"number of rows of result is not a multiple of vector length (arg 2)"
Warning message in cbind(parts\$left, ellip_h, parts\$right, deparse.level = OL):
"number of rows of result is not a multiple of vector length (arg 2)"
Warning message in cbind(parts\$left, ellip_h, parts\$right, deparse.level = OL):
"number of rows of result is not a multiple of vector length (arg 2)"
Warning message in cbind(parts\$left, ellip_h, parts\$right, deparse.level = OL):

"number of rows of result is not a multiple of vector length (arg 2)"

gene_id variant_id AA EAV1chrNcmpos our_counted A grouped_df: 0×28 <chr><chr><dbl><dbl><int><int><int><int><chr>

1.3 Session Info

```
[14]: Sys.time()
      proc.time()
      options(width = 120)
      sessioninfo::session_info()
     [1] "2022-03-08 19:36:23 EST"
                 system elapsed
         user
     4868.976
                268.288 1938.636
     $platform $version 'R version 4.1.2 (2021-11-01)'
          $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 '2022-03-08'
          $pandoc '2.14.1 @ /usr/bin/pandoc'
```

		package	ondiskversion	loadedversion	path
		<chr></chr>	<chr></chr>	<chr></chr>	<chr></chr>
	abind	abind	1.4.5	1.4-5	/home/jb
	assertthat	assertthat	0.2.1	0.2.1	/home/jb
	backports	backports	1.4.1	1.4.1	/home/jb
	base64enc	base64enc	0.1.3	0.1 - 3	/home/jb
	broom	broom	0.7.12	0.7.12	/home/jb
	cachem	cachem	1.0.6	1.0.6	/home/jb
	car	car	3.0.12	3.0 - 12	/home/jb
	$\operatorname{carData}$	carData	3.0.5	3.0-5	/home/jb
	cellranger	cellranger	1.1.0	1.1.0	/home/jb
	cli	cli	3.2.0	3.2.0	/home/jb
	colorspace	colorspace	2.0.3	2.0 - 3	/home/jb
	crayon	crayon	1.5.0	1.5.0	/home/jb
	data.table	data.table	1.14.2	1.14.2	/home/jb
	DBI	DBI	1.1.2	1.1.2	/home/jb
	dbplyr	dbplyr	2.1.1	2.1.1	/home/jb
	digest	digest	0.6.29	0.6.29	/home/jb
	dplyr	dplyr	1.0.8	1.0.8	/home/jb
	ellipsis	ellipsis	0.3.2	0.3.2	/home/jb
	evaluate	evaluate	0.15	0.15	/home/jb
	fansi	fansi	1.0.2	1.0.2	/home/jb
	farver	farver	2.1.0	2.1.0	/home/jb
	fastmap	fastmap	1.1.0	1.1.0	/home/jb
	forcats	forcats	0.5.1	0.5.1	/home/jb
	fs	fs	1.5.2	1.5.2	/home/jb
	generics	generics	0.1.2	0.1.2	/home/jb
	ggplot2	ggplot2	3.3.5	3.3.5	/home/jb
	ggpubr	ggpubr	0.4.0	0.4.0	/home/jb
	ggsci	ggsci	2.9	2.9	/home/jb
	ggsignif	ggsignif	0.6.3	0.6.3	/home/jb
\$packages A packages_info: 78×11	glue	glue	1.6.1	1.6.1	/home/jb
	purrr	purrr	0.3.4	0.3.4	/home/jb
	R.methodsS3	R.methodsS3	1.8.1	1.8.1	/home/jb
	R.oo	R.oo	1.24.0	1.24.0	/home/jb
	R.utils	R.utils	2.11.0	2.11.0	/home/jb
	R6	R6	2.5.1	2.5.1	/home/jb
	Rcpp	Rcpp	1.0.8	1.0.8	/home/jb
	readr	readr	2.1.2	2.1.2	/home/jb
	readxl	readxl	1.3.1	1.3.1	/home/jb
	repr	repr	1.1.4	1.1.4	/home/jb
	reprex	reprex	2.0.1	2.0.1	/home/jb
	rlang	rlang	1.0.1	1.0.1	/home/jb
	rstatix	rstatix	0.7.0	0.7.0	/home/jb
	rstudioapi	rstudioapi	0.13	0.13	/home/jb
	rvest	rvest	1.0.2	1.0.2	/home/jb
	scales	scales	1.1.1	1.1.1	/home/jb
	sessioninfo	sessioninfo	1.2.2	1.2.2	/home/jb
	st ri ngi	stringi	1.7.6	1.7.6	/home/jb
	stringr	stringr	1.4.0	1.4.0	/home/jb
	svglite	svglite	2.1.0	2.1.0	/home/jb
	systemfonts	systemfonts	1.0.4	1.0.4	/home/jb