

# GWAS\_vis\_vignette

*Arcadio*

*12/13/2018*

Loading up packages

```
library(rJava)
library(GenomicRanges)
```

```
## Loading required package: stats4
## Loading required package: BiocGenerics
## Loading required package: parallel
##
## Attaching package: 'BiocGenerics'
## The following objects are masked from 'package:parallel':
##
##   clusterApply, clusterApplyLB, clusterCall, clusterEvalQ,
##   clusterExport, clusterMap, parApply, parCapply, parLapply,
##   parLapplyLB, parRapply, parSapply, parSapplyLB
## The following objects are masked from 'package:rJava':
##
##   anyDuplicated, duplicated, sort, unique
## The following objects are masked from 'package:stats':
##
##   IQR, mad, sd, var, xtabs
## The following objects are masked from 'package:base':
##
##   anyDuplicated, append, as.data.frame, basename, cbind,
##   colMeans, colnames, colSums, dirname, do.call, duplicated,
##   eval, evalq, Filter, Find, get, grep, grepl, intersect,
##   is.unsorted, lapply, lengths, Map, mapply, match, mget, order,
##   paste, pmax, pmax.int, pmin, pmin.int, Position, rank, rbind,
##   Reduce, rowMeans, rownames, rowSums, sapply, setdiff, sort,
##   table, tapply, union, unique, unsplit, which, which.max,
##   which.min
## Loading required package: S4Vectors
##
## Attaching package: 'S4Vectors'
## The following object is masked from 'package:base':
##
##   expand.grid
## Loading required package: IRanges
## Loading required package: GenomeInfoDb
library(SummarizedExperiment)
```

```
## Loading required package: Biobase
## Welcome to Bioconductor
##
##     Vignettes contain introductory material; view with
##     'browseVignettes()'. To cite Bioconductor, see
##     'citation("Biobase")', and for packages 'citation("pkgname)".
## Loading required package: DelayedArray
## Loading required package: matrixStats
##
## Attaching package: 'matrixStats'
## The following objects are masked from 'package:Biobase':
##
##     anyMissing, rowMedians
## Loading required package: BiocParallel
##
## Attaching package: 'DelayedArray'
## The following objects are masked from 'package:matrixStats':
##
##     colMaxs, colMins, colRanges, rowMaxs, rowMins, rowRanges
## The following objects are masked from 'package:base':
##
##     aperm, apply
library(GWASpoly)
library(ggplot2)
library(ggrepel)
library(stringr)

multiple_trait_manhattan <-function(traitGWASresults){
  ggplot(traitGWASresults) + geom_point(aes(Position, markerLogPVal, col = trait), alpha=0.3) + geom_po
}
```

Set directory, loading up start functions and rTassel

```
is_experimental <- TRUE

#set workdir for rtassel
setwd("~/myBins/bucklerlabBitbucket/rtassel/")

path_exp_tassel <- paste0(getwd(), "/inst/java/sTASSEL.jar")
path_exp_tassel_libs <- paste0(getwd(), "/inst/java/lib")

## jinit
rJava::.jinit(parameters="-Xmx6g")
.jcall(.jnew("java/lang/Runtime"), "J", "totalMemory")

## [1] 257425408
.jcall(.jnew("java/lang/Runtime"), "J", "maxMemory")

## [1] 5726797824
```

```
## Add class paths
if(is_experimental == TRUE) {
  tasselpath <- path_exp_tassel
  tassellibs <- path_exp_tassel_libs
}

rJava::.jaddClassPath(tasselpath)
rJava::.jaddClassPath(tassellibs)
print(.jclassPath())

## [1] "/Users/jav246/myBins/R-packages/rJava/java"
## [2] "/Users/jav246/myBins/bucklerlabBitbucket/rtassel/inst/java/sTASSEL.jar"
## [3] "/Users/jav246/myBins/bucklerlabBitbucket/rtassel/inst/java/lib"

## Source files
source("R/AllGenerics.R")
source("R/AllClasses.R")
source("R/TasselPluginWrappers.R")
source("R/PullFunctions.R")
source("R/gwasPolyObjectCreator.R")
```

Load up genotypes implementing Tassel code through rJava

```
geno_fileName <- "/Users/jav246/myBins/bucklerlabBitbucket/rtassel/data/mdp_genotype.hmp.txt"
```

```
## Make genotype table from tassels sample data
tasGenoTable <- readGenotypeTable(geno_fileName)
```

```
## Make summarized experiment from genotypetable
tas_se <- summarizeExperimentFromGenotypeTable(tasGenoTable) # not working right now, but
```

```
## Extracting chromosome names for each position...
## ...is there a quicker way to get this? (~ Brandon)
```

```
tas_se
```

```
## class: RangedSummarizedExperiment
## dim: 3093 281
## metadata(0):
## assays(1): ''
## rownames: NULL
## rowData names(0):
## colnames: NULL
## colData names(1): Sample
```

Load phenotype data

```
####straight load as dataframe, skipping first two rows on tassels specific phenotype table format
pheno_fileName <- "/Users/jav246/myBins/bucklerlabBitbucket/rtassel/data/mdp_phenotype.txt"
phenos <- read.table(file = pheno_fileName, skip = 2, header = T, sep = "\t", na.strings = "NaN")
summary(phenos)
```

##	Taxa	location	EarHT	dpoll	EarDia
##	33-16	: 2	A:283	Min. : 6.40	Min. :52.60
##	38-11	: 2	B:280	1st Qu.: 48.50	1st Qu.:63.50
##	4226	: 2		Median : 60.20	Median :67.50
##	4722	: 2		Mean : 61.58	Mean :67.78

```
## A188 : 2 3rd Qu.: 72.50 3rd Qu.:71.50 3rd Qu.:40.09
## A214N : 2 Max. :138.80 Max. :85.80 Max. :49.30
## (Other):551 NA's :4 NA's :7 NA's :37
## Q1 Q2 Q3
## Min. :0.0010 Min. :0.0010 Min. :0.0000
## 1st Qu.:0.0020 1st Qu.:0.0050 1st Qu.:0.0020
## Median :0.0100 Median :0.5700 Median :0.0190
## Mean :0.1744 Mean :0.5011 Mean :0.3245
## 3rd Qu.:0.1205 3rd Qu.:0.9680 3rd Qu.:0.7940
## Max. :0.9990 Max. :0.9980 Max. :0.9980
##
```

```
#### select sinlge location, as GWASpoly requires single entries for taxa.
phenosOneLoc <- phenos[phenos$location == "A",]
rownames(phenosOneLoc) <- phenosOneLoc$Taxa
####remove location as it is now redundant.
####Also, GWASpoly expects all traits as initial columns, and fixed effect covariates last
phenosOneLoc <- phenosOneLoc[,-c(2)]
```

```
summary(phenosOneLoc)
```

```
## Taxa EarHT dpoll EarDia
## 33-16 : 1 Min. : 8.00 Min. :54.50 Min. :23.72
## 38-11 : 1 1st Qu.: 48.12 1st Qu.:64.00 1st Qu.:34.86
## 4226 : 1 Median : 60.50 Median :67.50 Median :37.32
## 4722 : 1 Mean : 61.75 Mean :67.75 Mean :37.20
## A188 : 1 3rd Qu.: 73.00 3rd Qu.:71.50 3rd Qu.:40.02
## A214N : 1 Max. :136.00 Max. :85.00 Max. :46.35
## (Other):277 NA's :1 NA's :3 NA's :33
## Q1 Q2 Q3
## Min. :0.0010 Min. :0.001 Min. :0.0000
## 1st Qu.:0.0020 1st Qu.:0.005 1st Qu.:0.0020
## Median :0.0090 Median :0.579 Median :0.0230
## Mean :0.1728 Mean :0.502 Mean :0.3253
## 3rd Qu.:0.1160 3rd Qu.:0.968 3rd Qu.:0.7940
## Max. :0.9990 Max. :0.998 Max. :0.9980
##
```

Run GWASpoly

```
## create GWASpoly object with coopted read.GWASpoly function
data_gwasPoly <- se_createGWASpolyObject(ploidy = 2, phenoDF = phenosOneLoc,
                                         SummarizedExperimentObject = tas_se,
                                         format = "numeric", n.traits = 3)
```

```
## Number of polymorphic markers: 3093
## Missing marker data imputed with population mode
## N = 264 individuals with phenotypic and genotypic information
## Detected following fixed effects:
## Q1
## Q2
## Q3
## Detected following traits:
## EarHT
## dpoll
## EarDia
```

```
## add kinship information to object
data_gwasPoly <- set.K(data_gwasPoly)

## set parameters for mixed model
params <- set.params(fixed=unlist(strsplit("Q1,Q2,Q3", ",")),
                     fixed.type=rep("numeric",3))

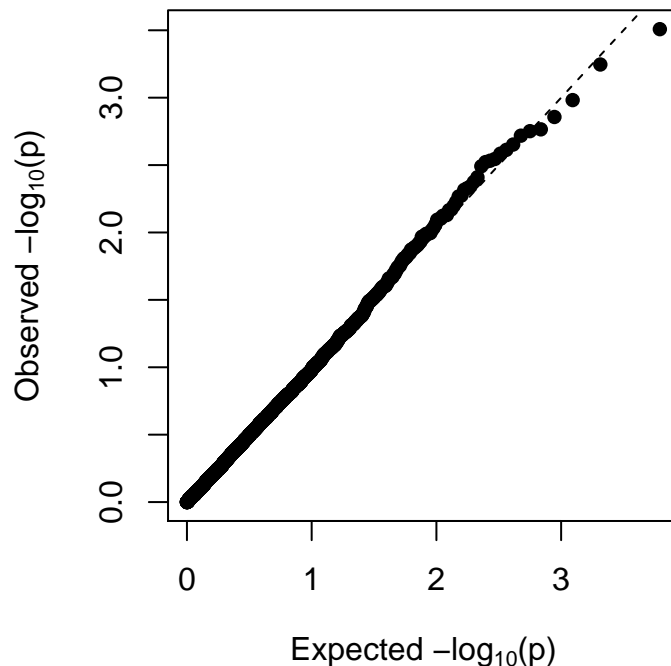
## run gwas
data_gwasPoly_res <- GWASpoly(data = data_gwasPoly, models = "additive",
                              params = params)
```

```
## Analyzing trait: EarHT
## P3D approach: Estimating variance components...Completed
## Testing markers for model: additive
## Analyzing trait: dpoll
## P3D approach: Estimating variance components...Completed
## Testing markers for model: additive
## Analyzing trait: EarDia
## P3D approach: Estimating variance components...Completed
## Testing markers for model: additive
```

Create GWASpoly plots and set thresholds to get QTLs

```
qq.plot(data_gwasPoly_res, trait = "EarDia", model = "additive")
```

### EarDia (additive)



```
## NULL
```

```
#can set Bonferroni and own pvalue
data_gwasPoly_res <- set.threshold(data_gwasPoly_res, method = "FDR", level=0.05)

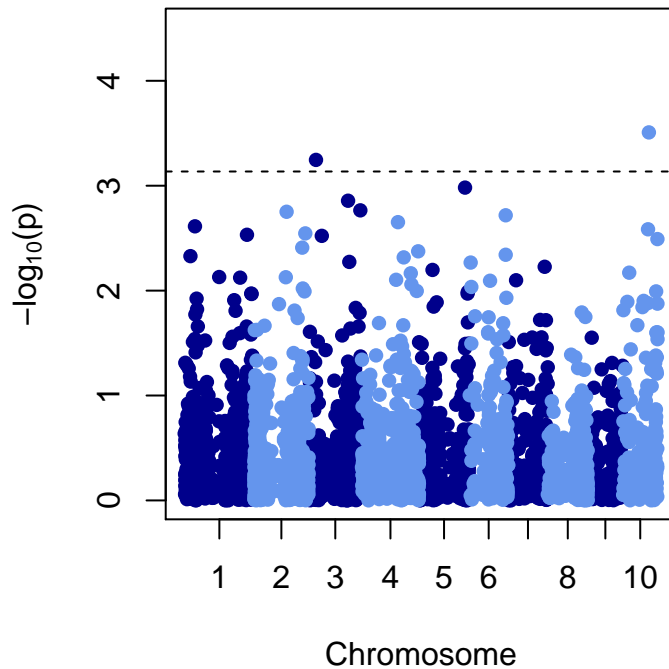
get.QTL(data = data_gwasPoly_res)
```

```
##      Trait      Model Threshold      Marker Chrom  Position Ref Alt Score
## 987 EarDia additive      3.14 dummy-987      3  31695534  0  1  3.25
## 2987 EarDia additive      3.14 dummy-2987    10 111608788  0  1  3.51
##      Effect
## 987    -1.08
## 2987   -1.15
```

*#can set any of the 3 traits*

```
manhattan.plot(data_gwasPoly_res, trait = "EarDia", model = "additive")
```

### EarDia (additive)



```
## NULL
```

Unwrap gwaspoly results class object

```
traitGWASresults <- data.frame()
for(trait in names(data_gwasPoly_res@scores)){
  message(paste("Getting results for:", trait))
  traitMarkerpScores <- as.data.frame(data_gwasPoly_res@scores[trait])
  Marker <- rownames(traitMarkerpScores)
  colnames(traitMarkerpScores) <- "markerLogPVal"
  traitMarkerpVals <- exp(traitMarkerpScores*-1)
  colnames(traitMarkerpVals) <- "markerpVal"
  traitMarkerEffects <- as.data.frame(data_gwasPoly_res@effects[trait])
  colnames(traitMarkerEffects) <- "markerEffect"
  sigTreshhold <- data_gwasPoly_res@threshold[rownames(data_gwasPoly_res@threshold)==trait]
  traitGWASresults <- rbind(traitGWASresults, data.frame(Marker, traitMarkerEffects, traitMarkerpVals,
})
```

```
## Getting results for: EarHT
## Getting results for: dpoll
## Getting results for: EarDia
```

```
summary(traitGWASresults)
```

```
##           Marker      markerEffect      markerpVal      markerLogPVal
## dummy-1      : 3   Min.      :-4.37836   Min.      :0.02396   Min.      :0.000007
## dummy-10     : 3   1st Qu.: -0.29295   1st Qu.: 0.54946   1st Qu.: 0.127346
## dummy-100    : 3   Median : 0.02879   Median : 0.73557   Median : 0.307111
## dummy-1000   : 3   Mean      : 0.07314   Mean      : 0.69642   Mean      : 0.434249
## dummy-1001   : 3   3rd Qu.: 0.36503   3rd Qu.: 0.88043   3rd Qu.: 0.598824
## dummy-1002   : 3   Max.      : 5.34599   Max.      : 0.99999   Max.      : 3.731254
## (Other)      :9261
##           trait      sigTreshold
## EarHT :3093   Min.      :3.135
## dpoll :3093   1st Qu.: 3.135
## EarDia:3093   Median : 4.023
##           Mean      :3.825
##           3rd Qu.: 4.317
##           Max.      :4.317
##
```

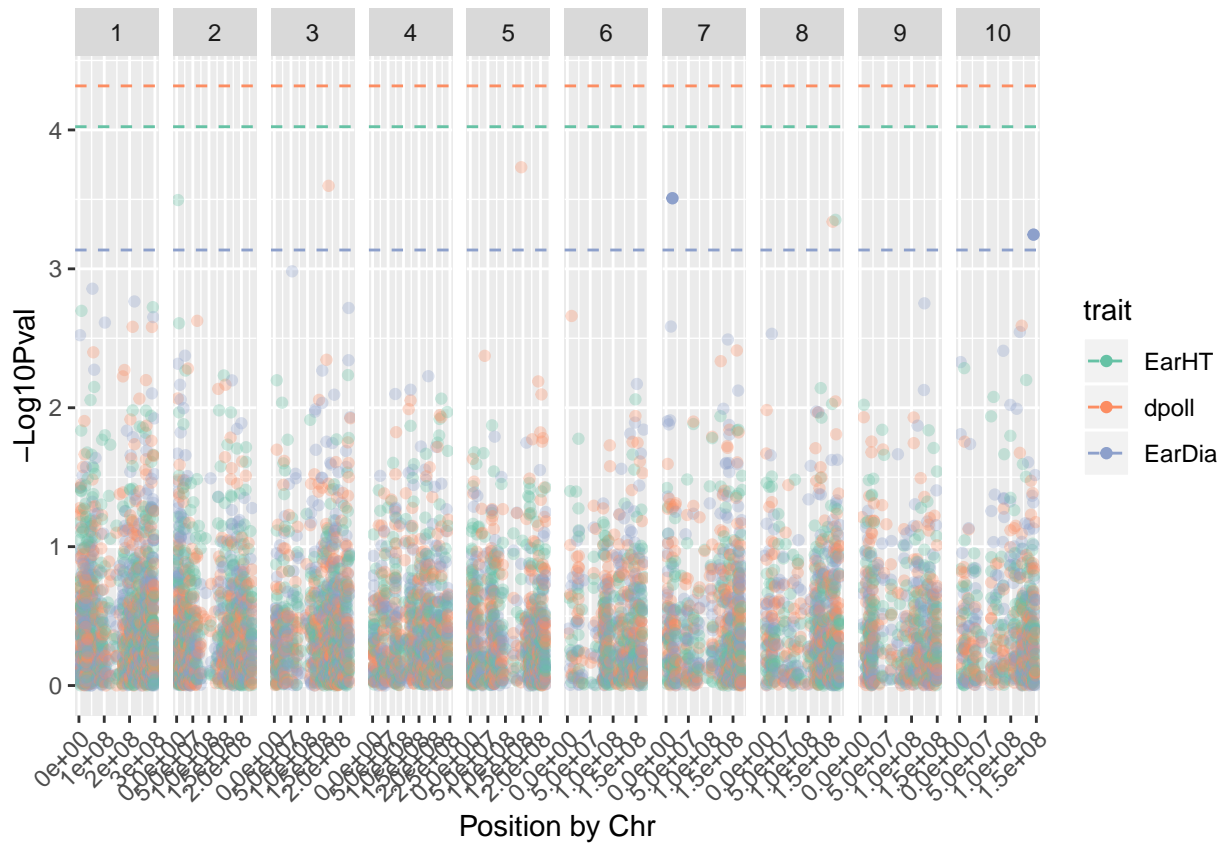
```
traitGWASresults <- merge(traitGWASresults, data_gwasPoly_res@map, by = "Marker")
```

```
summary(traitGWASresults)
```

```
##           Marker      markerEffect      markerpVal      markerLogPVal
## dummy-1      : 3   Min.      :-4.37836   Min.      :0.02396   Min.      :0.000007
## dummy-10     : 3   1st Qu.: -0.29295   1st Qu.: 0.54946   1st Qu.: 0.127346
## dummy-100    : 3   Median : 0.02879   Median : 0.73557   Median : 0.307111
## dummy-1000   : 3   Mean      : 0.07314   Mean      : 0.69642   Mean      : 0.434249
## dummy-1001   : 3   3rd Qu.: 0.36503   3rd Qu.: 0.88043   3rd Qu.: 0.598824
## dummy-1002   : 3   Max.      : 5.34599   Max.      : 0.99999   Max.      : 3.731254
## (Other)      :9261
##           trait      sigTreshold      Chrom      Position
## EarHT :3093   Min.      :3.135      1      :1620   Min.      : 139753
## dpoll :3093   1st Qu.: 3.135      2      :1179   1st Qu.: 43868122
## EarDia:3093   Median : 4.023      5      :1071   Median :128402775
##           Mean      :3.825      3      :1065   Mean      :119893324
##           3rd Qu.: 4.317      4      : 957   3rd Qu.:175159119
##           Max.      :4.317      8      : 768   Max.      :299170077
##           (Other):2619
##           Ref      Alt
## Min.      :0      Min.      :1
## 1st Qu.:0      1st Qu.:1
## Median :0      Median :1
## Mean      :0      Mean      :1
## 3rd Qu.:0      3rd Qu.:1
## Max.      :0      Max.      :1
##
```

Create simple manhattan like plot for all traits

```
multiple_trait_manhattan(traitGWASresults = traitGWASresults)
```



Parse GFF file to get genes and create GenomicRanges object

```
maizeGFF <- read.table("~/Box/projectMaize/PHG/cimmyt_assemblies_analy/b73/Zea_mays.AGPv4.40.chr.gff3",
  colnames(maizeGFF) <- c("Chr", "Source", "annotType", "Start", "End", "other", "Strand", "other2", "IDs")

maizeGFF$Chr <- as.factor(maizeGFF$Chr)

maizeGFFgenes <- maizeGFF[maizeGFF$annotType=="gene",]

maizeGFFgenes$Gene <- str_split(str_split(maizeGFFgenes$IDs, ";", simplify = T)[,1], ":", simplify = T)[,1]

head(maizeGFFgenes)
```

##	Chr	Source	annotType	Start	End	other	Strand	other2
## 2	1	gramene	gene	44289	49837	.	+	.
## 24	1	gramene	gene	50877	55716	.	-	.
## 170	1	gramene	gene	92299	95134	.	-	.
## 184	1	gramene	gene	111655	118312	.	-	.
## 217	1	gramene	gene	118683	119739	.	-	.
## 231	1	gramene	gene	122120	122614	.	+	.
##								
## 2	ID=gene:Zm00001d027230;biotype=protein_coding;description=Mitochondrial transcription termination							
## 24	ID=gene:Zm00001d027231;biotype=protein_coding;description=OSJNBa009300							
## 170	ID=gene:Zm00001d027232							
## 184	ID=gene:Zm00001d027233							
## 217	ID=gene:Zm00001d027234							
## 231	ID=gene:Zm00001d027235;biotype=protein_coding;description=Pentatricopeptide rep							
##	Gene							



```
## 2   Zm00001d027230
## 24  Zm00001d027231
## 170 Zm00001d027232
## 184 Zm00001d027233
## 217 Zm00001d027234
## 231 Zm00001d027235
```

```
maizeGFFgenesGR <- makeGRangesFromDataFrame(maizeGFFgenes, keep.extra.columns = T)
```

```
maizeGFFgenesGR
```

```
## GRanges object with 39179 ranges and 6 metadata columns:
```

```
##      seqnames      ranges strand |   Source annotType   other
##      <Rle>       <IRanges> <Rle> | <factor>  <factor> <factor>
##      2          1   44289-49837   + |   gramene    gene      .
##     24          1   50877-55716   - |   gramene    gene      .
##    170          1   92299-95134   - |   gramene    gene      .
##    184          1  111655-118312   - |   gramene    gene      .
##    217          1  118683-119739   - |   gramene    gene      .
##     ...      ...      ...      ... |   ...      ...      ...
## 2804827      Pt 134341-134862   - |   gramene    gene      .
## 2804831      Pt 134923-135222   - |   gramene    gene      .
## 2804835      Pt 138323-139807   + |   gramene    gene      .
## 2804849      Pt 139824-140048   + |   gramene    gene      .
## 2804853      Pt 140068-140361   + |   gramene    gene      .
```

```
##      other2
##      <factor>
##      2      .
##     24      .
##    170      .
##    184      .
##    217      .
##     ...      ...
## 2804827      .
## 2804831      .
## 2804835      .
## 2804849      .
## 2804853      .
```

```
##
##
##      2      ID=gene:Zm00001d027230;biotype=protein_coding;description=
##     24      ID=gene:Zm00001d027231;biotype=p
##    170
##    184
##    217
##     ...
## 2804827      ID=gene:GRMZM5G885905;Name=ycf73-A;biotype=protein_coding;description=Uncharacter
## 2804831 ID=gene:GRMZM5G866761;Name=ycf15-A;biotype=protein_coding;description=Putative uncharacter
## 2804835
## 2804849
## 2804853
##
##      Gene
##      <character>
##      2 Zm00001d027230
##     24 Zm00001d027231
```

```
##      170 Zm00001d027232
##      184 Zm00001d027233
##      217 Zm00001d027234
##      ...
## 2804827 GRMZM5G885905
## 2804831 GRMZM5G866761
## 2804835 GRMZM5G818111
## 2804849 GRMZM5G866064
## 2804853 GRMZM5G855343
## -----
## seqinfo: 12 sequences from an unspecified genome; no seqlengths
```

Make GenomicRanges object out of gwas results

```
traitGWASresultsGR <- makeGRangesFromDataFrame(traitGWASresults, seqnames.field = "Chrom", start.field =
traitGWASresultsGR
```

```
## GRanges object with 9279 ranges and 8 metadata columns:
##      seqnames      ranges strand |      Marker      markerEffect
##      <Rle> <IRanges> <Rle> | <factor>      <numeric>
##      [1]      1      157104      * | dummy-1      0.343306778643111
##      [2]      1      157104      * | dummy-1      0.454322818890773
##      [3]      1      157104      * | dummy-1      0.357036991759809
##      [4]      1      3206090     * | dummy-10     0.721473858204604
##      [5]      1      3206090     * | dummy-10    -0.0265087099725783
##      ...      ...      ...      ... .      ...      ...
## [9275]      3      43868043     * | dummy-998     0.571750799793797
## [9276]      3      43868043     * | dummy-998     2.09836224957921
## [9277]      3      43868067     * | dummy-999    -0.112171180072146
## [9278]      3      43868067     * | dummy-999     0.686141690548524
## [9279]      3      43868067     * | dummy-999     2.57171471115537
##      markerpVal      markerLogPVal      trait      sigThreshold
##      <numeric>      <numeric> <factor>      <numeric>
##      [1] 0.912009425744584 0.0921049537149598 EarHT 4.02323519245304
##      [2] 0.495647822750053 0.701889639229406 dpoll 4.31685670024708
##      [3] 0.528853881117141 0.637043102448693 EarDia 3.13549015264627
##      [4] 0.240756564892385 1.42396896017148 dpoll 4.31685670024708
##      [5] 0.968201579430578 0.032314970171761 EarDia 3.13549015264627
##      ...      ...      ...      ...
## [9275] 0.391523703112522 0.937709221035299 dpoll 4.31685670024708
## [9276] 0.438150807877916 0.825192117643301 EarHT 4.02323519245304
## [9277] 0.865945729907151 0.1439330399271 EarDia 3.13549015264627
## [9278] 0.292826155607051 1.22817617162555 dpoll 4.31685670024708
## [9279] 0.329427758725234 1.11039819422075 EarHT 4.02323519245304
##      Ref      Alt
##      <numeric> <numeric>
##      [1]      0      1
##      [2]      0      1
##      [3]      0      1
##      [4]      0      1
##      [5]      0      1
##      ...      ...
## [9275]      0      1
## [9276]      0      1
```

```
##      [9277]          0          1
##      [9278]          0          1
##      [9279]          0          1
##      -----
##      seqinfo: 10 sequences from an unspecified genome; no seqlengths

Subset geneRanges with gwasRanges. Finding genes with SNPs in them
geneGWAShits <- findOverlaps(maizeGFFgenesGR, traitGWASresultsGR)
geneGWAShits

## Hits object with 909 hits and 0 metadata columns:
##      queryHits subjectHits
##      <integer>  <integer>
##      [1]        145        2335
##      [2]        145        2336
##      [3]        145        2337
##      [4]        145        2668
##      [5]        145        2669
##      ...         ...         ...
##      [905]       38508        6254
##      [906]       38508        6255
##      [907]       38550        6271
##      [908]       38550        6272
##      [909]       38550        6273
##      -----
##      queryLength: 39179 / subjectLength: 9279

genesInGWASranges <- maizeGFFgenesGR[unique(queryHits(geneGWAShits)), ]
genesInGWASranges

## GRanges object with 157 ranges and 6 metadata columns:
##      seqnames          ranges strand | Source annotType
##      <Rle>             <IRanges> <Rle> | <factor> <factor>
##      25628             1      4832874-4838381      + | gramene      gene
##      25882             1      4909652-4914685      - | gramene      gene
##      51549             1      18907931-18909955     + | gramene      gene
##      56160             1      22599568-22606524     + | gramene      gene
##      57715             1      23265205-23270017     + | gramene      gene
##      ...             ...             ...     ... | ...         ...
##      2692010           9      109895426-109918891    + | gramene      gene
##      2717909           9      130889521-130890480    - | gramene      gene
##      2741832           9      142497782-142500049    - | gramene      gene
##      2752056           9      147127843-147139955    - | gramene      gene
##      2754518           9      148427300-148449087    + | gramene      gene
##      other            other2
##      <factor> <factor>
##      25628             .             .
##      25882             .             .
##      51549             .             .
##      56160             .             .
##      57715             .             .
##      ...             ...             ...
##      2692010           .             .
##      2717909           .             .
```

```

## 2741832 . .
## 2752056 . .
## 2754518 . .
##
##
## 25628 ID=gene:Zm00001d027415;Name=autophagy1a;biotype=protein_coding;description=Serine/
## 25882 ID=gene:Zm00001d027423;Name=flavone synthase typeI2;biotype=protein_coding;descrip
## 51549 ID=gene:Zm00001d027973;Name=flavone synthase typeI2;biotype=protein_coding;descrip
## 56160 ID=gene:Zm00001d028088;biotype=protein_coding;description=Ubiquitin
## 57715 ID=gene:Zm00001d028114;biotype=protein_coding;description=n
## ...
## 2692010 ID=gene:Zm00001d046905;biotype=prote
## 2717909 ID=gene:Zm00001d047460;biotype=protein_coding;description=Haloacid dehalogenase-like hydro
## 2741832 ID=gene:Zm00001d047814;biotype=protein_coding;desc
## 2752056 ID=gene:Zm00001d047983;biotype=protein_coding;description=LisH and Ran
## 2754518 ID=gene:Zm00001d048031;biotype=protein_coding;description=Calmodulin-bin
##
## Gene
## <character>
## 25628 Zm00001d027415
## 25882 Zm00001d027423
## 51549 Zm00001d027973
## 56160 Zm00001d028088
## 57715 Zm00001d028114
## ...
## 2692010 Zm00001d046905
## 2717909 Zm00001d047460
## 2741832 Zm00001d047814
## 2752056 Zm00001d047983
## 2754518 Zm00001d048031
## -----
## seqinfo: 12 sequences from an unspecified genome; no seqlengths

```

Annotating SNPs with their closest gene

```

genesNearestToGWASranges <- nearest(traitGWASresultsGR, maizeGFFgenesGR, ignore.strand = T)

distToNearest <- distanceToNearest(traitGWASresultsGR, maizeGFFgenesGR, ignore.strand = T)

genesNearestToGWASranges <- maizeGFFgenesGR[genesNearestToGWASranges]

genesNearestToGWASranges <- as.data.frame(genesNearestToGWASranges, row.names = NULL)

genesNearestToGWASranges$distanceToNearestAnnot <- distToNearest@elementMetadata$distance

traitGWASresults_temp <- traitGWASresults

colnames(traitGWASresults_temp) <- paste(colnames(traitGWASresults), "gwas", sep = "_")

genesNearestToGWASranges <- cbind(traitGWASresults_temp, genesNearestToGWASranges)

rm(traitGWASresults_temp)

genesNearestToGWASranges <- makeGRangesFromDataFrame(genesNearestToGWASranges, keep.extra.columns = T)

genesNearestToGWASranges

```

```
## GRanges object with 9279 ranges and 17 metadata columns:
##      seqnames      ranges strand | Marker_gwas
##      <Rle>        <IRanges> <Rle> | <factor>
##      [1]          1      138378-139043 - | dummy-1
##      [2]          1      138378-139043 - | dummy-1
##      [3]          1      138378-139043 - | dummy-1
##      [4]          1    3224423-3230647 + | dummy-10
##      [5]          1    3224423-3230647 + | dummy-10
##      ...          ...          ...   ...   ...
##      [9275]        3 43825329-43828852 - | dummy-998
##      [9276]        3 43825329-43828852 - | dummy-998
##      [9277]        3 43825329-43828852 - | dummy-999
##      [9278]        3 43825329-43828852 - | dummy-999
##      [9279]        3 43825329-43828852 - | dummy-999
##      markerEffect_gwas  markerpVal_gwas markerLogPVal_gwas
##      <numeric>          <numeric>          <numeric>
##      [1] 0.343306778643111 0.912009425744584 0.0921049537149598
##      [2] 0.454322818890773 0.495647822750053 0.701889639229406
##      [3] 0.357036991759809 0.528853881117141 0.637043102448693
##      [4] 0.721473858204604 0.240756564892385 1.42396896017148
##      [5] -0.0265087099725783 0.968201579430578 0.032314970171761
##      ...          ...          ...
##      [9275] 0.571750799793797 0.391523703112522 0.937709221035299
##      [9276] 2.09836224957921 0.438150807877916 0.825192117643301
##      [9277] -0.112171180072146 0.865945729907151 0.1439330399271
##      [9278] 0.686141690548524 0.292826155607051 1.22817617162555
##      [9279] 2.57171471115537 0.329427758725234 1.11039819422075
##      trait_gwas sigTreshold_gwas Chrom_gwas Position_gwas Ref_gwas
##      <factor>    <numeric>    <ordered>    <integer> <numeric>
##      [1] EarHT 4.02323519245304 1 157104 0
##      [2] dpoll 4.31685670024708 1 157104 0
##      [3] EarDia 3.13549015264627 1 157104 0
##      [4] dpoll 4.31685670024708 1 3206090 0
##      [5] EarDia 3.13549015264627 1 3206090 0
##      ...          ...          ...   ...   ...
##      [9275] dpoll 4.31685670024708 3 43868043 0
##      [9276] EarHT 4.02323519245304 3 43868043 0
##      [9277] EarDia 3.13549015264627 3 43868067 0
##      [9278] dpoll 4.31685670024708 3 43868067 0
##      [9279] EarHT 4.02323519245304 3 43868067 0
##      Alt_gwas Source annotType other other2
##      <numeric> <factor> <factor> <factor> <factor>
##      [1] 1 gramene gene . .
##      [2] 1 gramene gene . .
##      [3] 1 gramene gene . .
##      [4] 1 gramene gene . .
##      [5] 1 gramene gene . .
##      ...          ...          ...   ...   ...
##      [9275] 1 gramene gene . .
##      [9276] 1 gramene gene . .
##      [9277] 1 gramene gene . .
##      [9278] 1 gramene gene . .
##      [9279] 1 gramene gene . .
##
```

```
##
##      [1] ID=gene:Zm00001d027236;biotype=protein_coding;description=Di-glucose binding protein with K
##      [2] ID=gene:Zm00001d027236;biotype=protein_coding;description=Di-glucose binding protein with K
##      [3] ID=gene:Zm00001d027236;biotype=protein_coding;description=Di-glucose binding protein with K
##      [4] ID=gene:Zm00001d027337;biotype=protein_coding;description=Di-glucose binding protein with K
##      [5] ID=gene:Zm00001d027337;biotype=protein_coding;description=Di-glucose binding protein with K
##      ...
## [9275] ID=gene:Zm00001d040449;biotype=protein_coding;description=protein kinase C substrate l
## [9276] ID=gene:Zm00001d040449;biotype=protein_coding;description=protein kinase C substrate l
## [9277] ID=gene:Zm00001d040449;biotype=protein_coding;description=protein kinase C substrate l
## [9278] ID=gene:Zm00001d040449;biotype=protein_coding;description=protein kinase C substrate l
## [9279] ID=gene:Zm00001d040449;biotype=protein_coding;description=protein kinase C substrate l
##
##      Gene distanceToNearestAnnot
##      <character>      <integer>
##      [1] Zm00001d027236      18060
##      [2] Zm00001d027236      18060
##      [3] Zm00001d027236      18060
##      [4] Zm00001d027337      18332
##      [5] Zm00001d027337      18332
##      ...
## [9275] Zm00001d040449      39190
## [9276] Zm00001d040449      39190
## [9277] Zm00001d040449      39214
## [9278] Zm00001d040449      39214
## [9279] Zm00001d040449      39214
## -----
## seqinfo: 12 sequences from an unspecified genome; no seqlengths
```

Subset GWAS hits by significance for plotting with genes.

```
traitGWASresultsGR_sig <- traitGWASresultsGR[traitGWASresultsGR$markerLogPVal > traitGWASresultsGR$sigT]

traitGWASresultsGR_sig_df <- as.data.frame(traitGWASresultsGR_sig)

maxDistToAnnotation <- 3e5

genesInGWASranges_sigHits <- findOverlaps(genesInGWASranges, traitGWASresultsGR_sig, maxgap = maxDistToAnnotation)

genesInGWASranges_sigGenesGR <- genesInGWASranges[unique(queryHits(genesInGWASranges_sigHits)), ]

annotationPadding <- 1e3

annotationsDF <- data.frame()
for(seqName in as.character(seqnames(genesInGWASranges_sigGenesGR))){
  #working whole sequence present
  oneSeqRange <- genesInGWASranges_sigGenesGR[as.character(seqnames(genesInGWASranges_sigGenesGR)) == seqName]
  seqRangeStart <- start(oneSeqRange) - annotationPadding - maxDistToAnnotation
  seqRangeEnd <- end(oneSeqRange) + annotationPadding + maxDistToAnnotation

  annotationCount <- length(oneSeqRange)

  #working individual annotations
  annotationNum <- 0
  for(annotationName in oneSeqRange$Gene){
    annotationNum <- annotationNum + 1
  }
}
```

```

oneAnnotationRange <- oneSeqRange[oneSeqRange$Gene == annotationName,]

annotationStart <- start(oneAnnotationRange)
annotationEnd <- end(oneAnnotationRange)
annotationStrand <- strand(oneAnnotationRange)

annotationsDF <- rbind(annotationsDF, data.frame(seqname = seqName, start=seqRangeStart, end=seqRangeEnd, strand=annotationStrand))
}
ggplot(annotationsDF) + geom_rect(aes(xmin = seqRangeStart, xmax = seqRangeEnd, ymin = 0, ymax=1), fill="white", stroke="black")
}

annotationsDF_GR <- makeGRangesFromDataFrame(annotationsDF, keep.extra.columns = T)

traitGWASresultsGR_annotatons <- mergeByOverlaps(traitGWASresultsGR, annotationsDF_GR)

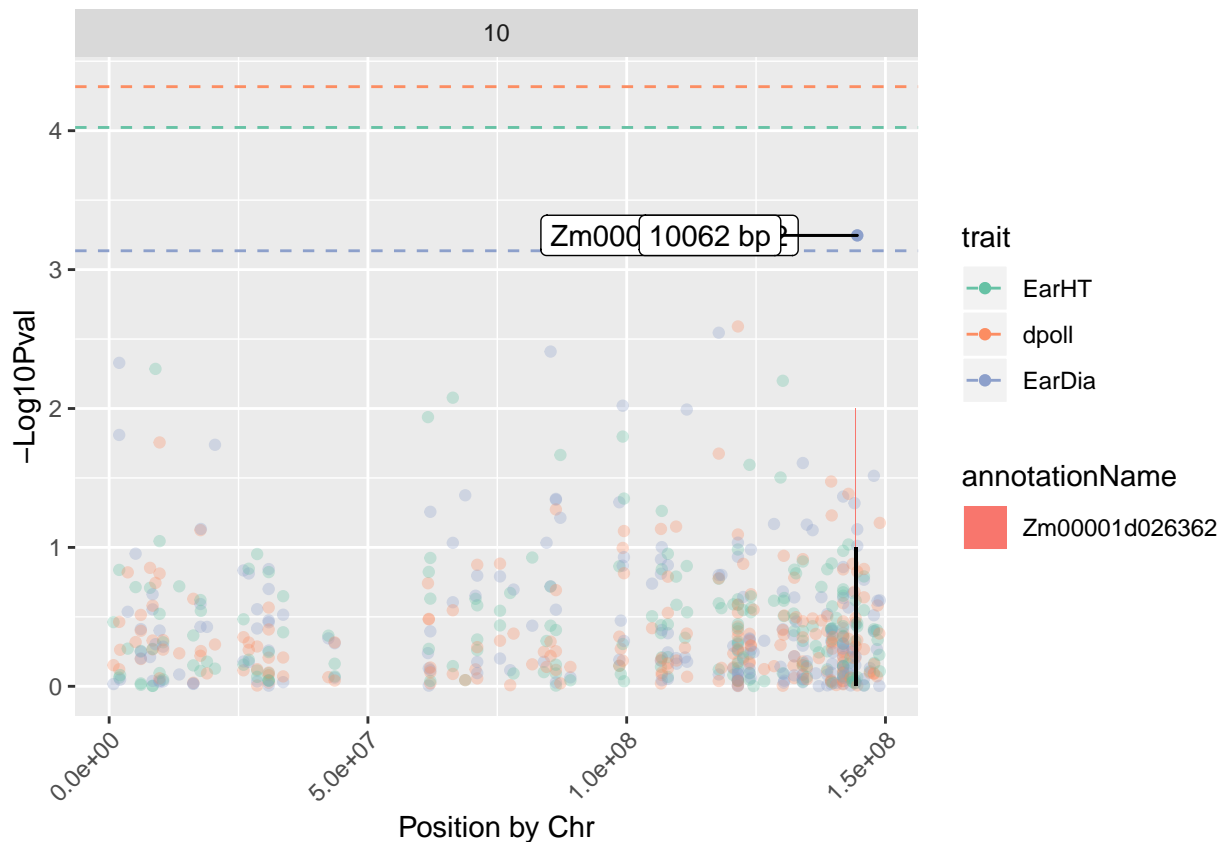
traitGWASresultsGR_annotatons_df <- as.data.frame(traitGWASresultsGR_annotatons)

traitGWASresultsGR_annotatons_df$MinDistToAnnotation <- min(abs(traitGWASresultsGR_annotatons_df$seqname - traitGWASresultsGR_annotatons_df$seqname))

traitGWASresultsGR_annotatons_df$MinDistToAnnotation[traitGWASresultsGR_annotatons_df$seqname == annotationName] <- 0

ggplot(subset(traitGWASresults, Chrom %in% traitGWASresultsGR_annotatons_df$traitGWASresultsGR.seqname))

```



Subset GWAS hits by significance for plotting with genes. Working to make multiple genes show up right

```

genesNearestToGWASranges$Gene_label <- as.character(genesNearestToGWASranges$Gene)

bestHitByAnnot <- data.frame()
for(annotationName in unique(genesNearestToGWASranges$Gene)){
  annotationData <- genesNearestToGWASranges[genesNearestToGWASranges$Gene == annotationName,]
  bestHit <- min(annotationData$markerpVal_gwas)
  bestHitByAnnot <- rbind(bestHitByAnnot, data.frame(Gene = annotationName, bestHitPval = bestHit))
}

genesNearestToGWASranges <- merge(genesNearestToGWASranges, bestHitByAnnot, by = "Gene")

#to only plot gene names for best gwas SNP
genesNearestToGWASranges$Gene_label <- as.character(genesNearestToGWASranges$Gene)
genesNearestToGWASranges$Gene_label[genesNearestToGWASranges$bestHitPval != genesNearestToGWASranges$ma
genesNearestToGWASranges$Gene_label <- as.factor(genesNearestToGWASranges$Gene_label)
summary(genesNearestToGWASranges$Gene_label)

```

```

##          Zm00001d001781 Zm00001d001798 Zm00001d001800 Zm00001d001802
##          7791          1          1          1          1
## Zm00001d001803 Zm00001d001887 Zm00001d001896 Zm00001d001912 Zm00001d001939
##          1          1          1          1          1
## Zm00001d001945 Zm00001d002004 Zm00001d002031 Zm00001d002058 Zm00001d002090
##          1          1          1          1          1
## Zm00001d002096 Zm00001d002109 Zm00001d002130 Zm00001d002135 Zm00001d002184
##          1          1          1          1          1
## Zm00001d002278 Zm00001d002307 Zm00001d002328 Zm00001d002336 Zm00001d002370
##          1          1          1          1          1
## Zm00001d002390 Zm00001d002396 Zm00001d002431 Zm00001d002456 Zm00001d002457
##          1          1          1          1          1
## Zm00001d002488 Zm00001d002524 Zm00001d002573 Zm00001d002621 Zm00001d002628
##          1          1          1          1          1
## Zm00001d002649 Zm00001d002706 Zm00001d002714 Zm00001d002735 Zm00001d002737
##          1          1          1          1          1
## Zm00001d002742 Zm00001d002762 Zm00001d002773 Zm00001d002778 Zm00001d002793
##          1          1          1          1          1
## Zm00001d002862 Zm00001d002873 Zm00001d002889 Zm00001d002937 Zm00001d002944
##          1          1          1          1          1
## Zm00001d002969 Zm00001d003033 Zm00001d003081 Zm00001d003093 Zm00001d003123
##          1          1          1          1          1
## Zm00001d003150 Zm00001d003188 Zm00001d003205 Zm00001d003231 Zm00001d003237
##          1          1          1          1          1
## Zm00001d003266 Zm00001d003292 Zm00001d003343 Zm00001d003357 Zm00001d003394
##          1          1          1          1          1
## Zm00001d003403 Zm00001d003423 Zm00001d003445 Zm00001d003452 Zm00001d003543
##          1          1          1          1          1
## Zm00001d003592 Zm00001d003646 Zm00001d003730 Zm00001d003735 Zm00001d003772
##          1          1          1          1          1
## Zm00001d003774 Zm00001d003794 Zm00001d003852 Zm00001d003855 Zm00001d003863
##          1          1          1          1          1
## Zm00001d003925 Zm00001d003929 Zm00001d003993 Zm00001d004105 Zm00001d004178
##          1          1          1          1          1
## Zm00001d004327 Zm00001d004342 Zm00001d004376 Zm00001d004560 Zm00001d004599
##          1          1          1          1          1

```



```
## Zm00001d004649 Zm00001d004667 Zm00001d004669 Zm00001d004689 Zm00001d004753
##           1           1           1           1           1
## Zm00001d004794 Zm00001d004800 Zm00001d004804 Zm00001d004829      (Other)
##           1           1           1           1           1390
```

```
genesNearestToGWASranges$distanceToNearestAnnot_label <- genesNearestToGWASranges$distanceToNearestAnnot_label[genesNearestToGWASranges$bestHitPval != genesNearestToGWASranges$bestHitPval]
```

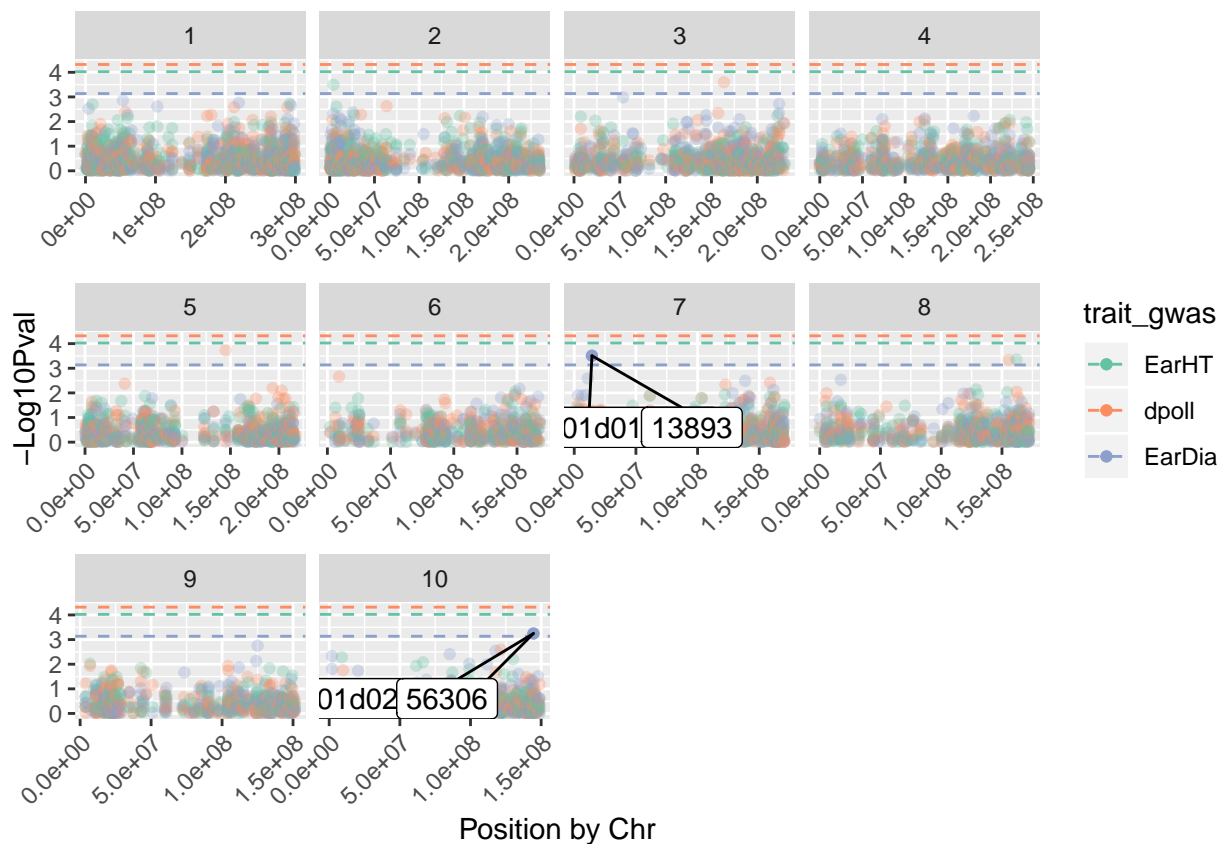
```
#to only plot gene names for significant SNPs
```

```
genesNearestToGWASranges$Gene_label <- as.character(genesNearestToGWASranges$Gene)
genesNearestToGWASranges$Gene_label[genesNearestToGWASranges$markerLogPVal_gwas < genesNearestToGWASranges$markerLogPVal_gwas] <- as.factor(genesNearestToGWASranges$Gene_label)
summary(genesNearestToGWASranges$Gene_label)
```

```
##           Zm00001d019055 Zm00001d026374
##           9277           1           1
```

```
genesNearestToGWASranges$distanceToNearestAnnot_label <- genesNearestToGWASranges$distanceToNearestAnnot_label[genesNearestToGWASranges$markerLogPVal_gwas < genesNearestToGWASranges$markerLogPVal_gwas]
```

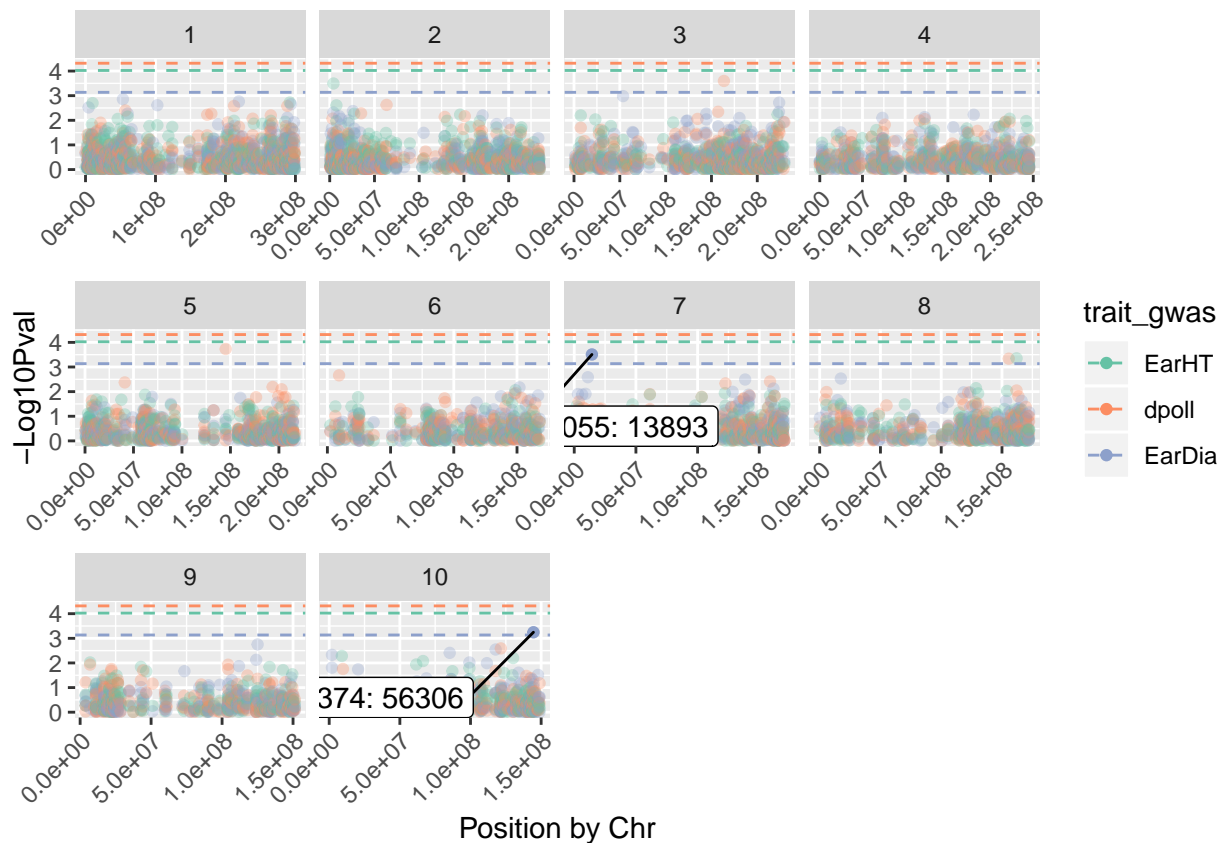
```
ggplot(genesNearestToGWASranges) + geom_point(aes(Position_gwas, markerLogPVal_gwas, col = trait_gwas),
```



```
genesNearestToGWASranges$single_label <- paste(genesNearestToGWASranges$Gene_label, genesNearestToGWASranges$single_label)
```

```
genesNearestToGWASranges$single_label <- gsub("^: $", "" ,genesNearestToGWASranges$single_label)
```

```
ggplot(genesNearestToGWASranges) + geom_point(aes(Position_gwas, markerLogPVal_gwas, col = trait_gwas),
```



```
#+ geom_rect(aes(xmin = seqRangeStart, xmax = seqRangeEnd, ymin = 0, ymax=1), fill = "black", data=anno
```

Subset GWAS hits by significance for plotting with genes. Working to make multiple genes show up right.  
Temp work not working

```
traitGWASresultsGR_sig <- traitGWASresultsGR[traitGWASresultsGR$markerLogPval > traitGWASresultsGR$sigT

traitGWASresultsGR_sig_df <- as.data.frame(traitGWASresultsGR_sig)

maxDistToAnnotation <- 3e5

genesInGWASranges_sigHits <- findOverlaps(genesInGWASranges, traitGWASresultsGR_sig, maxgap = maxDistTo

genesInGWASranges_sigGenesGR <- genesInGWASranges[unique(queryHits(genesInGWASranges_sigHits)), ]

annotationPadding <- 1e3

annotationsDF <- data.frame()
for(seqName in as.character(seqnames(genesInGWASranges_sigGenesGR))){
  #working whole sequence present
  oneSeqRange <- genesInGWASranges_sigGenesGR[as.character(seqnames(genesInGWASranges_sigGenesGR)) == s
  seqRangeStart <- start(oneSeqRange) - annotationPadding - maxDistToAnnotation
  seqRangeEnd <- end(oneSeqRange) + annotationPadding + maxDistToAnnotation

  annotationCount <- length(oneSeqRange)

  #working individual annotations
  annotationNum <- 0
```

```

for(annotationName in oneSeqRange$Gene){
  annotationNum <- annotationNum + 1
  oneAnnotationRange <- oneSeqRange[oneSeqRange$Gene == annotationName,]

  annotationStart <- start(oneAnnotationRange)
  annotationEnd <- end(oneAnnotationRange)
  annotationStrand <- strand(oneAnnotationRange)

  annotationsDF <- rbind(annotationsDF, data.frame(seqname = seqName, start=seqRangeStart, end=seqRangeEnd, strand=annotationStrand))
}
#ggplot(annotationsDF) + geom_rect(aes(xmin = seqRangeStart, xmax = seqRangeEnd, ymin = 0, ymax=1), fill = "black", data=annotationsDF)

annotationsDF_GR <- makeGRangesFromDataFrame(annotationsDF, keep.extra.columns = T)

traitGWASresultsGR_annotatons <- mergeByOverlaps(traitGWASresultsGR, annotationsDF_GR)

traitGWASresultsGR_annotatons_df <- as.data.frame(traitGWASresultsGR_annotatons)

traitGWASresultsGR_annotatons_df$MinDistToAnnotation <- min(abs(traitGWASresultsGR_annotatons_df$seqname - traitGWASresultsGR_annotatons_df$seqname))

traitGWASresultsGR_annotatons_df$MinDistToAnnotation[traitGWASresultsGR_annotatons_df$seqname == traitGWASresultsGR_annotatons_df$seqname] <- 0

bestHitByAnnot <- data.frame()
for(annotationName in unique(traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName)){
  message(annotationName)
  annotationData <- traitGWASresultsGR_annotatons_df[traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName == annotationName,]
  bestHit <- min(annotationData$traitGWASresultsGR.markerpVal)
  message(bestHit)
  bestHitByAnnot <- rbind(bestHitByAnnot, data.frame(annotationsDF_GR.annotationName = annotationName, bestHit = bestHit))
}

traitGWASresultsGR_annotatons_df <- merge(traitGWASresultsGR_annotatons_df, bestHitByAnnot, by = "annotationName", all=TRUE)

traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName_label <- as.character(traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName)

traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName_label[traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName == ""] <- NA

traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName_label <- as.factor(traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName_label)

summary(traitGWASresultsGR_annotatons_df$annotationsDF_GR.annotationName_label)

traitGWASresultsGR_annotatons_df$MinDistToAnnotation_label <- paste(traitGWASresultsGR_annotatons_df$seqname, traitGWASresultsGR_annotatons_df$MinDistToAnnotation, sep="_")

traitGWASresultsGR_annotatons_df$MinDistToAnnotation_label[traitGWASresultsGR_annotatons_df$bestHitPValue == 0] <- NA

#ggplot(subset(traitGWASresults, Chrom %in% traitGWASresultsGR_annotatons_df$traitGWASresultsGR.seqname)) +
ggplot(traitGWASresultsGR_annotatons_df) + geom_point(aes(traitGWASresultsGR.start, traitGWASresultsGR.end, traitGWASresultsGR.markerpVal)) +
# + geom_rect(aes(xmin = seqRangeStart, xmax = seqRangeEnd, ymin = 0, ymax=1), fill = "black", data=annotationsDF)

```