

# mir181 binding sites - union of mir181 enriched binding sites and Ago binding sites targeted by mir181

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## 1 Libraries and settings

```
# -----  
# libraries  
# -----  
library(tidyverse)  
library(GenomicRanges)  
library(colorspace)  
library(eulerr)  
library(gghalves)  
library(ggpubr)  
  
# -----  
# settings  
# -----  
out <- "/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/R_github/miR181_paper/Figure1/mir181  
source("/Users/melinaklostermann/Documents/projects/R_general_functions/theme_paper.R")  
source("/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/mirko_files/mirECLIP/DifferentialBin  
  
# farben  
farbeneg <- "#B4B4B4"  
farbe1 <- "#0073C2FF" #WT farbe  
farbe2 <- "#EFC000FF"
```

```

farbe3 <- "#CD534CFF" #mir181KO farbe
farbe4 <- "#7AA6DCFF"
farbe5 <- "#868686FF"
farbe6 <- "#003C67FF"
farbe7 <- "#8F7700FF"
farbe8 <- "#3B3B3BFF"
farbe9 <- "#A73030FF"
farbe10 <- "#4A6990FF"
farbe11 <- "#FF6F00FF"
farbe12 <- "#C71000FF"
farbe13 <- "#008EAOFF"
farbe14 <- "#8A4198FF"
farbe15 <- "#5A9599FF"
farbe16 <- "#FF6348FF"

```

## 2 What was done?

mir181 binding sites are defined as the union of - AGO binding sites that contain at least 2 chimirc mir181 crosslinks (from the IP\_WT chimeric reads or the IP\_mir181\_WT chimeric reads) in a window from 10nt before till 10nt after a the AGO binding site - binding sites defined on enriched mir181 data (IP\_mir181\_WT)

- the two subgroups are plotted as a venn diagram (figure 1 XX)
- this is compared to the differentially regulated AGO binding sites from the mir181 KO condition

## 3 Files

```

# -----
# mir181 enriched binding sites
# -----
mir181_enriched <- readRDS("/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/R_github/miR181_p

# -----
# chimeric reads
# -----

chimeric_reads <- readRDS("/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/R_github/miR181_p

# -----
# AGO binding sites
# -----
ago_bs <- readRDS("/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/R_github/miR181_paper/Figur

# -----
# BS downregulated in mir181 KO
# -----
diff <- readRDS("/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/R_github/miR181_paper/Figur
diff$down <- (diff$resBs.padj < 0.05) & (diff$resBs.log2FoldChange < -log2(2))

```

## 4 mir181 binding sites

### 4.1 Get AGO binding sites with chimeric mir181

Here we define mir181 AGO binding sites by overlapping the AGO binding sites (see script Methods/02\_AGO\_binding\_site\_definition) with the chimeric mir181 reads (see script Figure1/Ago\_targetome). AGO binding sites that contained at least 2 chimeric mir181 crosslinks in the binding site or within 10nt proximity to the binding site are selected as mir181 Ago binding sites.

```
# use region of bs +/-10nt for overlaps
ago_bs_10 <- ago_bs + 10

# use chimeric reads from both mir181 enriched and non-enriched data
chimeric_reads <- c(makeGRangesFromDataFrame(chimeric_reads$IP_WT, keep.extra.columns = T), makeGRangesFromDataFrame(chimeric_reads$IP_KO, keep.extra.columns = T))

# find overlaps of mir181 and AGO bs
idx <- findOverlaps(ago_bs_10, chimeric_reads)

# make a data frame from the ago bs
names(ago_bs) <- 1:NROW(ago_bs)
ago_bs <- as.data.frame(ago_bs)
ago_bs$BS_ID <- rownames(ago_bs)

# add mir info to ago bs
ago_bs_mir181_chi <- cbind(ago_bs[queryHits(idx),], mir_IP = chimeric_reads[subjectHits(idx),]$Name)

ago_bs_mir181_chi <- ago_bs_mir181_chi[grepl(ago_bs_mir181_chi$mir_IP,
                                           pattern = "miR-181"),]

# count chimerics
mir181_chi <- ago_bs_mir181_chi %>% group_by(BS_ID) %>%
  summarize(n_mir181 = sum(grepl(mir_IP, pattern = "miR-181")),
            n_mir181a = sum(grepl(mir_IP, pattern = "miR-181a")),
            n_mir181b = sum(grepl(mir_IP, pattern = "miR-181b")),
            n_mir181c = sum(grepl(mir_IP, pattern = "miR-181c")),
            n_mir181d = sum(grepl(mir_IP, pattern = "miR-181d")),
            .groups = "keep") %>% subset (n_mir181 > 0)

ago_bs_mir181_chi <- ago_bs_mir181_chi %>%
  subset(!duplicated(ago_bs_mir181_chi$BS_ID)) %>%
  left_join(., mir181_chi, by = "BS_ID") %>% makeGRangesFromDataFrame(keep.extra.columns = T)
```

### 4.2 Combine AGO binding sites with chimeric mir181 with mir181 enriched binding sites

I combine the mir181 Ago binding sites that we obtained above with the binding sites from the mir181 enriched Ago-eCLIP (see Methods/mir181-enriched\_binding\_site\_definition). In order to do that, I first select binding sites from both conditions that do not overlap with any binding site from the other set. For the binding sites that overlap between the two conditions, I select the AGO mir181 binding sites and tag them as occurring in both sets. Then I combine the three subsets. The obtained union of mir181 binding sites from both conditions are our final mir181 binding sites.

```
# -----
# combine mir181 Ago BS and mir181 enriched Bs
# -----
```

```

# get only Ago mir181 BS with now overlaps to enriched mir181 BS
only_ago_bs_mir181_chi <- subsetByOverlaps(ago_bs_mir181_chi, mir181_enriched, type = "any", invert = T)
only_ago_bs_mir181_chi$set <- "ago_bs_mir181_chi"

# get only enriched mir181 BS with now overlaps to Ago mir181 BS
only_mir181_enriched <- subsetByOverlaps(mir181_enriched, ago_bs_mir181_chi, type = "any", invert = T)
only_mir181_enriched$set <- "mir181_enriched"

# get only Ago mir181 BS overlapping with mir181 enriched BS
both_mir181_enriched_chi <- subsetByOverlaps(ago_bs_mir181_chi, mir181_enriched, type = "any")
both_mir181_enriched_chi$set <- "ago_bs_mir181_chi&mir181_enriched"

# combine all three sets
mir181_bs <- c(only_ago_bs_mir181_chi, only_mir181_enriched, both_mir181_enriched_chi)

```

## 5 Combine mir181 binding sites with differential binding sites

Next, I combine the obtained mir181 binding sites with the results we obtained from the differential binding between AGO binding sites and AGO binding sites with mir181 KO (see script Figure1/Differential\_Binding\_AGO\_BS\_mir181\_KO).

```

# -----
# combine with differential BS
# -----
# get overlaps with diff binding
diff_overlap <- findOverlaps( mir181_bs, makeGRangesFromDataFrame(diff, keep.extra.columns = T) , type = "any")
# add differential information to mir181 binding sites
d <- diff[,9:49]
mcols(mir181_bs) <- cbind(mcols(mir181_bs), d[diff_overlap,])

# add only diff bs (these are Ago binding sites but not mir181 Binding sites)
diff_only <- subsetByOverlaps( makeGRangesFromDataFrame(diff, keep.extra.columns = T), mir181_bs , type = "any")
mir181_bs_diff <- c(mir181_bs, diff_only)

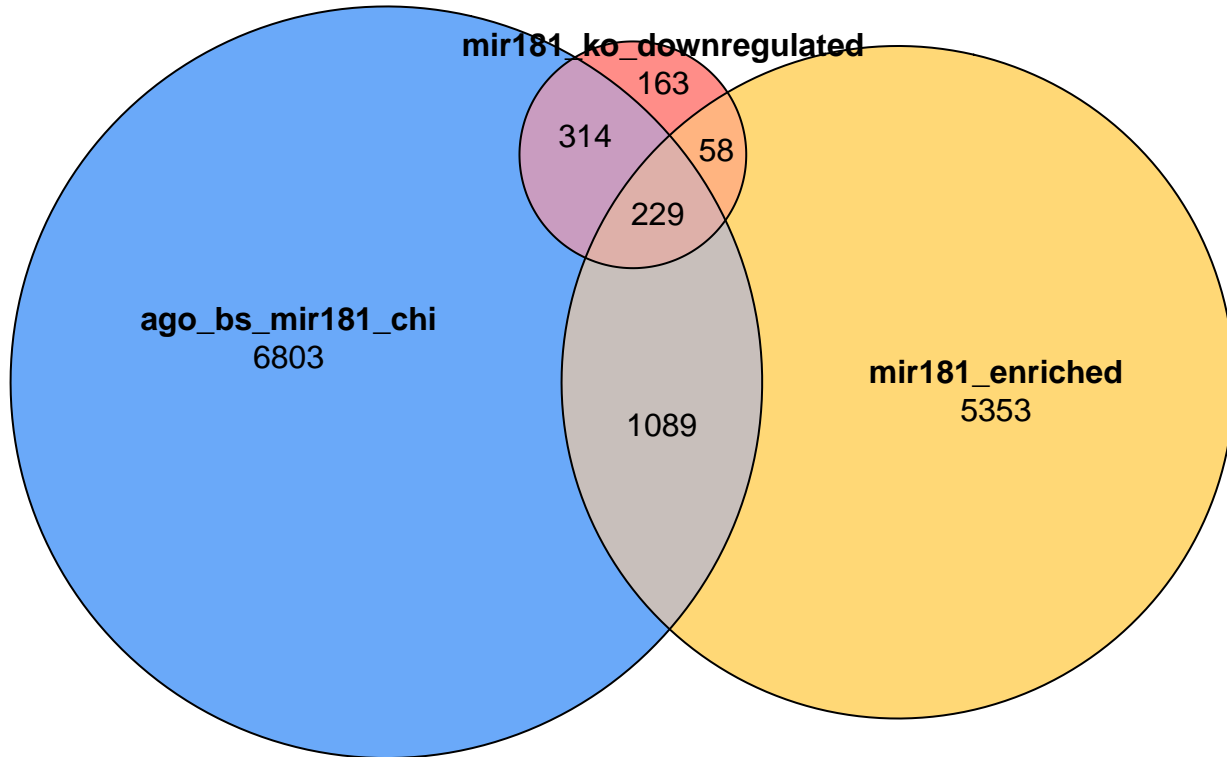
# -----
# make venn diagram
# -----

# select downregulated BS from differential BS
mir181_bs_diff_anygroup <- mir181_bs_diff %>% subset((down == T) | !is.na(set))

# make venn
venn_df <- data.frame(ago_bs_mir181_chi =
  (mir181_bs_diff_anygroup$set == "ago_bs_mir181_chi") | (mir181_bs_diff_anygroup$set == "mir181_enriched") &
  (mir181_bs_diff_anygroup$set == "mir181_enriched") | (mir181_bs_diff_anygroup$set == "mir181_ko_downregulated") &
  (mir181_ko_downregulated == (mir181_bs_diff_anygroup$down == T) )
) %>%
mutate(ago_bs_mir181_chi = case_when(is.na(ago_bs_mir181_chi) ~ F, T~ago_bs_mir181_chi),
  mir181_enriched = case_when(is.na(mir181_enriched) ~ F, T ~ mir181_enriched),
  mir181_ko_downregulated = case_when(is.na(mir181_ko_downregulated) ~ F, T ~ mir181_ko_downregulated))

```

```
venn <- eulerr::euler(venn_df)
p <- plot(venn, quantities = T, fills = c( lighten(farbe1, amount = 0.4), lighten(farbe2, amount = 0.4)
p
```



```
pdf(paste0(out, "Figure1I_Venn_overlaps_mirBs.pdf"), height = 3, width = 4 )
p
dev.off()
```

```
## pdf
## 2
```

## 5.1 Differential binding both sets

Here I look at the overall binding changes between mir181 KO and WT for the three subsets of the mir181 binding sites.

```
# -----
# Compare regulation of BS during mir181 KO of the different subgroups of mir181 binding sites
# -----
names(mir181_bs_diff) <- 1:NROW(mir181_bs_diff)

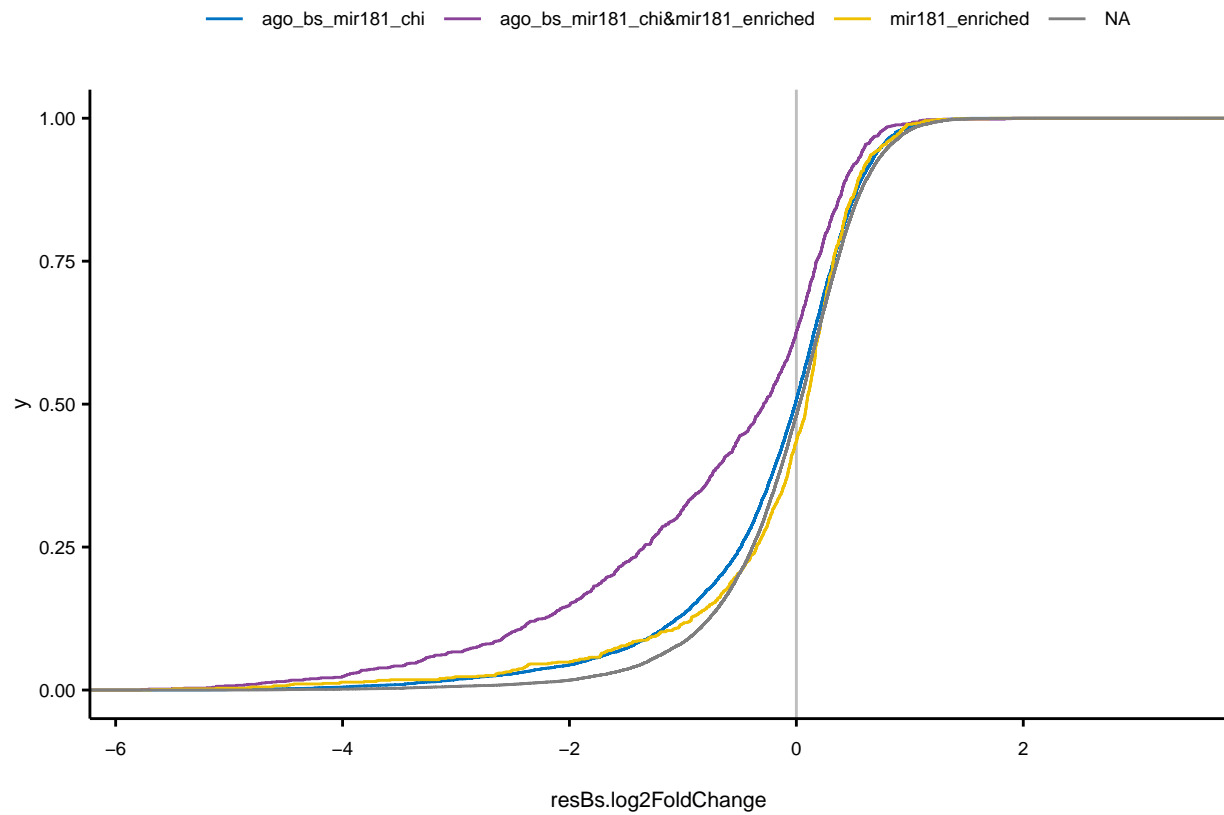
mir181_bs_diff <- as.data.frame(mir181_bs_diff)

p1 <- ggplot(mir181_bs_diff, aes(x = resBs.log2FoldChange, color = set))+
  geom_vline(xintercept = 0, color = "grey")+
  stat_ecdf()+
  scale_color_manual(values = c(farbe1, farbe14, farbe2, farbeneg))+
  theme_paper()+
```

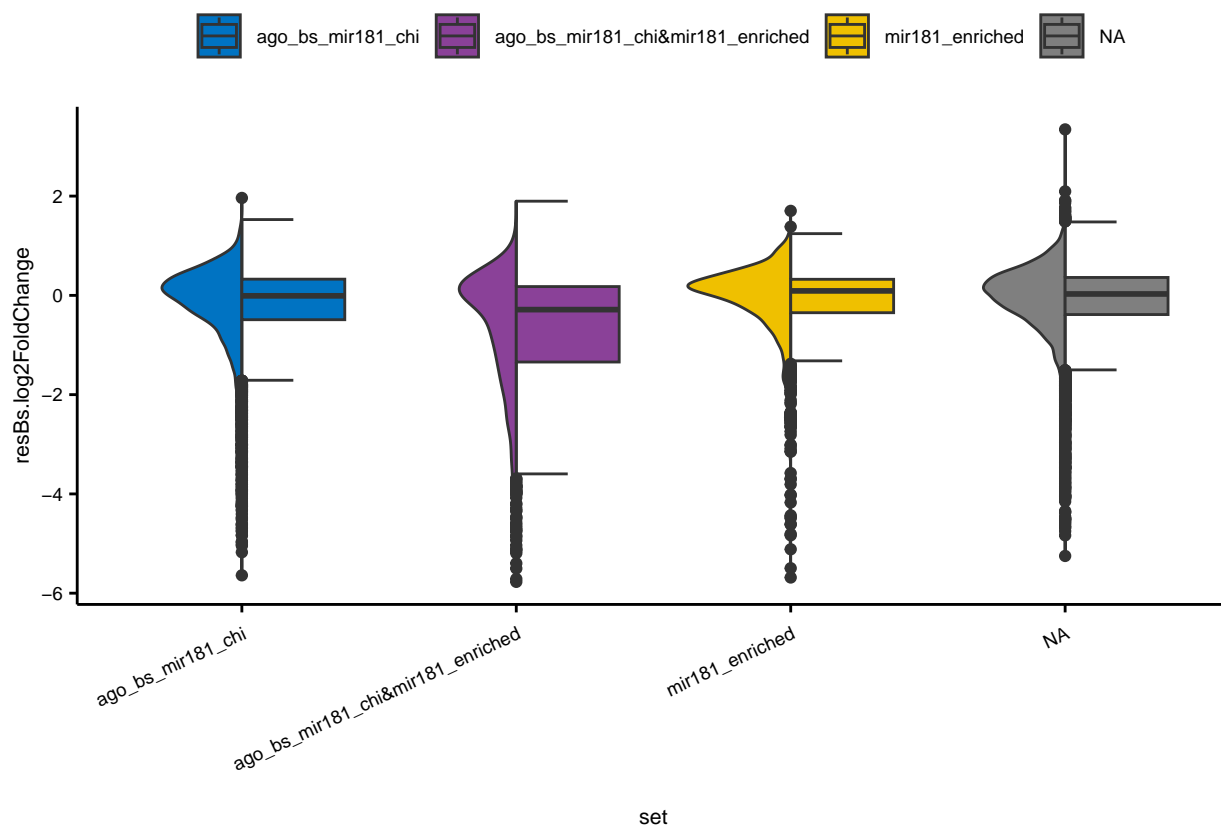
```
theme(legend.position = "top")
```

```
p2 <- ggplot(mir181_bs_diff, aes(y = resBs.log2FoldChange, x = set, fill = set))+
  geom_half_violin()+
  geom_half_boxplot(side = "r")+
  theme_paper()+
  scale_fill_manual(values = c(farbe1, farbe14, farbe2, farbeneg))+
  theme(legend.position = "top")+
  scale_x_discrete(guide = guide_axis(angle = 25))
```

p1



p2



```
ggsave(p1, filename = paste0(out, "Figure_1J_ecdf_differntial_binding_vs_mir181BS.pdf"), width = 6, height = 6)
ggsave(p2, filename = paste0(out, "violin_differntial_binding_vs_mir181BS.pdf"), width = 10, height = 10)
```

### 5.1.1 Statistical tests for differential binding changes

```
# t.tests of 3 sets against not bound
# -----
t1 <- t.test(x = mir181_bs_diff %>% subset(set == "mir181_enriched") %>% pull(resBs.lfcSE),
             y = mir181_bs_diff %>% subset(is.na(set)) %>% pull(resBs.lfcSE))
t1
```

#### 5.1.1.1 T-tests

```
##
## Welch Two Sample t-test
##
## data: mir181_bs_diff %>% subset(set == "mir181_enriched") %>% pull(resBs.lfcSE) and mir181_bs_diff %>% subset(is.na(set)) %>% pull(resBs.lfcSE)
## t = -15.992, df = 1000.5, p-value < 2.2e-16
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.1305064 -0.1019782
## sample estimates:
## mean of x mean of y
## 0.4005421 0.5167844

t.test(x = mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi") %>% pull(resBs.lfcSE),
       y = mir181_bs_diff %>% subset(is.na(set)) %>% pull(resBs.lfcSE))

##
```

```
## Welch Two Sample t-test
##
## data: mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi") %>% pull(resBs.lfcSE) and mir181_bs_diff
## t = -17.36, df = 10729, p-value < 2.2e-16
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.05261416 -0.04193797
## sample estimates:
## mean of x mean of y
## 0.4695083 0.5167844

t.test(x = mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi&mir181_enriched") %>% pull(resBs.lfcSE),
       y = mir181_bs_diff %>% subset(is.na(set)) %>% pull(resBs.lfcSE))

##
## Welch Two Sample t-test
##
## data: mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi&mir181_enriched") %>% pull(resBs.lfcSE) and
## t = -5.9647, df = 1276.1, p-value = 3.168e-09
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.06323341 -0.03193289
## sample estimates:
## mean of x mean of y
## 0.4692012 0.5167844

# --> the p-values are driven strongly by the number of binding sites per set, for this reason the high

# check power of test
# -----
pwr::pwr.t2n.test(n1 = mir181_bs_diff %>% subset(set == "mir181_enriched") %>% nrow(.),
                  n2 = mir181_bs_diff %>% subset(is.na(set)) %>% nrow(.),
                  d = abs(t1$estimate[1] - t1$estimate[2])/t1$stderr)

##
##      t test power calculation
##
##          n1 = 5411
##          n2 = 17499
##          d = 15.99168
##      sig.level = 0.05
##          power = 1
##      alternative = two.sided

# Kolmogorov-Smirnov Tests
# -----
ks.test(x = mir181_bs_diff %>% subset(set == "mir181_enriched") %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE),
        y = mir181_bs_diff %>% subset(is.na(set)) %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE))
```

#### 5.1.1.2 Kolmogorov-Smirnov Tests

```
##
## Asymptotic two-sample Kolmogorov-Smirnov test
##
## data: mir181_bs_diff %>% subset(set == "mir181_enriched") %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE)
```



```
## D = 0.4051, p-value < 2.2e-16
## alternative hypothesis: two-sided
ks.test(x = mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi") %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE),
        y = mir181_bs_diff %>% subset(is.na(set) ) %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE))

##
## Asymptotic two-sample Kolmogorov-Smirnov test
##
## data: mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi") %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE)
## D = 0.17649, p-value < 2.2e-16
## alternative hypothesis: two-sided
ks.test(x = mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi&mir181_enriched") %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE),
        y = mir181_bs_diff %>% subset(is.na(set) ) %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE))

##
## Asymptotic two-sample Kolmogorov-Smirnov test
##
## data: mir181_bs_diff %>% subset(set == "ago_bs_mir181_chi&mir181_enriched") %>% arrange(resBs.lfcSE) %>% pull(resBs.lfcSE)
## D = 0.3015, p-value < 2.2e-16
## alternative hypothesis: two-sided
```

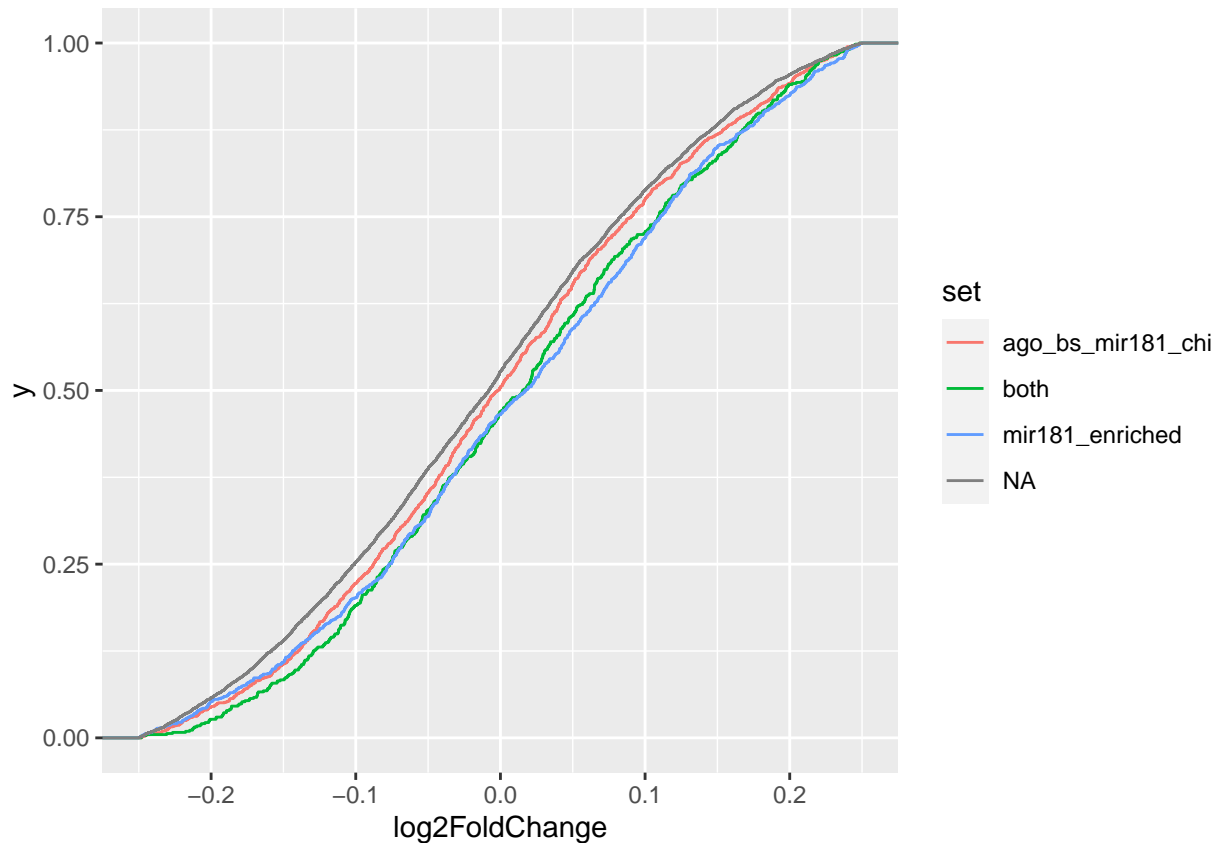
## 5.2 Venn bound genes from all sets

## 5.3 Ribofootprint both sets

```
# TODO need to be other script later
rpf <- read.csv("/Users/melinaklostermann/Documents/projects/AgoCLIP_miR181/files_RNAseq_RiboP/RPF_master.csv")
rpf <- mutate(rpf, Gene = substr(Gene,1,18))

rpf <- rpf %>% mutate(., set =
  case_when(Gene %in% both_mir181_enriched_chi$geneID ~ "both",
            (Gene %in% ago_bs_mir181_chi$geneID) & !(Gene %in% both_mir181_enriched_chi$geneID) ~ "ago",
            (Gene %in% mir181_enriched$geneID) & !(Gene %in% both_mir181_enriched_chi$geneID) ~ "mir181",
            TRUE ~ "other")
)

ggplot(rpf, aes(x = log2FoldChange, color = set))+
  stat_ecdf()+
  xlim(-0.25, 0.25)
```



## 6 Save output

```
saveRDS(mir181_bs, paste0(out, "mir181_bs.rds"))
```

## 7 Session Info

```
sessionInfo()

## R version 4.2.2 (2022-10-31)
## Platform: x86_64-apple-darwin17.0 (64-bit)
## Running under: macOS Big Sur ... 10.16
##
## Matrix products: default
## BLAS: /Library/Frameworks/R.framework/Versions/4.2/Resources/lib/libRblas.0.dylib
## LAPACK: /Library/Frameworks/R.framework/Versions/4.2/Resources/lib/libRlapack.dylib
##
## locale:
## [1] en_US.UTF-8/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-8/en_US.UTF-8
##
## attached base packages:
## [1] stats4      stats      graphics  grDevices  utils      datasets  methods
## [8] base
##
## other attached packages:
## [1] ggpubr_0.5.0      gghalves_0.1.4      eulerr_7.0.0
```

```

## [4] colorspace_2.1-0      GenomicRanges_1.50.2 GenomeInfoDb_1.34.7
## [7] IRanges_2.32.0        S4Vectors_0.36.1     BiocGenerics_0.44.0
## [10] forcats_0.5.2         stringr_1.5.0         dplyr_1.0.10
## [13] purrr_1.0.1           readr_2.1.3           tidyr_1.3.0
## [16] tibble_3.1.8          ggplot2_3.4.0         tidyverse_1.3.2
## [19] knitr_1.42
##
## loaded via a namespace (and not attached):
## [1] httr_1.4.4             jsonlite_1.8.4         carData_3.0-5
## [4] modelr_0.1.10          assertthat_0.2.1       highr_0.10
## [7] GenomeInfoDbData_1.2.9 googlesheets4_1.0.1    cellranger_1.1.0
## [10] yaml_2.3.7             pillar_1.8.1           backports_1.4.1
## [13] glue_1.6.2            digest_0.6.31          XVector_0.38.0
## [16] ggsignif_0.6.4         polyclip_1.10-4        rvest_1.0.3
## [19] htmltools_0.5.4        pkgconfig_2.0.3        broom_1.0.3
## [22] haven_2.5.1           zlibbioc_1.44.0        scales_1.2.1
## [25] tzdb_0.3.0            timechange_0.2.0       googledrive_2.0.0
## [28] farver_2.1.1          generics_0.1.3         car_3.1-1
## [31] ellipsis_0.3.2        withr_2.5.0            cli_3.6.0
## [34] magrittr_2.0.3        crayon_1.5.2           readxl_1.4.1
## [37] evaluate_0.20         fs_1.6.0              fansi_1.0.4
## [40] rstatix_0.7.1         xml2_1.3.3            textshaping_0.3.6
## [43] tools_4.2.2           hms_1.1.2             gargle_1.2.1
## [46] lifecycle_1.0.3       munsell_0.5.0          reprex_2.0.2
## [49] compiler_4.2.2        systemfonts_1.0.4      rlang_1.0.6
## [52] grid_4.2.2           Rcurl_1.98-1.9         rstudioapi_0.14
## [55] labeling_0.4.2        bitops_1.0-7           rmarkdown_2.20
## [58] gtable_0.3.1         abind_1.4-5            DBI_1.1.3
## [61] R6_2.5.1             lubridate_1.9.1        pwr_1.3-0
## [64] fastmap_1.1.0         utf8_1.2.2            ragg_1.2.5
## [67] polylabelr_0.2.0     stringi_1.7.12         Rcpp_1.0.10
## [70] vctrs_0.5.2          dbplyr_2.3.0          tidyselect_1.2.0
## [73] xfun_0.36

```