# Downstream analysis followig the image analysis to quantify Dvl2\_mEos codensates (Figure S8C)

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# 1 Load dependencies

```
knitr::opts_chunk$set(echo = TRUE)
library(tidyverse)
library(here)
library(cowplot)
library(ggpubr)
theme_set(theme_cowplot())
```

### 2 Preamble

The image analysis was carried out as follows: In a first step cells were identified by segmentation. Nuclei were identified based on a Hoechst staining (DAPI channel) and cell bodies based on a DyLight Phalloidin staining (Cy5 channel). After segmentation condensates inside cells were identified based on intensity-based thresholding (FITC channel) and counted. Condensate counts and cell counts (number of segmented cells) were saved for each image. An exemplary image analysis can be found with the script images\_analysis\_condesate\_quantification.Rmd in this repository. Two different cell lines and different conditions were compared. Cell lines: - HEK cells with a DVL1-mEOS tag - HEK cells with a DVL1-mEOS tag and a EVI k.o. (clone 1)

Conditions: - transfection reagent control (RNAiMax) - transfection reagent (RNAiMax) + non-targeting siRNA - transfection reagent (RNAiMax) + siCTNNB1

The time course experiment 4 separate xcperiments (i.e. on different days) were performed. Each experiment was carried out on one 384-well plates. Per condition, 42 wells with 4 fields of view per well were analyzed.

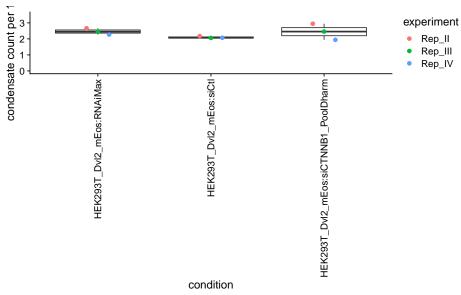
# 3 Load the data after the image analysis

Note: The raw data for Figure S8C was saved in one data frame with the war data for figure 4G.

```
raw_data <- read_rds(here("raw_data/raw_data_Fig4G","raw_data_Fig4G.rds"))</pre>
```

# 4 Prepare boxplot

```
boxplot <- raw_data %>% filter(plate_id == "Plate_IV") %>%
  filter(treatment %in% c("RNAiMax","siCTNNB1_PoolDharm","siCtl")) %>%
  mutate(norm_punct_count = (punct_count_adapt/cell_count)*100) %>%
  group_by(barcode,cell_line,plate_id,experiment,treatment) %>%
  summarise_if(is.numeric,mean) %>%
  ungroup() %>%
  ungroup() %>%
  unite(condition,cell_line,treatment,sep=":") %>%
  arrange(condition) %>%
  ggplot(aes(x = condition,y=norm_punct_count)) + geom_boxplot(outlier.shape = NA) +
```



# 5 Session info

```
sessionInfo()
## R version 4.1.2 (2021-11-01)
## Platform: x86_64-apple-darwin17.0 (64-bit)
## Running under: macOS Big Sur 10.16
##
## Matrix products: default
## BLAS: /Library/Frameworks/R.framework/Versions/4.1/Resources/lib/libRblas.0.dylib
## LAPACK: /Library/Frameworks/R.framework/Versions/4.1/Resources/lib/libRlapack.dylib
##
## locale:
```

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```
## [1] en_US.UTF-8/en_US.UTF-8/en_US.UTF-8/en_US.UTF-8
##
## attached base packages:
## [1] stats graphics grDevices utils datasets methods base
## other attached packages:
## [1] ggpubr_0.4.0 cowplot_1.1.1 here_1.0.1 forcats_0.5.1
## [5] stringr_1.4.0 dplyr_1.0.7 purrr_0.3.4 readr_2.1.1 ## [9] tidyr_1.1.4 tibble_3.1.6 ggplot2_3.3.5 tidyverse_1.3.1
## [13] BiocStyle_2.22.0
##
## loaded via a namespace (and not attached):
tidyselect_1.1.1
## [61] knitr_1.36 haven_2.4.3
```