

Downstream analysis followig the image analysis to quanttify Dvl2_mEos coden-sates (Figure 4F)

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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_condensate_quantification.Rmd` in this repository. Four different cell lines were compared: - HEK cells with a DVL1-mEOS tag - HEK cells with a DVL1-mEOS tag and a EVI k.o. (clone 1) - HEK cells with a DVL1-mEOS tag and a EVI k.o. (clone 3) - HEK cells with a DVL1-mEOS tag and a FZD1 k.o. (clone 1) - HEK wild-type

For each cell line three replicates were carried out (E01,E02,E03). Per replicate multiple 384-well plates were seeded, fixed/stained and imaged. For each replicate representative images were selected. The image-level data (see description above) was further averaged per plate and differences between the cell lines were compared using a box plot and statistical testing with the `ggbbubr` package (non-parametric Wilcoxon rank-sum test).

3 Load the data after the image analysis

For each replicate the results were saved in a separate data frame.

```
results_path <- here("./raw_data/raw_data_Fig4F/results_image_analysis/data_frames")
files <- dir(results_path, pattern = "*.rds")

raw_data <- files %>%
  map(~ read_rds(file.path(results_path, .))) %>%
  reduce(rbind) %>%
  mutate(cell_line=factor(cell_line,
                           levels = c("wt", "mEos", "Evi1", "Evi3", "mFZD")))
```

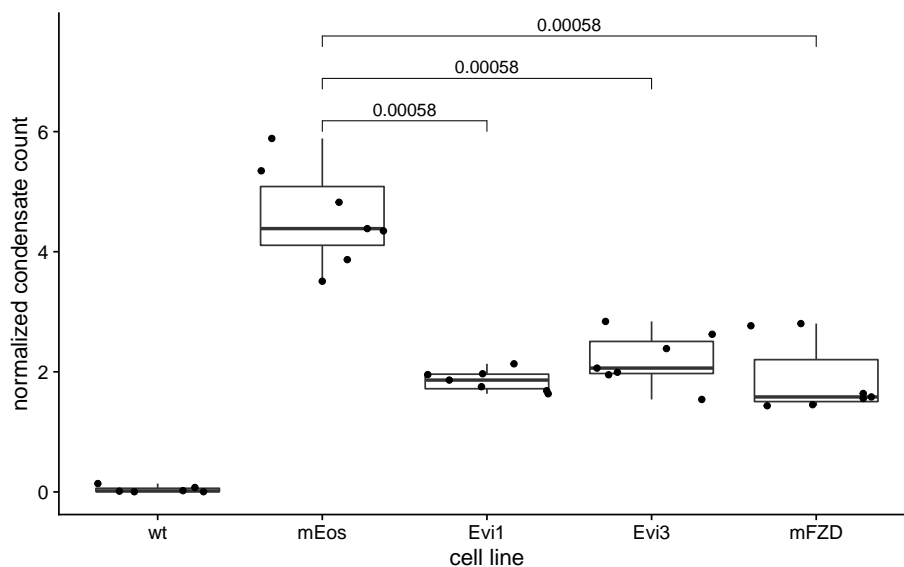
4 Prepare boxplot

```
boxplot <- raw_data %>%
  separate(well_id, c("well", "field")) %>%
  mutate(norm_punct_count = punct_count_dapt/(cell_count/100)) %>%
  group_by(experiment, plate_id, cell_line) %>%
```

Downstream analysis followig the image analysis to quanttify Dvl2_mEos condensates (Figure 4F)

```
summarise_if(is.numeric,mean) %>%
ungroup() %>%
  ggplot(aes(x = cell_line,y=norm_punct_count)) +
  geom_boxplot() +
  geom_jitter() +
  stat_compare_means(comparisons = list(c("Evi1","mEos"),
                                         c("Evi3","mEos"),
                                         c("mFZD","mEos")),
                    method = "wilcox.test") +
  ylab("normalized condensate count") +
  xlab("cell line")
```

boxplot



```
ggsave(here("graphics/Figure4F","boxplot_figure4f_raw.pdf"),
  plot=boxplot,
  width = 6,
  height = 4)
```

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
```

Downstream analysis following the image analysis to quantify Dvl2_mEos condensates (Figure 4F)

```
##
## 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] 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):
## [1] Rcpp_1.0.7        lubridate_1.8.0    assertthat_0.2.1
## [4] rprojroot_2.0.2   digest_0.6.29      utf8_1.2.2
## [7] R6_2.5.1          cellranger_1.1.0   backports_1.4.0
## [10] reprex_2.0.1      evaluate_0.14      http_1.4.2
## [13] pillar_1.6.4      rlang_0.4.12       readxl_1.3.1
## [16] rstudioapi_0.13   car_3.0-12         rmarkdown_2.11
## [19] labeling_0.4.2    munsell_0.5.0      broom_0.7.10
## [22] compiler_4.1.2    modelr_0.1.8       xfun_0.28
## [25] pkgconfig_2.0.3   htmltools_0.5.2    tidyselect_1.1.1
## [28] bookdown_0.24     fansi_0.5.0        crayon_1.4.2
## [31] tzdb_0.2.0        dbplyr_2.1.1       withr_2.4.3
## [34] grid_4.1.2        jsonlite_1.7.2     gtable_0.3.0
## [37] lifecycle_1.0.1   DBI_1.1.1          magrittr_2.0.1
## [40] scales_1.1.1      carData_3.0-4      cli_3.1.0
## [43] stringi_1.7.6     farver_2.1.0       ggsignif_0.6.3
## [46] fs_1.5.1          xml2_1.3.3         ellipsis_0.3.2
## [49] generics_0.1.1    vctrs_0.3.8        tools_4.1.2
## [52] glue_1.5.1        hms_1.1.1          abind_1.4-5
## [55] fastmap_1.1.0     yaml_2.2.1         colorspace_2.0-2
## [58] BiocManager_1.30.16 rstatix_0.7.0      rvest_1.0.2
## [61] knitr_1.36        haven_2.4.3
```