$MitoGraph_simplified_width_filtered.R$

mcleland

2022-05-26

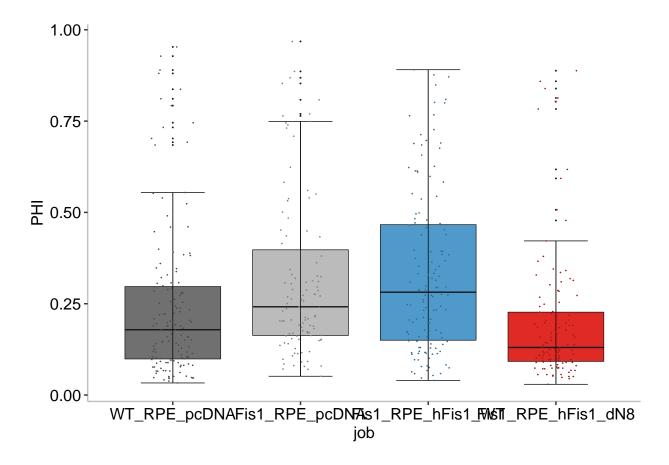
```
# Summarize MitoGraph Metrics using Box Plots
# First set working directory to source file location
# Run with command + option + R
library(tidyverse)
## -- Attaching packages ------ tidyverse 1.3.1 --
## v ggplot2 3.3.5 v purrr 0.3.4
## v tibble 3.1.6 v dplyr 1.0.7
## v tidyr 1.2.0 v stringr 1.4.0
## v readr 2.1.2 v forcats 0.5.1
## -- Conflicts ----- tidyverse_conflicts() --
## x dplyr::filter() masks stats::filter()
## x dplyr::lag()
                    masks stats::lag()
library(reshape2)
## Attaching package: 'reshape2'
## The following object is masked from 'package:tidyr':
##
##
      smiths
library(stringr)
library(formattable)
library(igraph)
## Attaching package: 'igraph'
## The following object is masked from 'package:formattable':
##
##
      normalize
```

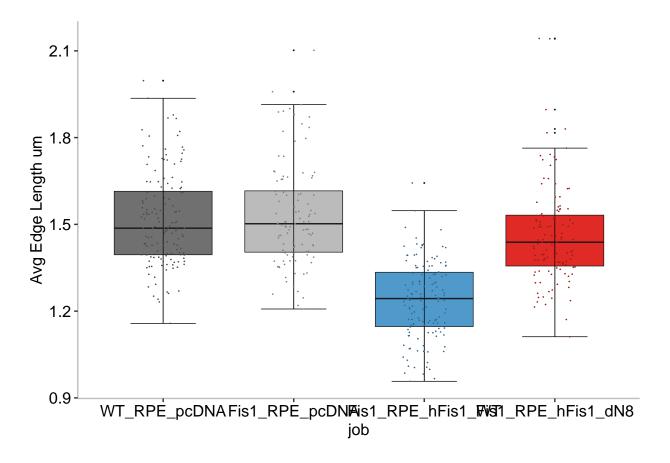
```
## The following objects are masked from 'package:dplyr':
##
       as_data_frame, groups, union
##
## The following objects are masked from 'package:purrr':
##
##
       compose, simplify
## The following object is masked from 'package:tidyr':
##
##
       crossing
## The following object is masked from 'package:tibble':
##
##
       as_data_frame
## The following objects are masked from 'package:stats':
##
##
       decompose, spectrum
## The following object is masked from 'package:base':
##
       union
# Import data from log files (*.txt)
datafolder1 <- "~/mount/home110/mcharwig/20210903_Ntermtrun_repeat4/mitoYFP_DrpIF/combined_drp1_days/Mi
# Import data
# with read csv command below
data <- read_csv(file = "output-summary_filtered.csv")</pre>
## Rows: 496 Columns: 19
## -- Column specification -----
## Delimiter: ","
## chr (4): file, job, isoform, treatment
## dbl (15): Total_Nodes, Total_Edges, Total_Length_um, Total_Connected_Compone...
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
data2 <- data %>%
  mutate(job = factor(job, levels = c("WT_RPE_pcDNA",
                                      "Fis1_RPE_pcDNA",
                                      "Fis1_RPE_hFis1_WT",
                                      "Fis1_RPE_hFis1_dN8"), ordered = T)) #RENAME with your sample ord
data3 <- data2 %>%
  mutate(treatment = factor(treatment, levels = c("pcDNA",
```

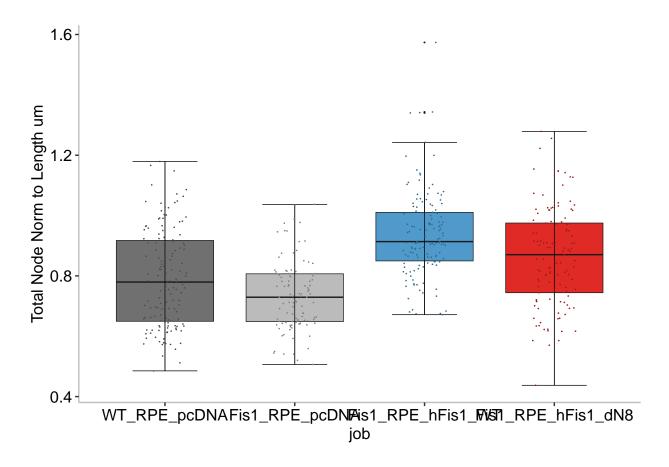
```
"hFis1 WT",
                                                   "hFis1_dN8"), ordered = T)) #RENAME with your sample
data3
## # A tibble: 496 x 19
##
     file
                   job
                          isoform treatment Total_Nodes Total_Edges Total_Length_um
##
      <chr>
                    <ord> <chr>
                                  <ord>
                                                   <dbl>
                                                               <dbl>
                                                                               <dbl>
## 1 /Users/mclel~ Fis1~ Fis1 R~ pcDNA
                                                                                240.
                                                     175
                                                                 144
## 2 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     460
                                                                 429
                                                                                716.
## 3 /Users/mclel~ Fis1~ Fis1 R~ pcDNA
                                                     716
                                                                 625
                                                                                942.
## 4 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     451
                                                                 386
                                                                                530.
## 5 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     269
                                                                 237
                                                                                426.
## 6 /Users/mclel~ Fis1~ Fis1 R~ pcDNA
                                                                                322.
                                                     193
                                                                 168
## 7 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     302
                                                                                424.
                                                                 276
## 8 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     284
                                                                 294
                                                                                442.
## 9 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     428
                                                                 363
                                                                                538.
## 10 /Users/mclel~ Fis1~ Fis1_R~ pcDNA
                                                     563
                                                                 439
                                                                                703.
## # ... with 486 more rows, and 12 more variables:
## #
       Total_Connected_Components <dbl>, PHI <dbl>, Avg_Edge_Length_um <dbl>,
## #
       Total_Edge_Norm_to_Length_um <dbl>, Total_Node_Norm_to_Length_um <dbl>,
## #
       Total_Connected_Components_Norm_to_Length_um <dbl>, Free_Ends <dbl>,
       three_way_junction <dbl>, four_way_junction <dbl>, Avg_Degree <dbl>,
## #
       MitoGraph_Connectivity_Score <dbl>, Average_width_um <dbl>
# Here is a loop that creates 11 graphs where the X and Y variable are listed accordingly.
PlotsToBeMade <- c("job",
                   "treatment",
                   "PHI",
                   "Avg_Edge_Length_um",
                   "Total_Node_Norm_to_Length_um",
                   "Total_Connected_Components_Norm_to_Length_um",
                   "Free Ends",
                   "three_way_junction",
                   "four_way_junction",
                   "Avg Degree",
                   "MitoGraph_Connectivity_Score",
                   "Average_width_um")
AxisLabels <- c("job",
                "treatment",
                "PHI",
                "Avg Edge Length um",
                "Total Node Norm to Length um",
                "Total Connected Components Norm to Length um",
                "Free Ends",
                "three way junction",
                "four way junction",
                "Avg Degree",
                "MitoGraph Connectivity Score",
                "Average width um")
Titles <- c("job",
```

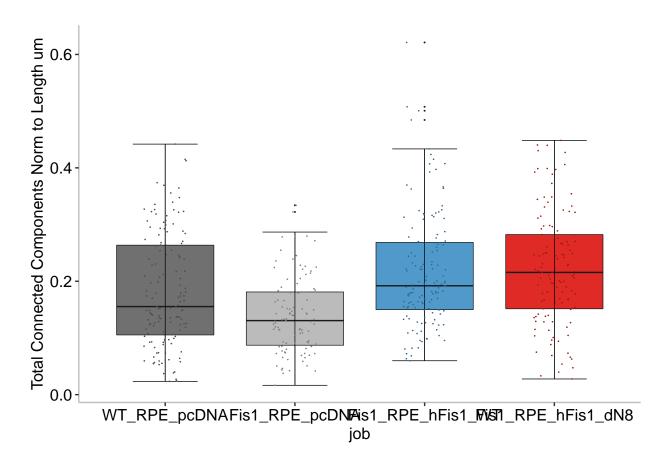
```
"treatment",
             "PHI",
             "Avg Edge Length um",
             "Total Node Norm to Length um",
             "Total Connected Components Norm to Length um",
             "Free Ends",
             "Three way junction",
             "Four way junction",
             "Avg Degree",
             "MitoGraph Connectivity Score",
             "Average width um")
# Uncomment to modify specific axis settings
yAxisMinimum <- c(0,
                   0,
                   0.
                  0,
                  0,
                  0,
                  0,
                  0,
                  0,
                  0,
                  0,
                  0)
# yAxisMaximum <- c(1,</pre>
                    1,
#
                    150,
#
                    150,
#
                    150,
#
                    150,
#
                    150,
#
                    1.5,
#
                    10000,
#
                    20000,
#
                    10000,
#
                    20000)
for (p in seq(3,length(PlotsToBeMade))) {
  yaxis <- PlotsToBeMade[p]</pre>
 xaxis <- PlotsToBeMade[1]</pre>
 fill <- PlotsToBeMade[1]</pre>
  # Adjust to modify error bar width
  # error <- c(0.03,0.05,
               0.03,0.025,
               0.02,0.05,
               8,0.005)
  plot(ggplot(data=data3,aes_string(x=xaxis, y=yaxis, fill=fill)) +
```

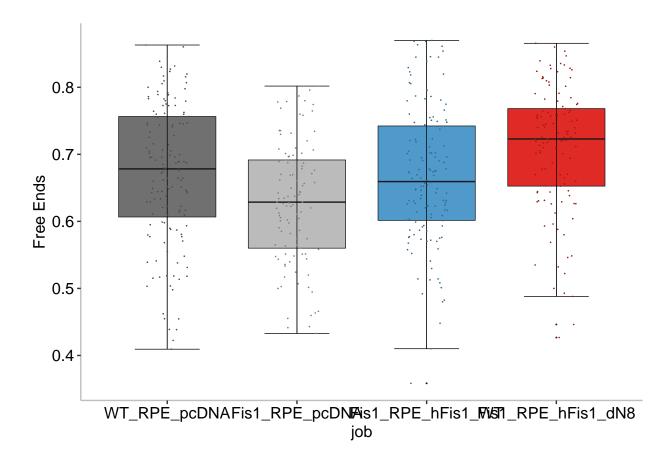
```
geom_boxplot(outlier.size = 0, colour = "grey10", position = position_dodge (width = 1), size =
       stat_boxplot(geom="errorbar", position = position_dodge (width = 1), width = 0.5, size = 0.25)
       scale_fill_manual(values = c("#707070","#BBBBBBB","#4F9ACB","#E02A26")) +
       geom_point(data=data3,aes_string(colour = fill, x=xaxis, y=yaxis), pch=20, size=0.01, position
       scale_colour_manual(values=c("#4e4e4e","#828282","#376b8e","#9c1d1a")) +
       labs(title = Titles[p],
            x = xaxis,
            y = AxisLabels[p]) +
       #ylab(AxisLabels[p]) +
       theme_bw() +
       theme(axis.text.x = element_text(size = 12, color = "black"),
             axis.title.x = element_text(size = 12, color = "black"),
             axis.text.y = element_text(size = 12, color = "black"),
             axis.title.y = element_text(size = 12, color = "black"),
            plot.title = element_blank(),
             #plot.title = element_text(size = 8, color = "black", hjust = 0.5, face = "bold"),
             panel.grid.major = element_blank(),
             panel.grid.minor = element_blank(),
             panel.border = element_blank(),
            panel.background = element_blank(),
             axis.line.x = element_line(color = "grey75", size = 0.5, linetype = 1),
            axis.line.y = element_line(color = "grey75", size = 0.5, linetype = 1),
            legend.position = "none"
             #legend.title = element_blank(),
             #legend.justification = c(0, 1),
             #legend.position = "right",
             #legend.text = element_text(size = 6, color = "black")
     # Add a "+" above and uncomment the below 2 lines to add custom axis scales.
     #ylim(yAxisMinimum[p], yAxisMaximum[p])
)
ggsave(paste(datafolder1, "Plot-filtered", yaxis, ".png", sep=""), width = 5.8, height = 8, units = "cm"
ggsave(paste(datafolder1, "Plot-filtered", yaxis, ".eps", sep=""), width = 5.8, height = 8, units = "cm"
```

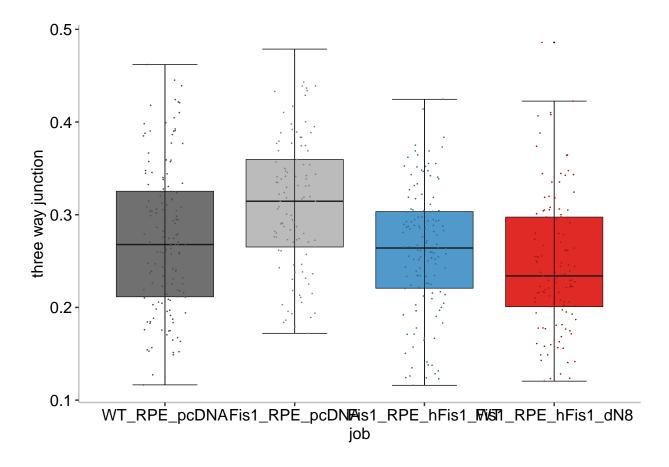


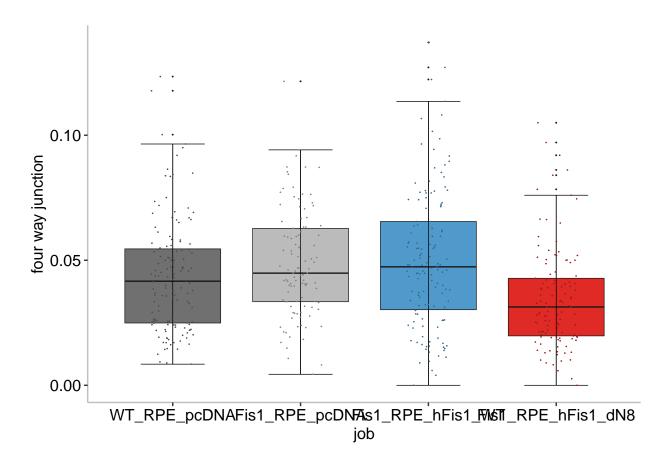


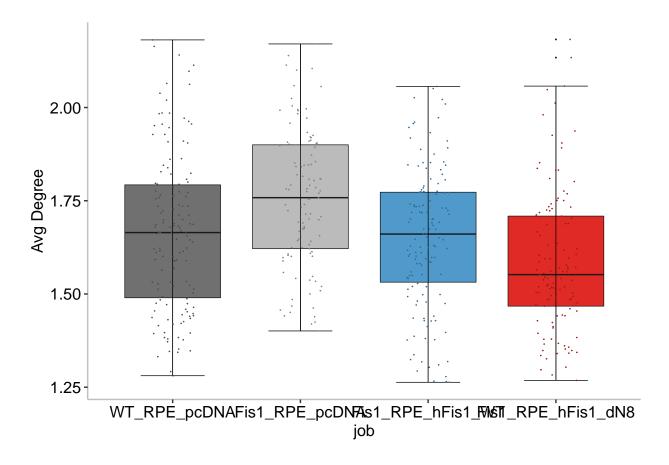


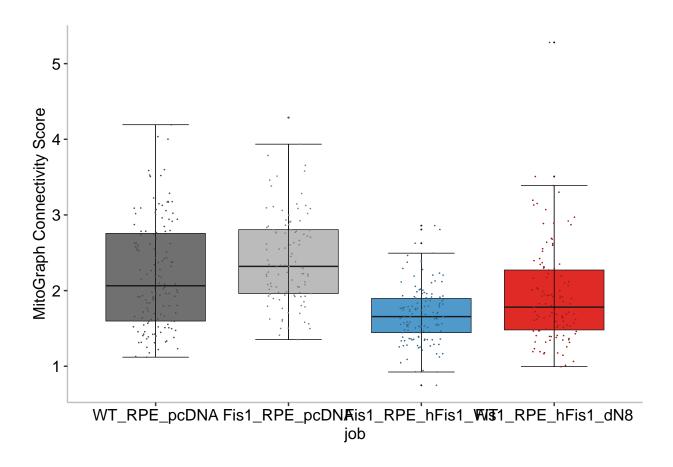


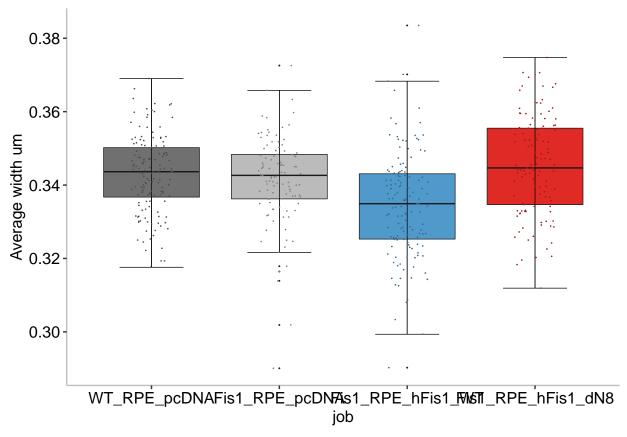






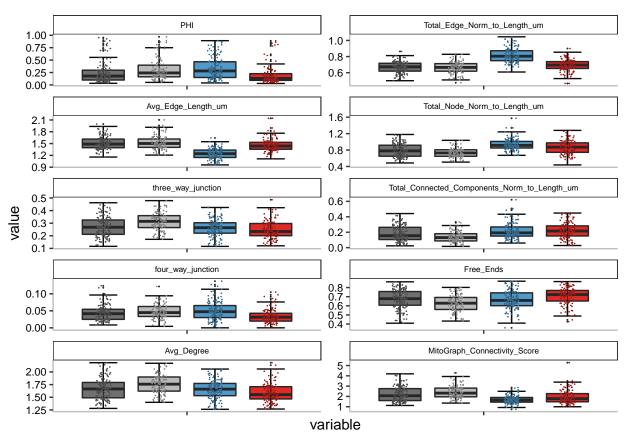






```
#test
data4 <- read_csv(file = "output-summary_filtered.csv")</pre>
## Rows: 496 Columns: 19
## -- Column specification
## Delimiter: ","
## chr (4): file, job, isoform, treatment
## dbl (15): Total_Nodes, Total_Edges, Total_Length_um, Total_Connected_Compone...
##
## i Use 'spec()' to retrieve the full column specification for this data.
## i Specify the column types or set 'show_col_types = FALSE' to quiet this message.
data5 <- data4 %>%
  mutate(job = factor(job, levels = c("WT_RPE_pcDNA",
                                       "Fis1_RPE_pcDNA",
                                       "Fis1_RPE_hFis1_WT",
                                       "Fis1_RPE_hFis1_dN8"), ordered = T)) #RENAME with your sample ord
data5_long <- reshape2::melt(data = data5, id.vars = "job", measure.vars = c("PHI",</pre>
                                                                               "Total_Edge_Norm_to_Length
                                                                               "Avg_Edge_Length_um",
                                                                               "Total_Node_Norm_to_Length
                                                                               "three_way_junction",
```

```
"Total_Connected_Component
                                                                              "four_way_junction",
                                                                              "Free Ends",
                                                                              "Avg Degree",
                                                                              "MitoGraph_Connectivity_Sc
                                                                    ))
ggplot(data5_long, aes(fill=job, x=variable, y=value)) +
  stat_boxplot(geom = "errorbar", colour = "grey15", width = 0.5, position = position_dodge (width = 1)
  geom_boxplot (outlier.size = 0, colour = "grey15", position = position_dodge (width = 1)) +
  stat_boxplot(geom="errorbar", position = position_dodge (width = 1), width = 0.5, size = 0.25) +
  scale_fill_manual(values = c( "#707070", "#BBBBBBB", "#4F9ACB", "#E02A26")) +
  facet_wrap(~variable, scales = "free", ncol = 2)+
  geom_point (aes(colour = job, x=variable, y=value), pch=20, position=position_jitterdodge(jitter.widt
  scale_colour_manual(values=c("#4e4e4e","#828282","#376b8e","#9c1d1a")) +
  theme_bw() +
  theme(axis.text.x = element_blank(),
        axis.text.y = element_text(size = 8, color = "black"),
        panel.grid.major = element_blank(),
       panel.grid.minor = element_blank(),
        panel.border = element_blank(),
        panel.background = element_blank(),
        axis.line.x = element_line(color = "grey75", size = 0.5, linetype = 1),
        axis.line.y = element_line(color = "grey75", size = 0.5, linetype = 1),
        legend.position = "none",
        #legend.title = element_blank(),
        \#legend.justification = c(0, 1),
        #legend.position = "right",
        #legend.text = element_text(size = 8, color = "black"),
        strip.background = element_rect(fill = "white"),
        strip.text.x = element_text(size = 6, colour = "black")
```



```
ggsave(paste("All_metrics-filtered.eps",sep=""), width = 9, height = 13, units = "cm", dpi = 300)
ggsave(paste("All_metrics-filtered.png",sep=""), width = 9, height = 13, units = "cm", dpi = 300)
#AOV_STATS (ANOVA and TUKEY Post Hoc analysis)
AOV_STATS <- TukeyHSD(aov(data=data, MitoGraph_Connectivity_Score-job))
AOV_Table <- (AOV_STATS)
AOV_Table_summary <- as.data.frame(AOV_Table[1:1])
write.csv(AOV_Table_summary, paste("MitoGraph_CS_AOV_stats.csv",sep=""))
formattable(AOV_Table_summary, Condition.p.adj=formatter("span", style = x-style(color=ifelse(x < 0.05
job.diff
job.lwr
job.upr
job.p.adj
Fisl_RPE_hFisl_WT-Fisl_RPE_hFisl_dN8</pre>
```

-0.2516827 -0.44550351 -0.05786193

4.854263 e-03

 $Fis1_RPE_pcDNA\text{-}Fis1_RPE_hFis1_dN8$

0.4916388

0.28571918

0.69755842

9.459815 e - 09

 $WT_RPE_pcDNA\text{-}Fis1_RPE_hFis1_dN8$

0.2807320

0.08885694

0.47260698

1.041522e-03

 $Fis1_RPE_pcDNA-Fis1_RPE_hFis1_WT$

0.7433215

0.54367547

0.94296757

9.724466e-11

 $WT_RPE_pcDNA\text{-}Fis1_RPE_hFis1_WT$

0.5324147

0.34728856

0.71754079

1.004776e-10

 $WT_RPE_pcDNA\text{-}Fis1_RPE_pcDNA$

-0.2109068

-0.40866445

-0.01314924

3.137726 e-02