Patient FLC and FLC Sphereoid miR correlation

Alaa R. Farghli

1/27/2021

```
library(dplyr)
library(ggpubr)
library(tibble)
library(ggrepel)
```

Load CSV

```
flc.counts <- read.csv("../NML-PRI-MET_noDups_avg-normalized-counts-by-group.csv")
flc.sphereoids <- read.csv("FLC-PDX_DESeq_miRs.csv")
flc.pri <- flc.counts %>%
    select(miR, avg_PRI)
flc.mets <- flc.counts %>%
    select(miR, avg_MET)
```

Workflow with primary patient FLC primary tumors

```
#Join intracellular sphereoids miRs with primary tumor miRs
flc.combined.pri <- full_join(flc.sphereoids, flc.pri, by = "miR")</pre>
#change is.na rows to 0
flc.combined.pri[is.na(flc.combined.pri)] <- 0</pre>
#Move the column to rowname. this is necessary for the final clean up step of removing rows that contai
flc.combined.pri <- column_to_rownames(flc.combined.pri, var = "miR")</pre>
flc.clean.100 <- flc.combined.pri %>%
  filter_all(any_vars(. > 100))
write.csv(x = flc.clean.100, "PrimaryVsSpheroids100count.csv")
#this is done again with reads at least 50 reads in either column
flc.clean.50 <- flc.combined.pri %>%
  filter_all(any_vars(. > 50))
write.csv(x = flc.clean.50, "PrimaryVsSpheroids50count.csv")
most.dev.pri.genes <- flc.clean.100 %>%
  rownames_to_column(var = "miR") %>%
  filter(miR %in% c("hsa-mir-122-5p", "hsa-mir-21-5p", "hsa-mir-320a", "hsa-mir-192-5p", "hsa-mir-126-3
```

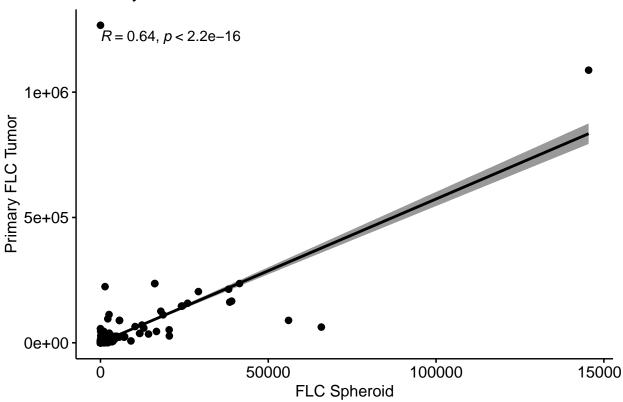
Plots for primary FLC primary tumors with and without log10 transformation

```
set.seed(1)
p.pri.noclean <- ggscatter(flc.combined.pri, x = "avg_Intra_FLC", y = "avg_PRI",</pre>
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "FLC Spheroid", ylab = "Primary FLC Tumor", title = "Primary FLC tumor miRS vs FLC-PDX
p.pri.100 <- ggscatter(flc.clean.100, x = "avg_Intra_FLC", y = "avg_PRI",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "FLC Spheroid", ylab = "Primary FLC Tumor", title = "Primary FLC tumor miRs vs FLC Sph
  geom_label_repel(data = most.dev.pri.genes,
                  aes(label = miR),
                  size = 3,
                  force = 100,
                  nudge_y = -30000)
p.pri.100.log10 <- ggscatter(flc.clean.100, x = "avg_Intra_FLC", y = "avg_PRI",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "log10 FLC Spheroid", ylab = "log 10 Primary FLC", title = "log10(Primary FLC tumor mi
  scale_x_log10() +
  scale_y_log10() +
  geom_label_repel(data = most.dev.pri.genes,
                  aes(label = miR),
                  size = 3) +
  geom label repel(data = most.dev.pri.genes,
                  aes(label = miR),
                  size = 3,
                  nudge_y = -1)
p.pri.50 <- ggscatter(flc.clean.50, x = "avg_Intra_FLC", y = "avg_PRI",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "FLC Spheroid", ylab = "Primary FLC Tumor", title = "Primary FLC tumor miRs vs FLC Sph
  geom_label_repel(data = most.dev.pri.genes,
                  aes(label = miR),
                  size = 3,
                  force = 100.
                  nudge_y = -30000)
p.pri.50.log10 <- ggscatter(flc.clean.50, x = "avg_Intra_FLC", y = "avg_PRI",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "log10 FLC Spheroid", ylab = "log 10 Primary FLC Tumor", title = "log10(Primary FLC tumor")
  scale x log10() +
  scale_y_log10() +
  geom_label_repel(data = most.dev.pri.genes,
                  aes(label = miR),
                  size = 3,
                  nudge_y = -1)
```

p.pri.noclean

'geom_smooth()' using formula 'y ~ x'

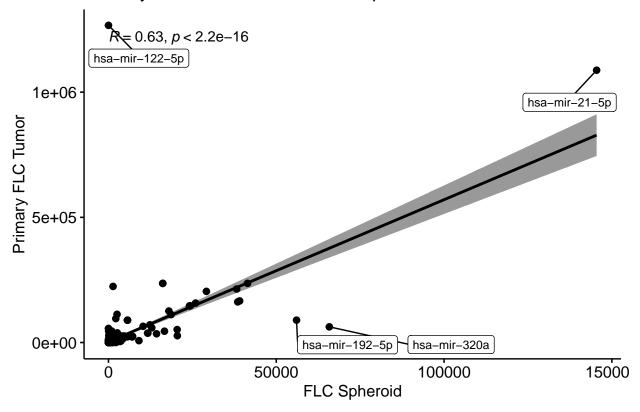
Primary FLC tumor miRS vs FLC-PDX miRs



p.pri.100

- ## 'geom_smooth()' using formula 'y ~ x'
- ## Warning: ggrepel: 9 unlabeled data points (too many overlaps). Consider
- ## increasing max.overlaps

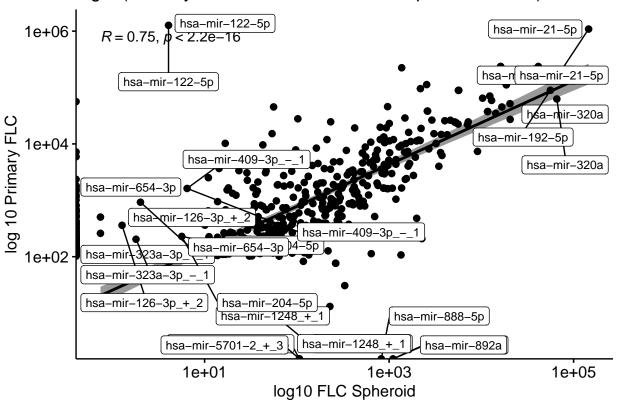
Primary FLC tumor miRs vs FLC Sphereoid miRs - 100+ reads in



p.pri.100.log10

Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous y-axis
"geom_smooth()" using formula "y ~ x"
Warning: Removed 119 rows containing non-finite values (stat_smooth).
Warning: Removed 119 rows containing non-finite values (stat_cor).

log10(Primary FLC tumor miRs vs FLC Sphereoid miRs) 100+ rea



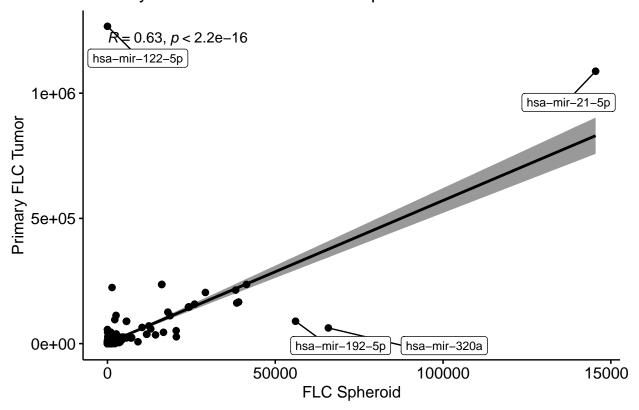
p.pri.50

```
## 'geom_smooth()' using formula 'y ~ x'
```

Warning: ggrepel: 9 unlabeled data points (too many overlaps). Consider

increasing max.overlaps

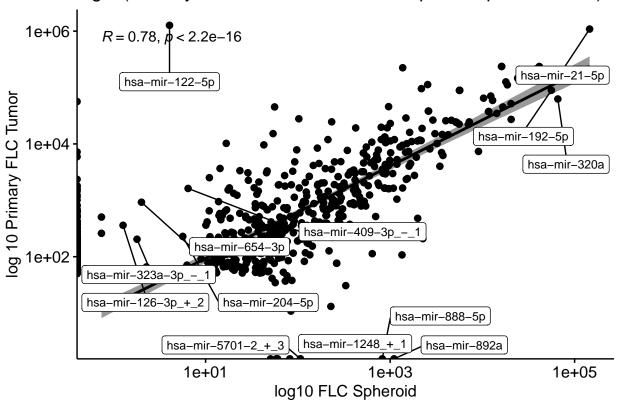
Primary FLC tumor miRs vs FLC Sphereoid miRs - 50+ reads in a



p.pri.50.log10

Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous y-axis
'geom_smooth()' using formula 'y ~ x'
Warning: Removed 213 rows containing non-finite values (stat_smooth).
Warning: Removed 213 rows containing non-finite values (stat_cor).

log10(Primary FLC tumor miRs vs DESeq_FLC Spheroid miRs) 50



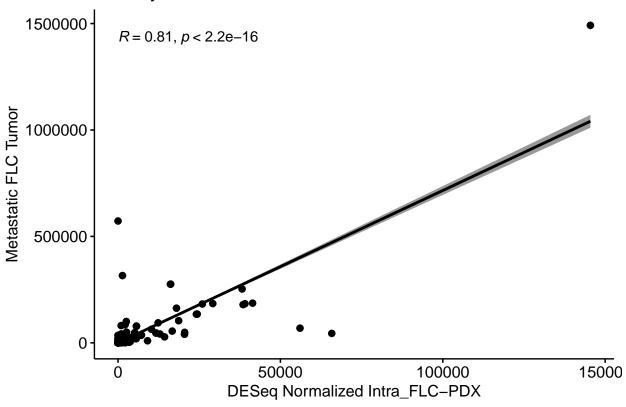
Workflow with primary patient FLC metastatic tumors

```
#The steps here are similar to that seen in code chunk 4
flc.combined.mets <- full_join(flc.mets, flc.sphereoids, by = "miR")</pre>
flc.combined.mets[is.na(flc.combined.mets)] <- 0</pre>
flc.combined.mets <- column_to_rownames(flc.combined.mets, var = "miR")
flc.clean.mets.100 <- flc.combined.mets %>%
  filter_all(any_vars(. > 100))
write.csv(x = flc.clean.mets.100, "MetastaticVsSpheroids100count.csv")
flc.clean.mets.50 <- flc.combined.mets %>%
  filter_all(any_vars(. > 50))
write.csv(x = flc.clean.mets.50, "MetastaticVsSpheroids50count.csv")
most.dev.met.genes <- flc.clean.mets.100 %>%
  rownames_to_column(var = "miR") %>%
  filter(miR %in% c("hsa-mir-21-5p", "hsa-mir-122-5p", "hsa-mir-143-3p", "hsa-mir-26a-1-5p", "hsa-mir-3
set.seed(1)
p.met.noclean <- ggscatter(flc.combined.mets, x = "avg_Intra_FLC", y = "avg_MET",</pre>
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "DESeq Normalized Intra_FLC-PDX", ylab = "Metastatic FLC Tumor", title = "Primary FLC
```

```
p.met.100 <- ggscatter(flc.clean.mets.100, x = "avg_Intra_FLC", y = "avg_MET",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "FLC Spheroid", ylab = "Metastatic FLC Tumor", title = "Metastatic FLC tumor miRs vs F
  geom_label_repel(data = most.dev.met.genes,
                  aes(label = miR),
                  size = 3,
                  force = 100,
                  nudge_x = 30000,
                  nudge_y = 300000)
p.met.100.log10 <- ggscatter(flc.clean.mets.100, x = "avg_Intra_FLC", y = "avg_MET",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "log10 FLC SpheroidC", ylab = "log10 Metastatic FLC Tumor", title = "log10(Metastatic Tumor")
  scale_x_log10() +
  scale_y_log10() +
  geom_text_repel(data = most.dev.met.genes,
                  aes(label = miR),
                  size = 3,
                  force = 100.
                  nudge_y = .5)
p.met.50 <- ggscatter(flc.clean.mets.50, x = "avg_Intra_FLC", y = "avg_MET",</pre>
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "FLC Spheroid", ylab = "Metastatic FLC Tumor", title = "Metastatic FLC tumor miRs vs F
  geom_label_repel(data = most.dev.met.genes,
                  aes(label = miR),
                  size = 3,
                  force = 100,
                  nudge_x = 30000,
                  nudge_y = 300000)
p.met.50.log10 <- ggscatter(flc.clean.mets.50, x = "avg_Intra_FLC", y = "avg_MET",
          add = "reg.line", conf.int = TRUE,
          cor.coef = TRUE, cor.method = "pearson",
          xlab = "log10 FLC Spheroid", ylab = "log10 Metastatic FLC Tumor", title = "log10 (Metastatic F.
  scale_x_log10() +
  scale_y_log10() +
  geom_text_repel(data = most.dev.met.genes,
                   aes(label = miR),
                   size = 3,
                   force = 100,
                  nudge_y = .5)
p.met.noclean
```

'geom_smooth()' using formula 'y ~ x'

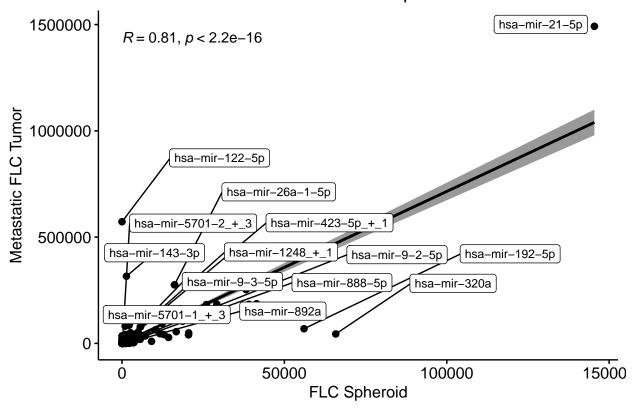
Primary FLC tumor miRS vs FLC miRs



p.met.100

'geom_smooth()' using formula 'y ~ x'

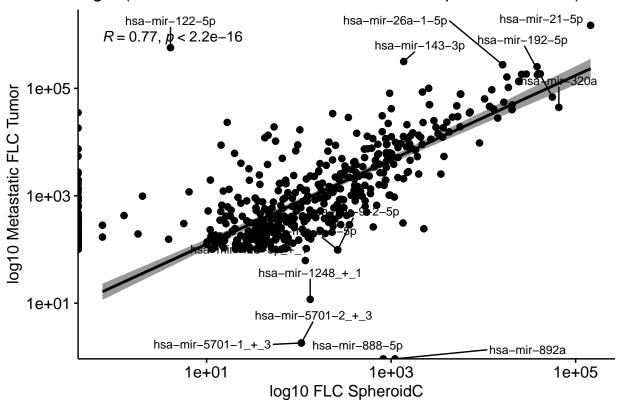
Metastatic FLC tumor miRs vs FLC sphereoids miRs - 100+ rea



p.met.100.log10

Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous y-axis
'geom_smooth()' using formula 'y ~ x'
Warning: Removed 120 rows containing non-finite values (stat_smooth).
Warning: Removed 120 rows containing non-finite values (stat_cor).

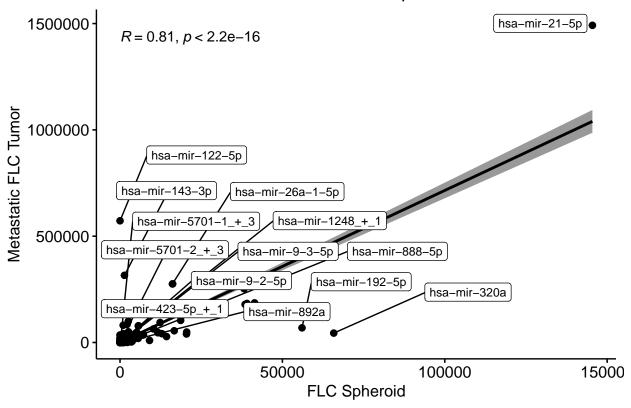
log10(Metastatic FLC tumor miRs vs _FLC sphereoids miRs) 100+



p.met.50

'geom_smooth()' using formula 'y ~ x'

Metastatic FLC tumor miRs vs FLC sphereoids miRs - 50+ read



p.met.50.log10

Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous x-axis
Warning: Transformation introduced infinite values in continuous y-axis
Warning: Transformation introduced infinite values in continuous y-axis
'geom_smooth()' using formula 'y ~ x'
Warning: Removed 203 rows containing non-finite values (stat_smooth).
Warning: Removed 203 rows containing non-finite values (stat_cor).

log10(Metastatic FLC tumor miRs vs _FLC sphereoids miRs) 50+

