R plotting

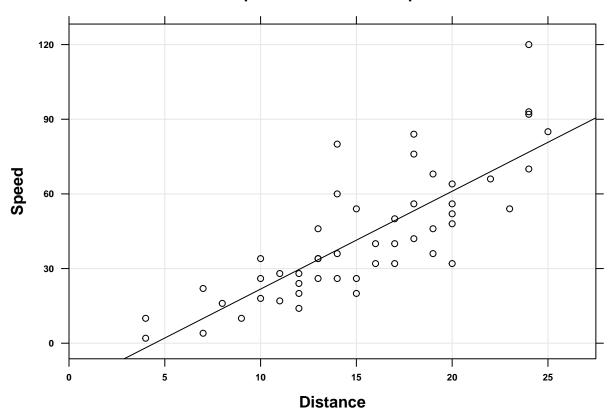
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Question 2

2a. Create a simple scatterplot using BoutrosLab.plotting.general

```
cars <- datasets::cars;</pre>
summary(cars);
##
       speed
                       dist
## Min. : 4.0 Min. : 2.00
## 1st Qu.:12.0 1st Qu.: 26.00
## Median: 15.0 Median: 36.00
## Mean :15.4 Mean : 42.98
## 3rd Qu.:19.0 3rd Qu.: 56.00
## Max. :25.0 Max. :120.00
create.scatterplot(
 formula = dist ~ speed,
 data = cars,
 main = "Scatter plot of distance and speed",
 xlab.label = "Distance",
 ylab.label = "Speed",
 yat = seq(0,150,30),
 xaxis.cex = 0.6,
 yaxis.cex = 0.6,
 xlab.cex = 1,
 ylab.cex = 1,
 main.cex = 1,
 pch = 1,
 col= "black",
 type = c ( "p" , "g" , "r" )
)
```

Scatter plot of distance and speed



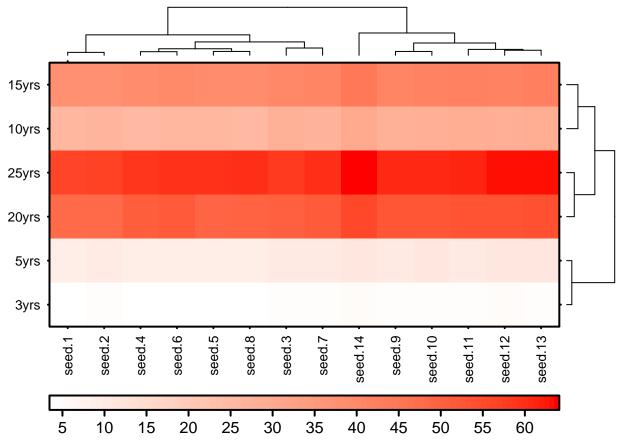
2b. Create a heatmap displaying data found in the 'Loblolly' dataset

```
loblolly <- datasets::Loblolly;</pre>
loblolly <- reshape(data = loblolly, idvar = "Seed",</pre>
                    v.names = "height",
                    timevar = "age",
                    direction = "wide");
loblolly$Seed <- as.numeric(loblolly$Seed);</pre>
loblolly.sort <- loblolly[order(loblolly$Seed),];</pre>
loblolly.matrix <- as.matrix(loblolly.sort[,-1]);</pre>
row.names(loblolly.matrix) <- paste("seed", 1:14, sep = ".");</pre>
colnames(loblolly.matrix) <- c("3yrs", "5yrs", "10yrs", "15yrs", "20yrs", "25yrs");</pre>
create.heatmap(
  x = loblolly.matrix,
  # format the colour key
  colourkey.cex = 1,
  colourkey.labels.at = seq(0, 70, 5),
  # set labels to NA -- results in default labels
  xaxis.lab = NA,
  yaxis.lab = NA,
  xaxis.cex = 0.8,
  yaxis.cex = 0.8,
```

```
# set font style (default is bold, 1 is roman)
xaxis.fontface = 1,
yaxis.fontface = 1,

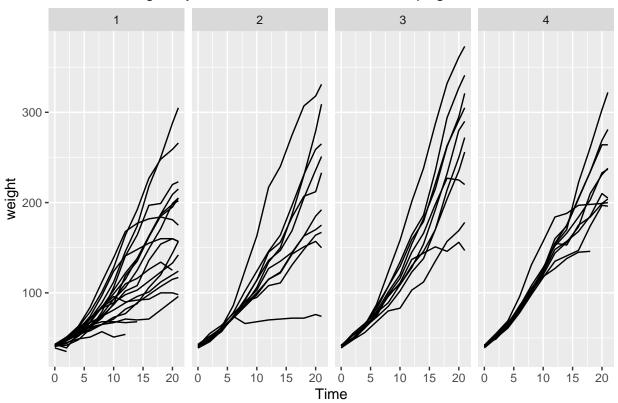
# specify clustering method
# if no clustering is desired, set this to "none"
clustering.method = "complete",

# select distance measure
rows.distance.method = "euclidean",
cols.distance.method = "manhattan"
);
```



2c. Take a look at the 'ChickWeight' dataset

Chicken Weight By Time Under Different Diet, Spaghetti Plot

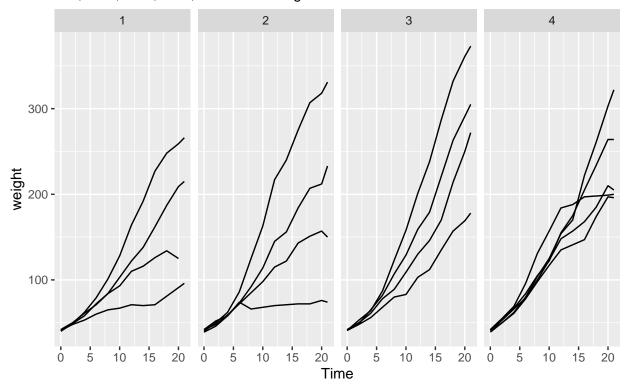


From the spaghetti plot, chicken tend to have a bigger increasement in weight under Diet 3.

```
chick.residual <- chickweight %>%
  group_by(Time,Diet) %>%
  mutate(meanweight = mean(weight)) %>%
  ungroup() %>%
  mutate(residual = weight - meanweight) %>%
  group_by(Chick,Diet) %>%
  mutate(median.residual = median(residual)) %>%
  ungroup();
chick.stat <- chick.residual %>%
  group by(Diet) %>%
  mutate(min = min(median.residual),
         q1 = quantile(median.residual,c(.25)),
         q2 = quantile(median.residual,c(.5)),
        q3 = quantile(median.residual,c(.75)),
       max = max(median.residual)) %>%
  ungroup()%>%
  select(Diet,min, q1, q2, q3, max) %>%
  unique();
chick.id <- chick.residual %>%
  filter(median.residual %in% as.matrix(chick.stat[,-1]));
ggplot(chick.id, aes(x = Time, y = weight, group = Chick)) +
```

Weight By Time Under Different Diet, Spaghetti Plot

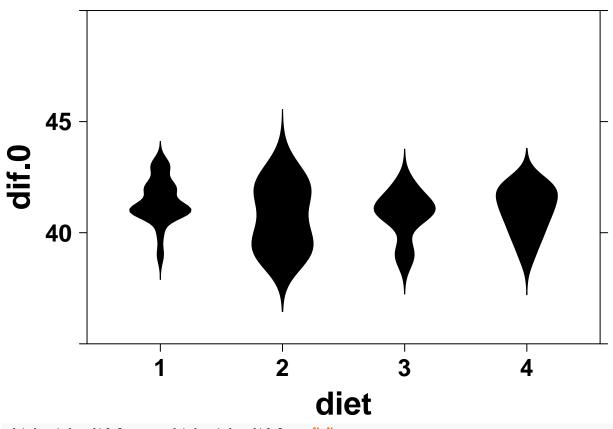
min, 25%, 50%, 75%, max of the weight residuals



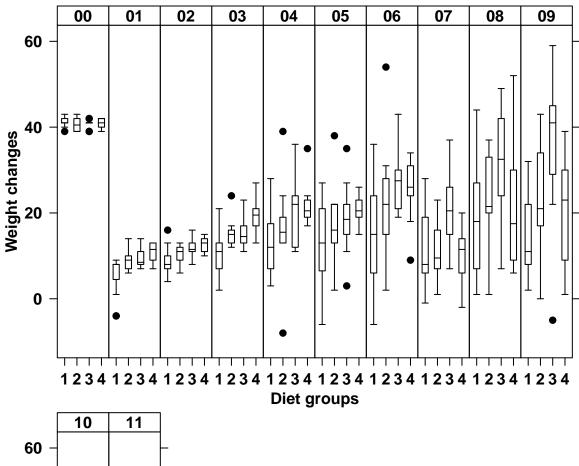
• Chicken tend to have higher vairation in weight as time goes by under Diet2. Least variation in weight was shown in chicken under diet4.

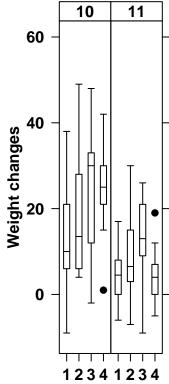
```
##Input variables:
#number of row, number of columns, column data type, and column names
#Output variables:
#empty dataset
#Description:
#Function that create empty dataframe
emptydf <- function(numrow, numcol, type, name){</pre>
 df <- data.frame(matrix(NA, nrow <- numrow, ncol <- numcol));</pre>
 for (i in 1:numcol){
   print(type[i])
   if('numeric' == type[i]) {df[,i] <- as.numeric(df[,i])</pre>
   colnames(df)[i] <- name[i]};</pre>
   if('character' == type[i]) {df[,i] <- as.character(df[,i])</pre>
   colnames(df)[i] <- name[i]};</pre>
   if('logical' == type[i]) {df[,i] <- as.logical(df[,i])</pre>
   colnames(df)[i] <- name[i]};</pre>
   if('factor' == type[i]) {df[,i] <- as.factor(df[,i])</pre>
   colnames(df)[i] <- name[i]};</pre>
```

```
return(df);
# chicken weight difference by time, box plot
chickweight.dif <- emptydf(nlevels(as.factor(chickweight$Chick)),</pre>
                           nlevels(as.factor(chickweight$Time))+2 ,
                       c('character', 'character', rep('numeric',nlevels(as.factor(chickweight$Time)) ))
                         c('chick', 'diet', paste("dif", 0 : nlevels(as.factor(chickweight$Time)) , sep
## [1] "character"
## [1] "character"
## [1] "numeric"
chickweight$Time <- as.character(chickweight$Time);</pre>
chickweight.wide <- chickweight %>%
  group_by(Chick, Diet) %>%
  spread(Time, weight, fill=NA, sep = ".");
chickweight.wide < chickweight.wide[, c(01,02,03,09,12,13,14,04,05,06,07,08,10,11)];
# Calculate weight difference between each measurement
for(i in 1: (nlevels(as.factor(chickweight$Time))-1) ){
  chickweight.dif [,i+3]<- chickweight.wide[,i+3]-chickweight.wide[,i+2];</pre>
}
chickweight.dif[,1:3] <- chickweight.wide[,1:3];</pre>
chickweight.dif.long <- chickweight.dif %>%
  gather(interval,dif,dif.0:dif.11 );
create.violinplot(formula = dif.0 ~ diet,
               data = chickweight.dif,
               ylimits = c(35,50)
);
```



```
chickweight.dif.long = chickweight.dif.long %>%
  mutate ( interval.num = sprintf("%02d", as.numeric(substr(interval,5,6))));
diet.colour <- recode.vector(</pre>
    chickweight.dif.long$diet,
    list(
        blue = 1,
        hazel = 2,
       green = 3,
       brown = 4
        )
    );
create.boxplot(formula = dif ~ diet | interval.num,
    data = chickweight.dif.long,
    xaxis.cex = 1,
    yaxis.cex = 1,
    ylab.label = "Weight changes",
    xlab.label = "Diet groups",
    xlab.cex = 1,
    ylab.cex = 1,
    layout = c(10,1)
    );
```





Diet groups

- From the boxplot, we can find out that there is not a big difference in weight between 4 groups at time0.
- Chicken on Diet 1 tend to have the least weight changes, while chicken on Diet 3 have the most weight

changes during the time.

- Chicken on Diet 3 tend to have the most weight changes in the last few weeks.
- In the last few weeks the weight changes tend to have higher vairance among diet groups.

Question 3

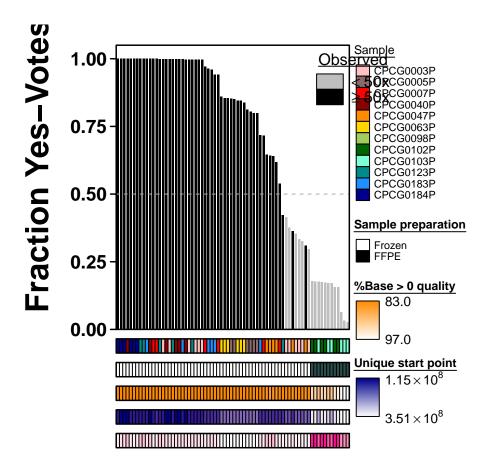
```
seq.control <- read.table("/cloud/project/Q3_SeqControl_data", header = T);</pre>
# Ste 1 Reorder data
seq.control1 <- seq.control[order(seq.control$yes.votes, decreasing = T),];</pre>
# Step 2 Create the CPCG bars
colour.scheme.large <- c(</pre>
  'rosybrown1',
  'rosybrown4',
  'red',
  'darkred',
  'darkorange',
  'gold',
  'darkolivegreen3',
  'darkgreen',
  'aquamarine',
  'cyan4',
  'dodgerblue',
  'darkblue',
  'plum',
  'magenta',
  'darkorchid',
  'purple4',
  'gray70',
  'gray30'
);
##Input variables:
# data that need to create a bar plot, colour scheme
#Output variables:
# covariate bars
#Description:
#Function that create covariate bars
heatmap <- function(data,colour){</pre>
 plot <- create.heatmap(</pre>
   x <- t(as.matrix(as.numeric(data))),
   clustering.method = "none",
   scale.data = FALSE,
   colour.scheme = colour,
   total.col = 12,
   force.grid.col = TRUE,
   grid.col = TRUE,
   print.colour.key = FALSE,
    # remove y-axis ticks
```

```
yaxis.tck = 0,
    height = 1,
    xaxis.lab = NULL,
    yaxis.lab = NULL,
    yat = 1,
    yaxis.cex = 1);
  return(plot);
gene.heatmap <- heatmap(</pre>
  data = seq.control1$CPCG,
  colour = colour.scheme.large[1:12]);
average.reads.start.heatmap <- heatmap(</pre>
  data = seq.control1$Average.reads.start,
  colour = c("white", "deeppink"));
unique.start.points.heatmap <- heatmap(</pre>
  data = seq.control1$Unique.start.points,
  colour = c("white", "darkblue"));
x.base.O.quality.heatmap <- heatmap(</pre>
  data = seq.control1$X..Bases...0.quality,
          colour = c("white", "darkorange"));
# Create FFPE bar
seq.control1 <- seq.control1 %>%
  dplyr::mutate(FFPE = ifelse("CPCG0102P" == CPCG | "CPCG0103P" == CPCG,1,0) );
FFPE.heatmap <- heatmap(data = seq.control1$FFPE,
                        colour = c("white", "darkslategrey"));
# Step 5 Create the barplot
barplot.colour.choice <- c("grey", "black");</pre>
barplot.colour <- barplot.colour.choice[factor(seq.control1$outcome, levels <- c(0,1))];</pre>
yes.votes.barplot <- create.barplot(</pre>
 formula = yes.votes ~ c(1:72),
  border.col = 'transparent',
 data = seq.control1,
  col = barplot.colour,
  right.padding = 2,
  xaxis.cex = 0.5,
  abline.h = 0.5,
 abline.lty = 2,
  abline.col = 'darkgrey'
);
# Step 6 Create a legend for each of the covariates
# create legend for cpcgene sample heatmap
  sample.legend <- list(</pre>
    legend = list(
```

```
colours = colour.scheme.large[1:12],
    title = expression(underline("Sample")),
    labels = levels(seq.control1$CPCG),
    continuous = FALSE
)
);
# create legend for outcome heatmap
 prep.legend <- list(</pre>
    legend = list(
      colours = c('white', 'black'),
      labels = c('Frozen', 'FFPE'),
      title = expression(bold(underline('Sample preparation'))),
      continuous = FALSE
    )
);
# create legend for x..base....O.quality heatmap
qual.legend <- list(
 legend = list(
  colours = c("white", "darkorange"),
 labels = c("97.0", "83.0"),
 title = expression(bold(underline('%Base > 0 quality'))),
 continuous = TRUE
 )
);
# create legend for unique.start.point.heatmap
1.s <- scientific.notation(x=min(seq.control1$Unique.start.points),digits=2);</pre>
h.s <- scientific.notation(x=max(seq.control1$Unique.start.points),digits=2);
  uni.start.legend <- list(
    legend = list(
      colours = c("white", "darkblue"),
      labels = c(h.s, l.s),
      title = expression(bold(underline('Unique start point'))),
      continuous = TRUE
    )
  );
# create legend for average.reads.start.heatmap
1.r <- round(min(seq.control1$Average.reads.start),2);</pre>
h.r <- round(max(seq.control1$Average.reads.start),2);</pre>
  ave.start.legend <- list(</pre>
    legend <- list(</pre>
      colours <- c("white", "deeppink"),</pre>
      labels \leftarrow c(h.r, l.r),
      title <- expression(bold(underline('Unique start point'))),</pre>
      continuous <- TRUE
```

```
)
  );
#combine all covariates legends
covariate.legends <- c(</pre>
  sample.legend,
  prep.legend,
  qual.legend,
  uni.start.legend,
  ave.start.legend
);
legends1 <- BoutrosLab.plotting.general::legend.grob(</pre>
  legends = covariate.legends,
 title.cex = 0.75,
  title.just = 'left',
  label.cex = 0.65,
  size = 1.5,
  between.row = 1.0,
 between.col = 0.5,
layout = c(1,6)
);
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
# create legend for barplot
barplot.legend <- list(</pre>
legend = list(
    colours = c('grey', 'black'),
    labels = c(
      as.expression(substitute(x < '50x',list(x = ''))),
      as.expression(substitute(x >= '50x',list(x = '')))
    ),
    title = expression(underline('Observed'))
  )
);
legends2 <- BoutrosLab.plotting.general::legend.grob(barplot.legend);</pre>
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
# Multiplot
  plot.objects <- list(</pre>
    average.reads.start.heatmap,
    unique.start.points.heatmap,
    x.base.O.quality.heatmap,
    FFPE.heatmap,
    gene.heatmap,
    yes.votes.barplot
  );
```

```
# identify where plotting objects should be placed in the multiplot
 yat.vals <- list();</pre>
  for (n in 1:(length(plot.objects)-1)) {
   yat.vals <- c(yat.vals, list(NULL));</pre>
  }
# combine plots
  create.multiplot(
   plot.objects = plot.objects,
   #filename = 'testing_votes.tiff',
   panel.heights = c(1, rep(0.05, length(plot.objects)-1)),
   yat = c(yat.vals, list(seq(0,1,0.25))),
   yaxis.cex = 1.15,
   ylab.label = c(' ', 'Fraction Yes-Votes', ' ', ' ', ' '),
   ylab.padding = 6.5,
   right.padding = 35,
   bottom.padding = -5,
   xat = NULL,
   y.spacing = 0.5,
   ylimits = list(c(0.9,1), c(0.9,1), c(0.9,1), c(0.9,1), c(0.9,1), c(0.0,1.05)),
   legend = list(
      inside = list(
       fun = legends1,
       x = 1.02,
       y = 1
     ),
      inside = list(
       x = 0.85
       y = 0.98,
       fun = legends2
      )
   ),
     print.new.legend = TRUE,
   width = 12,
   height = 6,
   resolution = 1200,
 );
## Warning in formals(fun): argument is not a function
## Warning in formals(fun): argument is not a function
## Warning in formals(fun): argument is not a function
## Warning in formals(fun): argument is not a function
## Warning in formals(fun): argument is not a function
```



Question 4

```
het <- read.table("/cloud/project/Q4 HetStudy data.txt", header=T);</pre>
# create covariate table
covariate <-as.data.frame(colnames(het[,1:28]));</pre>
names(covariate) <- c("id");</pre>
covariate$sample <- substr(covariate$id, 1,8);</pre>
covariate$cohort <- c(rep("Bx",5),rep("Sx",23));</pre>
# convert gleason score 3+4 as 1, 4+3 as 2, 4+4 as 3
covariate$gleason.score <- c(1,1,2,1,2,1,2,2,3,3,1,1,1,1,1,2,1,2,2,2,1,1,1,3,2,2,2,1);
covariate$gleason.score.plus <- c(1,rep(NA,14),rep(1,4),rep(NA,9));</pre>
covariate$tissue.type <- substr(covariate$id, 9,10);</pre>
# create matrix for gleason score positive
gleason.score.plus.matrix <- data.frame(matrix(NA, nrow <- 28, ncol <- 2));</pre>
gleason.score.plus.matrix <- cbind(gleason.score.plus.matrix,covariate$gleason.score.plus);</pre>
gleason.score.plus.matrix <- cbind(gleason.score.plus.matrix,data.frame(matrix(NA, nrow <- 28, ncol <-
covariate.numeric <- data.frame(lapply(covariate, as.character),stringsAsFactors = FALSE);</pre>
covariate.numeric <- covariate.numeric[,2:6];</pre>
covariate.numeric <- covariate.numeric %>%
  dplyr::select(-c("gleason.score.plus"));
```

```
# convert covariate data to numeric
# create sample id
covariate.numeric <- covariate.numeric %>%
  dplyr::mutate(sample.id = as.integer(as.factor(sample)));
covariate.numeric$sample <- as.character(covariate.numeric$sample.id);</pre>
covariate.numeric <- covariate.numeric[,-5];</pre>
#convert Bx (biopsy) as 11, and Sx (surgery) as 12
covariate.numeric$cohort["Bx" == covariate.numeric$cohort] = 11;
covariate.numeric$cohort["Sx" == covariate.numeric$cohort] = 12;
#convert gleason score
covariate.numeric$gleason.score[1 == covariate.numeric$gleason.score] <- 13;</pre>
covariate.numeric$gleason.score[2 == covariate.numeric$gleason.score] <- 14;</pre>
covariate.numeric$gleason.score[3 == covariate.numeric$gleason.score] <- 15;</pre>
# concert "FO" samples are Frozen(2), while all other samples are FPPE(1)
covariate.numeric$tissue.type["F0" != covariate.numeric$tissue.type] <- 17;</pre>
covariate.numeric$tissue.type["F0" == covariate.numeric$tissue.type] <- 16;</pre>
# set colour scheme
sample.colour <- c( 'blue', 'purple', 'green', 'orange', 'yellow', 'black', 'wheat4', 'green4', 'grey', 'red4');</pre>
cohort.colour <- c('royalblue', 'pink');</pre>
gleason.score.colour <- c("yellow1", "orange", "red");</pre>
tissue.type.colour <- c('Frozen' = colours()[532],'FFPE' = colours()[557]);
# create covairate bar on the right
covariate.bar <- create.heatmap(x = t(data.matrix(covariate.numeric)),</pre>
                                 clustering.method = "none",
                                 print.colour.key = FALSE,
                                 total.colours = 17,
                                 colour.scheme = c(sample.colour,cohort.colour,gleason.score.colour,tiss
                                 at=seq(0.5,17.5,1),
                                 # add row lines
                                 force.grid.col = TRUE,
                                 grid.col = TRUE,
                                 grid.row = TRUE,
                                 row.colour = "black",
                                 col.colour = "black",
                                 row.pos = which(1 == t(gleason.score.plus.matrix[1:28,1:4]), arr.ind = '
                                 col.pos = which(1 == t(gleason.score.plus.matrix[1:28,1:4]), arr.ind = '
                                 cell.text =rep("+", 5),
                                 text.cex = 1,
                                 xaxis.tck = 0,
                                 yaxis.tck = 0
);
# create fraction plot
het.frac <- het %>%
  dplyr::select(-Baca, -Berger, -Weischenfeldt);
het.frac$frac <- rowSums(het.frac != 0)/28;
het.frac <- het.frac%>%
```

```
dplyr::mutate(
    ends = sapply( strsplit(rownames(het),"-", fixed = TRUE),tail, 1),
    chr = sapply( strsplit(rownames(het), ":", fixed = TRUE), head, 1),
    number = c(1:3113);
het.frac<- het.frac%>%
  dplyr::mutate(chr.num = gsub("chr", "", chr));
frac.plot <- create.barplot(formula = frac ~ number,</pre>
                            data = het.frac,
                            main = NULL,
                            stack = FALSE,
                            xlab.top.label = NULL,
                            xlab.label = NULL,
                            ylab.label = "Fraction",
                            ylab.cex = 0.8,
                            xaxis.tck = 0,
                            xaxis.lab = rep('',3113),
                            ylimits = c(0, 0.5),
                            yat = seq(0,0.6,0.25),
                            yaxis.cex = 0.6,
                            yaxis.tck = c(1,0)
);
# create literature covariate bars
het.literature <- cbind(het.frac[,30:32],het[,29:31]);</pre>
het.literature <- data.frame(lapply(het.literature, as.character),stringsAsFactors = FALSE);</pre>
# set colour scheme
literature.colour <- c('white', 'darkred', 'cornflowerblue', 'darkolivegreen4');</pre>
# create
publication.bar <- create.heatmap(x =data.matrix(het.literature[,4:6]),</pre>
                                   clustering.method = "none",
                                   print.colour.key = FALSE,
                                   #otal.colours = 4,
                                   colour.scheme = c( literature.colour ),
                                   # add row lines
                                   at=seq(-0.5,3.5,1),
                                   col.lines = which("1000000" == het.literature$ends),
                                   force.grid.col = TRUE,
                                   grid.col = TRUE,
                                   force.grid.row = TRUE,
                                   grid.row = TRUE,
                                   row.colour = "black",
                                   yaxis.tck = 0
);
# create main heatmap
literature.colour <- c('white', 'cornflowerblue', 'darkolivegreen4', 'darkred');</pre>
```

```
# create x axis location
het.frac.plot <- het.frac%>%
  dplyr::group by(chr.num) %>%
  dplyr::mutate(x.location = ifelse (number == round(mean(number), digits = 0),1,0));
het.frac.plot <- data.frame(lapply(het.frac.plot, as.character),stringsAsFactors = FALSE);</pre>
# create y axis location
covariate <- covariate %>%
  dplyr::group_by(sample) %>%
  dplyr::mutate(first = row_number() == min( row_number() ));
# create main heat map
main.heatmap <- create.heatmap(x = data.matrix(het.frac.plot[,1:28]),</pre>
                               clustering.method = "none",
                               print.colour.key = FALSE,
                               total.colours = 4,
                               colour.scheme = c(literature.colour),
                               at = seq(-0.5, 3.5, 1),
                               # add row lines
                               row.lines = which(TRUE == covariate$first)+0.5,
                               grid.row = TRUE,
                               # add col lines
                               col.lines = which("1000000" == het.literature$ends),
                               force.grid.col = TRUE,
                               grid.col = TRUE,
                               \#set\ labels\ for\ x\ and\ y\ axis
                               xaxis.lab = c(1:22, "X", "Y"),
                               xat = which(1 == het.frac.plot$x.location),
                               xaxis.cex = 0.6,
                               xaxis.rot = 0,
                               yaxis.lab = covariate$tissue.type,
                               yaxis.tck = c(1,0),
                               yaxis.cex = 0.6
                               #row.colour = "black",
                               #col.colour = "black"
);
# create the bottom notation heatmap
notation <- matrix(1:4, nrow = 1);</pre>
notation.heatmap <- create.heatmap(x = notation,
                                   clustering.method = "none",
                                   print.colour.key = FALSE,
                                   total.colours = 4,
                                   colour.scheme = c('white', 'cornflowerblue', 'darkolivegreen4', 'darkre
                                   at = seq(0.5, 4.5, 1),
                                   force.grid.col = TRUE,
                                   grid.col = TRUE,
```

```
xaxis.lab = c("", "None", "CTX", "ITX","INV"),
                                   xaxis.cex = 0.6,
                                   xaxis.rot = 0,
                                   yaxis.lab = NULL,
                                   yaxis.cex = 0.6,
                                   yaxis.tck = 0,
                                   right.padding = 0
);
#create legend on the left
# create legend for samples
pid.legend <- list(</pre>
 legend = list(
    colours = c( 'blue', 'purple', 'green', 'orange', 'yellow', 'black', 'wheat4', 'green4', 'grey', 'red4'),
    labels = unique(covariate$sample),
    title = expression(bold(underline('Patient ID'))),
    continuous = FALSE
  )
);
# create legend for cohort
cohort.legend <- list(</pre>
  legend = list(
    colours = c('pink', 'royalblue'),
    labels = c('Sx', 'Bx'),
    title = expression(bold(underline('Cohort'))),
    continuous = FALSE
  )
);
# create legend for Gleason score
gs.legend <- list(</pre>
  legend = list(
    colours = c('yellow', 'orange','red'),
    labels = c('3+4', '4+3', '4+4'),
    title = expression(bold(underline('Gleason score'))),
    continuous = FALSE
  )
);
# create legend for Tissue type
tissue.legend <- list(</pre>
  legend = list(
    colours = c(colours()[532],colours()[557]),
    labels = c('FFPE', 'Frozen'),
    title = expression(bold(underline('Tissue type'))),
    continuous = FALSE
  )
);
# create legend for Publication
```

```
pub.legend <- list(</pre>
  legend = list(
    colours = c('darkred', 'cornflowerblue', 'darkolivegreen4'),
    labels = c('Baca', 'Berger', "Weischenfeldt"),
    title = expression(bold(underline('Publication'))),
    continuous = FALSE
  )
);
left.legends <- legend.grob(</pre>
  legends = c(pid.legend, cohort.legend, gs.legend, tissue.legend, pub.legend),
  title.cex = 0.75,
 title.just = 'left',
 label.cex = 0.65,
  size = 1,
  between.row = 1.0,
  between.col = 0.5
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
## Warning in FUN(X[[i]], ...): 'x' is NULL so the result will be NULL
# create multiplot
plot.objects <- list(frac.plot,publication.bar, main.heatmap,covariate.bar,notation.heatmap );</pre>
create.multipanelplot(
  plot.objects = plot.objects,
  #filename = 'HetStudy.tiff',
  plot.objects.heights = c(0.6, 0.3, 1.4, 0.2),
  plot.objects.widths = c(1.1, 0.1),
  layout.skip = c(FALSE, TRUE, FALSE, TRUE, FALSE, FALSE, TRUE),
  layout.height = 4,
  layout.width = 2,
  #yaxis.cex = 1.15,
  ylab.axis.padding = 3.5,
  y.spacing = c(-1,-1,-1),
  x.spacing = 0,
  legend = list(
   left = list(
     fun = left.legends,
     x = 1.02,
      y = 1
    )
  ),
```

```
left.legend.padding = 2,

width = 12,
height = 6,
resolution = 1200
```

Warning in rbind(plot.objects.heights, y.spacing): number of columns of
result is not a multiple of vector length (arg 2)

