

## Coronavirus Liver and Blood Capillary Samples

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These samples are the headers added from three Gene Expression Omnibus studies at

- [ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE89166](https://ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE89166)
- [ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE89160](https://ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE89160)
- [ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE100509](https://ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE100509)

The first two studies are part of the same study that used human liver tumor samples in vitro to compare the effects of the coronavirus over time. The third study used human microvascular blood capillaries in vitro to study the effects of the coronavirus over time.

In the first two studies that used the liver tumor samples to examine the effects of the coronavirus in vitro, there were four groups inoculated or treated with the active coronavirus and four groups not inoculated with the active coronavirus, and two samples that were treated with heat inactivated coronavirus, and two samples that were treated with active coronavirus and IL-1alpha to see the gene expression changes over one hour's time.

In the the third study that used blood capillaries, there were five samples followed over a 0,12,24,36, and 48 hour time intervals in groups A,B,C,D, and E that compared the time interval values of screening for changes in microarray analysis with a mock group of the same.

This following data is the data of all genes in common between these three studies, cleaned to remove missing values and with the attached gene symbols from the GEO platform for the probe IDs.

---

The libraries used for this script are tidyr, dplyr, and ggplot2.

```
library(ggplot2)
library(dplyr)
library(tidyr)
```

---

```
both <- read.csv('both_clean_liver_capillary_CoV.csv', sep=',', header=TRUE,
                 na.strings=c('', ' '))
```

```
dim(both)
```

```
## [1] 21754    63
```

colnames(both)

```
## [1] "GENE_SYMBOL"
## [2] "LiverTumorSamples.GSM2359851_CoV1"
## [3] "LiverTumorSamples.GSM2359853_CoV2"
## [4] "LiverTumorSamples.GSM2359910_CoV3"
## [5] "LiverTumorSamples.GSM2359913_CoV4"
## [6] "LiverTumorSamples.GSM2359850_ctrl1"
## [7] "LiverTumorSamples.GSM2359852_ctrl2"
## [8] "LiverTumorSamples.GSM2359911_ctrl3"
## [9] "LiverTumorSamples.GSM2359914_ctrl4"
## [10] "LiverTumorSamples.GSM2359912_IL1"
## [11] "LiverTumorSamples.GSM2359917_IL2"
## [12] "LiverTumorSamples.GSM2359915_inactiveHeatCoV1"
## [13] "LiverTumorSamples.GSM2359916_inactiveHeatCoV2"
## [14] "capillarySamples.GSM2685693_MERS_CoV_0hr_A"
## [15] "capillarySamples.GSM2685694_MERS_CoV_0hr_B"
## [16] "capillarySamples.GSM2685695_MERS_CoV_0hr_C"
## [17] "capillarySamples.GSM2685696_MERS_CoV_0hr_D"
## [18] "capillarySamples.GSM2685697_MERS_CoV_0hr_E"
## [19] "capillarySamples.GSM2685698_ctrl_0hr_A"
## [20] "capillarySamples.GSM2685699_ctrl_0hr_B"
## [21] "capillarySamples.GSM2685700_ctrl_0hr_C"
## [22] "capillarySamples.GSM2685701_ctrl_0hr_D"
## [23] "capillarySamples.GSM2685702_ctrl_0hr_E"
## [24] "capillarySamples.GSM2685703_MERS_CoV_12hr_A"
## [25] "capillarySamples.GSM2685704_MERS_CoV_12hr_B"
## [26] "capillarySamples.GSM2685705_MERS_CoV_12hr_C"
## [27] "capillarySamples.GSM2685706_MERS_CoV_12hr_D"
## [28] "capillarySamples.GSM2685707_MERS_CoV_12hr_E"
## [29] "capillarySamples.GSM2685708_ctrl_12hr_A"
## [30] "capillarySamples.GSM2685709_ctrl_12hr_B"
## [31] "capillarySamples.GSM2685710_ctrl_12hr_C"
## [32] "capillarySamples.GSM2685711_ctrl_12hr_D"
## [33] "capillarySamples.GSM2685712_ctrl_12hr_E"
## [34] "capillarySamples.GSM2685713_MERS_CoV_24hr_A"
## [35] "capillarySamples.GSM2685714_MERS_CoV_24hr_B"
## [36] "capillarySamples.GSM2685715_MERS_CoV_24hr_C"
## [37] "capillarySamples.GSM2685716_MERS_CoV_24hr_D"
## [38] "capillarySamples.GSM2685717_MERS_CoV_24hr_E"
## [39] "capillarySamples.GSM2685718_ctrl_24hr_A"
## [40] "capillarySamples.GSM2685719_ctrl_24hr_B"
## [41] "capillarySamples.GSM2685720_ctrl_24hr_C"
## [42] "capillarySamples.GSM2685721_ctrl_24hr_D"
## [43] "capillarySamples.GSM2685722_ctrl_24hr_E"
## [44] "capillarySamples.GSM2685723_MERS_CoV_36hr_A"
## [45] "capillarySamples.GSM2685724_MERS_CoV_36hr_B"
## [46] "capillarySamples.GSM2685725_MERS_CoV_36hr_C"
## [47] "capillarySamples.GSM2685726_MERS_CoV_36hr_D"
## [48] "capillarySamples.GSM2685727_MERS_CoV_36hr_E"
```

```
## [49] "capillarySamples.GSM2685728_ctrl_36hr_A"
## [50] "capillarySamples.GSM2685729_ctrl_36hr_B"
## [51] "capillarySamples.GSM2685730_ctrl_36hr_C"
## [52] "capillarySamples.GSM2685731_ctrl_36hr_D"
## [53] "capillarySamples.GSM2685732_ctrl_36hr_E"
## [54] "capillarySamples.GSM2685733_MERS_CoV_48hr_A"
## [55] "capillarySamples.GSM2685734_MERS_CoV_48hr_B"
## [56] "capillarySamples.GSM2685735_MERS_CoV_48hr_C"
## [57] "capillarySamples.GSM2685736_MERS_CoV_48hr_D"
## [58] "capillarySamples.GSM2685737_MERS_CoV_48hr_E"
## [59] "capillarySamples.GSM2685738_ctrl_48hr_A"
## [60] "capillarySamples.GSM2685739_ctrl_48hr_B"
## [61] "capillarySamples.GSM2685740_ctrl_48hr_C"
## [62] "capillarySamples.GSM2685741_ctrl_48hr_D"
## [63] "capillarySamples.GSM2685742_ctrl_48hr_E"
```

---

Lets group the samples that are our columns with descriptive and GEO ID names into their respective groups, get the fold change between the controls from those groups, attach to the original data table, both, as a different names, then order by the genes that have the most fold change then the least fold change. Take the first 100 genes from both lists, combine into one table of 200 genes and the samples with their fold change values ordered, make into a transposed data frame so that the samples are the rows, the stats removed, and the 200 genes are the header columns to save as a machine learning ready file. \*\*\*

Liver tumor study control and CoV treated. Also, the IL-alpha treated and the inactive CoV treated tables are in this code block.

```
names <- both$GENE_SYMBOL

liverCtrl <- both[,c(6:9)]
row.names(liverCtrl) <- names

liverCoV <- both[,c(2:5)]
row.names(liverCoV) <- names

liverIL <- both[,10:11]
row.names(liverIL) <- names

liverIACoV <- both[,12:13]
row.names(liverIACoV) <- names
```

Get the row means of those liver samples groups each.

```
liverCtrl$CtrlMeanLvr <- rowMeans(liverCtrl)
liverCoV$CoVMeanLvr <- rowMeans(liverCoV)
liverIL$ILMeanLvr <- rowMeans(liverIL)
liverIACoV$IACoVMeanLvr <- rowMeans(liverIACoV)
```

Get the fold change values of those states as a ratio to the control group values.

```
fold1 <-  
as.data.frame(cbind(liverCtrl$CtrlMeanLvr,liverCoV$CoVMeanLvr,liverIL$ILMeanLvr,  
                    liverIACoV$IACoVMeanLvr))  
row.names(fold1) <- names  
colnames(fold1) <- c('CtrlMeanLvr','CoVMeanLvr','ILMeanLvr','IACoVMeanLvr')  
  
fold1$FC_CoV <- fold1$CoVMeanLvr/fold1$CtrlMeanLvr  
fold1$FC_IL <- fold1$ILMeanLvr/fold1$CtrlMeanLvr  
fold1$FC_IACoV <- fold1$IACoVMeanLvr/fold1$CtrlMeanLvr
```

Most expressed in liver samples by fold change of the Coronavirus, inactive CoronaVirus, and the IL-alpha treated Coronavirus as tables.

```
mostCoV <- fold1[order(fold1$FC_CoV, decreasing = TRUE)[0:100],]  
mostIL <- fold1[order(fold1$FC_IL, decreasing = TRUE)[0:100],]  
mostIACoV <- fold1[order(fold1$FC_IACoV, decreasing = TRUE)[0:100],]
```

Least expressed in liver samples by fold change of the Coronavirus, inactive CoronaVirus, and the IL-alpha treated Coronavirus as tables.

```
leastCoV <- fold1[order(fold1$FC_CoV, decreasing = FALSE)[0:100],]  
leastIL <- fold1[order(fold1$FC_IL, decreasing = FALSE)[0:100],]  
leastIACoV <- fold1[order(fold1$FC_IACoV, decreasing = FALSE)[0:100],]
```

Gene Expressions with most changes in the liver samples.

```
changes <- rbind(mostCoV,mostIL,mostIACoV,leastCoV,leastIL,leastIACoV)  
Changes <- changes[!duplicated(row.names(changes)),]  
length(unique(row.names(Changes)))  
## [1] 600
```

Get the magnitude of the fold change genes' row means.

```
Changes$MagnitudeFCs <- abs(rowMeans(Changes[,5:7]))
```

Combine this to the samples data for the liver tumor group.

```
Changes$Gene <- row.names(Changes)  
combined1 <- merge(both, Changes, by.x='GENE_SYMBOL', by.y='Gene')  
  
combined2 <- combined1[order(combined1$MagnitudeFCs, decreasing=TRUE),]  
  
CombinedLiver <- combined2[c(0:100,354:453),]
```

Machine Learning data for liver samples with 200 genes in the group of most gene expression changes.

```

names1 <- CombinedLiver$GENE_SYMBOL
names2 <- colnames(CombinedLiver)
row.names(CombinedLiver) <- names1

Combo_lvr_ML <- as.data.frame(t(CombinedLiver))

colnames(Combo_lvr_ML) <- gsub('-', '_', colnames(Combo_lvr_ML))
Combo1 <- Combo_lvr_ML[c(2:63),] #remove stats of fold change values and gene symbol row

```

Lets add a class field called Class\_Type to use machine learning on predicting class with these 200 genes and 62 mixed samples of capillary and liver tumor both inoculated with Coronavirus.

```

a <- rep('liver_CoV', 4)
b <- rep('liver_Ctrl', 4)
c <- rep('liver_CoV_IL', 2)
d <- rep('liver_IA_CoV', 2)
e <- rep('capillary_CoV_0hr', 5)
f <- rep('capillary_Ctrl_0hr', 5)
g <- rep('capillary_Cov_12hr', 5)
h <- rep('capillary_Ctrl_12hr', 5)
i <- rep('capillary_Cov_24hr', 5)
j <- rep('capillary_Ctrl_24hr', 5)
k <- rep('capillary_Cov_36hr', 5)
l <- rep('capillary_Ctrl_36hr', 5)
m <- rep('capillary_Cov_48hr', 5)
n <- rep('capillary_Ctrl_48hr', 5)

type <- as.data.frame(c(a,b,c,d,e,f,g,h,i,j,k,l,m,n))
colnames(type) <- 'Class_Type'
row.names(type) <- row.names(Combo1)

Combo2 <- cbind(type, Combo1)

```

Write this ML ready file to csv.

```
write.csv(Combo2, 'ML_ready_CoV_14_classes.csv', row.names=TRUE)
```

Make a separate ML ready file with a smaller set of classes to classify by liver or capillary and control or CoronaVirus

```

a <- rep('liver', 4)
b <- rep('liver', 4)
c <- rep('liver', 2)
d <- rep('liver', 2)
e <- rep('capillary', 5)
f <- rep('capillary', 5)
g <- rep('capillary', 5)
h <- rep('capillary', 5)
i <- rep('capillary', 5)

```

```

j <- rep('capillary',5)
k <- rep('capillary',5)
l <- rep('capillary',5)
m <- rep('capillary',5)
n <- rep('capillary',5)

type <- as.data.frame(c(a,b,c,d,e,f,g,h,i,j,k,l,m,n))
colnames(type) <- 'Class_Type'
row.names(type) <- row.names(Combo1)

Combo3 <- cbind(type,Combo1)

write.csv(Combo3, 'ML_ready_CoV_2_classes.csv', row.names=TRUE)

a <- rep('CoV', 4)
b <- rep('Ctrl',4)
c <- rep('CoV_IL',2)
d <- rep('IA_CoV',2)
e <- rep('CoV',5)
f <- rep('Ctrl',5)
g <- rep('Cov',5)
h <- rep('Ctrl',5)
i <- rep('Cov',5)
j <- rep('Ctrl',5)
k <- rep('Cov',5)
l <- rep('Ctrl',5)
m <- rep('Cov',5)
n <- rep('Ctrl',5)

type <- as.data.frame(c(a,b,c,d,e,f,g,h,i,j,k,l,m,n))
colnames(type) <- 'Class_Type'
row.names(type) <- row.names(Combo1)

Combo4 <- cbind(type,Combo1)

write.csv(Combo4, 'ML_ready_CoV_4_classes.csv', row.names=TRUE)

```

---

We didn't do any fold change or stat measures on the capillary samples, but we can plot them by using ggplot2 and group the sets by timed intervals for each group A through E and picking a handful of genes to compare over the 0,12,24,36, and 48 hour time intervals for the control group and the Coronavirus inoculated groups.

---

When the values are a ratio like this, it is easier to see the larger changes as in 9 compared to a low change like 0.0005, but this just means that compared to the control samples the inoculated Coronavirus had 9 times the gene expression values or had downregulated or suppressed gene expression values to 1/5000th the amount of the normal range of gene expression values respectively. \*\*\*

It makes sense to use some genes we already know have a higher magnitude of change, and we have a column for that in the CombinedLiver table called MagnitudeFCs that was already sorted from largest to smallest when made. We'll just select the first five of those genes to compare in these capillary samples over time.

```
mostChanged <- CombinedLiver[1:5,c(1,71)]
mostSuppressed <- CombinedLiver[196:200,c(1,71)]
row.names(mostChanged)

## [1] "NEURL3" "DUSP1" "ATF3" "PCLO" "LHB"

row.names(mostSuppressed)

## [1] "RASSF7" "LOC100335030" "C2orf78" "DEFB1" "ZNF610"

capillary <- merge(mostChanged, CombinedLiver, by.x='GENE_SYMBOL',
by.y='GENE_SYMBOL')
capillary1 <- merge(mostSuppressed, CombinedLiver, by.x='GENE_SYMBOL',
by.y='GENE_SYMBOL')
capillaries <- rbind(capillary, capillary1)
Capillaries <- capillaries[,c(1,15:64)]
row.names(Capillaries) <- Capillaries$GENE_SYMBOL

Capillaries2 <- as.data.frame(t(Capillaries))
Capillaries2 <- Capillaries2[-1,]
row.names(Capillaries2) <-
gsub('capillarySamples.', '', row.names(Capillaries2))
row.names(Capillaries2) <- gsub('GSM[0-9][0-9][0-9][0-9][0-9][0-9][0-9][0-9]',
'9_', '', row.names(Capillaries2))
row.names(Capillaries2) <- gsub('MERS_', '', row.names(Capillaries2))

CoV <- grep('CoV', row.names(Capillaries2))
ctrl <- grep('ctrl', row.names(Capillaries2))

Capillaries2$Class <- 'CoV or ctrl'

Capillaries2[CoV,11] <- 'Coronavirus'
Capillaries2[ctrl,11] <- 'control'

A <- grep('_A', row.names(Capillaries2))
B <- grep('_B', row.names(Capillaries2))
C <- grep('_C', row.names(Capillaries2))
D <- grep('_D', row.names(Capillaries2))
E <- grep('_E', row.names(Capillaries2))
```

```

Capillaries2$Group <- 'group'

Capillaries2[A,12] <- 'A'
Capillaries2[B,12] <- 'B'
Capillaries2[C,12] <- 'C'
Capillaries2[D,12] <- 'D'
Capillaries2[E,12] <- 'E'

hr0 <- grep('0hr', row.names(Capillaries2))
hr12 <- grep('12hr', row.names(Capillaries2))
hr24 <- grep('24hr', row.names(Capillaries2))
hr36 <- grep('36hr', row.names(Capillaries2))
hr48 <- grep('48hr', row.names(Capillaries2))

Capillaries2$TimeInterval <- 'time'

Capillaries2[hr0,13] <- '0 hr'
Capillaries2[hr12,13] <- '12 hr'
Capillaries2[hr24,13] <- '24 hr'
Capillaries2[hr36,13] <- '36 hr'
Capillaries2[hr48,13] <- '48 hr'

write.csv(Capillaries2, 'FC_10_capillaries_CoV.csv', row.names=TRUE)

```

---

The above table has 10 genes as the columns with the added Class (Coronavirus or control), Group (A,B,C,D,E), and TimeInterval (0,12,24,36,48 hours) fields to filter by and plot.

Lets make these group tables for the corona virus and see how they compare over time.

```

A_group <- filter(Capillaries2, Group=='A' & Class == 'Coronavirus')
B_group <- filter(Capillaries2, Group=='B' & Class == 'Coronavirus')
C_group <- filter(Capillaries2, Group=='C' & Class == 'Coronavirus')
D_group <- filter(Capillaries2, Group=='D' & Class == 'Coronavirus')
E_group <- filter(Capillaries2, Group=='E' & Class == 'Coronavirus')

```

We will do this for the A\_group table and ignore the Group and Class fields, because we made it only the A group of the Coronavirus class.

```

A_group2 <- A_group[,c(1,3,5,7,9,11:13)]
A_tidy <- gather(A_group2, 'Gene', 'GeneExpression', 1:5)

## Warning: attributes are not identical across measure variables;
## they will be dropped

A_tidy$GeneExpression <- round(as.numeric(A_tidy$GeneExpression),1)
A_tidy$TimeInterval <- as.factor(A_tidy$TimeInterval)

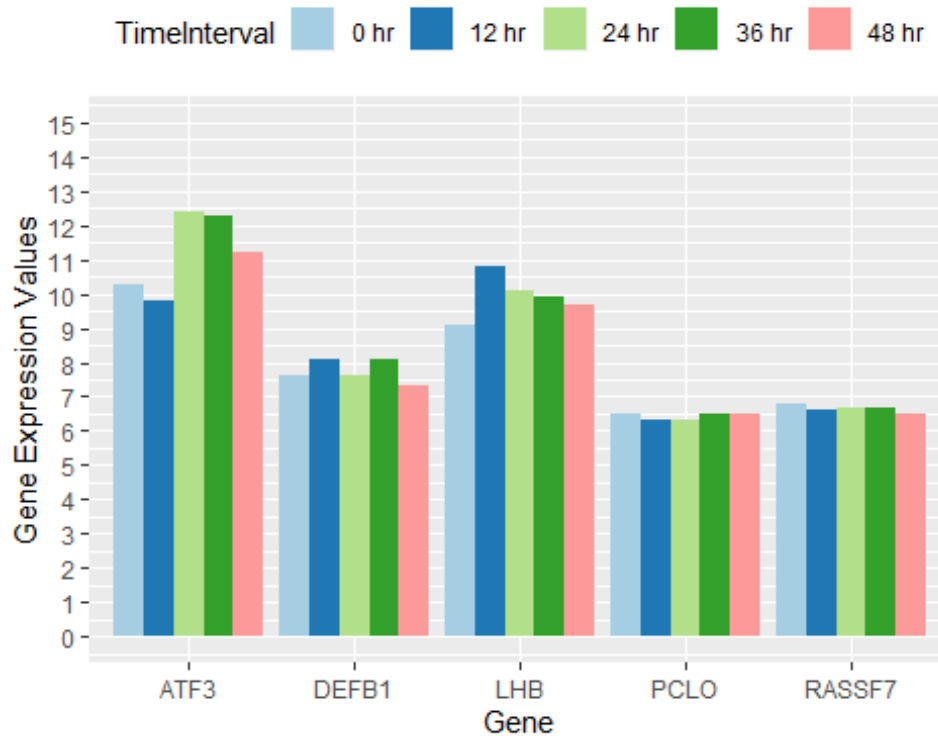
```



```
A_tidy$Gene <- as.factor(A_tidy$Gene)
A_tidy
```

##		Class	Group	TimeInterval	Gene	GeneExpression
## 1	Coronavirus	A		0 hr	ATF3	10.3
## 2	Coronavirus	A		12 hr	ATF3	9.8
## 3	Coronavirus	A		24 hr	ATF3	12.4
## 4	Coronavirus	A		36 hr	ATF3	12.3
## 5	Coronavirus	A		48 hr	ATF3	11.2
## 6	Coronavirus	A		0 hr	LHB	9.1
## 7	Coronavirus	A		12 hr	LHB	10.8
## 8	Coronavirus	A		24 hr	LHB	10.1
## 9	Coronavirus	A		36 hr	LHB	9.9
## 10	Coronavirus	A		48 hr	LHB	9.7
## 11	Coronavirus	A		0 hr	PCLO	6.5
## 12	Coronavirus	A		12 hr	PCLO	6.3
## 13	Coronavirus	A		24 hr	PCLO	6.3
## 14	Coronavirus	A		36 hr	PCLO	6.5
## 15	Coronavirus	A		48 hr	PCLO	6.5
## 16	Coronavirus	A		0 hr	DEFB1	7.6
## 17	Coronavirus	A		12 hr	DEFB1	8.1
## 18	Coronavirus	A		24 hr	DEFB1	7.6
## 19	Coronavirus	A		36 hr	DEFB1	8.1
## 20	Coronavirus	A		48 hr	DEFB1	7.3
## 21	Coronavirus	A		0 hr	RASSF7	6.8
## 22	Coronavirus	A		12 hr	RASSF7	6.6
## 23	Coronavirus	A		24 hr	RASSF7	6.7
## 24	Coronavirus	A		36 hr	RASSF7	6.7
## 25	Coronavirus	A		48 hr	RASSF7	6.5

```
ggplot(data = A_tidy, aes(x=Gene, y=GeneExpression, fill=TimeInterval)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous(breaks = seq(0, 15, by=1), limits=c(0,15))+
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="top")+
  #ggtitle('Group A with Coronavirus for Selected Genes Part 1')+
  xlab('Gene')+
  ylab('Gene Expression Values')
```



The genes above for Part 1 of the group A samples of coronavirus in blood capillaries show some variation in gene expression values for some of these genes that had the most change in the liver tumor samples. Starting at the initial hour up to 48 hours after being inoculated in vitro, there is an increase then decrease for ATF3 and LHB genes, while a decrease then increase close to initial value with PCLO and slightly with RASSF7. For DEFB1, it has a cyclical increase, decrease, increase, then decrease to stabilize closer to the initial gene expressio value.

Now lets find the other five genes in the group A set of ten genes found to have the most change in the liver tumor samples, and examined here in the blood capillary samples.

```
A_group3 <- A_group[,c(2,4,6,8,10,11:13)]
A_tidy1 <- gather(A_group3, 'Gene', 'GeneExpression', 1:5)

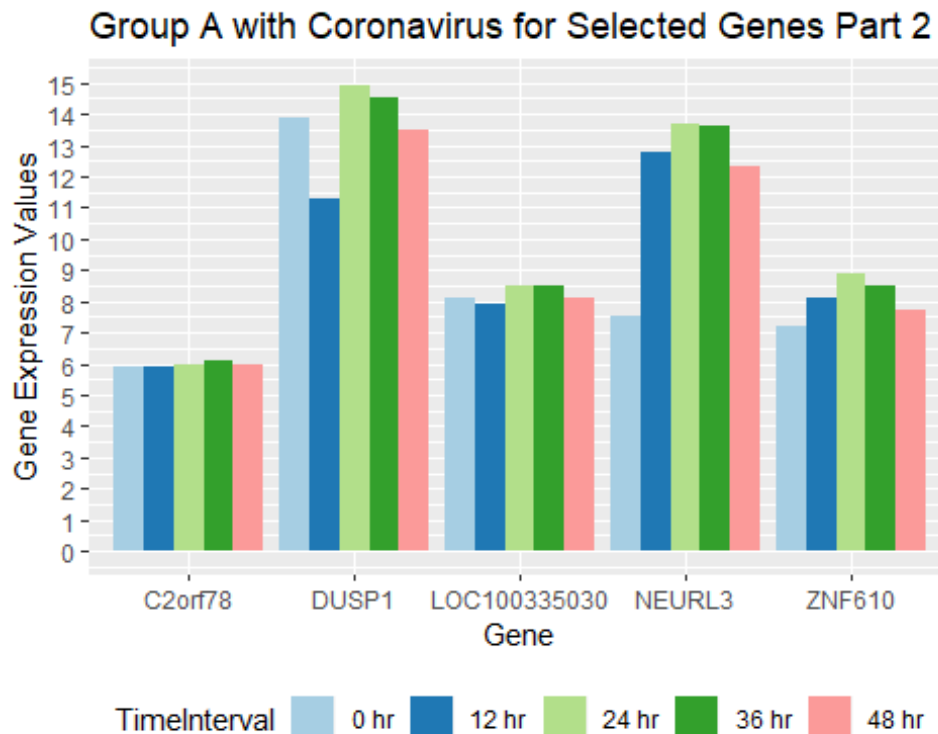
## Warning: attributes are not identical across measure variables;
## they will be dropped

A_tidy1$GeneExpression <- round(as.numeric(A_tidy1$GeneExpression),1)
A_tidy1$TimeInterval <- as.factor(A_tidy1$TimeInterval)
A_tidy1$Gene <- as.factor(A_tidy1$Gene)
```

---

```
ggplot(data = A_tidy1, aes(x=Gene, y=GeneExpression, fill=TimeInterval)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous(breaks = seq(0, 15, by=1), limits=c(0,15))+
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="bottom")+
  ggtitle('Group A with Coronavirus for Selected Genes Part 2')+
  xlab('Gene')+
  ylab('Gene Expression Values')
```

---



The above genes in part 2 of the group A Coronavirus samples over 48 hours, shows that the gene expression values increase up to 24 hours then decrease to 48 hours for most of the genes above.

---

Lets look back at the platforms and the features removed. The sequence field is an interesting field because it can show copy number variants of genes by the genes that are duplicates at other probes from the samples.

---

```
Platform13497 <- read.csv('GPL13497-9755-forSequenceFeature-GSE100509.csv',
sep=',',
```

---

```

na.strings=c('', ' '), header=TRUE)
Platform16699 <- read.csv('GPL16699-forSequenceFeatureGSE89166_GSE89160.csv',
sep=',',
na.strings=c('', ' '), header=TRUE)

```

The features in Platform13497 and the first five listed ID values for this platform:

```

colnames(Platform13497)

## [1] "ID" "SPOT_ID" "CONTROL_TYPE"
## [4] "REFSEQ" "GB_ACC" "GENE"
## [7] "GENE_SYMBOL" "GENE_NAME" "UNIGENE_ID"
## [10] "ENSEMBL_ID" "TIGR_ID" "ACCESSION_STRING"
## [13] "CHROMOSOMAL_LOCATION" "CYTOBAND" "DESCRIPTION"
## [16] "GO_ID" "SEQUENCE"

```

The features in Platform16699 and the first five listed ID values of that platform:

```

colnames(Platform16699)

## [1] "ID" "COL" "ROW"
## [4] "NAME" "SPOT_ID" "CONTROL_TYPE"
## [7] "REFSEQ" "GB_ACC" "LOCUSLINK_ID"
## [10] "GENE_SYMBOL" "GENE_NAME" "UNIGENE_ID"
## [13] "ENSEMBL_ID" "ACCESSION_STRING" "CHROMOSOMAL_LOCATION"
## [16] "CYTOBAND" "DESCRIPTION" "GO_ID"
## [19] "SEQUENCE"

```

Lets keep only the ID (Platform16699) or SPOT\_ID (Platform13497), GENE\_SYMBOL, DESCRIPTION, and SEQUENCE features of both platforms.

```

P16699 <- Platform16699[,c(1,10,17,19)]
P13497 <- Platform13497[,c(2,7,15,17)]

```

Lets also remove the incomplete cases in both platforms.

```

work16699 <- P16699[complete.cases(P16699),]
work13497 <- P13497[complete.cases(P13497),]

```

Now merge these data sets to their corresponding samples by SPOT\_ID. First read in the samples data for each platform and series.

```

GSE89166_89160 <- read.csv('GSE89166_GSE89160.csv', sep=',', na.strings=c('', ' '),
header=TRUE)
GSE100509 <- read.csv('GSE100509.csv', sep=',', header=TRUE,
na.strings=c('', ' '))

```

Now merge the series data sets to their respective platforms of gene informational meta features.

```

Series100509 <- merge(work13497, GSE100509, by.x='SPOT_ID', by.y='ID_REF')
Series89166_89160 <- merge(work16699, GSE89166_89160, by.x='ID', by.y='ID_REF')

```

Rename the columns of the Series100509 to the 5 groups for each of CoV and Ctrl over 0,12,24,36, and 48 hours.

```
colnames(Series100509)
```

```
## [1] "SPOT_ID" "GENE_SYMBOL"
## [3] "DESCRIPTION" "SEQUENCE"
## [5] "GSM2685693_MERS_CoV_0hr" "GSM2685694_MERS_CoV_0hr"
## [7] "GSM2685695_MERS_CoV_0hr" "GSM2685696_MERS_CoV_0hr"
## [9] "GSM2685697_MERS_CoV_0hr" "GSM2685698_ctrl_0hr"
## [11] "GSM2685699_ctrl_0hr" "GSM2685700_ctrl_0hr"
## [13] "GSM2685701_ctrl_0hr" "GSM2685702_ctrl_0hr"
## [15] "GSM2685703_MERS_CoV_12hr" "GSM2685704_MERS_CoV_12hr"
## [17] "GSM2685705_MERS_CoV_12hr" "GSM2685706_MERS_CoV_12hr"
## [19] "GSM2685707_MERS_CoV_12hr" "GSM2685708_ctrl_12hr"
## [21] "GSM2685709_ctrl_12hr" "GSM2685710_ctrl_12hr"
## [23] "GSM2685711_ctrl_12hr" "GSM2685712_ctrl_12hr"
## [25] "GSM2685713_MERS_CoV_24hr" "GSM2685714_MERS_CoV_24hr"
## [27] "GSM2685715_MERS_CoV_24hr" "GSM2685716_MERS_CoV_24hr"
## [29] "GSM2685717_MERS_CoV_24hr" "GSM2685718_ctrl_24hr"
## [31] "GSM2685719_ctrl_24hr" "GSM2685720_ctrl_24hr"
## [33] "GSM2685721_ctrl_24hr" "GSM2685722_ctrl_24hr"
## [35] "GSM2685723_MERS_CoV_36hr" "GSM2685724_MERS_CoV_36hr"
## [37] "GSM2685725_MERS_CoV_36hr" "GSM2685726_MERS_CoV_36hr"
## [39] "GSM2685727_MERS_CoV_36hr" "GSM2685728_ctrl_36hr"
## [41] "GSM2685729_ctrl_36hr" "GSM2685730_ctrl_36hr"
## [43] "GSM2685731_ctrl_36hr" "GSM2685732_ctrl_36hr"
## [45] "GSM2685733_MERS_CoV_48hr" "GSM2685734_MERS_CoV_48hr"
## [47] "GSM2685735_MERS_CoV_48hr" "GSM2685736_MERS_CoV_48hr"
## [49] "GSM2685737_MERS_CoV_48hr" "GSM2685738_ctrl_48hr"
## [51] "GSM2685739_ctrl_48hr" "GSM2685740_ctrl_48hr"
## [53] "GSM2685741_ctrl_48hr" "GSM2685742_ctrl_48hr"
```

```
group <- rep(1:5,10)
```

```
Group <- gsub('1','Group_A',group)
```

```
Group <- gsub('2','Group_B',Group)
```

```
Group <- gsub('3','Group_C',Group)
```

```
Group <- gsub('4','Group_D',Group)
```

```
Group <- gsub('5','Group_E',Group)
```

```
names <- colnames(Series100509)[5:54]
```

```
Names <- paste(names,Group,sep='_')
```

```
newNames <- gsub('_MERS','', Names)
```

```
newNames
```

```
## [1] "GSM2685693_CoV_0hr_Group_A" "GSM2685694_CoV_0hr_Group_B"
## [3] "GSM2685695_CoV_0hr_Group_C" "GSM2685696_CoV_0hr_Group_D"
## [5] "GSM2685697_CoV_0hr_Group_E" "GSM2685698_ctrl_0hr_Group_A"
```

```
## [7] "GSM2685699_ctrl_0hr_Group_B" "GSM2685700_ctrl_0hr_Group_C"
## [9] "GSM2685701_ctrl_0hr_Group_D" "GSM2685702_ctrl_0hr_Group_E"
## [11] "GSM2685703_CoV_12hr_Group_A" "GSM2685704_CoV_12hr_Group_B"
## [13] "GSM2685705_CoV_12hr_Group_C" "GSM2685706_CoV_12hr_Group_D"
## [15] "GSM2685707_CoV_12hr_Group_E" "GSM2685708_ctrl_12hr_Group_A"
## [17] "GSM2685709_ctrl_12hr_Group_B" "GSM2685710_ctrl_12hr_Group_C"
## [19] "GSM2685711_ctrl_12hr_Group_D" "GSM2685712_ctrl_12hr_Group_E"
## [21] "GSM2685713_CoV_24hr_Group_A" "GSM2685714_CoV_24hr_Group_B"
## [23] "GSM2685715_CoV_24hr_Group_C" "GSM2685716_CoV_24hr_Group_D"
## [25] "GSM2685717_CoV_24hr_Group_E" "GSM2685718_ctrl_24hr_Group_A"
## [27] "GSM2685719_ctrl_24hr_Group_B" "GSM2685720_ctrl_24hr_Group_C"
## [29] "GSM2685721_ctrl_24hr_Group_D" "GSM2685722_ctrl_24hr_Group_E"
## [31] "GSM2685723_CoV_36hr_Group_A" "GSM2685724_CoV_36hr_Group_B"
## [33] "GSM2685725_CoV_36hr_Group_C" "GSM2685726_CoV_36hr_Group_D"
## [35] "GSM2685727_CoV_36hr_Group_E" "GSM2685728_ctrl_36hr_Group_A"
## [37] "GSM2685729_ctrl_36hr_Group_B" "GSM2685730_ctrl_36hr_Group_C"
## [39] "GSM2685731_ctrl_36hr_Group_D" "GSM2685732_ctrl_36hr_Group_E"
## [41] "GSM2685733_CoV_48hr_Group_A" "GSM2685734_CoV_48hr_Group_B"
## [43] "GSM2685735_CoV_48hr_Group_C" "GSM2685736_CoV_48hr_Group_D"
## [45] "GSM2685737_CoV_48hr_Group_E" "GSM2685738_ctrl_48hr_Group_A"
## [47] "GSM2685739_ctrl_48hr_Group_B" "GSM2685740_ctrl_48hr_Group_C"
## [49] "GSM2685741_ctrl_48hr_Group_D" "GSM2685742_ctrl_48hr_Group_E"
```

Change the column names in Series100509 to the new column names identifying which group the samples is from in A:E.

```
colnames(Series100509)[5:54] <- newNames
colnames(Series100509)
```

```
## [1] "SPOT_ID" "GENE_SYMBOL"
## [3] "DESCRIPTION" "SEQUENCE"
## [5] "GSM2685693_CoV_0hr_Group_A" "GSM2685694_CoV_0hr_Group_B"
## [7] "GSM2685695_CoV_0hr_Group_C" "GSM2685696_CoV_0hr_Group_D"
## [9] "GSM2685697_CoV_0hr_Group_E" "GSM2685698_ctrl_0hr_Group_A"
## [11] "GSM2685699_ctrl_0hr_Group_B" "GSM2685700_ctrl_0hr_Group_C"
## [13] "GSM2685701_ctrl_0hr_Group_D" "GSM2685702_ctrl_0hr_Group_E"
## [15] "GSM2685703_CoV_12hr_Group_A" "GSM2685704_CoV_12hr_Group_B"
## [17] "GSM2685705_CoV_12hr_Group_C" "GSM2685706_CoV_12hr_Group_D"
## [19] "GSM2685707_CoV_12hr_Group_E" "GSM2685708_ctrl_12hr_Group_A"
## [21] "GSM2685709_ctrl_12hr_Group_B" "GSM2685710_ctrl_12hr_Group_C"
## [23] "GSM2685711_ctrl_12hr_Group_D" "GSM2685712_ctrl_12hr_Group_E"
## [25] "GSM2685713_CoV_24hr_Group_A" "GSM2685714_CoV_24hr_Group_B"
## [27] "GSM2685715_CoV_24hr_Group_C" "GSM2685716_CoV_24hr_Group_D"
## [29] "GSM2685717_CoV_24hr_Group_E" "GSM2685718_ctrl_24hr_Group_A"
## [31] "GSM2685719_ctrl_24hr_Group_B" "GSM2685720_ctrl_24hr_Group_C"
## [33] "GSM2685721_ctrl_24hr_Group_D" "GSM2685722_ctrl_24hr_Group_E"
## [35] "GSM2685723_CoV_36hr_Group_A" "GSM2685724_CoV_36hr_Group_B"
## [37] "GSM2685725_CoV_36hr_Group_C" "GSM2685726_CoV_36hr_Group_D"
## [39] "GSM2685727_CoV_36hr_Group_E" "GSM2685728_ctrl_36hr_Group_A"
## [41] "GSM2685729_ctrl_36hr_Group_B" "GSM2685730_ctrl_36hr_Group_C"
```

```
## [43] "GSM2685731_ctrl_36hr_Group_D" "GSM2685732_ctrl_36hr_Group_E"
## [45] "GSM2685733_CoV_48hr_Group_A"  "GSM2685734_CoV_48hr_Group_B"
## [47] "GSM2685735_CoV_48hr_Group_C"  "GSM2685736_CoV_48hr_Group_D"
## [49] "GSM2685737_CoV_48hr_Group_E"  "GSM2685738_ctrl_48hr_Group_A"
## [51] "GSM2685739_ctrl_48hr_Group_B" "GSM2685740_ctrl_48hr_Group_C"
## [53] "GSM2685741_ctrl_48hr_Group_D" "GSM2685742_ctrl_48hr_Group_E"
```

Remove the ID and SPOT\_ID fields of the probe labels that won't be needed for the analysis.

```
Series89 <- Series89166_89160[, -1]
Series100 <- Series100509[, -c(1,3,4)]
```

Now combine the series together by genes in common.

```
ComboLiverCapillarySequences <- merge(Series89, Series100, by.x='GENE_SYMBOL',
                                       by.y='GENE_SYMBOL')
```

There should be different genotypes or copy number variations in the SEQUENCE feature column, which will identify which nucleotide jumps, is deleted, rearranged in the gene expressions. This time around let's group by SEQUENCE to see how many genotypes there are in all the genes. This could give more information in the analysis of any genotypes of genes could be more susceptible to pathogenesis of CoV or immunity to it. \*\*\*

This next portion of this analysis shows the top five genes in the data, the number of genotypes or copy number variations of the nucleotide sequences are in each gene, and the gene name. This could be useful to determine how well the genotypes within these five genes that were expressed more than the other 65k genes were when analyzing the effects of CoV, inactive CoV, CoV treated with an interleukin, and the control samples.

```
SeqGroup <- ComboLiverCapillarySequences %>% group_by(GENE_SYMBOL) %>%
count(n=n())
SeqGroup <- SeqGroup[order(SeqGroup$n, decreasing=TRUE), -3]

SeqGroup5 <- SeqGroup[1:5,]

genes5sequences <- merge(SeqGroup5, ComboLiverCapillarySequences,
                        by.x='GENE_SYMBOL', by.y='GENE_SYMBOL')

genotypes5 <- genes5sequences %>% group_by(SEQUENCE) %>% count(n=n)

genotypes5$n <- as.factor(genotypes5$n)
SeqGroup5$n <- as.factor(SeqGroup5$n)

genotypes5_1 <- merge(genotypes5, SeqGroup5, by.x='n', by.y='n')
colnames(genotypes5_1)[c(1,3)] <- c('geneCount', 'genotypeCount')
head(genotypes5_1)
```

```
##      geneCount                                     SEQUENCE
## 1      255 AAACCTTACTCCAGAGCTCCTTGTGCATCTGACCAGCACCATCGACAGAATAAACACAGAA
## 2      255 AACAGAGTCCTCAGGGAAGAAAATCGAAGACTTCAGGCTCAACTGAGTCATGTTTCCAGA
## 3      255 GCACCTGTGTTCTTTGAGTTACATCATGAATGTGGTGATTTCAGATACCATCTCAGG
## 4      255 ATGGGGTGCTCTGGGGAAAATATTGGAGGGTCATCCATTCCACATTAAGAGCAAGTTGT
## 5      255 AAGTGCTTGGAATACTTGGGTGAATGTTACCAGACTCCTTCTCTCTCAGCTTACAGCCT
## 6      255 AATCTACGAGGCACTTTATGGCAATTCCAAGAAGGGGCTGAAAGGTATGTGTTCTTCTCC
##      genotypeCount GENE_SYMBOL
## 1      15      PDE4DIP
## 2      15      PDE4DIP
## 3      15      PDE4DIP
## 4      15      PDE4DIP
## 5      15      PDE4DIP
## 6      15      PDE4DIP
```

Write the files above out to csv.

```
write.csv(ComboLiverCapillarySequences,
'SequencesBothCleaned.csv',row.names=FALSE)
```

Write this large file into three subsets of three tiers because it is more than 57mb in size, and I want to load it into github. They would be read in separately, then they would be row binded in order of tier1, tier2, and then tier 3: - **ComboLiverCapillarySequences <- rbind(SequencesBothCleaned\_tier1,SequencesBothCleaned\_tier2,SequencesBothCleaned\_tier3)**

```
write.csv(ComboLiverCapillarySequences[1:21000,],
'SequencesBothCleaned_tier1.csv',row.names=FALSE)
write.csv(ComboLiverCapillarySequences[21001:42000,],
'SequencesBothCleaned_tier2.csv',row.names=FALSE)
write.csv(ComboLiverCapillarySequences[42001:65505,],
'SequencesBothCleaned_tier3.csv',row.names=FALSE)

write.csv(genotypes5_1, 'Genotypes_5_1.csv', row.names=FALSE)
```

```
ComboCNV <- ComboLiverCapillarySequences[,-c(1,2)]
copynumbers <- merge(genotypes5_1,ComboCNV,
                     by.x='SEQUENCE', by.y='SEQUENCE')
CNV <- copynumbers[!duplicated(copynumbers$SEQUENCE),]
write.csv(CNV, 'copyNumbers.csv', row.names=FALSE)
```

```
colnames(CNV)
```

```
## [1] "SEQUENCE" "geneCount"
## [3] "genotypeCount" "GENE_SYMBOL"
## [5] "GSM2359851_CoV1" "GSM2359853_CoV2"
## [7] "GSM2359910_CoV3" "GSM2359913_CoV4"
## [9] "GSM2359850_ctrl1" "GSM2359852_ctrl2"
## [11] "GSM2359911_ctrl3" "GSM2359914_ctrl4"
## [13] "GSM2359912_IL1" "GSM2359917_IL2"
## [15] "GSM2359915_inactiveHeatCoV1" "GSM2359916_inactiveHeatCoV2"
```



```
## [17] "GSM2685693_CoV_0hr_Group_A" "GSM2685694_CoV_0hr_Group_B"
## [19] "GSM2685695_CoV_0hr_Group_C" "GSM2685696_CoV_0hr_Group_D"
## [21] "GSM2685697_CoV_0hr_Group_E" "GSM2685698_ctrl_0hr_Group_A"
## [23] "GSM2685699_ctrl_0hr_Group_B" "GSM2685700_ctrl_0hr_Group_C"
## [25] "GSM2685701_ctrl_0hr_Group_D" "GSM2685702_ctrl_0hr_Group_E"
## [27] "GSM2685703_CoV_12hr_Group_A" "GSM2685704_CoV_12hr_Group_B"
## [29] "GSM2685705_CoV_12hr_Group_C" "GSM2685706_CoV_12hr_Group_D"
## [31] "GSM2685707_CoV_12hr_Group_E" "GSM2685708_ctrl_12hr_Group_A"
## [33] "GSM2685709_ctrl_12hr_Group_B" "GSM2685710_ctrl_12hr_Group_C"
## [35] "GSM2685711_ctrl_12hr_Group_D" "GSM2685712_ctrl_12hr_Group_E"
## [37] "GSM2685713_CoV_24hr_Group_A" "GSM2685714_CoV_24hr_Group_B"
## [39] "GSM2685715_CoV_24hr_Group_C" "GSM2685716_CoV_24hr_Group_D"
## [41] "GSM2685717_CoV_24hr_Group_E" "GSM2685718_ctrl_24hr_Group_A"
## [43] "GSM2685719_ctrl_24hr_Group_B" "GSM2685720_ctrl_24hr_Group_C"
## [45] "GSM2685721_ctrl_24hr_Group_D" "GSM2685722_ctrl_24hr_Group_E"
## [47] "GSM2685723_CoV_36hr_Group_A" "GSM2685724_CoV_36hr_Group_B"
## [49] "GSM2685725_CoV_36hr_Group_C" "GSM2685726_CoV_36hr_Group_D"
## [51] "GSM2685727_CoV_36hr_Group_E" "GSM2685728_ctrl_36hr_Group_A"
## [53] "GSM2685729_ctrl_36hr_Group_B" "GSM2685730_ctrl_36hr_Group_C"
## [55] "GSM2685731_ctrl_36hr_Group_D" "GSM2685732_ctrl_36hr_Group_E"
## [57] "GSM2685733_CoV_48hr_Group_A" "GSM2685734_CoV_48hr_Group_B"
## [59] "GSM2685735_CoV_48hr_Group_C" "GSM2685736_CoV_48hr_Group_D"
## [61] "GSM2685737_CoV_48hr_Group_E" "GSM2685738_ctrl_48hr_Group_A"
## [63] "GSM2685739_ctrl_48hr_Group_B" "GSM2685740_ctrl_48hr_Group_C"
## [65] "GSM2685741_ctrl_48hr_Group_D" "GSM2685742_ctrl_48hr_Group_E"
```

This feature, SEQUENCE, provides the single nucleotide polymorphisms (SNP)s or copy number variants of each gene's DNA, many are duplicated. Here are the first few SNPs.

```
head(CNV$SEQUENCE)
```

```
## [1] AAACCTTACTCCAGAGCTCCTTGTGCATCTGACCAGCACCATCGACAGAATAAACACAGAA
## [2] AACAGAGTCCTCAGGGAAGAAAATCGAAGACTTCAGGCTCAACTGAGTCATGTTTCCAGA
## [3] AAGGATTTGCTTATAAGGGTTCCTGCTTTACAAAATTATTCCAGGTTTTATGTGTCAGG
## [4] AAGGATTTGTTTGTAAAGGGCTCCCGCTTTACAGAAATTATTCCAGGGTTTATGTGTCAGG
## [5] AAGTGCTTGAAATACTTGGGTGAATGTTACCAGACTCCTTCTCTCTCAGCTTACAGCCT
## [6] AATCTACGAGGCACTTTATGGCAATTCCAAGAAGGGGCTGAAAGGTATGTGTTCTTCTCC
## 50571 Levels: AAACAAAAAACAGGTTAAGAAAATTACTTGGGTGGGCAGACTTAGGAACGCTCTACTCGG
...

```

---

It would be interesting in the fold change values between genotypes of the genes expressed in comparing the capillary samples all within 1 hour after being inoculated with CoV, inactive CoV, CoV and an interleukin alpha, or control group. Then compare the liver tumor samples of each group A through E that was monitored after being inoculated with CoV at 0,12,24,36, and 48 hours side by side with the control groups of groups A through E.

We could also detect patterns to analyze those sequences of copy number variations within the top 5 genes expressed the most number of times in this data. Comparing networks of genes associated with processes in the body like immune response, pathogenesis of disease onset, networks of human processes in the body associated with cancer or subsequent diseases like autoimmune, celiac disease, hemochromatosis, anemia, etc. To see how well any of those genotypes fair.

We have the five genes that had the highest copy number variations within them and merged it with our liver tumor and capillary CoV and control samples in the CNV data table.

I joined [genecards.org](http://genecards.org) and found that this site has some useful information on gene networks for analyzing genes that play a role in a network of genes that function at some level on the human anatomy. Such as anemia, cannabidiol, celiac disease, diabetes type 1 and 2, hemochromatosis, immunity, pain, and tumorigenesis. I downloaded the csv files for each of these networks just mentioned. There is a ranking of each gene as importance the higher the ranked order in the network functional role on the human body.

We can import each of those now.

[illegible]

---

We could then take our combined data from the liver and capillary datasets to look at the copy number variants of genotypes within these networks separately to understand how or if these genes have a high number of copy number in control versus diseased and diseased/treated states.

Lets take the top five genes from each network, add a column feature to label the gene network, then combine those tables. They are already ordered each by Relevance.score.

```
anemia5 <- anemiaNetwork[1:5,]
cannabidiol5 <- cannabidiolNetwork[1:5,]
diabetesOne5 <- diabetesType1Network[1:5,]
diabetesTwo5 <- diabetesType2Network[1:5,]
hemochromatosis5 <- hemochromatosisNetwork[1:5,]
immunity5 <- immunityNetwork[1:5,]
pain5 <- painNetwork[1:5,]
tumorigenesis5 <- tumorigenesisNetwork[1:5,]
```

Lets merge each network disease pathway with the ComboLiverCapillarySequences data table.

```
anemia5_path <- merge(anemia5, ComboLiverCapillarySequences,
  by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
cannabidiol5_path <- merge(cannabidiol5, ComboLiverCapillarySequences,
  by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
diabetesOne5_path <- merge(diabetesOne5, ComboLiverCapillarySequences,
  by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
diabetesTwo5_path <- merge(diabetesTwo5, ComboLiverCapillarySequences,
  by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
hemochromatosis5_path <- merge(hemochromatosis5,
  ComboLiverCapillarySequences, by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
immunity5_path <- merge(immunity5, ComboLiverCapillarySequences,
  by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
pain5_path <- merge(pain5, ComboLiverCapillarySequences, by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
tumorigenesis5_path <- merge(tumorigenesis5, ComboLiverCapillarySequences,
  by.x='Gene.Symbol',
  by.y='GENE_SYMBOL')
```

Combine the above data tables together.

```
Network <- as.data.frame(c(rep('anemia',length(anemia5_path$Gene.Symbol)),
  rep('cannabidiol',length(cannabidiol5_path$Gene.Symbol)),
```

```

        rep('diabetes
1',length(diabetesOne5_path$Gene.Symbol)),
        rep('diabetes 2',
length(diabetesTwo5_path$Gene.Symbol)),
        rep('hemochromatosis',
length(hemochromatosis5_path$Gene.Symbol)),
        rep('immunity', length(immunity5_path$Gene.Symbol)),
        rep('pain', length(pain5_path$Gene.Symbol)),

rep('tumorigenesis',length(tumorigenesis5_path$Gene.Symbol))))
colnames(Network) <- 'geneNetwork'

networkPathwayGenes <-
rbind(anemia5_path,cannabidiol5_path,diabetesOne5_path,
diabetesTwo5_path,hemochromatosis5_path,immunity5_path,
      pain5_path,tumorigenesis5_path)
PathwayGenes <- cbind(Network,networkPathwayGenes)

```

Now group by gene for the count and then by sequence for the copy number variant count within the gene.

```

pathGenesCount <- PathwayGenes %>% group_by(Gene.Symbol) %>% count(n())
pathGenotypeCount <- PathwayGenes %>% group_by(SEQUENCE) %>% count(n())

pathGenes <- pathGenesCount[, -2]
colnames(pathGenes)[2] <- 'GeneCount'
pathGenos <- pathGenotypeCount[, -2]
colnames(pathGenos)[2] <- 'GenotypeCount'

pG <- merge(pathGenos, PathwayGenes, by.x='SEQUENCE', by.y='SEQUENCE')
pG1 <- merge(pathGenes, pG, by.x='Gene.Symbol', by.y='Gene.Symbol')
colnames(pG1)[c(13,14,17,18)] <-
paste('Study1',colnames(pG1)[c(13,14,17,18)] ,
      sep='_')
write.csv(pG1, 'genes-genotypes-networks.csv', row.names=FALSE)

```

I want to remove and hold it separate for one of the liver tumor samples studies that uses CoV1, CoV2, ctrl1, and ctrl2, because the values are not scaled the same and these numbers will ruin the analysis included as is. They were already log scaled but seem to be much larger than the other values after scaling. The previous block of code added 'Study1' to those larger scaled samples to extract for separate analysis.

```

scaleSet <- pG1[,c(1:14,17,18)]
otherSet <- pG1[,c(1:12,15,16,19:74)]

```

---

Lets add in that the scaleSet columns are from study1 and leave the study2 column names in the otherSet as is. Combine and make a separate data set. I have previously examined cannabidiol in sebum/acne and brain tumor samples to briefly determine the CBD effects

on hormone and immune response genes. I added this cannabidiol network to get the top five CBD genes by relative score in the CBD network. We can look at this scale set of CoV and ctrl samples from liver tumors to see if there are any obvious effects to this network that has shown to treat chronic pain in some folks.

Lets get the data set of cannabidiol, CBD, to compare CoV to mock or control liver tumor samples from the larger scaled data set.

```
CBD <- subset(scaleSet, scaleSet$geneNetwork == 'cannabidiol')
orderCBD <- CBD[order(CBD$Gene.Symbol, decreasing=TRUE),]
```

We can see how many geno types are unique to a gene by adding up the genotypes to equal the genes in count. For instance, the PPARG gene has 22 counts of that gene in the data with one genotype having 22 counts, and the other genotype for that gene having only 2 counts. So that there are two unique genotypes for the gene PPARG. Lets go ahead and use dplyr to group by SEQUENCE then take the mean of that sequence but for the CoV and ctrl groups separately.

The two separate CoV and Ctrl datasets for the five CBD genes with the highest count of occurence in the data.

```
CBD_CoV <- CBD[, -c(15,16)]

CBD_ctrl <- CBD[, -c(13,14)]

samples <- as.vector(colnames(CBD_CoV)[13:14])
samples1 <- as.vector(colnames(CBD_ctrl)[13:14])

CBD_CovMeans <- CBD_CoV %>% group_by(SEQUENCE) %>%
  summarise_at(vars(samples),
                                     mean)

CBD_CtrlMeans <- CBD_ctrl %>% group_by(SEQUENCE) %>%
  summarise_at(vars(samples1),
                                     mean)
```

It seems a small data set, but with this network, CBD network, there was only one gene with more than one genotype. This is why there are only six genotypes when we selected five genes.

```
unique(CBD$SEQUENCE)

## [1] TGTACTAGGCCTACTGGGGATCAGAGTTCCCAAGAAAGGAAACCTTTTCTTGTATCTGGA
## [2] AGCATCTGCTCCCTCTACCAGCTGGAGAATACTGCAACTAGACGCAGCCCGCAGGCAGC
## [3] CCAAGGCTTCATGACAAGGGAGTTTCTAAAGAGCCTGCGAAAGCCTTTTGGTGACTTTAT
## [4] ACAATCAGATTGAAGCTTATCTATGACAGATGTGATCTTAAGTGTGCGATCCACAAAAA
## [5] TGAGATATTTAAGGTTGAATGTTTGTCTTAGGATAGGCCTATGTGCTAGCCACAAAGA
## [6] GGGGTATCTGGGGGACCCAATGTAGGAGCTGCCTTGCTCAGACATGTTTTCCGTGAAA
## 50571 Levels: AAACAAAAACAGGTTAAGAAAATTACTTGGGTGGGCAGACTTAGGAACGCTCTACTCGG
...
```

Lets combine the data means for each group by genotype or unique sequence with the genes they belong to from two features of the CBD data set, Gene.Symbol and SEQUENCE.

```
geneSeq <- CBD[,c(1:5)]
geneSeq1 <- merge(geneSeq, CBD_CovMeans, by.x='SEQUENCE', by.y='SEQUENCE')
geneSeq2 <- merge(geneSeq1, CBD_CtrlMeans, by.x='SEQUENCE', by.y='SEQUENCE')

geneSeq3 <- geneSeq2[!duplicated(geneSeq2),]
geneSeq3
```

##	Gene.Symbol	SEQUENCE
## 1	ACAATCAGATTGAAGCTTATCTATGACAGATGTGATCTTAAGTGTGCGGATCCACAAAAA	PPARG
## 3	AGCATCTGCTCCCTCTACCAGCTGGAGAACTACTGCAACTAGACGCAGCCCGCAGGCAGC	INS
## 4	CCAAGGCTTCATGACAAGGGAGTTTCTAAAGAGCCTGCGAAAGCCTTTTGGTGACTTTAT	PPARG
## 24	GGGGTATCCTGGGGGACCCAATGTAGGAGCTGCCTTGGCTCAGACATGTTTTCCGTGAAA	TNF
## 25	TGAGATATTTAAGGTTGAATGTTTGTCTTAGGATAGGCCTATGTGCTAGCCCACAAAGA	PTGS2
## 35	TGTACTAGGCCTACTGGGGATCAGAGTTCCCAAGAAAGGAAACCTTTTCTTGTATCTGGA	CNR1
##	GeneCount	GenotypeCount
## 1	22	2 cannabidiol
## 3	3	3 cannabidiol
## 4	22	20 cannabidiol
## 24	4	4 cannabidiol
## 25	10	10 cannabidiol
## 35	1	1 cannabidiol
##	Study1_GSM2359851_CoV1	
## 1	302.82500	
## 3	8.99750	
## 4	1076.81550	
## 24	11.87500	
## 25	12.35425	
## 35	65.61000	
##	Study1_GSM2359853_CoV2	Study1_GSM2359850_ctrl1
## 1	246.56250	393.17250
## 3	14.20750	15.63000
## 4	856.73738	1321.94000
## 24	15.31250	15.35750
## 25	10.88063	11.85675
## 35	84.29750	44.82750
##		Study1_GSM2359852_ctrl2
## 1		380.69500
## 3		20.21500
## 4		1184.44325
## 24		8.09750
## 25		10.14488
## 35		68.61000

The above table shows the mean value of each genotype in the four samples, where two samples are liver tumor inoculated with CoV, and the other two are the liver tumor samples without CoV with both having been screened after 1 hour.

Since we want the fold change values within each geno type of the CBD network, lets get the means of each type of either CoV or ctrl. This is now the collective sample means of the genotype means per unique sample of either CoV or ctrl.

```

geneSeq3$CoV_Genotype_Mean <- rowMeans(geneSeq3[,6:7])
geneSeq3$Ctrl_Genotype_Mean <- rowMeans(geneSeq3[,8:9])
geneSeq3[,c(1,2,10,11)]

##                               SEQUENCE
Gene.Symbol
## 1  ACAATCAGATTGAAGCTTATCTATGACAGATGTGATCTTAAGTGTGCGATCCACAAAAAA
PPARG
## 3  AGCATCTGCTCCCTCTACCAGCTGGAGAACTACTGCAACTAGACGCAGCCCGCAGGCAGC
INS
## 4  CCAAGGCTTCATGACAAGGGAGTTTCTAAAGAGCCTGCGAAAGCCTTTTGGTGACTTTAT
PPARG
## 24 GGGGTATCCTGGGGGACCCAATGTAGGAGCTGCCTTGGCTCAGACATGTTTTCCGTGAAA
TNF
## 25 TGAGATATTTAAGGTTGAATGTTTGTCTTAGGATAGGCCTATGTGCTAGCCCACAAAGA
PTGS2
## 35 TGTACTAGGCCTACTGGGGATCAGAGTTCCCAAGAAAGGAAACCTTTTCTTGATCTGGA
CNR1
##      CoV_Genotype_Mean Ctrl_Genotype_Mean
## 1           274.69375           386.93375
## 3             11.60250             17.92250
## 4           966.77644          1253.19162
## 24            13.59375             11.72750
## 25            11.61744             11.00081
## 35            74.95375             56.71875

```

Looking at the above, both genotypes of PPARG decrease in CoV treated samples as well as insulin, INS. But tumor necrosis factor or TNF, prostaglandin-Endoperoxide Synthase 2 or PTGS2 and Cannabinoid Receptor 1 or CNR1 all increase in CoV treated samples. This could mean that in liver tumor samples that are inoculated with Coronavirus, more tumor suppressant or destroyer called TNF is activated, more pain reliever activation in the increased CNR1, and less insulin is produced. Insulin is a diabetic factor that plays a role in glucose processing for energy. If less insulin is produced after one hour, this could mean the body doesn't want to feed a contaminant it is recognizing in the body by lessening the need for the body to consume more glucose. An aside on the insulin and glucose relationship in regulating the body: Insulin and glucose are supposed to be balanced, where the more glucose is burned by the body for energy will lead to more insulin produced to compensate energy needs. The two types of diabetes controlled by insulin is either an overproduction of insulin that leads to an irregular need to eat more than normal amounts of food to get more glucose in the blood, or an underproduction of insulin that leads to an inadequate amount of glucose the body needs, loss of appetite and irregularly small amounts of food consumed.

Immediately, some more analysis to answer questions of these assertions on the results above can be further confirmed by:

1.) Analyzing the **time sequence of the blood capillary samples** inoculated with CoV and their respective control groups. Examining how these CBD genes are effected over time.

2.) Confirming that insulin is effected by CoV by looking at the **diabetic gene networks** for type 1 and type 2 diabetes.

3.) Also, looking at the **tumorigenesis gene network** to see how the genes that are involved in tumor growth in the body react to CoV in the liver tumor samples and the blood capillary samples.

---

---

Lets put an interesting bar chart into this script to show what we have so far in the CBD genes network for liver tumor samples inoculated over one hour's time. Lets use the mutate function of dplyr to create a new feature that will name each gene with more than one genotype or copy number variant in this data. Then use tidyr to gather the sample means per genotype. And finally, plot the CoV and ctrl means using ggplot2.

```
CBD_plot <- geneSeq3 %>% mutate(geneClass = paste(Gene.Symbol, GenotypeCount,
                                                    sep='_'))

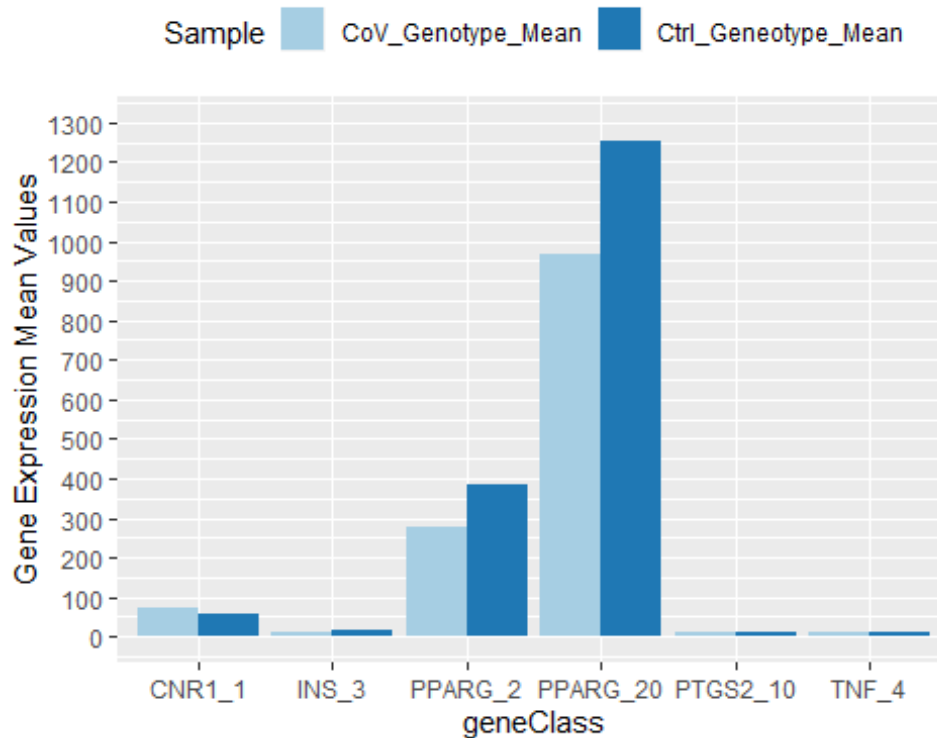
CBD_plot2 <- gather(CBD_plot, 'Sample', 'MeanValue', 10:11)
```

---

---

```
ggplot(data = CBD_plot2, aes(x=geneClass, y=MeanValue, fill=Sample)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous(breaks = seq(0, 1300, by=100), limits=c(0,1300)) +
  scale_fill_brewer(palette='Paired') +
  #ggtitle('CBD Network Gene Expression Values Treated with Coronavirus') +
  theme(legend.position="top") +
  #xlab('Gene and Genotype or Single Nucleotide Polymorphism or Copy Number
  Variant') +
  ylab('Gene Expression Mean Values')
```





The lower values aren't as easy to distinguish as the scale of change in Mean values is significantly greater for some genes. We could fix this by making the plot fold change and add that field, or adding a log2 field of the means to put on the same scale for visualization purposes. Or cut off the scale for those less than 100 in Mean Values and those greater than 100.

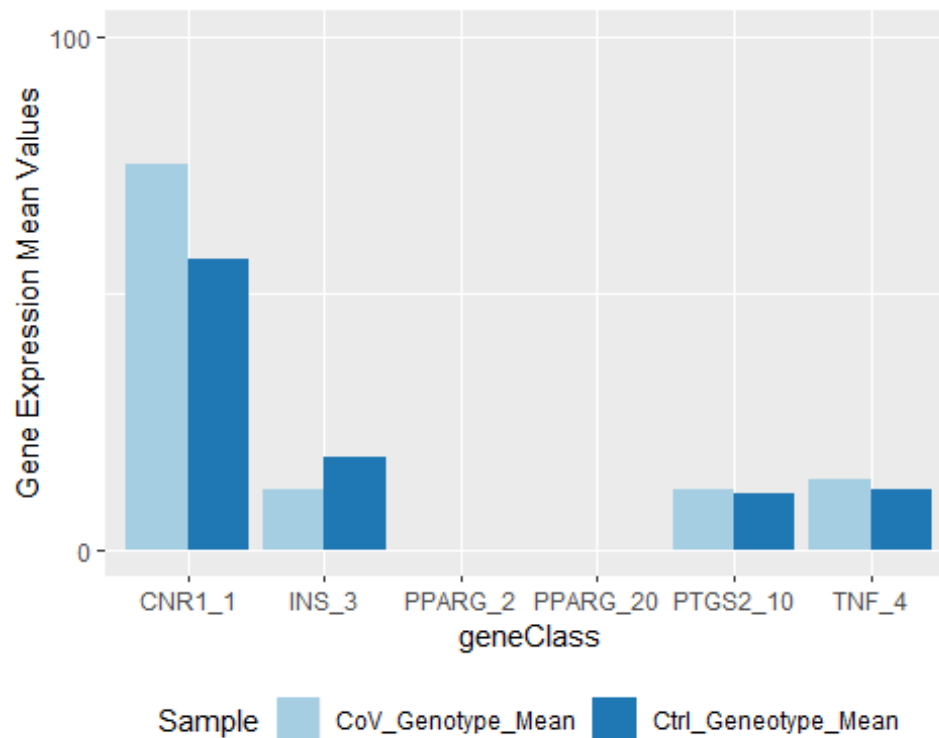
---



---

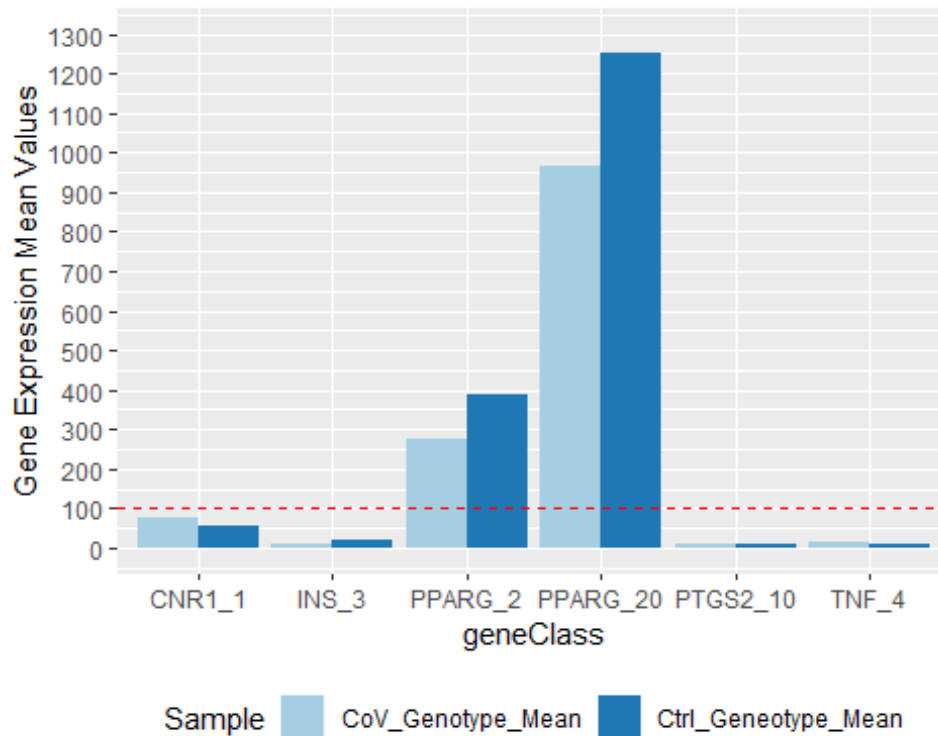
The following is the genes less than 100 for Mean genotype expression values.

```
ggplot(data = CBD_plot2, aes(x=geneClass, y=MeanValue, fill=Sample)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous(breaks = seq(0, 100, by=100), limits=c(0,100)) +
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="bottom") +
  #ggtitle('CBD Network Gene Expression Values Less than 100 Treated with
  CoV') +
  ylab('Gene Expression Mean Values')
```



The following is the genes greater than 100 for Mean genotype expression values:

```
ggplot(data = CBD_plot2, aes(x=geneClass, y=MeanValue, fill=Sample)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous(breaks = seq(0, 1300, by=100), limits=c(0,1300)) +
  scale_fill_brewer(palette='Paired') +
  geom_hline(yintercept=100, linetype="dashed", color = "red") +
  theme(legend.position="bottom") +
  #ggtitle('CBD Network Gene Levels > 100 Treated with CoV') +
  ylab('Gene Expression Mean Values')
```

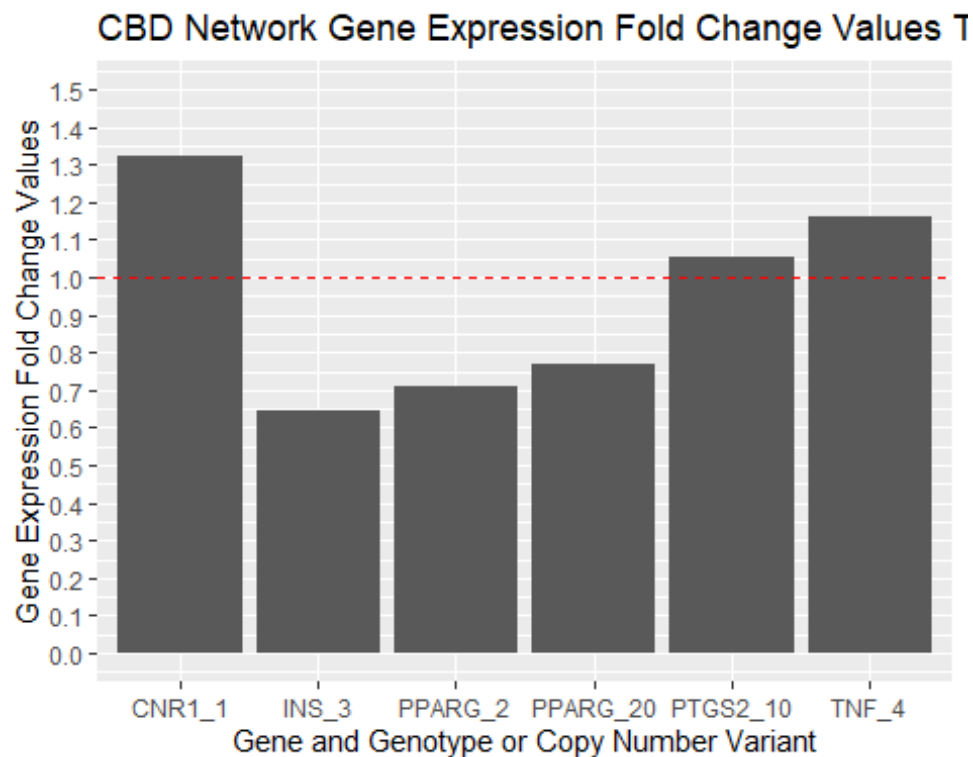


The following is the fold change chart.

```
CBD_plot$FoldChange <-
CBD_plot$CoV_Genotype_Mean/CBD_plot$Ctrl_Genotype_Mean
CBD_plot3 <- CBD_plot[,c(1:9,12,10,11,13)] #fold change plot data
CBD_plot2$logMean <- log2(CBD_plot2$MeanValue) # log2 scaled plot data
```

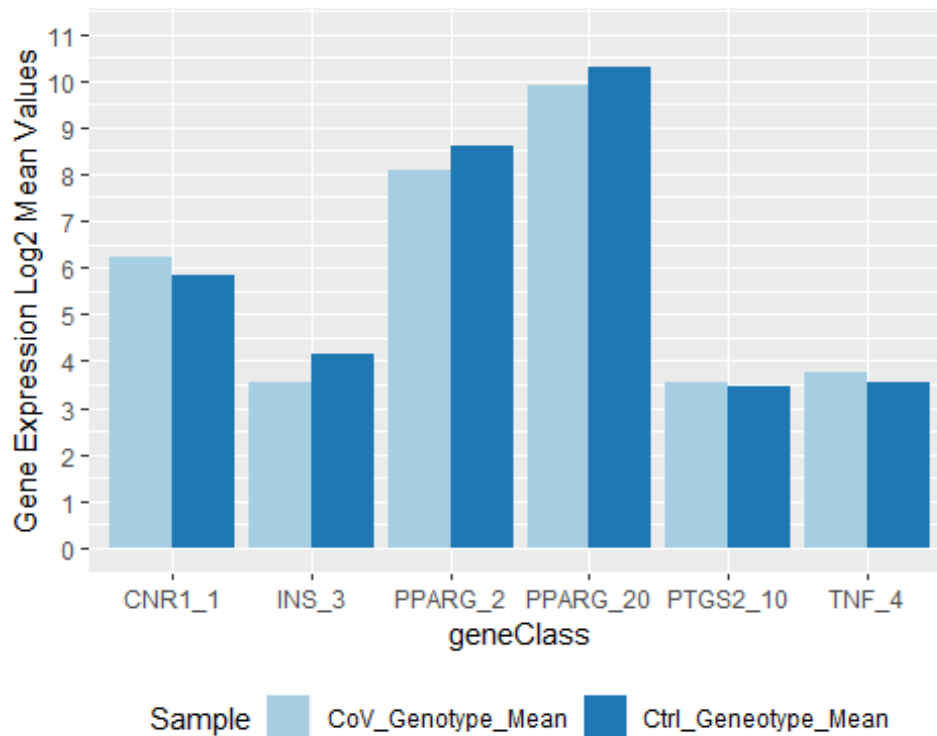
The chart of fold change values of the genotype or single nucleotide polymorphism (SNP) or copy number variant of the CBD gene network top five listed genes by Relevance.score.with an added dashed red line to show that genes below the line decreased and those above increased in percentage of gene expression of CoV/ctrl.

```
ggplot(data = CBD_plot3, aes(x=geneClass, y=FoldChange)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous(breaks = seq(0, 1.5, by=.1), limits=c(0,1.5))+
  scale_fill_brewer(palette='Paired') +
  geom_hline(yintercept=1, linetype="dashed", color = "red")+
  ggtitle('CBD Network Gene Expression Fold Change Values Treated with CoV')+
  xlab('Gene and Genotype or Copy Number Variant')+
  ylab('Gene Expression Fold Change Values')
```



The next chart is the log2 scaled Mean values for the CBD genotypes.

```
ggplot(data = CBD_plot2, aes(x=geneClass, y=logMean, fill=Sample)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous(breaks = seq(0, 11, by=1), limits=c(0,11)) +
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="bottom") +
  #ggtitle('CBD Network Gene Expression Log2 Mean Values Treated with
  Coronavirus') +
  ylab('Gene Expression Log2 Mean Values')
```



Those visuals help establish that the PPARG has two copy number variants of the gene that are expressed much more in both the control and CoV inoculated samples by mean genotype expression values, fold change values, and log2 scaled values. We can see that CoV infected samples after 1 hour in liver tumor samples increases genotype expressions of CNR\_1, PTGS2, and TNF. And that INS and PPARG are decreased in the first hour of inoculation with CoV.

---

Now, let's start working on the data that confirms or better asserts what is going on with these genes. Recall that we wanted to examine three possibilities to understand why we made the observation above on the what the data provided for the cannabidiol genes network.

- 1.) Analyzing the **time sequence of the blood capillary samples** inoculated with CoV and their respective control groups. Examining how these CBD genes are effected over time.
  - 2.) Confirming that insulin is effected by CoV by looking at the **diabetic gene networks** for type 1 and type 2 diabetes.
  - 3.) Also, looking at the **tumorigenesis gene network** to see how the genes that are involved in tumor growth in the body react to CoV in the liver tumor samples and the blood capillary samples.
-

---

Lets look at 1.) for the time sequence of the blood capillary samples for each of 0, 12, 24, 36, and 48 hours for the mean values of CoV and ctrl samples.

Time 0: We have to go back to the beginning when adding the platform sequence values to the probe ideas of the series. We have the cleaned series for this data, Series100509. Otherwise, there are duplicate gene values, and the SNP values of the CBD gene with 2 SNPs will be the same.

```
series100509 <- Series100509[, -c(1,3)]
s100 <- series100509[, c(1,2, grep('0hr', colnames(series100509)))]

CBD5 <- cannabidiol5[, 1:2]
CBD5_s100 <- merge(CBD5, s100, by.x='Gene.Symbol', by.y='GENE_SYMBOL')

seqN_CBD <- CBD5_s100 %>% group_by(SEQUENCE) %>% count(n=n())
seqN_CBD <- seqN_CBD[, -3]
colnames(seqN_CBD)[2] <- 'GenotypeCount'

gen_CBD <- CBD5_s100 %>% group_by(Gene.Symbol) %>% count(n=n())
gen_CBD <- gen_CBD[, -3]
colnames(gen_CBD)[2] <- 'GeneCount'

CBD5_s100_2 <- merge(seqN_CBD, CBD5_s100, by.x='SEQUENCE', by.y='SEQUENCE')
CBD5_s100_3 <- merge(gen_CBD, CBD5_s100_2, by.x='Gene.Symbol',
by.y='Gene.Symbol')

CBD5_s100_3$geneClass <- ifelse(CBD5_s100_3$GeneCount>1,
paste(as.character(CBD5_s100_3$Gene.Symbol),
CBD5_s100_3$GeneCount, sep='_'),
as.character(CBD5_s100_3$Gene.Symbol))
CBD5_s100_3$geneClass[3] <- 'PPARG_1'
CBD5_s100_3 <- CBD5_s100_3[, c(1:2, 16, 3:15)]

covSamples <- as.vector(colnames(CBD5_s100_3)[7:11])
CBD_seqCoV_t0 <- CBD5_s100_3 %>% group_by(SEQUENCE) %>%
summarise_at(vars(covSamples), mean)
CBD_seqCoV_t0$CoV_t0_SeqMeans <- rowMeans(CBD_seqCoV_t0[2:6])

CoV_t0 <- CBD5_s100_3[, 1:6]
CoV_t0_Means <- merge(CoV_t0, CBD_seqCoV_t0, by.x='SEQUENCE',
by.y='SEQUENCE')
CoV_t0_seqMean <- CoV_t0_Means[, c(1:6, 12)]

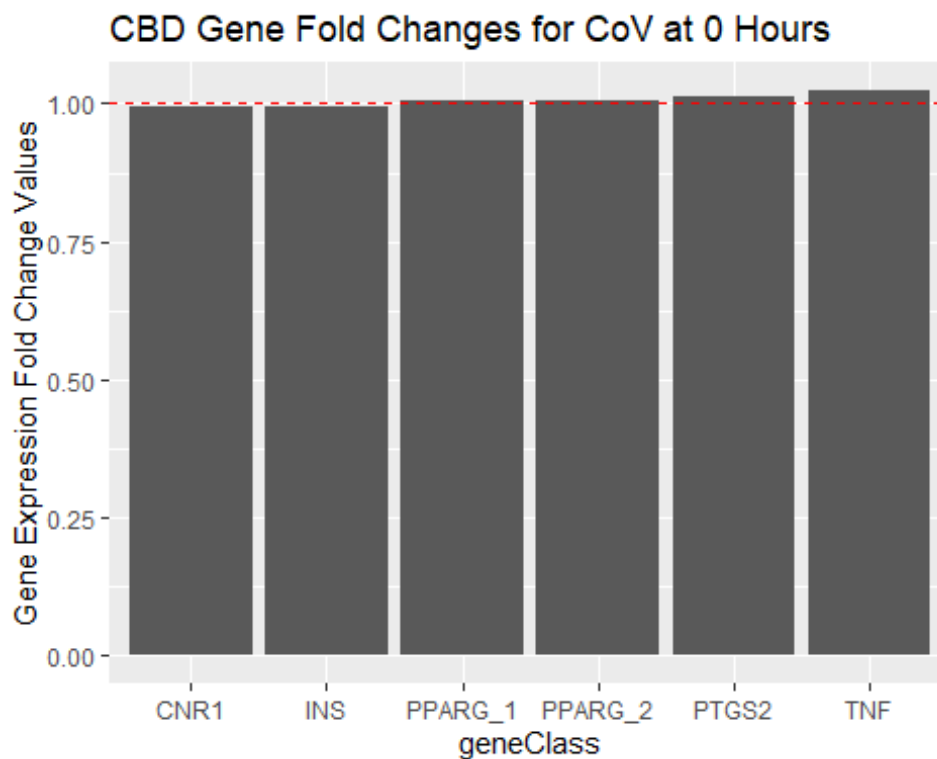
ctrlSamples <- as.vector(colnames(CBD5_s100_3)[12:16])
CBD_seqCtrl_t0 <- CBD5_s100_3 %>% group_by(SEQUENCE) %>%
summarise_at(vars(ctrlSamples), mean)
CBD_seqCtrl_t0$Ctrl_t0_SeqMeans <- rowMeans(CBD_seqCtrl_t0[2:6])
```

```
Ctrl_t0_Means <- merge(CoV_t0, CBD_seqCtrl_t0, by.x='SEQUENCE',
by.y='SEQUENCE')
Ctrl_t0_seqMean <- Ctrl_t0_Means[,c(1,12)]

t0_CBD <- merge(CoV_t0_seqMean, Ctrl_t0_seqMean, by.x='SEQUENCE',
by.y='SEQUENCE')

t0_CBD$FoldChange <- t0_CBD$CoV_t0_SeqMeans/t0_CBD$Ctrl_t0_SeqMeans
```

```
ggplot(data = t0_CBD, aes(x=geneClass, y=FoldChange)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+
  scale_fill_brewer(palette='Paired') +
  geom_hline(yintercept=1, linetype="dashed", color = "red")+
  theme(legend.position="bottom")+
  ggtitle('CBD Gene Fold Changes for CoV at 0 Hours')+
  ylab('Gene Expression Fold Change Values')
```

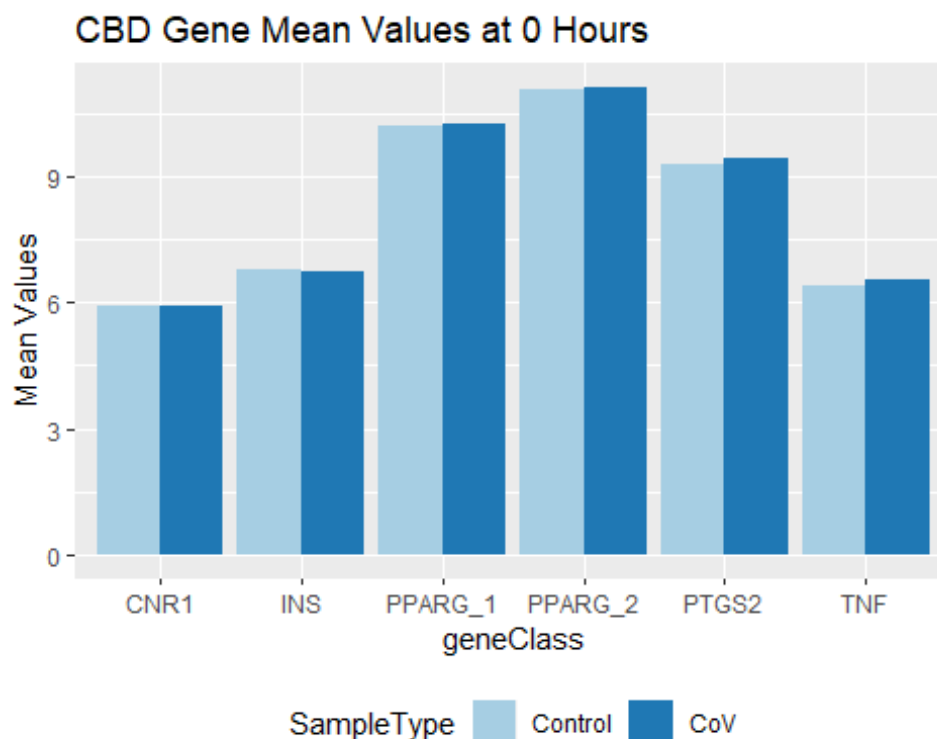


The above chart is one of the fold change values around 1, with those less than 1 insulin and the cannabidiol receptor 1 genes.

Now we will look at the mean values of those CBD genes on the blood capillary of the CoV and control sample means.

```
tidyCBD <- as.data.frame(gather(t0_CBD, 'SampleType', 'MeanValue', 7:8))
tidyCBD$SampleType <- gsub('CoV_t0_SeqMeans', 'CoV', tidyCBD$SampleType)
tidyCBD$SampleType <- gsub('Ctrl_t0_SeqMeans', 'Control', tidyCBD$SampleType)

ggplot(data = tidyCBD, aes(x=geneClass, y=MeanValue, fill=SampleType)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous() +
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="bottom") +
  ggtitle('CBD Gene Mean Values at 0 Hours') +
  ylab('Mean Values')
```



As we see in the chart above the tumor necrosis factor, TNF, gene increases in Coronavirus samples compared to control samples in blood capillaries as it did in liver tumor samples, as well as insulin, INS, decreasing in both tissue types. But that was at zero hours time for blood capillary tissue, closest to the one hour time of the liver tumor samples previously analyzed.

---

Now, let's look at what happens after 24 hours, skipping 12 hours for these blood capillary samples inoculated with CoV.



```

series100509_24 <- Series100509[, -c(1,3)]
s100_24 <- series100509[, c(1,2,grep('24hr', colnames(series100509)))]

CBD5_24 <- cannabidiol5[,1:2]
CBD5_s100_24 <- merge(CBD5, s100_24, by.x='Gene.Symbol', by.y='GENE_SYMBOL')

seqN_CBD_24 <- CBD5_s100_24 %>% group_by(SEQUENCE) %>% count(n=n())
seqN_CBD_24 <- seqN_CBD_24[, -3]
colnames(seqN_CBD_24)[2] <- 'GenotypeCount'

gen_CBD_24 <- CBD5_s100_24 %>% group_by(Gene.Symbol) %>% count(n=n())
gen_CBD_24 <- gen_CBD_24[, -3]
colnames(gen_CBD_24)[2] <- 'GeneCount'

CBD5_s100_2_24 <- merge(seqN_CBD_24, CBD5_s100_24, by.x='SEQUENCE',
by.y='SEQUENCE')
CBD5_s100_3_24 <- merge(gen_CBD_24, CBD5_s100_2_24, by.x='Gene.Symbol',
by.y='Gene.Symbol')

CBD5_s100_3_24$geneClass <- ifelse(CBD5_s100_3_24$GeneCount>1,
paste(as.character(CBD5_s100_3_24$Gene.Symbol),
      CBD5_s100_3_24$GeneCount, sep='_'),
      as.character(CBD5_s100_3_24$Gene.Symbol))
CBD5_s100_3_24$geneClass[3] <- 'PPARG_1'
CBD5_s100_3_24 <- CBD5_s100_3_24[, c(1:2,16,3:15)]

covSamples_24 <- as.vector(colnames(CBD5_s100_3_24)[7:11])
CBD_seqCoV_t_24 <- CBD5_s100_3_24 %>% group_by(SEQUENCE) %>%
summarise_at(vars(covSamples_24), mean)
CBD_seqCoV_t_24$CoV_t_24_SeqMeans <- rowMeans(CBD_seqCoV_t_24[2:6])

CoV_t_24 <- CBD5_s100_3_24[, 1:6]
CoV_t_24_Means <- merge(CoV_t_24, CBD_seqCoV_t_24, by.x='SEQUENCE',
by.y='SEQUENCE')
CoV_t_24_seqMean <- CoV_t_24_Means[, c(1:6,12)]

ctrlSamples_24 <- as.vector(colnames(CBD5_s100_3_24)[12:16])
CBD_seqCtrl_t_24 <- CBD5_s100_3_24 %>% group_by(SEQUENCE) %>%
summarise_at(vars(ctrlSamples_24), mean)
CBD_seqCtrl_t_24$Ctrl_t_24_SeqMeans <- rowMeans(CBD_seqCtrl_t_24[2:6])

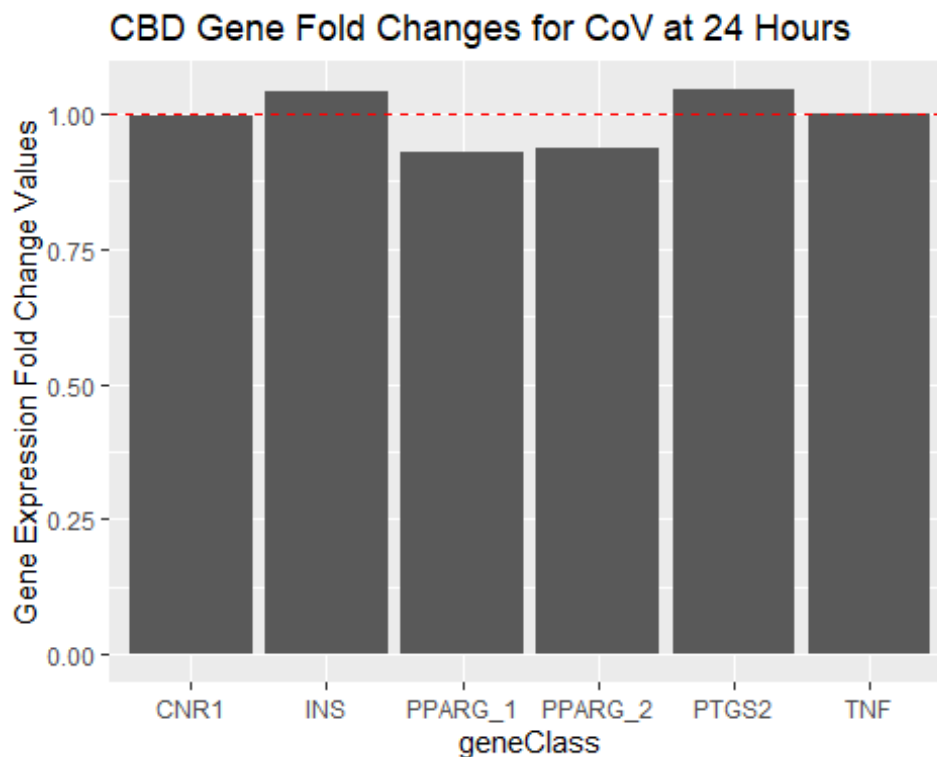
Ctrl_t_24_Means <- merge(CoV_t_24, CBD_seqCtrl_t_24, by.x='SEQUENCE',
by.y='SEQUENCE')
Ctrl_t_24_seqMean <- Ctrl_t_24_Means[, c(1,12)]

t_24_CBD <- merge(CoV_t_24_seqMean, Ctrl_t_24_seqMean, by.x='SEQUENCE',
by.y='SEQUENCE')

```

```
t_24_CBD$FoldChange <- t_24_CBD$CoV_t_24_SeqMeans/t_24_CBD$Ctrl_t_24_SeqMeans

ggplot(data = t_24_CBD, aes(x=geneClass, y=FoldChange)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+
  scale_fill_brewer(palette='Paired') +
  geom_hline(yintercept=1, linetype="dashed", color = "red")+
  theme(legend.position="bottom")+
  ggtitle('CBD Gene Fold Changes for CoV at 24 Hours')+
  ylab('Gene Expression Fold Change Values')
```



Looking at the chart above on the fold change values of the blood capillaries over 24 hours after having been inoculated with CoV compared to a control group, there is a switch or change in those genes that were previously up-regulated and down-regulated.

TNF and CNR1 have negligible changes, INS is now up-regulated after 24 hours of being inoculated, and both genotypes or copy number variants of PPARG have decreased. Previously, at 0 hours in blood capillaries we saw that INS and CNR1 were down-regulated, both PPARG genotypes were negligibly close to no change at all, and TNF and PTGS2 were up-regulated.

---

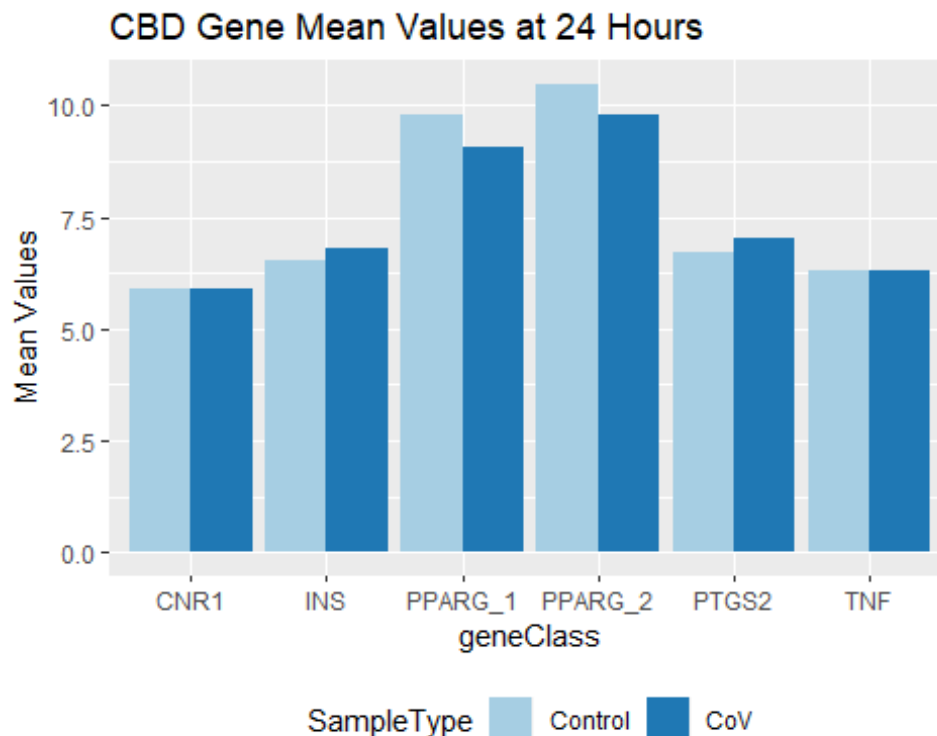
```
tidyCBD_24 <- as.data.frame(gather(t_24_CBD, 'SampleType', 'MeanValue', 7:8))
tidyCBD_24$SampleType <- gsub('CoV_t_24_SeqMeans', 'CoV',
tidyCBD_24$SampleType)
```

```

tidyCBD_24$SampleType <- gsub('Ctrl_t_24_SeqMeans', 'Control',
tidyCBD_24$SampleType)

ggplot(data = tidyCBD_24, aes(x=geneClass, y=MeanValue, fill=SampleType)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous() +
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="bottom") +
  ggtitle('CBD Gene Mean Values at 24 Hours') +
  ylab('Mean Values')

```



The above bar chart shows the mean values of the control and CoV inoculated samples after 24 hours in the blood capillary samples. In CoV inoculation, the gene expression values for both PPARG genotypes decrease, in INS and PTGS2 there is an up-regulation, and CNR1 and TNF show negligible changes, with a slight decrease in CNR1 gene expression.

---

Now, let's look at after 48 hours of being inoculated with CoV in these blood capillary samples.

```

series100509_48 <- Series100509[, -c(1,3)]
s100_48 <- series100509[, c(1,2,grep('48hr', colnames(series100509)))]

CBD5_48 <- cannabidiol5[, 1:2]
CBD5_s100_48 <- merge(CBD5, s100_48, by.x='Gene.Symbol1', by.y='GENE_SYMBOL')

```

```

seqN_CBD_48 <- CBD5_s100_48 %>% group_by(SEQUENCE) %>% count(n=n())
seqN_CBD_48 <- seqN_CBD_48[, -3]
colnames(seqN_CBD_48)[2] <- 'GenotypeCount'

gen_CBD_48 <- CBD5_s100_48 %>% group_by(Gene.Symbol) %>% count(n=n())
gen_CBD_48 <- gen_CBD_48[, -3]
colnames(gen_CBD_48)[2] <- 'GeneCount'

CBD5_s100_2_48 <- merge(seqN_CBD_48, CBD5_s100_48, by.x='SEQUENCE',
by.y='SEQUENCE')
CBD5_s100_3_48 <- merge(gen_CBD_48, CBD5_s100_2_48, by.x='Gene.Symbol',
by.y='Gene.Symbol')

CBD5_s100_3_48$geneClass <- ifelse(CBD5_s100_3_48$GeneCount>1,
paste(as.character(CBD5_s100_3_48$Gene.Symbol),
      CBD5_s100_3_48$GeneCount, sep='_'),
      as.character(CBD5_s100_3_48$Gene.Symbol))
CBD5_s100_3_48$geneClass[3] <- 'PPARG_1'
CBD5_s100_3_48 <- CBD5_s100_3_48[, c(1:2, 16, 3:15)]

covSamples_48 <- as.vector(colnames(CBD5_s100_3_48)[7:11])
CBD_seqCoV_t_48 <- CBD5_s100_3_48 %>% group_by(SEQUENCE) %>%
summarise_at(vars(covSamples_48), mean)
CBD_seqCoV_t_48$CoV_t_48_SeqMeans <- rowMeans(CBD_seqCoV_t_48[2:6])

CoV_t_48 <- CBD5_s100_3_48[, 1:6]
CoV_t_48_Means <- merge(CoV_t_48, CBD_seqCoV_t_48, by.x='SEQUENCE',
by.y='SEQUENCE')
CoV_t_48_seqMean <- CoV_t_48_Means[, c(1:6, 12)]

ctrlSamples_48 <- as.vector(colnames(CBD5_s100_3_48)[12:16])
CBD_seqCtrl_t_48 <- CBD5_s100_3_48 %>% group_by(SEQUENCE) %>%
summarise_at(vars(ctrlSamples_48), mean)
CBD_seqCtrl_t_48$Ctrl_t_48_SeqMeans <- rowMeans(CBD_seqCtrl_t_48[2:6])

Ctrl_t_48_Means <- merge(CoV_t_48, CBD_seqCtrl_t_48, by.x='SEQUENCE',
by.y='SEQUENCE')
Ctrl_t_48_seqMean <- Ctrl_t_48_Means[, c(1, 12)]

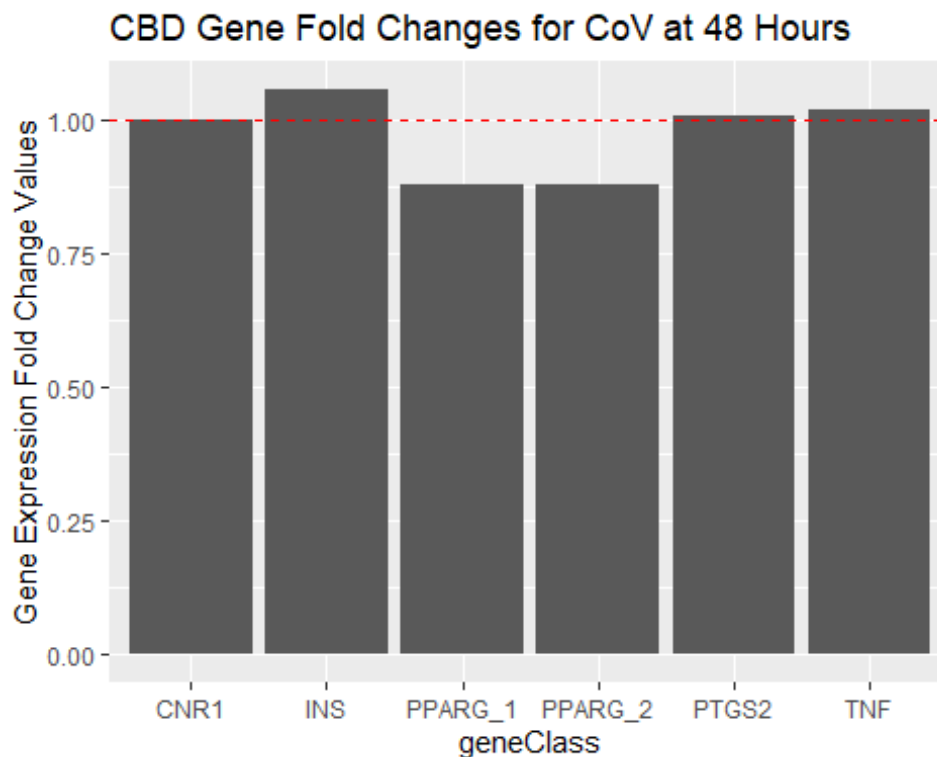
t_48_CBD <- merge(CoV_t_48_seqMean, Ctrl_t_48_seqMean, by.x='SEQUENCE',
by.y='SEQUENCE')

t_48_CBD$FoldChange <- t_48_CBD$CoV_t_48_SeqMeans/t_48_CBD$Ctrl_t_48_SeqMeans

ggplot(data = t_48_CBD, aes(x=geneClass, y=FoldChange)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+

```

```
scale_fill_brewer(palette='Paired') +
geom_hline(yintercept=1, linetype="dashed", color = "red")+
theme(legend.position="bottom")+
ggtitle('CBD Gene Fold Changes for CoV at 48 Hours')+
ylab('Gene Expression Fold Change Values')
```



From the above chart of fold changes after 48 hours, we see that not much has changed compared to after 24 hours, except the TNF has now become up-regulated as it was at 0 hours but not after 24 hours.

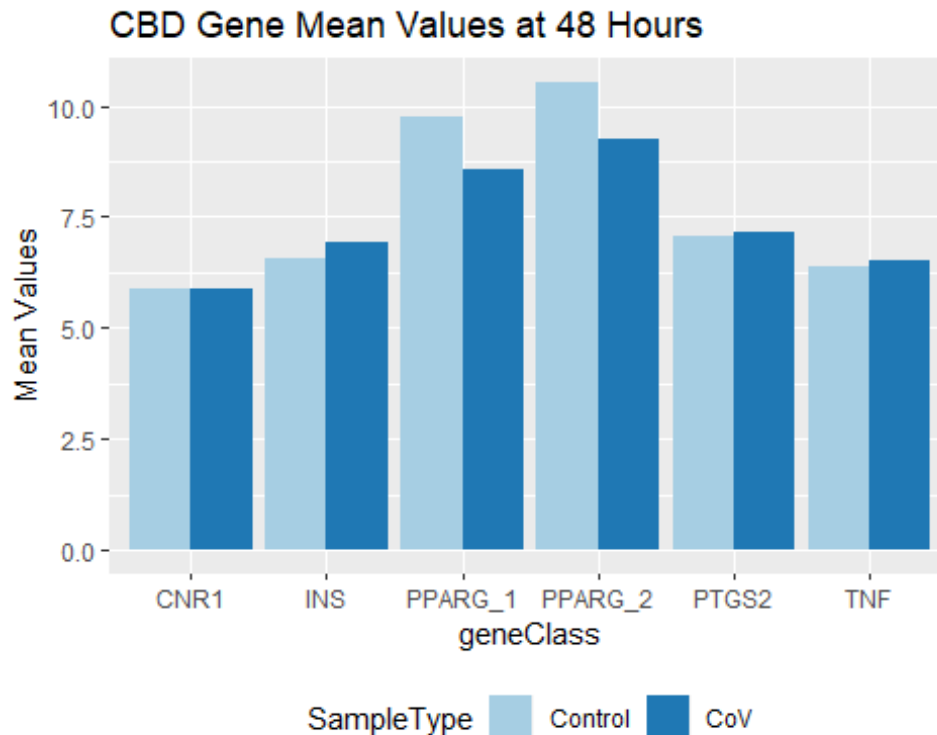
CNR1 remains having negligible changes as was the case after 24 hours, INS, TNF, and PTGS2 are up-regulated after 48 hours of being inoculated, and both genotypes or copy number variants of PPARG have decreased.

Previously, at 0 hours in blood capillaries we saw that INS and CNR1 were down-regulated, both PPARG genotypes were negligibly close to no change at all, and TNF and PTGS2 were up-regulated.

---

```
tidyCBD_48 <- as.data.frame(gather(t_48_CBD, 'SampleType', 'MeanValue', 7:8))
tidyCBD_48$SampleType <- gsub('CoV_t_48_SeqMeans', 'CoV',
tidyCBD_48$SampleType)
tidyCBD_48$SampleType <- gsub('Ctrl_t_48_SeqMeans', 'Control',
tidyCBD_48$SampleType)
```

```
ggplot(data = tidyCBD_48, aes(x=geneClass, y=MeanValue, fill=SampleType)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous() +
  scale_fill_brewer(palette='Paired') +
  theme(legend.position="bottom") +
  ggtitle('CBD Gene Mean Values at 48 Hours') +
  ylab('Mean Values')
```



The above bar chart shows the mean values of the control and CoV inoculated samples after 48 hours in the blood capillary samples. In CoV inoculation, the gene expression values for both PPARG genotypes decrease. In INS, TNF, and PTGS2 there is an up-regulation, and CNR1 shows negligible changes, with a slight decrease in CNR1 gene expression.

3.) Lets look at the tumorigenesis genes and the liver tumor samples that were inoculated with the Coronavirus.

Get the table with the five genes that were the highest relevance.scoring made earlier to be combined with all our samples. Then select only those liver tumor samples except for the overly scaled samples 1 and 2.Include the heat inactivated CoV and the added IL-alpha1 to CoV samples.

```
tumorigenesis5_path_1 <- tumorigenesis5_path[,c(1,9,12,13,16:21)]
describe <-
```

```
tumorigenesis5_path[!duplicated(tumorigenesis5_path$Gene.Symbol),c(1,2,8)]
describe
```

```
##      Gene.Symbol      Description
## 1      CTNNB1      Catenin Beta 1
## 23     EGFR      Epidermal Growth Factor Receptor
## 32     IGF2      Insulin Like Growth Factor 2
## 36     KRAS      KRAS Proto-Oncogene, GTPase
## 40     TP53      Tumor Protein P53
##
DESCRIPTION
## 1 Homo sapiens catenin (cadherin-associated protein), beta 1, 88kDa
(CTNNB1), transcript variant 1, mRNA [NM_001904]
## 23 Homo sapiens epidermal growth factor receptor
(EGFR), transcript variant 2, mRNA [NM_201282]
## 32 Homo sapiens insulin-like growth factor 2 (somatomedin A)
(IGF2), transcript variant 1, mRNA [NM_000612]
## 36 Homo sapiens v-Ki-ras2 Kirsten rat sarcoma viral oncogene homolog
(KRAS), transcript variant a, mRNA [NM_033360]
## 40 Homo sapiens tumor protein p53
(TP53), transcript variant 1, mRNA [NM_000546]
```

```
colnames(tumorigenesis5_path_1)
```

```
## [1] "Gene.Symbol"      "SEQUENCE"
## [3] "GSM2359910_CoV3"  "GSM2359913_CoV4"
## [5] "GSM2359911_ctrl3" "GSM2359914_ctrl4"
## [7] "GSM2359912_IL1"   "GSM2359917_IL2"
## [9] "GSM2359915_inactiveHeatCoV1" "GSM2359916_inactiveHeatCoV2"
```

Now, group by SEQUENCE to see how many genotypes there are in this set of five tumor creating network of genes.

```
seqTumor <- tumorigenesis5_path_1 %>% group_by(SEQUENCE) %>% count(n=n())
Seq_tumor <- as.data.frame(seqTumor[, -3])
```

```
colnames(Seq_tumor)[2] <- 'genoTypeCount'
```

```
Seq_Tumor <- merge(Seq_tumor, tumorigenesis5_path_1,
  by.x='SEQUENCE', by.y='SEQUENCE')
```

```
Seq_Tumor$CoV_Means <- rowMeans(Seq_Tumor[4:5])
Seq_Tumor$Ctrl_Means <- rowMeans(Seq_Tumor[6:7])
Seq_Tumor$IL_A_Means <- rowMeans(Seq_Tumor[8:9])
Seq_Tumor$IA_CoV_Means <- rowMeans(Seq_Tumor[10:11])
```

```
seqTumor1 <- Seq_Tumor %>% group_by(Gene.Symbol) %>% count(n=n())
Seq_tumor1 <- as.data.frame(seqTumor1[, -3])
colnames(Seq_tumor1)[2] <- 'geneCount'
Seq_Tumor1 <- merge(Seq_tumor1, Seq_Tumor, by.x='Gene.Symbol',
  by.y='Gene.Symbol')
```

```

Seq_Tumor1$FoldChange_CoV <- Seq_Tumor1$CoV_Means/Seq_Tumor1$Ctrl_Means
Seq_Tumor1$FoldChange_IL_a <- Seq_Tumor1$IL_A_Means/Seq_Tumor1$Ctrl_Means
Seq_Tumor1$FoldChange_IA_CoV <- Seq_Tumor1$IA_CoV_Means/Seq_Tumor1$Ctrl_Means

EGFR1 <- grep('^GGAAATC', Seq_Tumor1$SEQUENCE)
EGFR2 <- grep('^TATCAA', Seq_Tumor1$SEQUENCE)
EGFR3 <- grep('^AGAACT', Seq_Tumor1$SEQUENCE)

IGF2_1 <- grep('^CTCAACTC', Seq_Tumor1$SEQUENCE)
IGF2_2 <- grep('^TGCTTCC', Seq_Tumor1$SEQUENCE)

CTNNB1_1 <- grep('^TGATCAA', Seq_Tumor1$SEQUENCE)
CTNNB1_2 <- grep('^ATGATGGA', Seq_Tumor1$SEQUENCE)

KRAS_1 <- grep('^TCAGGACT', Seq_Tumor1$SEQUENCE)
KRAS_2 <- grep('^CTGAGTCA', Seq_Tumor1$SEQUENCE)

TP53_1 <- grep('^CTGTGAGG', Seq_Tumor1$SEQUENCE)
TP53_2 <- grep('^CAGCTACG', Seq_Tumor1$SEQUENCE)

Seq_Tumor1$Genotype_Class <- 'class'

Seq_Tumor1$Genotype_Class[EGFR1] <- 'EGFR_1'
Seq_Tumor1$Genotype_Class[EGFR2] <- 'EGFR_2'
Seq_Tumor1$Genotype_Class[EGFR3] <- 'EGFR_3'
Seq_Tumor1$Genotype_Class[IGF2_1] <- 'IGF2_1'
Seq_Tumor1$Genotype_Class[IGF2_2] <- 'IGF2_2'
Seq_Tumor1$Genotype_Class[CTNNB1_1] <- 'CTNNB1_1'
Seq_Tumor1$Genotype_Class[CTNNB1_2] <- 'CTNNB1_2'
Seq_Tumor1$Genotype_Class[KRAS_1] <- 'KRAS_1'
Seq_Tumor1$Genotype_Class[KRAS_2] <- 'KRAS_2'
Seq_Tumor1$Genotype_Class[TP53_1] <- 'TP53_1'
Seq_Tumor1$Genotype_Class[TP53_2] <- 'TP53_2'

```

Make the data tidy.

```

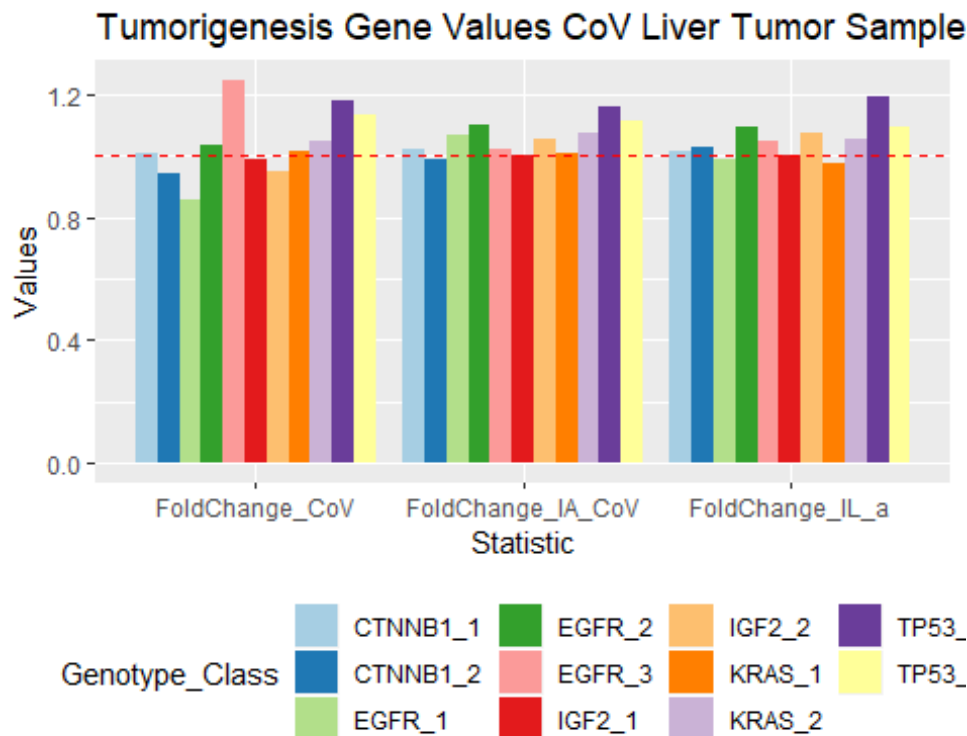
tidy_SeqTumor1 <- gather(Seq_Tumor1, 'Statistic', 'Value', 13:19)
foldChanges <- subset(tidy_SeqTumor1,
tidy_SeqTumor1$Statistic=="FoldChange_IL_a" |
tidy_SeqTumor1$Statistic=="FoldChange_IA_CoV" |
tidy_SeqTumor1$Statistic=="FoldChange_CoV")

ggplot(data = foldChanges, aes(x=Statistic, y=Value, fill=Genotype_Class)) +
  geom_bar(stat='identity', position=position_dodge()) +
  scale_y_continuous() +
  scale_fill_brewer(palette='Paired') +
  geom_hline(yintercept=1, linetype="dashed", color = "red") +
  theme(legend.position="bottom") +

```



```
ggtitle('Tumorigenesis Gene Values CoV Liver Tumor Samples')+
ylab('Values')
```



The above bar chart shows the fold change values of the tumorigenesis genes in the liver tumor samples inoculated with Coronavirus, Coronavirus and interleukin alpha1, and heat inactive CoV compared to the control group and over a one hour time span in vitro or cell culture. The genes closest to the dashed line represent negligible change. IGF2 genotype 1 of 3 and the 1st genotype of KRAS are negligible for changes after being inoculated with either of the three inoculations over one hour.

The EGFR for two genotypes out of three, both genotypes of TP53, and the second genotype of KRAS are increased as they are above the line in fold change values as the chart above shows.

The genotype of the three for EGFR that isn't increased is decreased in the CoV inoculated samples, but increased in the heat inactivated CoV samples and negligible change in the IL-A mixed with CoV sample. These could be clues to how CoV effects tumor producing genes or the network of genes responsible for causing tumors. However, this study is only done for a one hour time span, not like the blood capillary samples that were done over a 48 hour time span.

These five tumorigenesis genes are shown in the following table that was made earlier:

describe

```
##      Gene.Symbol              Description
## 1      CTNNB1              Catenin Beta 1
## 23      EGFR Epidermal Growth Factor Receptor
## 32      IGF2      Insulin Like Growth Factor 2
## 36      KRAS      KRAS Proto-Oncogene, GTPase
## 40      TP53              Tumor Protein P53
##
DESCRIPTION
## 1 Homo sapiens catenin (cadherin-associated protein), beta 1, 88kDa
(CTNNB1), transcript variant 1, mRNA [NM_001904]
## 23      Homo sapiens epidermal growth factor receptor
(EGFR), transcript variant 2, mRNA [NM_201282]
## 32      Homo sapiens insulin-like growth factor 2 (somatomedin A)
(IGF2), transcript variant 1, mRNA [NM_000612]
## 36 Homo sapiens v-Ki-ras2 Kirsten rat sarcoma viral oncogene homolog
(KRAS), transcript variant a, mRNA [NM_033360]
## 40      Homo sapiens tumor protein p53
(TP53), transcript variant 1, mRNA [NM_000546]
```

We just analyzed the tumorigenesis genes in the liver tumor samples of CoV, but let's look at the blood capillary samples as well to see how these genotypes behave. Are the genes expressed the same or do they differ when looking at blood capillary tissue instead of liver tumor tissue infected with CoV.

We will use the first four columns and the last one of the Seq\_Tumor1 table that has the count of the genes and genotypes in it as well as the corresponding given name by which genotype it was.

```
Seq_cap_tg <- Seq_Tumor1[,c(1:4,20)]
Seq_cap_tg <- Seq_cap_tg[!duplicated(Seq_cap_tg$Genotype_Class),]
colnames(Seq_cap_tg)

## [1] "Gene.Symbol"      "geneCount"      "SEQUENCE"      "genoTypeCount"
## [5] "Genotype_Class"

Seq_cap_tg$Genotype_Class

## [1] "CTNNB1_1" "CTNNB1_2" "EGFR_1"  "EGFR_2"  "EGFR_3"  "IGF2_1"
## [7] "IGF2_2"  "KRAS_1"   "KRAS_2"  "TP53_1"  "TP53_2"
```

Now use the Series100 table that has the blood capillary samples by gene name of all genes with gene names and sequence copy variant of each gene. Combine the tumorigenesis table Seq\_cap\_tg with it.

```
tg_genes_bcs <- merge(Seq_cap_tg, Series100509, by.x='SEQUENCE',
by.y='SEQUENCE')
```

Let's take the row means of each subset of samples in our blood capillary samples of CoV by 0, 12, 24, 36, and 48 hours.

```
colnames(tg_genes_bcs)
```

```
## [1] "SEQUENCE" "Gene.Symbol"
## [3] "geneCount" "genoTypeCount"
## [5] "Genotype_Class" "SPOT_ID"
## [7] "GENE_SYMBOL" "DESCRIPTION"
## [9] "GSM2685693_CoV_0hr_Group_A" "GSM2685694_CoV_0hr_Group_B"
## [11] "GSM2685695_CoV_0hr_Group_C" "GSM2685696_CoV_0hr_Group_D"
## [13] "GSM2685697_CoV_0hr_Group_E" "GSM2685698_ctrl_0hr_Group_A"
## [15] "GSM2685699_ctrl_0hr_Group_B" "GSM2685700_ctrl_0hr_Group_C"
## [17] "GSM2685701_ctrl_0hr_Group_D" "GSM2685702_ctrl_0hr_Group_E"
## [19] "GSM2685703_CoV_12hr_Group_A" "GSM2685704_CoV_12hr_Group_B"
## [21] "GSM2685705_CoV_12hr_Group_C" "GSM2685706_CoV_12hr_Group_D"
## [23] "GSM2685707_CoV_12hr_Group_E" "GSM2685708_ctrl_12hr_Group_A"
## [25] "GSM2685709_ctrl_12hr_Group_B" "GSM2685710_ctrl_12hr_Group_C"
## [27] "GSM2685711_ctrl_12hr_Group_D" "GSM2685712_ctrl_12hr_Group_E"
## [29] "GSM2685713_CoV_24hr_Group_A" "GSM2685714_CoV_24hr_Group_B"
## [31] "GSM2685715_CoV_24hr_Group_C" "GSM2685716_CoV_24hr_Group_D"
## [33] "GSM2685717_CoV_24hr_Group_E" "GSM2685718_ctrl_24hr_Group_A"
## [35] "GSM2685719_ctrl_24hr_Group_B" "GSM2685720_ctrl_24hr_Group_C"
## [37] "GSM2685721_ctrl_24hr_Group_D" "GSM2685722_ctrl_24hr_Group_E"
## [39] "GSM2685723_CoV_36hr_Group_A" "GSM2685724_CoV_36hr_Group_B"
## [41] "GSM2685725_CoV_36hr_Group_C" "GSM2685726_CoV_36hr_Group_D"
## [43] "GSM2685727_CoV_36hr_Group_E" "GSM2685728_ctrl_36hr_Group_A"
## [45] "GSM2685729_ctrl_36hr_Group_B" "GSM2685730_ctrl_36hr_Group_C"
## [47] "GSM2685731_ctrl_36hr_Group_D" "GSM2685732_ctrl_36hr_Group_E"
## [49] "GSM2685733_CoV_48hr_Group_A" "GSM2685734_CoV_48hr_Group_B"
## [51] "GSM2685735_CoV_48hr_Group_C" "GSM2685736_CoV_48hr_Group_D"
## [53] "GSM2685737_CoV_48hr_Group_E" "GSM2685738_ctrl_48hr_Group_A"
## [55] "GSM2685739_ctrl_48hr_Group_B" "GSM2685740_ctrl_48hr_Group_C"
## [57] "GSM2685741_ctrl_48hr_Group_D" "GSM2685742_ctrl_48hr_Group_E"
```

Add in the row means for each group of CoV or Ctrl over 0,12,24,36, or 48 hours.

```
tg_genes_bcs$Mean_0hr_Cov <- rowMeans(tg_genes_bcs[,9:13])
tg_genes_bcs$Mean_12hr_Cov <- rowMeans(tg_genes_bcs[,19:23])
tg_genes_bcs$Mean_24hr_Cov <- rowMeans(tg_genes_bcs[,29:33])
tg_genes_bcs$Mean_36hr_Cov <- rowMeans(tg_genes_bcs[,39:43])
tg_genes_bcs$Mean_48hr_Cov <- rowMeans(tg_genes_bcs[,49:53])

tg_genes_bcs$Mean_0hr_Ctrl <- rowMeans(tg_genes_bcs[,14:18])
tg_genes_bcs$Mean_12hr_Ctrl <- rowMeans(tg_genes_bcs[,24:28])
tg_genes_bcs$Mean_24hr_Ctrl <- rowMeans(tg_genes_bcs[,34:38])
tg_genes_bcs$Mean_36hr_Ctrl <- rowMeans(tg_genes_bcs[,44:48])
tg_genes_bcs$Mean_48hr_Ctrl <- rowMeans(tg_genes_bcs[,54:58])
```

Now, add in the fold change values of the CoV/Ctrl for each of 0,12,24,36, or 48 hours.

```
tg_genes_bcs$FC_0hr <- tg_genes_bcs$Mean_0hr_Cov/tg_genes_bcs$Mean_0hr_Ctrl
tg_genes_bcs$FC_12hr <-
tg_genes_bcs$Mean_12hr_Cov/tg_genes_bcs$Mean_12hr_Ctrl
```

```

tg_genes_bcs$FC_24hr <-
tg_genes_bcs$Mean_24hr_Cov/tg_genes_bcs$Mean_24hr_Ctrl
tg_genes_bcs$FC_36hr <-
tg_genes_bcs$Mean_36hr_Cov/tg_genes_bcs$Mean_36hr_Ctrl
tg_genes_bcs$FC_48hr <-
tg_genes_bcs$Mean_48hr_Cov/tg_genes_bcs$Mean_48hr_Ctrl

```

Lets look at all the columns of sample and statistical information.

```
head(tg_genes_bcs[,1:4])
```

```

##                               SEQUENCE Gene.Symbol
## 1 AGAACTCTGAGTGCATACAGTGCCACCCAGAGTGCCTGCCTCAGGCCATGAACATCACCT      EGFR
## 2 ATGATGGAACATGAGATGGGTGGCCACCACCCTGGTGCTGACTATCCAGTTGATGGGCTG      CTNNB1
## 3 CAGCTACGGTTTCCGTCTGGGCTTCTTGCATTCTGGGACAGCCAAGTCTGTGACTTGCAC      TP53
## 4 CTCAACTCAGCTCCTTTAACGCTAATATTTCCGGCAAAATCCCATGCTTGGGTTTTGTCT      IGF2
## 5 CTGAGTCACACTGCATAGGAATTTAGAACCTAACTTTATAGGTTATCAAACTGTTGTC      KRAS
## 6 CTGTGAGGGATGTTTGGGAGATGTAAGAAATGTTCTTGCAGTTAAGGGTTAGTTTACAAT      TP53
##   geneCount genoTypeCount
## 1         9             3
## 2        22             2
## 3        22             2
## 4         4             2
## 5         4             2
## 6        22            20

```

```
tail(tg_genes_bcs[,69:73])
```

```

##      FC_0hr  FC_12hr  FC_24hr  FC_36hr  FC_48hr
## 6  1.0035557 0.8901734 0.9371630 0.9166376 0.9227073
## 7  1.0037408 0.9708363 0.9652115 1.2515678 1.5938946
## 8  1.0035569 1.0252020 1.0569149 1.0924454 1.1058086
## 9  0.9957301 0.9123630 0.9475070 0.9431493 0.9034454
## 10 1.0015810 0.9618076 0.9400880 0.9736967 0.9659112
## 11 0.9983990 1.0491453 1.0020517 0.9762646 0.9773549

```

Lets use tidyr to gather the fold change columns and mean columns into separate tables to plot.

```

bc_fc <- as.data.frame(gather(tg_genes_bcs,
'TimeInterval_FCs', 'FC_Value', 69:73))
bc_means <- as.data.frame(gather(tg_genes_bcs, 'TimeInterval_Means',
'MeanValue', 59:68))

```

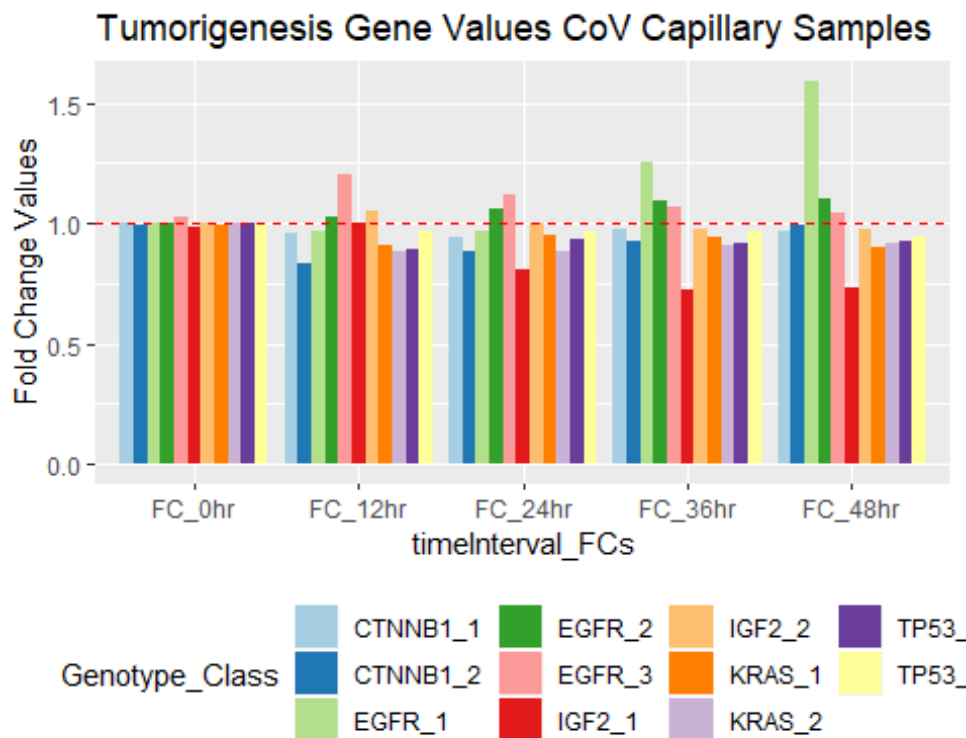
Now, we will plot the blood capillary tumorigenesis genes and their genotypes by fold change (FC) of each time interval and compare to the liver tumor samples of tumorigenesis fold change values.

```

ggplot(data = bc_fc, aes(x=TimeInterval_FCs, y=FC_Value, fill=Genotype_Class))
+
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+

```

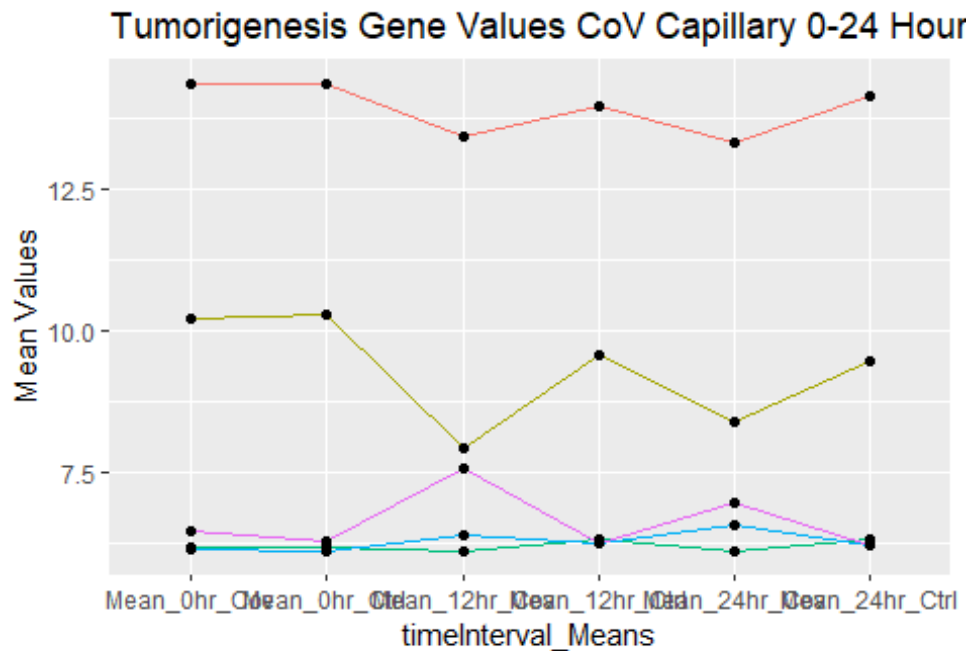
```
scale_fill_brewer(palette='Paired') +
geom_hline(yintercept=1, linetype="dashed", color = "red")+
theme(legend.position="bottom")+
ggtitle('Tumorigenesis Gene Values CoV Capillary Samples')+
ylab('Fold Change Values')
```



```
means_0_24 <- subset(bc_means, bc_means$timeInterval_Means=='Mean_0hr_Cov' |
  bc_means$timeInterval_Means=='Mean_0hr_Ctrl' |
  bc_means$timeInterval_Means=='Mean_12hr_Cov' |
  bc_means$timeInterval_Means=='Mean_12hr_Ctrl' |
  bc_means$timeInterval_Means=='Mean_24hr_Cov' |
  bc_means$timeInterval_Means=='Mean_24hr_Ctrl' )
means_0_24_a <- subset(means_0_24, means_0_24$Genotype_Class=='CTNNB1_1' |
  means_0_24$Genotype_Class=='CTNNB1_2' |
  means_0_24$Genotype_Class=='EGFR_1' |
  means_0_24$Genotype_Class=='EGFR_2' |
  means_0_24$Genotype_Class=='EGFR_3' )
ggplot(data = means_0_24_a, aes(x=timeInterval_Means,
y=MeanValue,group=Genotype_Class)) +
  geom_line(aes(color=Genotype_Class))+
  geom_point()+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
```

```
ggtitle('Tumorigenesis Gene Values CoV Capillary 0-24 Hours')+
ylab('Mean Values')
```

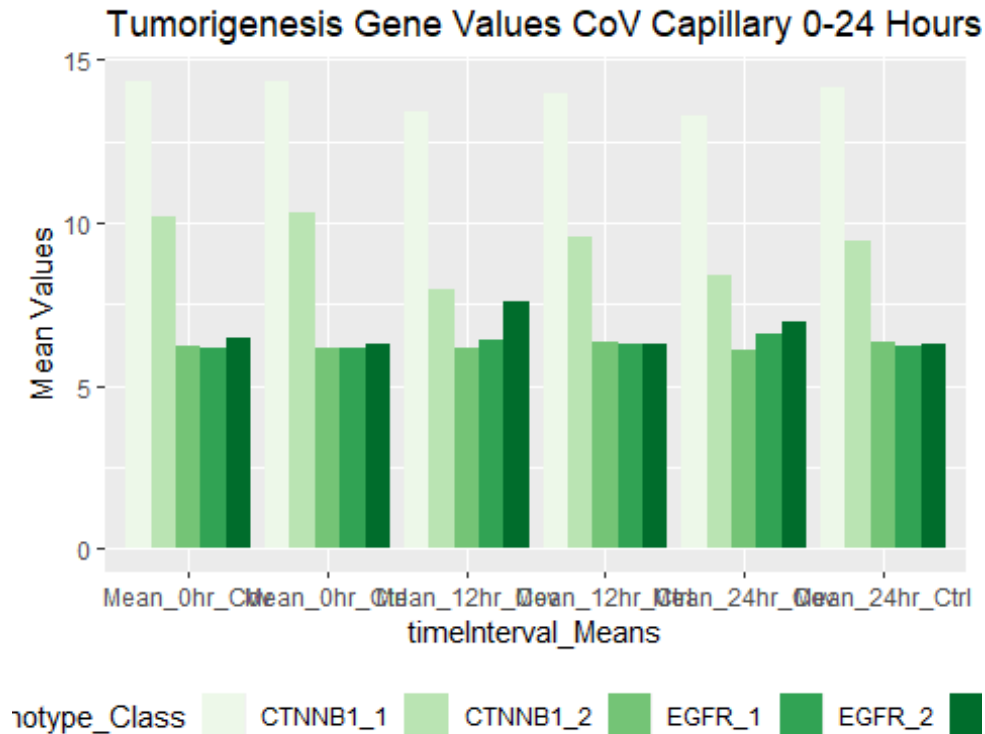
```
## Warning in pal_name(palette, type): Unknown palette paired
```



```
notype_Class CTNNB1_1 CTNNB1_2 EGFR_1 EGFR_2
```

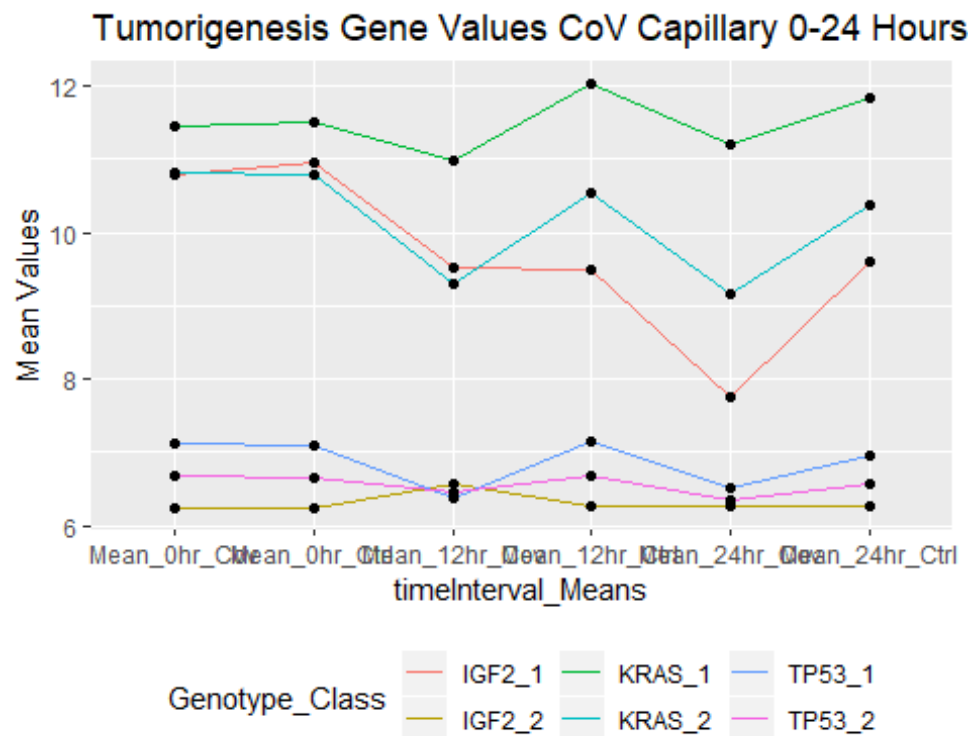
```
ggplot(data = means_0_24_a, aes(x=timeInterval_Means,
y=MeanValue,fill=Genotype_Class)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 0-24 Hours')+
  ylab('Mean Values')
```

```
## Warning in pal_name(palette, type): Unknown palette paired
```



```
means_0_24_b <- subset(means_0_24, means_0_24$Genotype_Class=='IGF2_1' |
  means_0_24$Genotype_Class=='IGF2_2' |
  means_0_24$Genotype_Class=='TP53_1' |
  means_0_24$Genotype_Class=='TP53_2' |
  means_0_24$Genotype_Class=='KRAS_1' |
  means_0_24$Genotype_Class=='KRAS_2')
ggplot(data = means_0_24_b, aes(x=timeInterval_Means,
y=MeanValue,group=Genotype_Class)) +
  geom_line(aes(color=Genotype_Class))+
  geom_point()+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 0-24 Hours')+
  ylab('Mean Values')
```

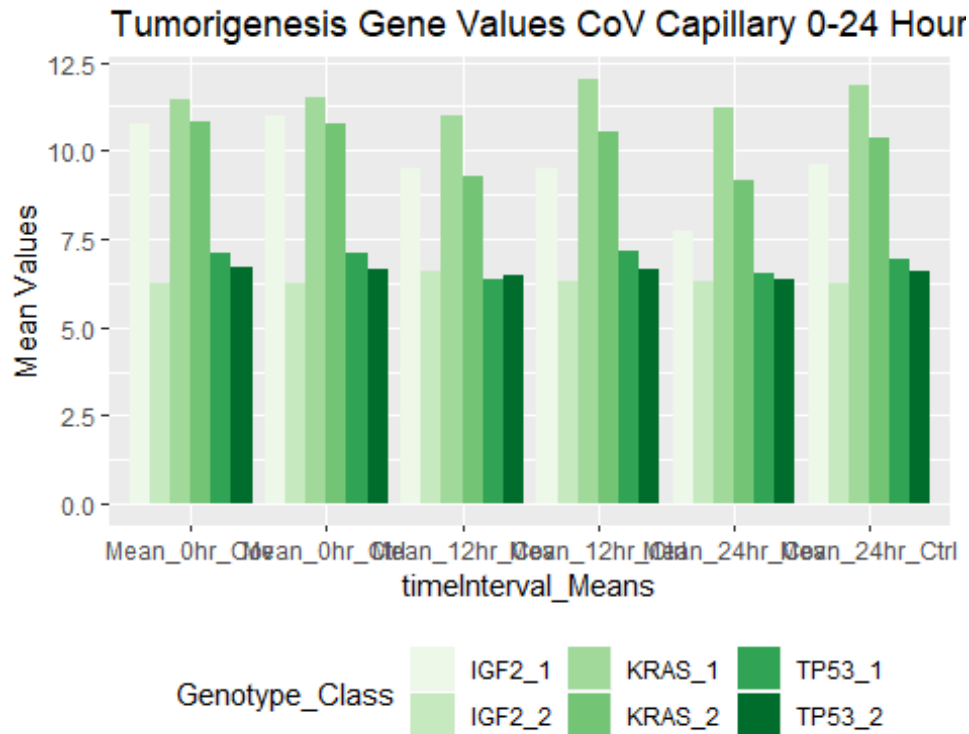
```
## Warning in pal_name(palette, type): Unknown palette paired
```



```
ggplot(data = means_0_24_b, aes(x=timeInterval_Means,
y=MeanValue,fill=Genotype_Class)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 0-24 Hours')+
  ylab('Mean Values')

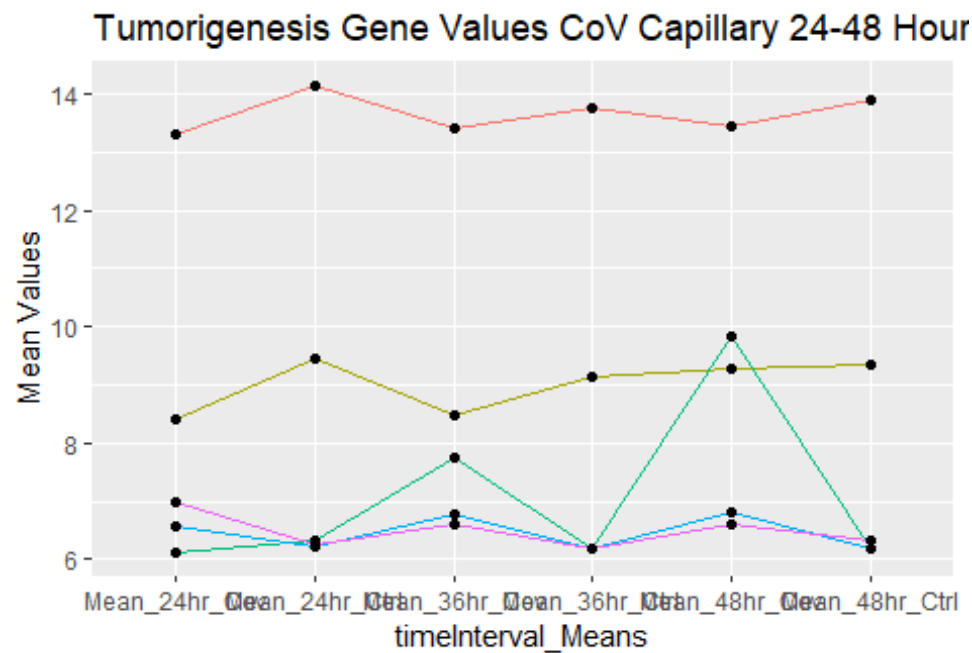
## Warning in pal_name(palette, type): Unknown palette paired
```





Now look at these genes split up into sets of genotypes and in the next 24 hours from time interval 24-48 hours.

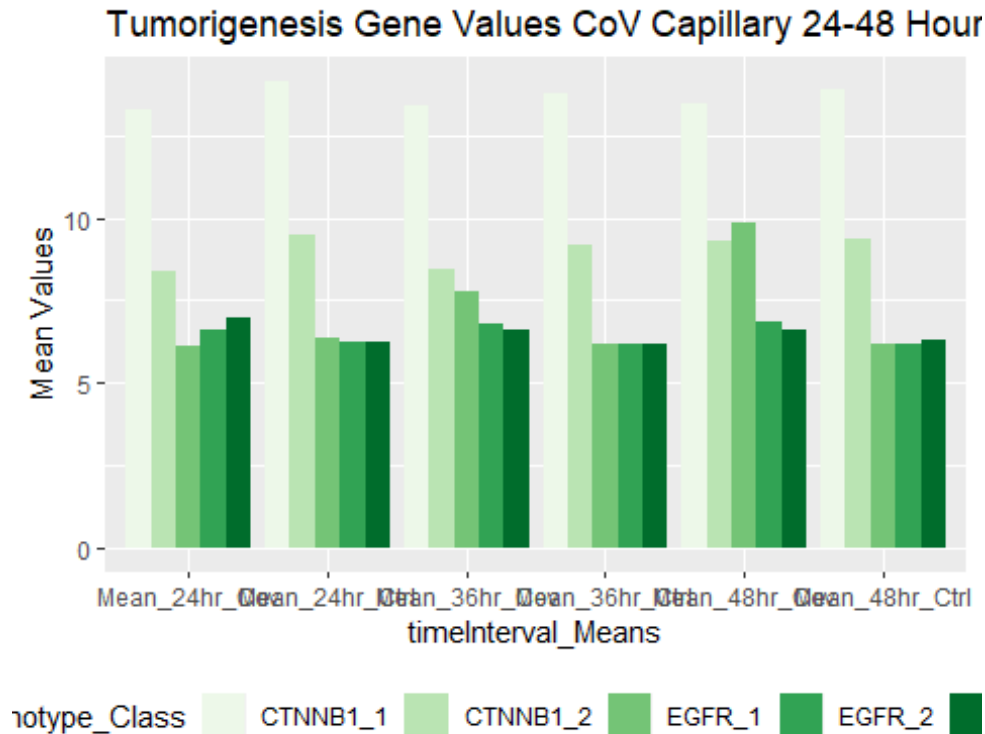
```
means_24_48 <- subset(bc_means, bc_means$timeInterval_Means=='Mean_24hr_Cov' |
  bc_means$timeInterval_Means=='Mean_24hr_Ctrl' |
  bc_means$timeInterval_Means=='Mean_36hr_Cov' |
  bc_means$timeInterval_Means=='Mean_36hr_Ctrl' |
  bc_means$timeInterval_Means=='Mean_48hr_Cov' |
  bc_means$timeInterval_Means=='Mean_48hr_Ctrl' )
means_24_48_a <- subset(means_24_48, means_24_48$Genotype_Class=='CTNNB1_1' |
  means_24_48$Genotype_Class=='CTNNB1_2' |
  means_24_48$Genotype_Class=='EGFR_1' |
  means_24_48$Genotype_Class=='EGFR_2' |
  means_24_48$Genotype_Class=='EGFR_3' )
ggplot(data = means_24_48_a, aes(x=timeInterval_Means,
y=MeanValue,group=Genotype_Class)) +
  geom_line(aes(color=Genotype_Class))+
  geom_point()+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 24-48 Hours')+
  ylab('Mean Values')
## Warning in pal_name(palette, type): Unknown palette paired
```



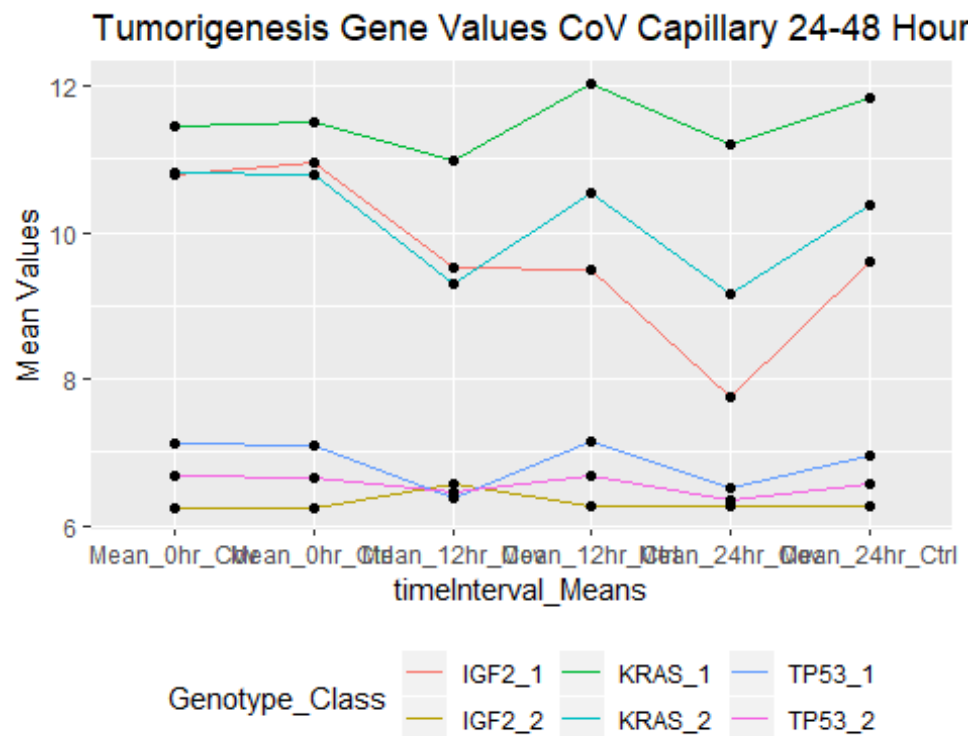
Genotype\_Class — CTNNB1\_1 — CTNNB1\_2 — EGFR\_1 — EGFR\_2 —

```
ggplot(data = means_24_48_a, aes(x=timeInterval_Means,
y=MeanValue,fill=Genotype_Class)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 24-48 Hours')+
  ylab('Mean Values')

## Warning in pal_name(palette, type): Unknown palette paired
```

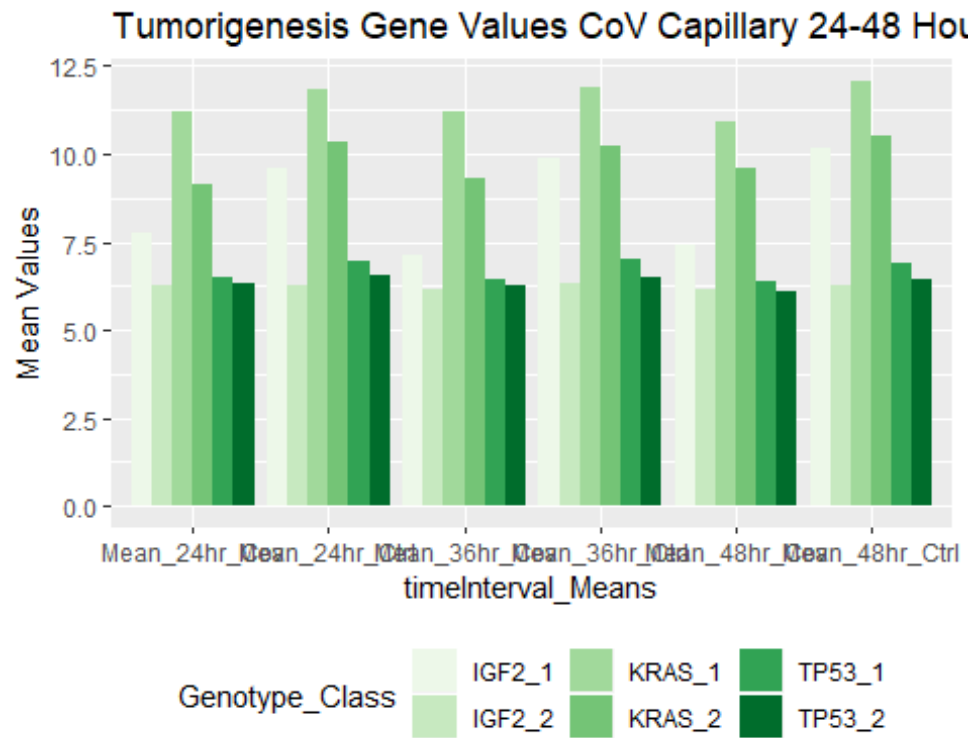


```
means_24_48_b <- subset(means_24_48, means_24_48$Genotype_Class=='IGF2_1' |
  means_24_48$Genotype_Class=='IGF2_2' |
  means_24_48$Genotype_Class=='TP53_1' |
  means_24_48$Genotype_Class=='TP53_2' |
  means_24_48$Genotype_Class=='KRAS_1' |
  means_24_48$Genotype_Class=='KRAS_2')
ggplot(data = means_0_24_b, aes(x=timeInterval_Means,
y=MeanValue,group=Genotype_Class)) +
  geom_line(aes(color=Genotype_Class))+
  geom_point()+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 24-48 Hours')+
  ylab('Mean Values')
## Warning in pal_name(palette, type): Unknown palette paired
```



```
ggplot(data = means_24_48_b, aes(x=timeInterval_Means,
y=MeanValue,fill=Genotype_Class)) +
  geom_bar(stat='identity', position=position_dodge())+
  scale_y_continuous()+
  scale_fill_brewer(palette="paired") +
  theme(legend.position="bottom")+
  ggtitle('Tumorigenesis Gene Values CoV Capillary 24-48 Hours')+
  ylab('Mean Values')

## Warning in pal_name(palette, type): Unknown palette paired
```



**updated**

**2/16/2020, analysis to follow as well as the diabetes type 1 and type 2 comparisons**