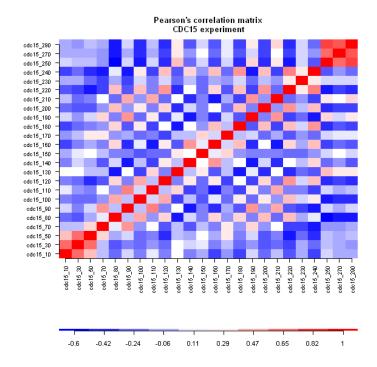
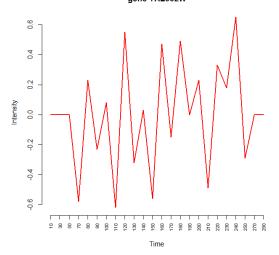
```
Lab #2
Data visualization
Solutions
2.)
dat <- read.table("C:\\Users\\higgsb\\Dropbox\\JHU\\Microarrays\\Data\\
spellman.txt",header=T,row.names=1)
3) dim(dat)
[1] 6178 77
4.)
> dat <- dat[,23:46]
> names(dat)
[1] "cdc15 10" "cdc15 30" "cdc15 50" "cdc15 70" "cdc15 80" "cdc15 90"
[7] "cdc15 100" "cdc15 110" "cdc15 120" "cdc15 130" "cdc15 140" "cdc15 150"
[13] "cdc15 160" "cdc15 170" "cdc15 180" "cdc15 190" "cdc15 200" "cdc15 210"
[19] "cdc15 220" "cdc15 230" "cdc15 240" "cdc15 250" "cdc15 270" "cdc15 290"
5.)
library(gplots)
dat.cor <- cor(dat,use="pairwise.complete.obs")
layout(matrix(c(1,1,1,1,1,1,1,2,2), 5, 2, byrow = TRUE))
par(oma=c(5,7,1,1))
cx <- rev(colorpanel(25,"red","white","blue"))
leg <- seq(min(dat.cor,na.rm=T),max(dat.cor,na.rm=T),length=10)
image(dat.cor,main="Pearson's correlation matrix\nCDC15 experiment",axes=F,col=cx)
axis(1.at=seq(0.1.length=ncol(dat.cor)),label=dimnames(dat.cor)[[2]],cex.axis=0.9.las=2)
axis(2,at=seq(0,1,length=ncol(dat.cor)),label=dimnames(dat.cor)[[2]],cex.axis=0.9,las=2)
image(as.matrix(leg),col=cx,axes=F)
tmp <- round(leg,2)
axis(1,at=seq(0,1,length=length(leg)),labels=tmp,cex.axis=1)
```



- 6.)
  > dat.m <- mean(as.numeric(as.matrix(dat["YAL002W",])), na.rm=T)
  > dat.m
  [1] 3.857953e-18
  > dat["YAL002W",is.na(dat["YAL002W",])] <- dat.m
- 7.) plot(as.numeric(dat["YAL002W",]),type='l',lwd=2,col='red', main="Spellman data set (CDC15 experiment)\ngene YAL002W",xlab="Time", ylab="Intensity",axes=F) axis(1,at=c(1:ncol(dat)),labels=sub("cdc15\_","",names(dat)),las=2,cex.axis=0.7) axis(2)

## Spellman data set (CDC15 experiment) gene YAL002W



```
8.)
library(shiny)
server <- function(input, output) {</pre>
         # Combine the selected variables into a new data frame
         selectedData <- reactive({</pre>
               dat[, c(input$xcol, input$ycol)]
         })
        output$plot1 <- renderPlot({
               par(mar = c(5.1, 4.1, 0, 1))
               plot(selectedData(), col=1, bg = input$colx, pch = 21, cex = 0.5)
         })
       fluidPage(
ui <-
               sidebarLayout(
               sidebarPanel(
               selectInput('xcol', 'X Variable', dimnames(dat)[[2]]),
               selectInput('ycol', 'Y Variable', dimnames(dat)[[2]],
                                       selected=dimnames(dat)[[2]][1]),
               selectInput('colx', 'Point color', c("red", "blue", "green", "black", "orange"))
         ),
         mainPanel(
               plotOutput('plot1')
       )
)
```

```
shinyApp(ui = ui, server = server)
```

## Appendix

(alternative means to obtain yeast cell cycle dastaset using R built-in datasets)

- > library(Biobase)
- > library(annotate)
- > library(yeastCC)
- > data(yeastCC)
- > dat <- exprs(yeastCC) # expression data