Lab 7 | Jon Lee

Code ▼

Lab 7 | BINF 6310 | Spring 2020 | Jon Lee

1. Consider a tournament with the following prize structure:

Number of Wins Payoff (\$) 0 1.45 1 1.72 2 2.24 3 2.76 4 3.55 5 4.60 6 5.65 7 6.75

You play in the tournament until you get three losses or 7 wins.

Make a graph of the expected value of the tournament (on the y-axis) vs. the probability of winning each game (on the x-axis). If the tournament costs \$3.75, at what win percentage does the expected value exceed the cost of the tournament? (Plot 3.75 as a horizontal line ranging from 0 to 1).

(hint: Calculate the probability of 7 wins a 1 – sum(prob of all other outcomes)).

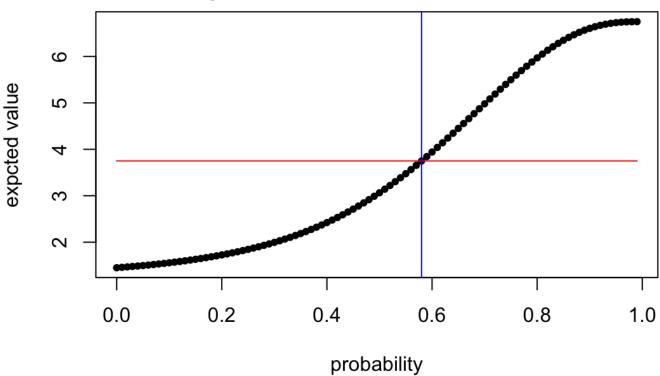
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```
#graph of expected values for tourney
#assuming the probability of a win is 0.5
entryPrice <- 3.75
numberOfWins <- 0:7</pre>
payoffPerWin <- c(1.45, 1.72, 2.24, 2.76, 3.55, 4.60, 5.65, 6.75)
probOfWin < - seq(0, 0.99, 0.01)
#did not include 1 becuase the dnbinom function's probability is not defined at a probab
ility of 0
expectVals <- vector()
for(i in 1:length(probOfWin))
 probs <- vector()</pre>
  expect <- vector()</pre>
  for(j in 1:(length(payoffPerWin)-1))
    probs[j] \leftarrow dnbinom((j-1), 3, (1-probOfWin[i]))
    expect[j] <- probs[j]*payoffPerWin[j]</pre>
  }
 probs[8] <- 1-sum(probs[1:7])</pre>
  expect[8] <- probs[8]*payoffPerWin[8]</pre>
  expectVals[i] <- sum(expect)</pre>
}
plot(probOfWin, expectVals, xlab = "probability", ylab = "expcted value", main = "expect
ed value of the tournament", pch = 20)
lines(probOfWin, rep(3.75, length(probOfWin)), col = "red")
```

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```
abline(v = 0.58, col= "blue")
```

## expected value of the tournament



Answer: At a win percentage (rate) of about 0.58 or 58% do we see the epected value of the tournament exceeding the cost.

Collabroated with David B. and Conor F. on Question 1

2. Generate a simulated dataset in which the variance is a function of the mean but the null hypothesis of differential expression is always true. The following code will accomplish this:

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```
rm(list=ls())
numRows = 3000
numCols = 20
for( i in 1:numCols)
myFrame <- data.frame(1:numRows)</pre>
#initiate the data.frame with the correct # of rows to suppress error messages.
#likely, there are much better ways to do this!
names(myFrame)[1] <- "tempColumn"</pre>
for( i in 1: numCols)
        vals <- vector(length=numRows)</pre>
        for( j in 1:numRows)
             aMean = j /10
             aMean = max(aMean, 5)
             aVar = aMean + 5* aMean
             aVal = round( max( rnorm(1, mean=aMean, sd=sqrt(aVar)), 1))
             vals[j] = aVal
        }
        colName <- paste( "sample" , i ,sep="")</pre>
        myFrame[[colName]] = vals
}
myFrame["tempColumn"] <- NULL</pre>
row.names(myFrame) <- paste("Gene ",1:numRows,sep="")</pre>
```

For each row in the spreadsheet, we will consider the first 10 columns to be "case" and the last 10 to be "control". For each row in the spreadsheet, run a t-test and generate a p-value for the null hypothesis that the case and control samples follow the same distribution (see slide 5 of lecture 11 for code to do this). Show your code for all answers.

```
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allPValsTTest <- vector()

for(i in 1:nrow(myFrame))
{
   vals1 <- as.numeric(myFrame[i,1:10])
   vals2 <- as.numeric(myFrame[i,11:20])
   allPValsTTest[i] <- t.test(vals1, vals2)$p.value
}</pre>
```

```
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```

```
hist(allPValsTTest, xlab = "t-test p values", main = "Histogram of t-test")
```

A. Use a simple threshold of p <0.05. How many significant hits would you expect to find if the null hypothesis is always true? How many hits did you actually find in your run?

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```
#how many expected hits
totalNumOfPVals <- length(allPValsTTest)
expectNumoOfPVals <- 0.05*totalNumOfPVals
print(expectNumoOfPVals)

#how many actual hits
countPVals <- 0
for(i in 1:length(allPValsTTest))
{
   if(allPValsTTest[i] < 0.05)
   {
     countPVals = countPVals + 1
   }
}
print(countPVals)</pre>
```

Answer: If the null hypothesis is true, we would expect to see 5% of the p-values to be significant, because the resulting p-values should be uniform. From the data we saw 156 p-values were significant, which is approximately 50, which confirms what we expected.

B. What is the Bonferroni adjusted p-value threshold? What percentage of the time would you expect to see a significant gene under this threshold? How many genes in fact do you see significant at this threshold?

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```
#Bonferroni adjusted p-value threshold
adjustedPVal <- 0.05/totalNumOfPVals
print(adjustedPVal)

#how many expected hits
adjustedExpectedPVals <- adjustedPVal*totalNumOfPVals
print(adjustedExpectedPVals)

#how many actual hits
countAdjustPVal <- 0
for(i in 1:length(allPValsTTest))
{
   if(allPValsTTest[i] < adjustedPVal)
   {
     countAdjustPVal = countAdjustPVal + 1
   }
}
print(countAdjustPVal)</pre>
```

Answer: Our new adjusted p-value threshold via Bonferroni is 1.67e-05, and we would expect to see none of our p-values to be below this threshold, and when looking at the data we see that this is true.

C. Next use a BH FDR corrected threshold of p < 0.05 (which R can do for you with p.adjust(pvals,method="BH") if pvals is a vector that holds the unadjusted p-values). How many hits do you

find with the BH adjusted p-values.

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```
#adjust p-values via BH method
adjustedPVals <- p.adjust(allPValsTTest, method = "BH")

#how many actual hits
countAdjustPVals <- 0
for(i in 1:length(allPValsTTest))
{
   if(allPValsTTest[i] < adjustedPVal)
   {
      countAdjustPVals = countAdjustPVals + 1
   }
}
print(countAdjustPVals)</pre>
```

Answer: After adjusting our p-values using the BH method, we find that none of our p-value lie beyondthe 005 threshold and therefore are significant.