STATS 201 Assignment 2

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Due Date: 2021/11/7

Loading required package: s20x

Question 1

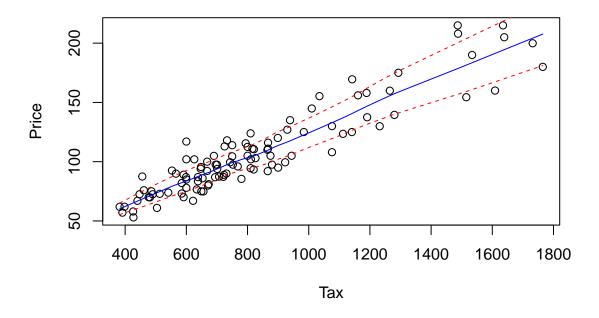
Question of interest/goal of the study

We want to build a model to explain the sale price of houses using their annual city tax bill (similar idea to rates in New Zealand) for houses in Albuquerque, New Mexico. In particular, we are interested in estimating the effect on sales price for houses which differ in city tax bills by 1% and 50%.

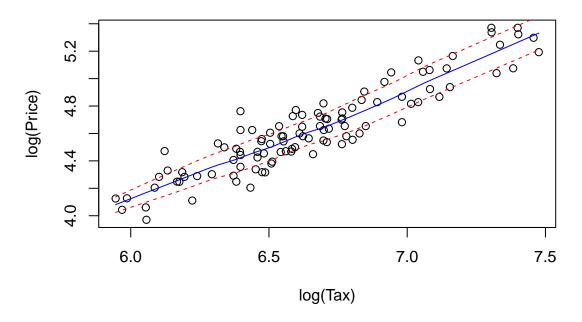
Read in and inspect the data:

```
hometax.df=read.csv("hometax.csv")
trendscatter(Price~Tax,main="Price vs Tax",data=hometax.df)
```

Price vs Tax



log(Price) vs log(Tax)



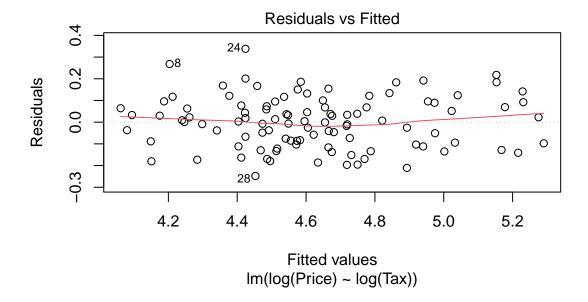
For initial plot, we can see the data is right-skewed. Because we find that some variables which have large explain variables have large response variables, it has characteristics of power law model.

Justify why a log-log (power) model is appropriate here.

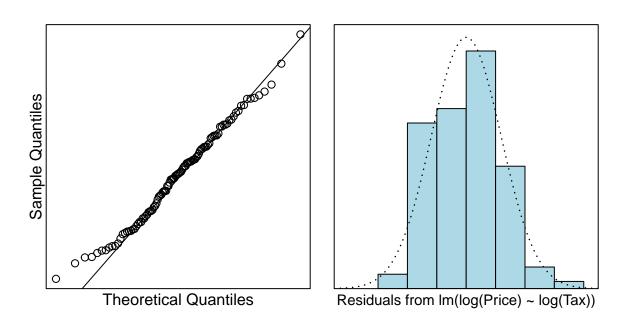
From the plot Price vs Tax, we could find that some points have a large explain variable value while they also have a large response variable value. It leads to the whole data set presents a right-skewed trend. It matches the characteristics of power law model. And when we add log() to each of the variables, the trendscatter plot seems to be perfect to fit using linear model. So we consider adopt power law model to fit this data set.

Fit model and check assumptions.

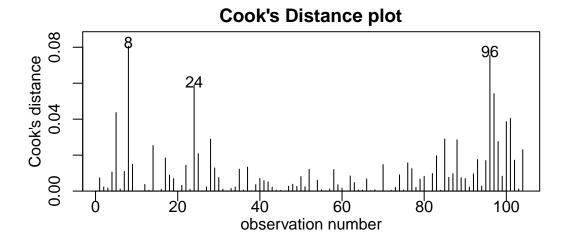
```
Price.lm = lm(log(Price) ~ log(Tax), data = hometax.df)
plot(Price.lm, which = 1)
```



normcheck(Price.lm)



cooks20x(Price.lm)



```
##
## Call:
## lm(formula = log(Price) ~ log(Tax), data = hometax.df)
##
## Residuals:
##
       Min
                  1Q
                       Median
                                     3Q
                                             Max
   -0.24820 -0.09519
                      0.00380
                               0.07994
##
##
##
  Coefficients:
##
               Estimate Std. Error t value Pr(>|t|)
   (Intercept) -0.71348
                           0.21679
                                    -3.291
                                            0.00137 **
                0.80311
                           0.03257
                                    24.660
                                            < 2e-16 ***
##
  log(Tax)
                     '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
## Signif. codes:
## Residual standard error: 0.1194 on 102 degrees of freedom
## Multiple R-squared: 0.8564, Adjusted R-squared: 0.855
## F-statistic: 608.1 on 1 and 102 DF, p-value: < 2.2e-16
confint(Price.lm)
                    2.5 %
##
                              97.5 %
## (Intercept) -1.1434829 -0.2834689
```

Methods and assumption checks

0.7385139 0.8677080

log(Tax)

summary(Price.lm)

We used the power law model to find out the relationship between sale price of houses and their annual city tax bill because we found that the data is right-skewed distributed. It has clarified that data was collected

from a random sample of 104 houses sold in Albuquerque. So there was no worries about the independence. The residual plot showed a patternless scatter with quite constant variability. The normcheck did not find serious problem. And there was no influential points. In conclusion, our model satisfies the problem.

Our model is:

$$log(Price_i) = \beta_0 + \beta_1 log(Tax_i) + \epsilon_i$$
 where $\epsilon \sim iid\ N(0, \sigma^2)$

Our model explains 86% response variables, which means it is resonable for prediction.

Executive Summary

In order to specify the relation ship between sale price of houses and their annual city tax bill, we used a power law model to fit this data set. We have strong evidence that there exists some relationship between Tax and Price. And we estimate that a 1% increase in the Tax results in a 0.74% to a 0.87% increase in the median value of Price.

Our model explains 86% data in the data set. So it is reliable to predict using this model.

Question 2

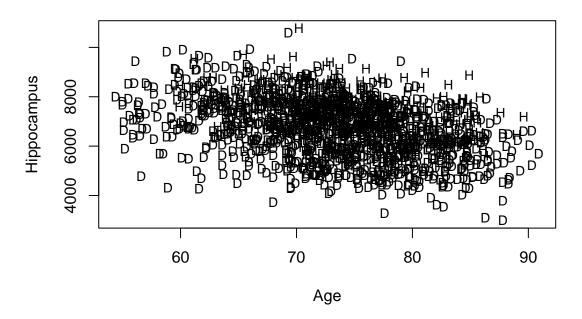
Question of interest/goal of the study

We want to explore the relationship between hippocampus size and age. In particular, we are interested in whether the relationship differs between healthy individuals and individuals with dementia related symptoms.

Read in and inspect the data:

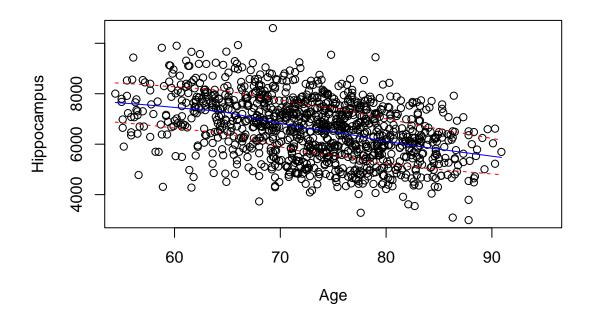
```
Hippocampus.df<-read.csv("Hippocampus.csv")
plot(Hippocampus~Age,main="Hippocampus Size versus Age",type="n",data=Hippocampus.df)
text(Hippocampus.df$Age, Hippocampus.df$Hippocampus, Hippocampus.df$AD, cex=.8)</pre>
```

Hippocampus Size versus Age

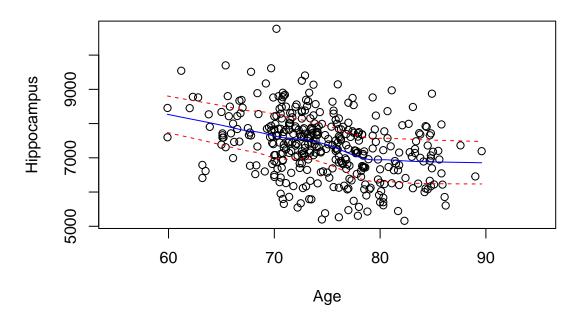


trendscatter(Hippocampus~Age, data=Hippocampus.df[Hippocampus.df\$AD=="D",],xlim=c(55,95),main="Dementia"

Dementia



Healthy



Generally speaking, the Hippocampus volume is decreasing with the increase of Age for both dementia and healthy volunteers. By comparing the two plots, we could find the average Hippocampus volume is quite different between these two groups. Healthy volunteers have larger Hippocampus volume than Dementia volunteers. And it seems that the decreasing speeds for these two groups are different.

Fit model and check assumptions.

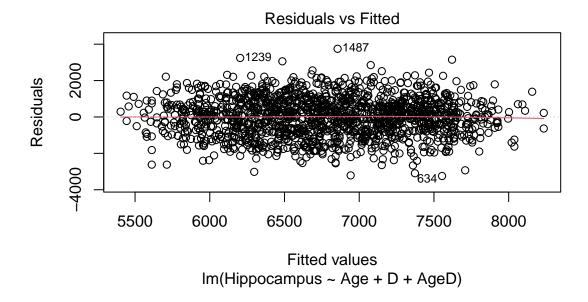
We define dummy variable D to represent healthy volunteers when D=1 while baseline represents those volunteers who are dementia.

```
Hippocampus.df$D = as.numeric(Hippocampus.df$AD == "H")
table(Hippocampus.df$AD, Hippocampus.df$D)
```

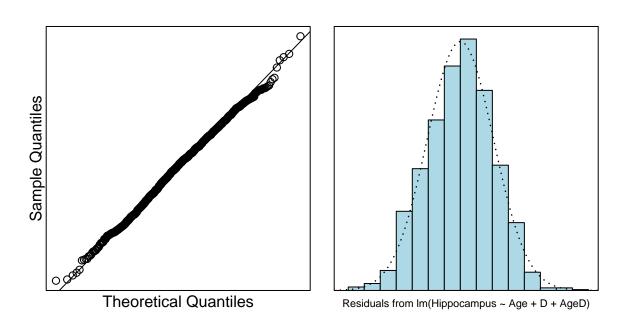
```
##
## 0 1
## D 1116 0
## H 0 373

Hippocampus.df$AgeD = with(Hippocampus.df, {AgeD = D * Age})
AgeHippo.fit = lm(Hippocampus ~ Age + D + AgeD, data = Hippocampus.df)

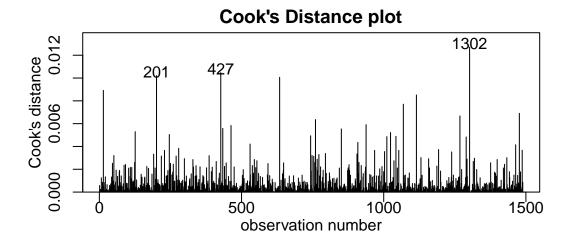
plot(AgeHippo.fit, which = 1)
```



normcheck(AgeHippo.fit)



cooks20x(AgeHippo.fit)



```
##
## lm(formula = Hippocampus ~ Age + D + AgeD, data = Hippocampus.df)
##
## Residuals:
##
       Min
                1Q Median
                                3Q
                                       Max
                             701.9 3746.6
## -3245.4 -729.8
                      52.1
##
## Coefficients:
##
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 11513.168
                            303.741 37.905
                                              <2e-16 ***
                 -67.212
                              4.132 -16.266
                                              <2e-16 ***
## Age
## D
                 291.487
                            787.293
                                     0.370
                                               0.711
## AgeD
                   7.617
                             10.546
                                      0.722
                                               0.470
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
```

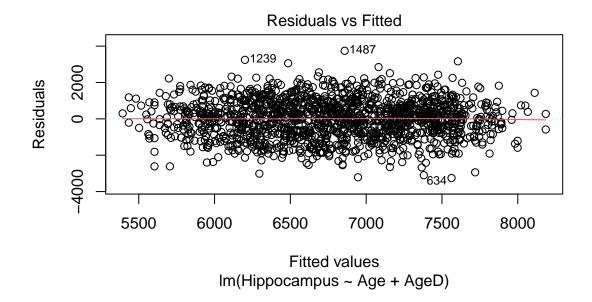
summary(AgeHippo.fit)

However, some coefficients seems may equal to 0. Let us drop some of the variables in the model.

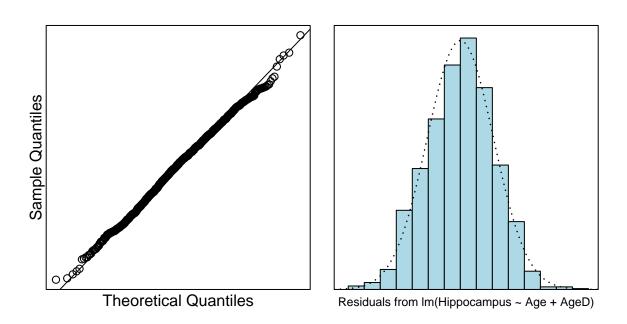
Fit model and check assumptions AGAIN (DROP D).

Residual standard error: 1039 on 1485 degrees of freedom
Multiple R-squared: 0.2328, Adjusted R-squared: 0.2313
F-statistic: 150.2 on 3 and 1485 DF, p-value: < 2.2e-16</pre>

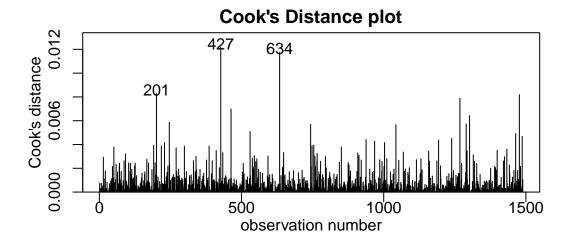
```
AgeHippo.fit2 = lm(Hippocampus ~ Age + AgeD, data = Hippocampus.df)
plot(AgeHippo.fit2, which = 1)
```



normcheck(AgeHippo.fit2)



cooks20x(AgeHippo.fit2)



summary(AgeHippo.fit2)

```
##
## Call:
## lm(formula = Hippocampus ~ Age + AgeD, data = Hippocampus.df)
## Residuals:
##
      Min
               1Q Median
                               3Q
                                       Max
                            709.7 3743.9
## -3254.2 -733.8
                     55.0
##
## Coefficients:
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 11556.5541
                           280.1440
                                       41.25
                                              <2e-16 ***
## Age
                -67.7991
                              3.8146 -17.77
                                               <2e-16 ***
## AgeD
                 11.5088
                              0.8359
                                       13.77
                                               <2e-16 ***
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## Residual standard error: 1039 on 1486 degrees of freedom
## Multiple R-squared: 0.2327, Adjusted R-squared: 0.2317
## F-statistic: 225.4 on 2 and 1486 DF, p-value: < 2.2e-16
```

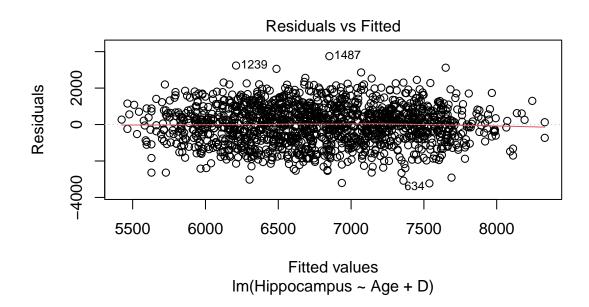
confint(AgeHippo.fit2)

```
## 2.5 % 97.5 %
## (Intercept) 11007.034397 12106.07381
## Age -75.281680 -60.31645
## AgeD 9.869105 13.14855
```

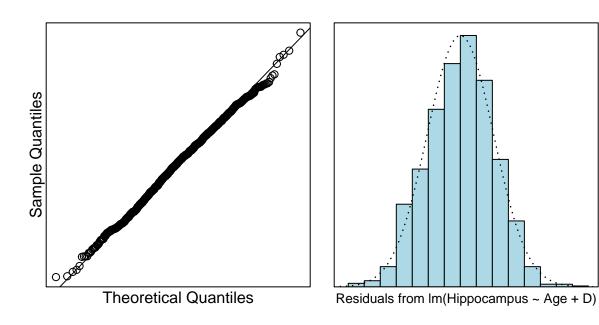
Fit model and check assumptions AGAIN AND AGAIN (DROP AGED).

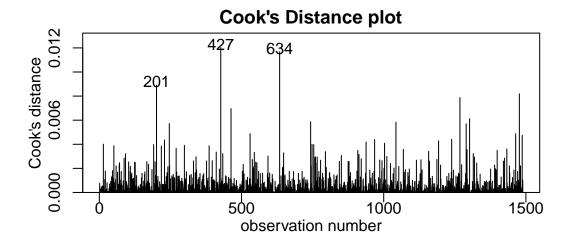
```
AgeHippo.fit3 = lm(Hippocampus ~ Age + D, data = Hippocampus.df)

plot(AgeHippo.fit3, which = 1)
```



normcheck(AgeHippo.fit3)



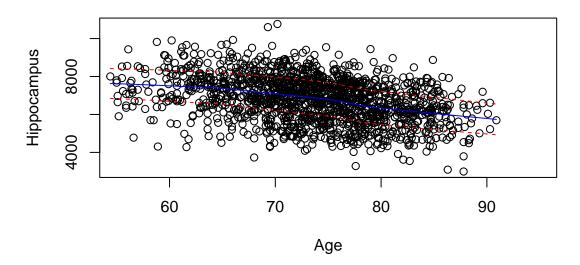


summary(AgeHippo.fit3)

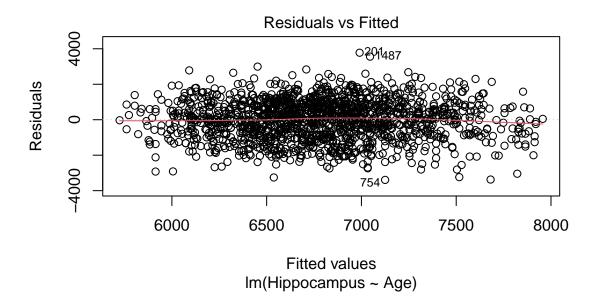
```
##
## Call:
## lm(formula = Hippocampus ~ Age + D, data = Hippocampus.df)
##
## Residuals:
##
       Min
                1Q Median
                               3Q
                                      Max
## -3228.8 -727.2
                     54.5
                            705.0 3751.1
##
## Coefficients:
               Estimate Std. Error t value Pr(>|t|)
## (Intercept) 11427.664
                            279.674
                                    40.86
                                             <2e-16 ***
## Age
                -66.043
                             3.801 -17.37
                                             <2e-16 ***
## D
                 858.307
                            62.413
                                     13.75
                                             <2e-16 ***
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## Residual standard error: 1039 on 1486 degrees of freedom
## Multiple R-squared: 0.2325, Adjusted R-squared: 0.2315
## F-statistic: 225.1 on 2 and 1486 DF, p-value: < 2.2e-16
```

Fit model and check assumptions AGAIN AND AGAIN AND AGAIN (DROP D AND AGED).

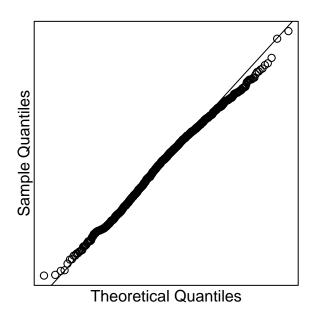
Plot of Hippocampus vs. Age (lowess+/-sd)

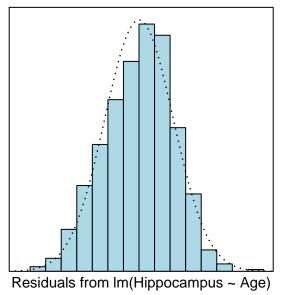


```
AgeHippo.fit4 = lm(Hippocampus ~ Age, data = Hippocampus.df)
plot(AgeHippo.fit4, which = 1)
```

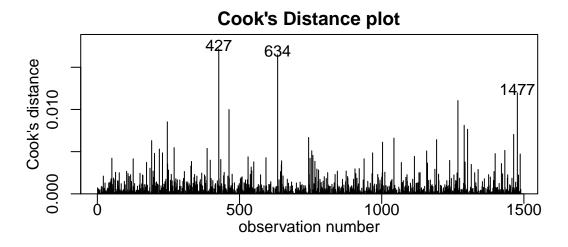


normcheck(AgeHippo.fit4)





cooks20x(AgeHippo.fit4)



summary(AgeHippo.fit4)

Call:

```
## lm(formula = Hippocampus ~ Age, data = Hippocampus.df)
##
## Residuals:
##
      Min
               1Q Median
                               ЗQ
                                     Max
## -3393.9 -769.2
                          788.1 3778.7
                     76.1
##
## Coefficients:
               Estimate Std. Error t value Pr(>|t|)
##
## (Intercept) 11283.689
                          296.630 38.04 <2e-16 ***
                           4.017 -15.23 <2e-16 ***
## Age
                -61.159
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
## Residual standard error: 1103 on 1487 degrees of freedom
## Multiple R-squared: 0.1349, Adjusted R-squared: 0.1343
## F-statistic: 231.8 on 1 and 1487 DF, p-value: < 2.2e-16
```

Plot the data with your appropriate model superimposed over it

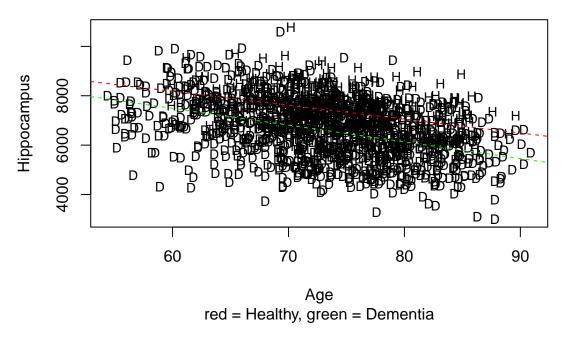
```
plot(Hippocampus~Age,main="Hippocampus Size versus Age",sub="red = Healthy, green = Dementia",type="n",text(Hippocampus.df$Age, Hippocampus.df$Hippocampus.df$AD, cex=.8)

b = coef(AgeHippo.fit2)
b

## (Intercept) Age AgeD
## 11556.55410 -67.79906 11.50883

abline(b[1:2], lty = 2, col = "green")
abline(b[1], b[2] + b[3], lty = 2, col = "red")
```

Hippocampus Size versus Age



We finally choose AgeHippo.fit2 as our fit model.

Methods and assumption checks

In order to study the relationship between hippocampus size, age and whether the relationship differs between healthy individuals and individuals with dementia related symptoms, we used linear model with one dummy variable to fit the data set. We find that Hippocampus volume is decreasing as the increase of Age. The average volumes of Healthy volunteers' Hippocampus are much larger than those who Dementia. And Dementia volunteers' decreasing speed is faster than Healthy volunteers. The main problem we encountered is when we used Hippocampus ~ Age + D + AgeD to fit the data set, we found some coefficients with dummy variable D could equal to 0. Then we are trying to remove some variable in our model to make all the coefficients have evidence existing. After removing D, AgeD and both D and AgeD, we found the plan which removes D variable can get the highest Multiple R-squared value. However, compared to not removing variable D, the Multiple R-squared value decreased about 0.01%, this does not affect the correctness of our model because dropping one useless variable makes our model more general. So we choose this linear model to fit. As the description of the question has stated that we can treat the data as if it came from random samples of subjects. So we do not have worries about independence. The residual plot showed a patternless scatter with quite constant variability. The normcheck did not find serious problem. And there was no influential points. In conclusion, our model can explain the data set.

Our model is:

$$Hippocampus_i = \beta_0 + \beta_1 \times Age_i + \beta_2 \times D_i \times Age_i + \epsilon_i$$

where $D_i = 1$ if the *i*th subject is healthy and 0 if they have signs of dementia, and $\epsilon_i \sim iid \ N(0, \sigma^2)$ Our model explains 23.27% data in the initial data set.

Executive Summary

We are going to study the relationship between hippocampus size, age and whether the relationship differs between healthy individuals and individuals with dementia related symptoms. We use linear model with numerical explain variable and categorical variable and we have extremely strong evidence that Hippocampus volume is decreasing as the increase of Age, the average volumes of Healthy volunteers' Hippocampus are much larger than those who Dementia, and Dementia volunteers' decreasing speed is faster than Healthy volunteers.

We estimate that for Dementia volunteers, their Hippocampus volumes decrease about $60 \sim 75$ units per year on average. For Healthy volunteers, their Hippocampus volumes decreasing speed could decrease $10 \sim 13$ units compared with Dementia volunteers per year on average.

Our model only explains about 23.27% variability in the initial data set, which means this model may not be very good for prediction. We estimate the reason may be the too scattered data points.

Question 3

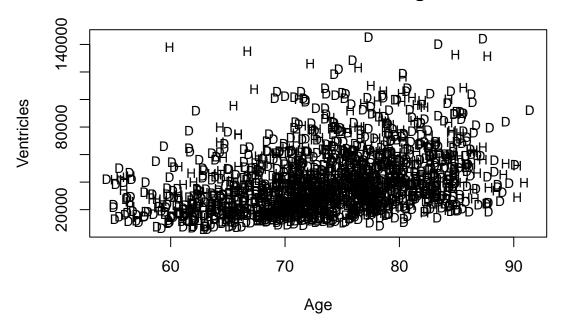
Question of interest/goal of the study

It is of interest to study the relationship between ventricles and age. In particular, we are interested in whether the relationship varies between healthy individuals and individuals with dementia related symptoms.

Read in and inspect the data:

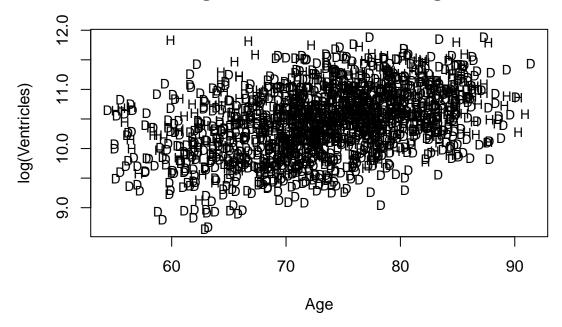
```
Ventricles.df=read.csv("Ventricles.csv")
plot(Ventricles~Age,main="Ventricles Size versus Age",type="n",data=Ventricles.df)
text(Ventricles.df$Age, Ventricles.df$Ventricles, Ventricles.df$AD, cex=.8)
```

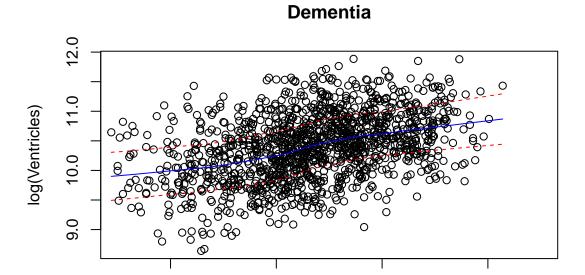
Ventricles Size versus Age



plot(log(Ventricles)~Age,main="log Ventricles Size versus Age",type="n",data=Ventricles.df) text(Ventricles.df\$Age, log(Ventricles.df\$Ventricles), Ventricles.df\$AD, cex=.8)

log Ventricles Size versus Age

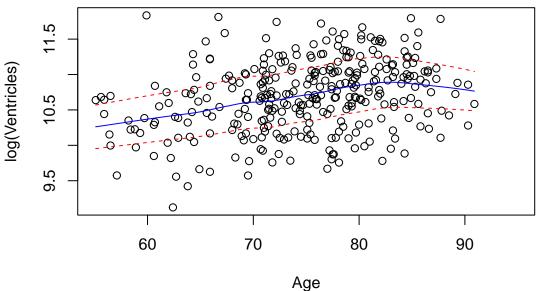




trendscatter(log(Ventricles)~Age,data=Ventricles.df[Ventricles.df\$AD=="H",],xlim=c(55,95),main="Healthy

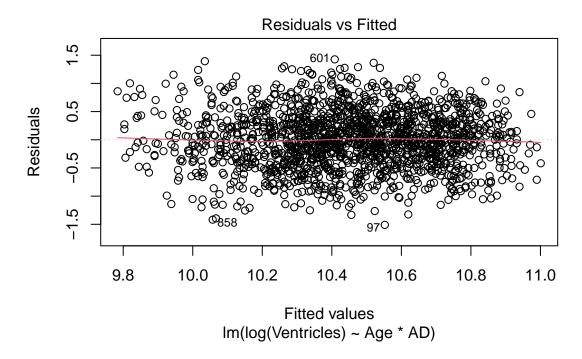


Age

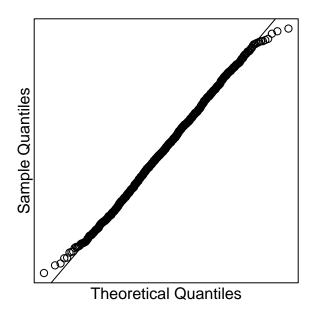


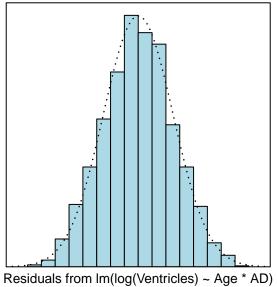
In the trendscatter plots above, we could find that the initial data presents a trend that right-skewed. So we can use multiplicative model to fit this model. After transforming response variable into its logarithm, the data presents a linear relationship. However, in Healthy group, when age larger than 83, the trend seems to be different. It could be ignored because we have too little data in the interval.

Ventriclesfit1=lm(log(Ventricles)~Age*AD,data=Ventricles.df)
plot(Ventriclesfit1,which=1)

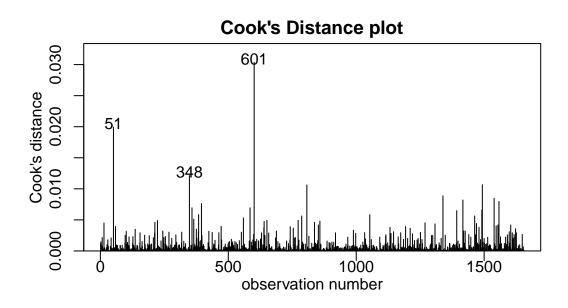


normcheck(Ventriclesfit1)





cooks20x(Ventriclesfit1)



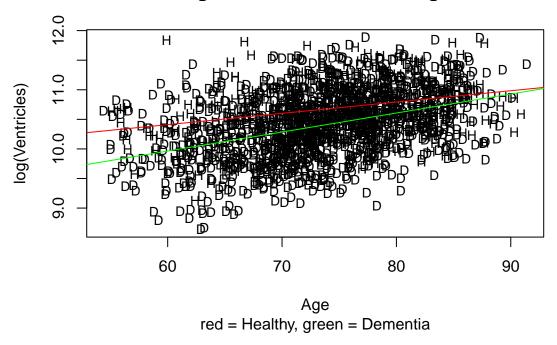
```
##
## Call:
## lm(formula = log(Ventricles) ~ Age * AD, data = Ventricles.df)
## Residuals:
       Min
                1Q
                   Median
                                3Q
## -1.51040 -0.34077 0.00086 0.33883 1.42693
## Coefficients:
              Estimate Std. Error t value Pr(>|t|)
## (Intercept) 8.034934 0.145792 55.112 < 2e-16 ***
## Age
             ## ADH
## Age:ADH
             ## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## Residual standard error: 0.5053 on 1651 degrees of freedom
## Multiple R-squared: 0.1877, Adjusted R-squared: 0.1862
## F-statistic: 127.2 on 3 and 1651 DF, p-value: < 2.2e-16
confint(Ventriclesfit1)
                   2.5 %
                             97.5 %
## (Intercept) 7.74897653 8.320891845
## Age
             0.02827412 0.036030020
              0.61838254 1.838252182
## ADH
## Age:ADH
             -0.02117928 -0.004891143
exp(confint(Ventriclesfit1))
                   2.5 %
                              97.5 %
## (Intercept) 2319.1975659 4108.8228069
## Age
                1.0286776
                           1.0366870
## ADH
                1.8559237
                            6.2855427
                0.9790434
## Age:ADH
                           0.9951208
(exp(confint(Ventriclesfit1))-1)*100
                    2.5 %
                                97.5 %
## (Intercept) 231819.756593 4.107823e+05
## Age
                 2.867762 3.668697e+00
                85.592372 5.285543e+02
## ADH
## Age:ADH
                -2.095657 -4.879201e-01
# rotate factor
Ventricles.df=within(Ventricles.df,{ADflip=factor(AD,levels=c("H","D"))})
Ventriclesfit2=lm(log(Ventricles)~Age*ADflip,data=Ventricles.df)
summary(Ventriclesfit2)
```

summary(Ventriclesfit1)

```
##
## Call:
## lm(formula = log(Ventricles) ~ Age * ADflip, data = Ventricles.df)
## Residuals:
##
       \mathtt{Min}
                  1Q
                     Median
                                    3Q
                                            Max
## -1.51040 -0.34077 0.00086 0.33883 1.42693
##
## Coefficients:
##
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 9.263252 0.274675 33.724 < 2e-16 ***
                           0.003651
                                     5.236 1.85e-07 ***
                0.019117
## ADflipD
               -1.228317
                           0.310969 -3.950 8.15e-05 ***
                           0.004152
                                    3.139 0.00172 **
## Age:ADflipD 0.013035
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## Residual standard error: 0.5053 on 1651 degrees of freedom
## Multiple R-squared: 0.1877, Adjusted R-squared: 0.1862
## F-statistic: 127.2 on 3 and 1651 DF, p-value: < 2.2e-16
confint(Ventriclesfit2)
                      2.5 %
##
                                 97.5 %
## (Intercept) 8.724504197 9.80199889
                0.011955341 0.02627837
## Age
## ADflipD
               -1.838252182 -0.61838254
## Age: ADflipD 0.004891143 0.02117928
exp(confint(Ventriclesfit2))
                      2.5 %
                                  97.5 %
##
## (Intercept) 6151.8258140 1.806983e+04
## Age
                  1.0120271 1.026627e+00
## ADflipD
                  0.1590953 5.388152e-01
## Age:ADflipD
                  1.0049031 1.021405e+00
(\exp(\text{confint}(\text{Ventriclesfit2}))-1)*100
##
                       2.5 %
                                    97.5 %
## (Intercept) 6.150826e+05 1.806883e+06
## Age
                1.202709e+00 2.662669e+00
## ADflipD
               -8.409047e+01 -4.611848e+01
## Age:ADflipD 4.903124e-01 2.140515e+00
```

Plot the data with your appropriate model superimposed over it

log Ventricles Size versus Age



or abline(Ventriclesfit2\$coef[1], Ventriclesfit2\$coef[2], col="red")

Methods and assumption checks

As the size of the ventricles increased the variability also increased so we logged the Ventricles data, this evened out the scatter. We have two explanatory variables, a grouping explanatory variable with two levels and a numeric explanatory variable, so have fitted a linear model with both variables and included an interaction term. The test for the interaction term proved to be significant, so the interaction term was kept and the model could not be simplified further.

Checking the assumptions there are no problems with assuming constant variability; looking at normality we see no issues and the Cook's plot doesn't reaveal any points of concern; as we have assumed the people were randomly sampled, independence is satisfied. The model assumptions are satisfied.

Our model is: $log(Ventricles_i) = \beta_0 + \beta_1 \times Age_i + \beta_2 \times ADH_i + \beta_3 \times Age_i \times ADH_i + \epsilon_i$ where $ADH_i = 1$ if the *i*th subject is healthy and 0 if they have signs of dementia, and $\epsilon_i \sim iid\ N(0, \sigma^2)$

Our model only explained 19% of the variability in the data.

In terms of slopes and/or intercepts, explain what the coefficient of Age:ADH is estimating.

For slopes, Age: ADH means Healthy volunteers could get additional times of $e^{Age:ADH}$ median increasement every year than Dementia volunteers.

For each of the following, either write a sentence interpreting a confidence interval to estimate the requested information or state why we cannot answer this from the R-output given:

-in general, the difference in size of ventricles between healthy people and those exhibiting dementia symptoms.

We can see in the output of (exp(confint(Ventriclesfit1))-1)*100, ADH is estimated between 85.59 and 528.55, which means the Healthy volunteers' median of Ventricles volumes could increased by 85.59% to 528.55% than Dementia volunteers. Though Age*ADH is minus number which means the Ventricles volumes of Dementia could exceed the Ventricles volumes of Healthy, the intersect point is outside the data range. So the Ventricles volumes of Healthy is always much larger than the Ventricles volumes of Dementia.

-the effect on the size of ventricles for each additional years aging on healthy people.

We can see in the output of (exp(confint(Ventriclesfit1))-1)*100, Age is estimated between 2.87 and 3.67, which means the Dementia volunteers' median of Ventricles volumes for each additional years could increased by 2.87% to 3.67%. And Age: ADH is estimated between -2.10 and -0.49, which means the Healthy volunteers' median of Ventricles volumes for each additional years could decreased by 2.10% to 0.49% than the estimated median of Ventricles volumes of Dementia volunteers.

So, the effect on the size of ventricles for each additional years aging on healthy people could be estimated by $2.87 - 2.10 \sim 3.67 - 0.49$. That is, the Healthy volunteers' median of Ventricles volumes for each additional years could increased by about 0.7% to 3.2%.

-the effect on the size of ventricles for each additional years aging on people exhibiting dementia symptoms.

We can see in the output of (exp(confint(Ventriclesfit1))-1)*100, Age is estimated between 2.87 and 3.67, which means the Healthy volunteers' median of Ventricles volumes could increased by 2.87% to 3.67% every year.

Looking at the plot with the model superimposed, describe what seems to be happening.

For Healthy volunteers, they usually have larger Ventricles volumes when they are young. However, when they getting old, they have a low speed increment of Ventricles volumes. For Dementia volunteers, they usually have less Ventricles volumes when they are young. However, when they getting old, they have a high speed increment of Ventricles volumes.

However, Healthy volunteers' Ventricles volumes are always lager than Dementia volunteers' Ventricles volumes even though Dementia volunteers have a high speed increment.