

CS 6363.001 Statistical Methods for Data Science (Spring 2017)

Mini Project #5

April 20, 2017

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Contribution.

Both of us discussed and completed the entire project together.

- psa is the response variable.
- all other variables are predictors.
- subject ID is not a predictor.
- Quantitative variables are: cancervol, weight, age, benpros, capspen
- vesinv is a qualitative variable with no ordering.
- gleason is a qualitative variable with ordering.

The simplest linear regression model looks like this: $y = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \dots + \beta_p x_p + \varepsilon$ (1)

The regression model comes with two main assumptions:

1. Linear relationship between response and predictor.
2. $\varepsilon \sim N(0, \sigma^2)$

-UNIVARIATE ANALYSIS:

#We begin by doing a univariate analysis(one predictor at a time) and see each variable's performance.

#Import the data set

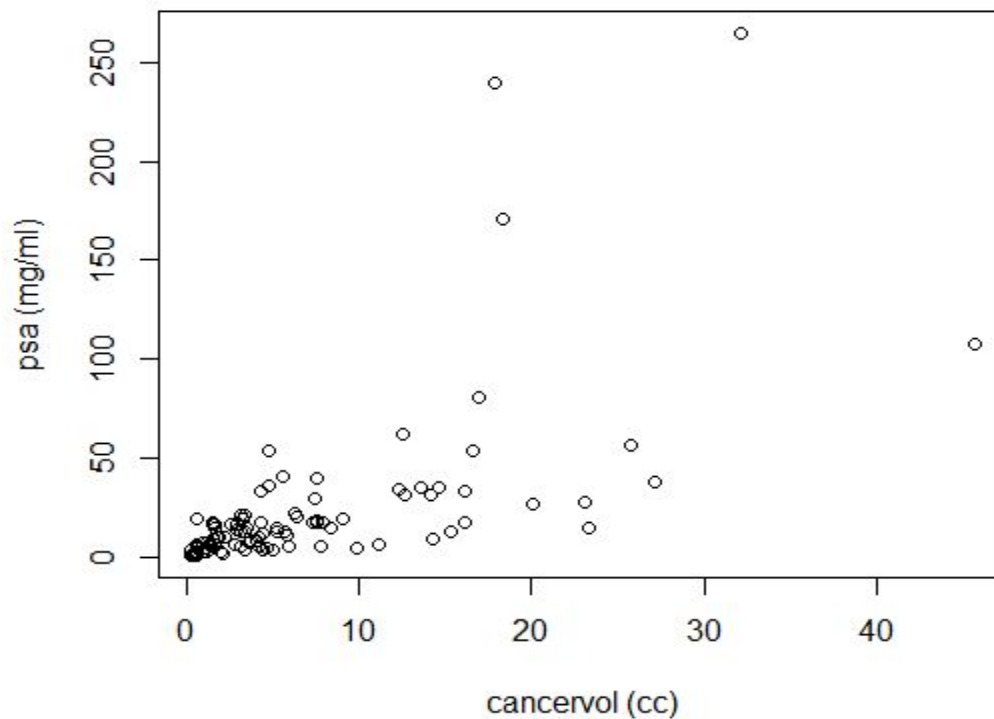
```
prostate1 <- read.table(file="prostate_cancer.csv", sep="," , header=T)
```

#Attach the data set values

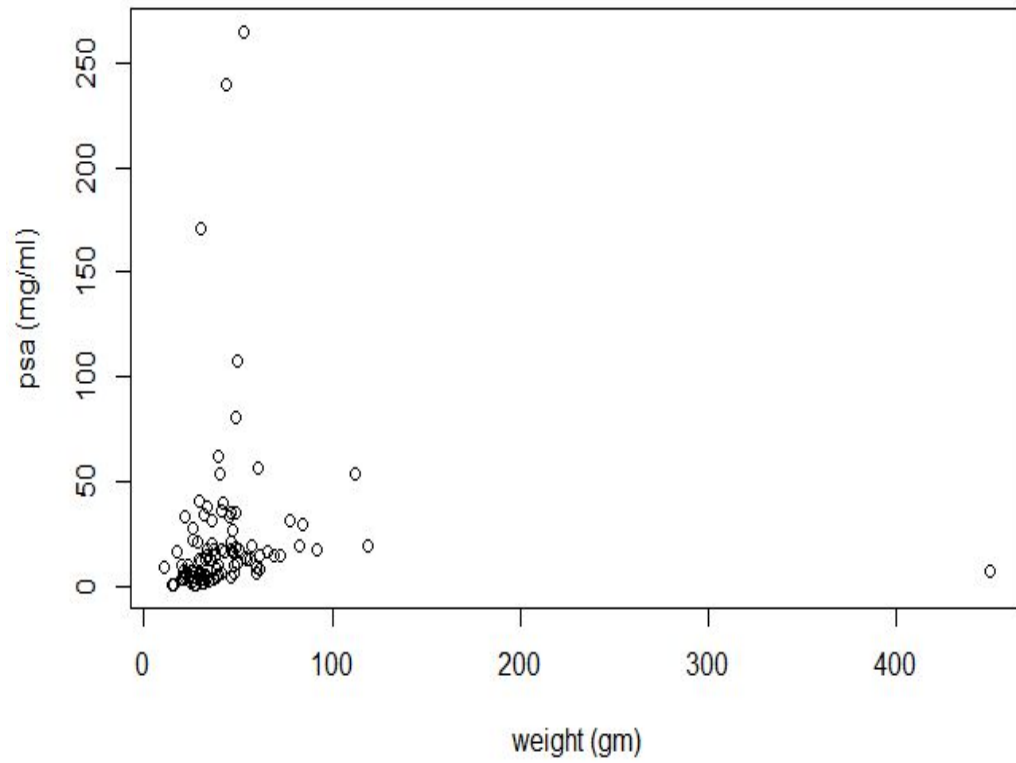
```
attach(prostate1)
```

#Scatterplot - psa vs cancervol

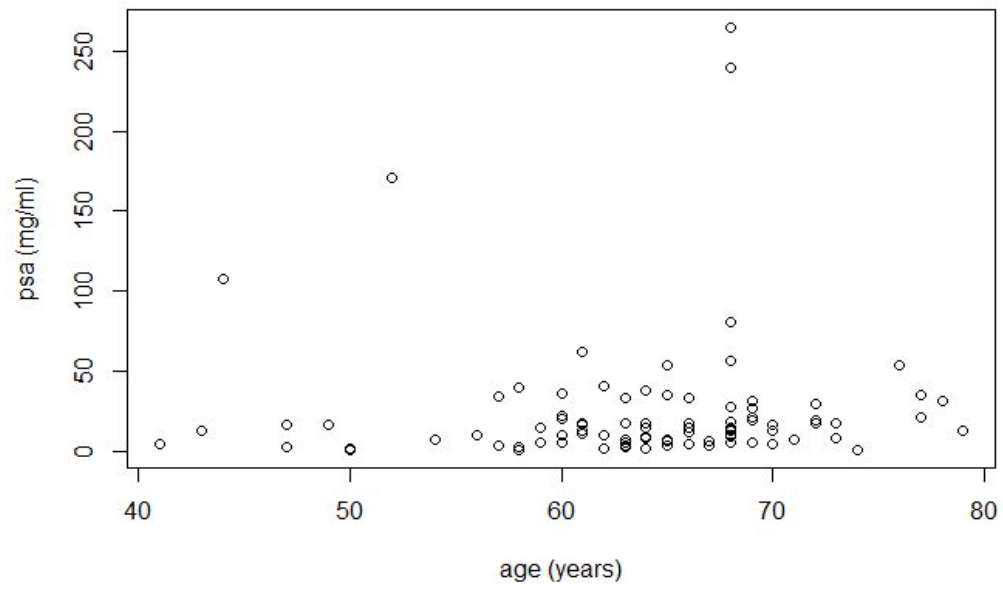
```
plot(cancervol, psa, xlab="cancervol (cc)", ylab="psa (mg/ml)")
```



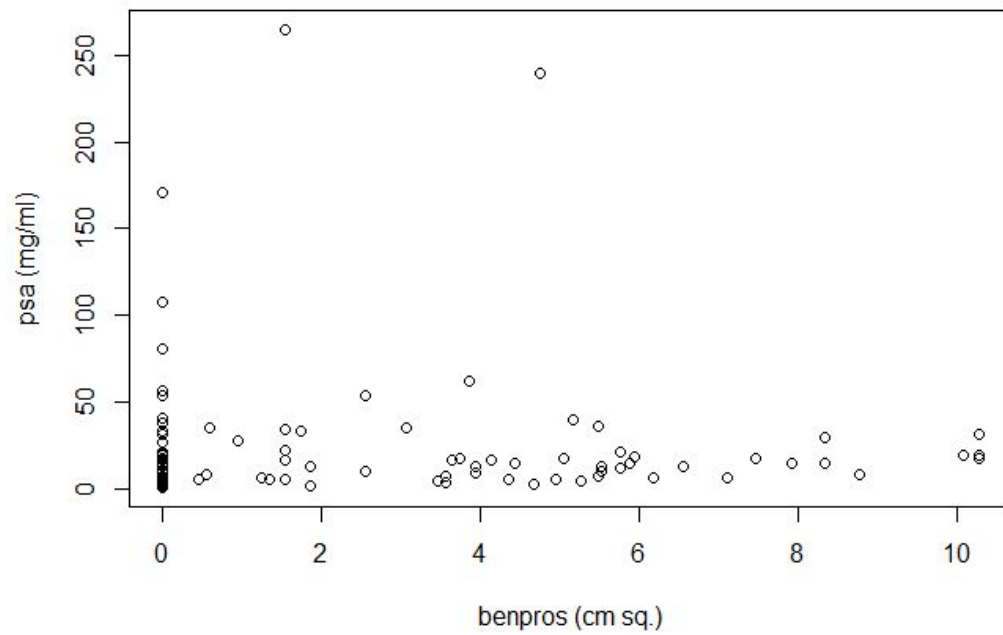
```
#Scatterplot - psa vs weight  
plot(weight, psa, xlab="weight (gm)", ylab="psa (mg/ml)")
```



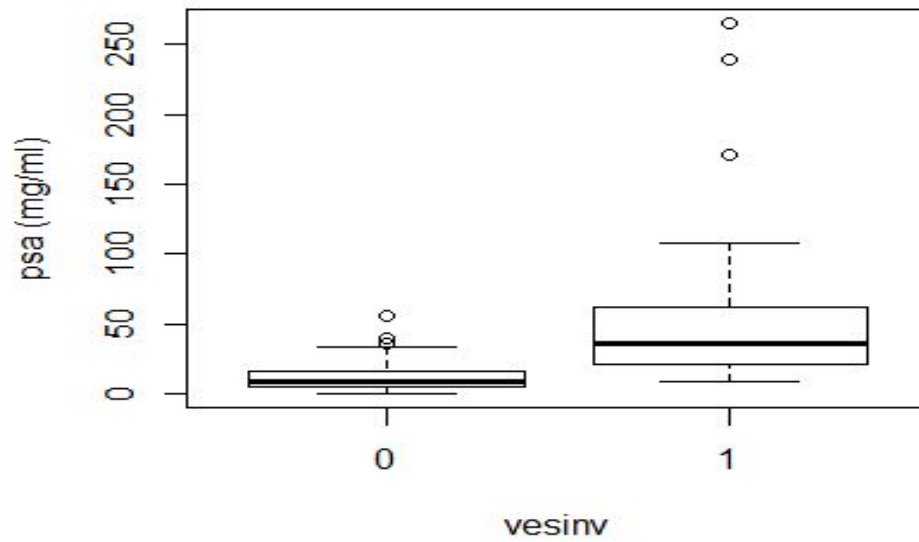
```
#Scatterplot - psa vs age  
plot(age, psa, xlab="age (years)", ylab="psa (mg/ml)")
```



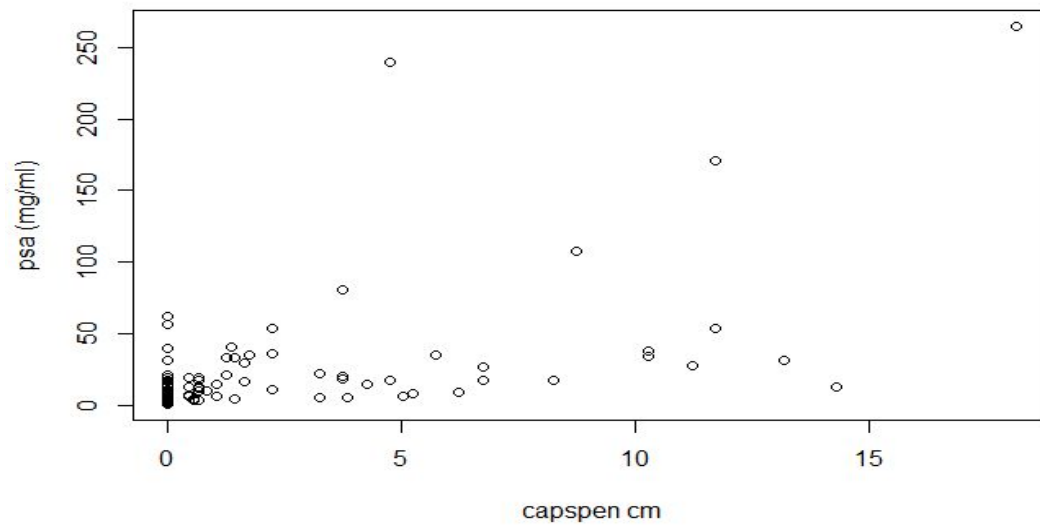
```
#Scatterplot - psa vs benpros  
plot(benpros, psa, xlab="benpros (cm sq.)", ylab="psa (mg/ml)")
```



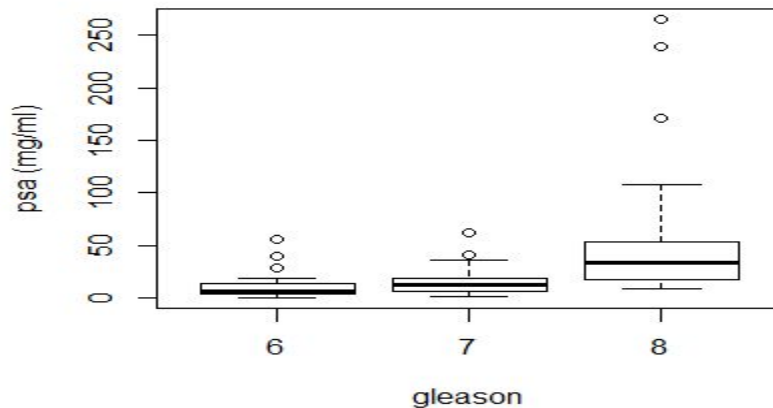
```
#Scatterplot - psa vs vesinv  
plot(factor(vesinv), psa, xlab="vesinv ", ylab="psa (mg/ml)")
```



```
#Scatterplot - psa vs capspen  
plot(capspen, psa, xlab="capspen cm", ylab="psa (mg/ml)")
```



```
#Scatterplot - psa vs gleason
plot(factor(gleason), psa, xlab="gleason", ylab="psa (mg/ml)")
```



```
#to generate correlations between psa and cancervol
cor(psa,cancervol)
[1] 0.6241506
```

#similarly, we find the correlation between psa and other variables

```
>cor(psa,weight)
[1] 0.02621343
```

```
>cor(psa,age)
[1] 0.01719938
```

```
>cor(psa,benpros)
[1] -0.01648649
```

```
>cor(psa,capspen)
[1] 0.5507925
```

- We can see the correlation is positive and moderately strong for the quantitative variable cancervol.
- So, from the univariate analysis, our predictor for the data is cancervol as the correlation between the PSA level and cancervol is a linear positive relationship from the scatter plot.

MODEL BUILDING:

-Multivariate Analysis:

#A sensible solution to picking a good model is to Minimize AIC.

#We wish to find which ones (if any) of the explanatory variables are important.

#We'll start off with including every variable as-is

#To do this, we first fit the full linear model:

```

> fit.lm = lm(psa~.,data=prostate1)
> summary(fit.lm)
> fit.lm = lm(psa~.,data=prostate)
> summary(fit.lm)

Call:
lm(formula = psa ~ ., data = prostate)

Residuals:
    Min       1Q   Median       3Q      Max
-51.856 -10.605   0.309   6.916 167.586

Coefficients:
              Estimate Std. Error t value
(Intercept)  1.0677412  39.7450228   0.027
subject      0.4377515   0.1665658   2.628
cancervol    1.3355863   0.6329973   2.110
weight      -0.0004541   0.0717523  -0.006
age         -0.5530385   0.4608701  -1.200
benpros      0.3740731   1.2156657   0.308
vesinv      9.6762126  11.2042175   0.864
capspen     1.6000523   1.3056066   1.226
gleason     2.9915123   5.2630095   0.568
Pr(>|t|)
(Intercept)  0.9786
subject      0.0101 *
cancervol    0.0377 *
weight       0.9950
age          0.2334
benpros      0.7590
vesinv       0.3901
capspen      0.2236
gleason      0.5712
---
Signif. codes:
  0 '***' 0.001 '**' 0.01 '*' 0.05 '.'
  0.1 ' ' 1

Residual standard error: 30.18 on 88 degrees of freedom
Multiple R-squared:  0.4979,    Adjusted R-squared:  0.4523
F-statistic: 10.91 on 8 and 88 DF,  p-value: 1.46e-10

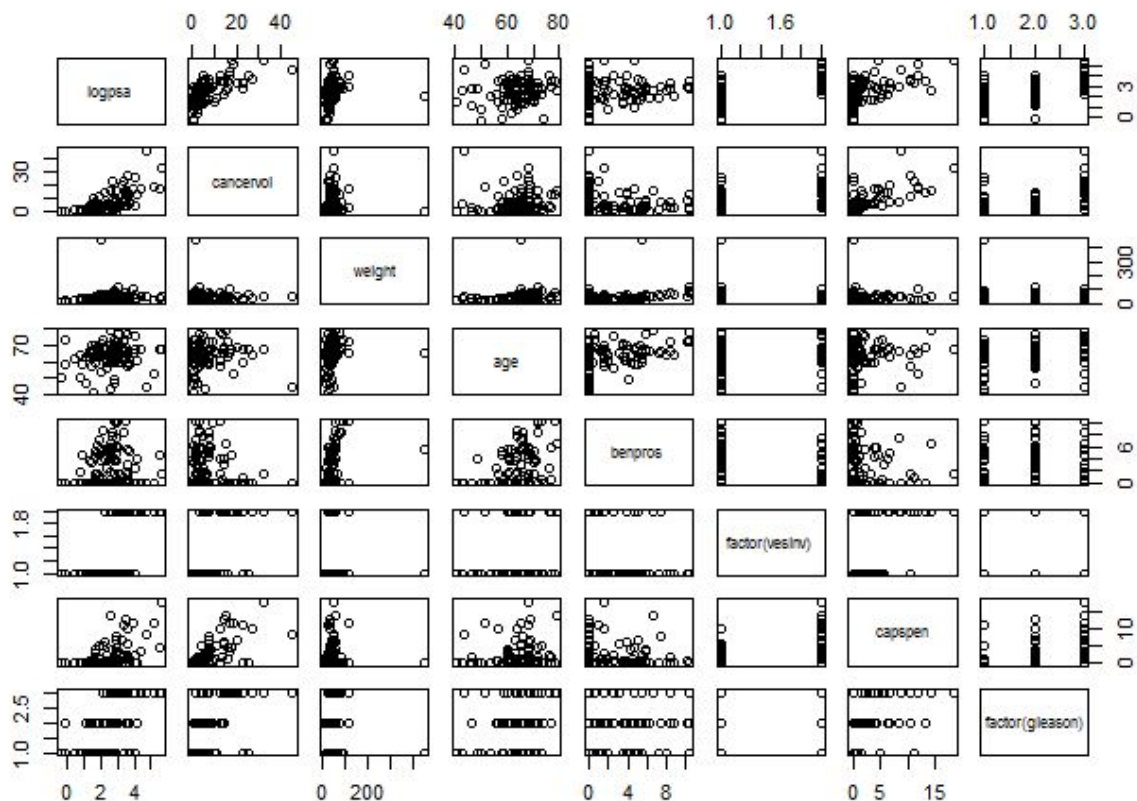
```

Observation:

Here, some of the predictors are unimportant.
(But there are too many parameters)

Transformations:

Often it is a good idea to transform the data to make it fit the assumptions of the model. Most commonly this means transforming the data to make the relationships more linear. In this case, let's try a log transformation on the dependent variable psa



```
> logpsa <- log(psa)
> psa = NULL
> pairs(logpsa ~ cancervol + weight + age + benpros + factor(vesinv) + capspen + factor(gleason), data = prostate1)
```

#Let's fit the model again

```
> transformedfit <- lm(logpsa ~ ., data = prostate1)
```

```
> summary(transformedfit)
```

Call:

```
lm(formula = logpsa ~ ., data = prostate)
```

Residuals:

Min	1Q	Median	3Q	Max
-1.11781	-0.09685	0.04960	0.17660	0.24607

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	0.6574253	0.3490985	1.883	0.063 .
subject	0.0334434	0.0015193	22.012	< 2e-16 ***
psa	0.0063728	0.0009363	6.806	1.22e-09 ***
cancervol	0.0032788	0.0056988	0.575	0.567
weight	0.0004085	0.0006302	0.648	0.519
age	-0.0005843	0.0040810	-0.143	0.886
benpros	0.0085855	0.0106834	0.804	0.424
vesinv	-0.1012339	0.0988275	-1.024	0.309
capspen	0.0047744	0.0115651	0.413	0.681


```
gleason    0.0024009 0.0463120 0.052 0.959
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Residual standard error: 0.2651 on 87 degrees of freedom
Multiple R-squared: 0.9521, Adjusted R-squared: 0.9472
F-statistic: 192.3 on 9 and 87 DF, p-value: < 2.2e-16

Observation:

The adjusted R-squared increased from 0.4523 to 0.9472, which is a huge increase. Based on these results, the log transformation of the dependent variables is definitely a good idea.

#But let us also try and see a Transformation using sqrt:

```
> sqrtpsa <-sqrt(prostate$psa)
> prostate$psa <- NULL
> pairs(sqrtpsa ~ cancervol + weight + age + benpros + factor(vesinv) + capspen + factor(gleason), data = prostate1)
> sqrtfit = lm(sqrtpsa ~ ., data = prostate1)

> summary(sqrtfit)
```

Call:

```
lm(formula = sqrtpsa ~ ., data = prostate)
```

Residuals:

```
    Min     1Q  Median     3Q    Max
-2.4953 -0.5348  0.0312  0.3088  7.2946
```

Coefficients:

```
            Estimate Std. Error t value Pr(>|t|)
(Intercept)  1.2813319  1.7877952   0.717  0.4755
subject      0.0576572  0.0074924  7.695 1.93e-11 ***
cancervol    0.0719255  0.0284732  2.526  0.0133 *
weight       0.0002907  0.0032275   0.090  0.9284
age          -0.0295024  0.0207307  -1.423  0.1582
benpros      0.0143057  0.0546826   0.262  0.7942
vesinv       0.5118460  0.5039838   1.016  0.3126
capspen      0.0551802  0.0587283   0.940  0.3500
gleason      0.1557187  0.2367387   0.658  0.5124
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

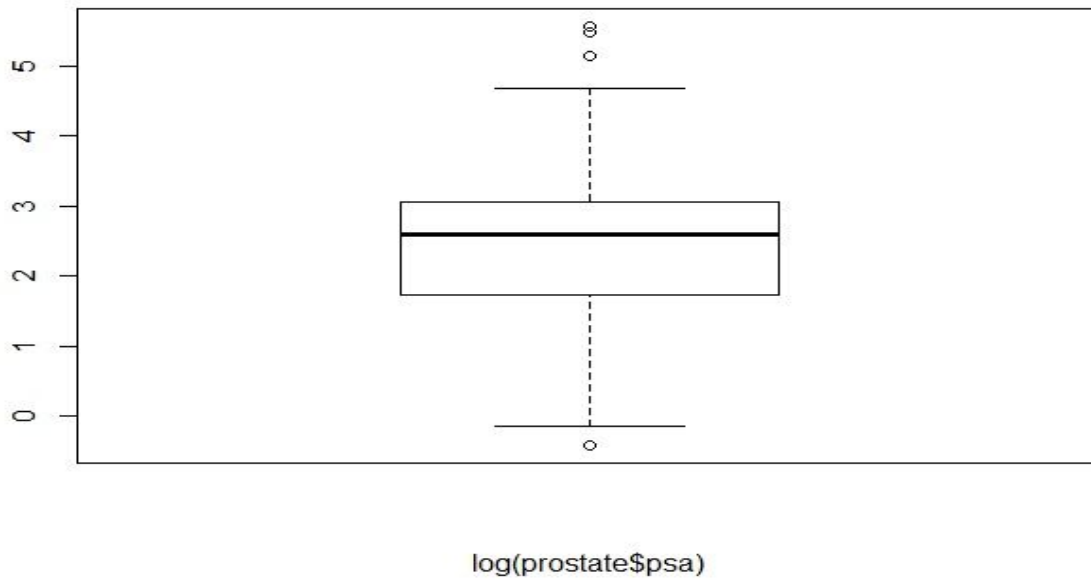
Residual standard error: 1.358 on 88 degrees of freedom
Multiple R-squared: 0.7638, Adjusted R-squared: 0.7423
F-statistic: 35.56 on 8 and 88 DF, p-value: < 2.2e-16

Observation:

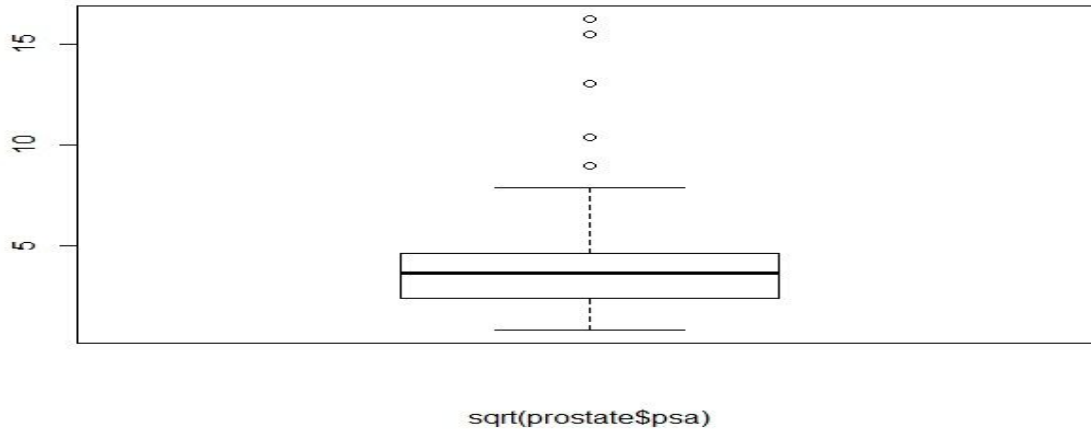
The adjusted R-squared increased from 0.4523 to 0.7423, which is a reasonable increase.

#Let's see how the boxplot of the log and sqrt transformation look like:

```
> boxplot(log(prostate1$psa),xlab="log(prostate1$psa)")
```



```
> boxplot(sqrt(prostate1$psa), xlab="sqrt(prostate1$psa)")
```



Observations of Box Plots:

We see that sqrt has more outliers than log transformation. So we can choose log as a better pick for transformation.

```
y = log(prostate1$psa)
```

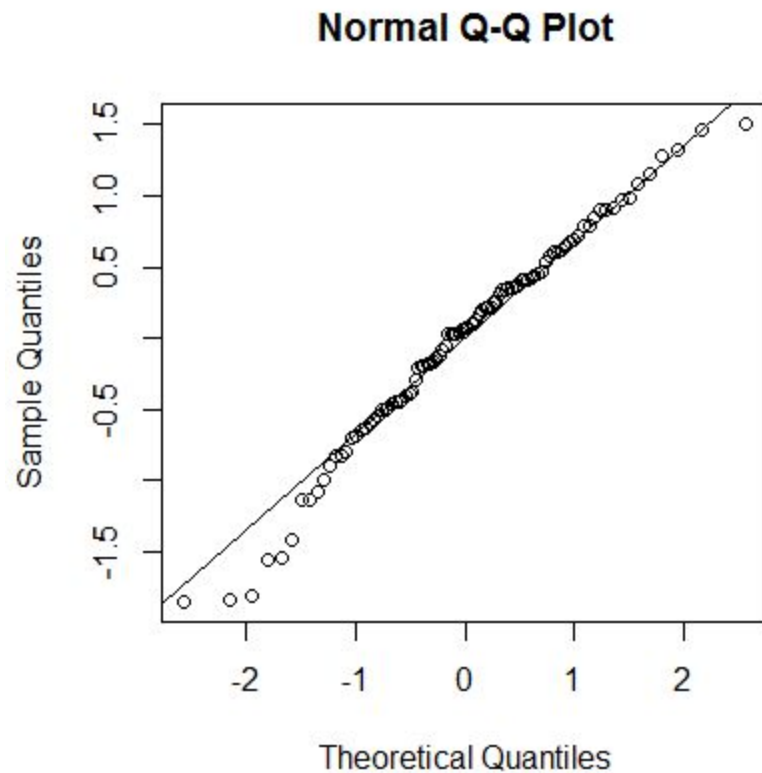
#check for the residulas in the fit

```
fit.check <- lm(y ~ cancervol + weight + age + benpros + factor(vesinv) + capspen + factor(gleason), data = prostate1)
```

#residuals in fit

```
fit.res <- resid(fit.check)
```

```
#qqplot for residuals
qqnorm(fit.res)
qqline(fit.res)
```



```
#find the outliers using IQR values
qnt = quantile(fit.res, probs=c(.25,.75))
>qnt
      25%      75%
-0.4540944 0.4554852
H = 1.5 * IQR(fit.res)
> H
>[1] 1.364369
fit.out <- fit.res
fit.out[fit.res < (qnt[1] - H)] <- NA
fit.out[fit.res > (qnt[2] + H)] <- NA
```

```
> fit.out
```

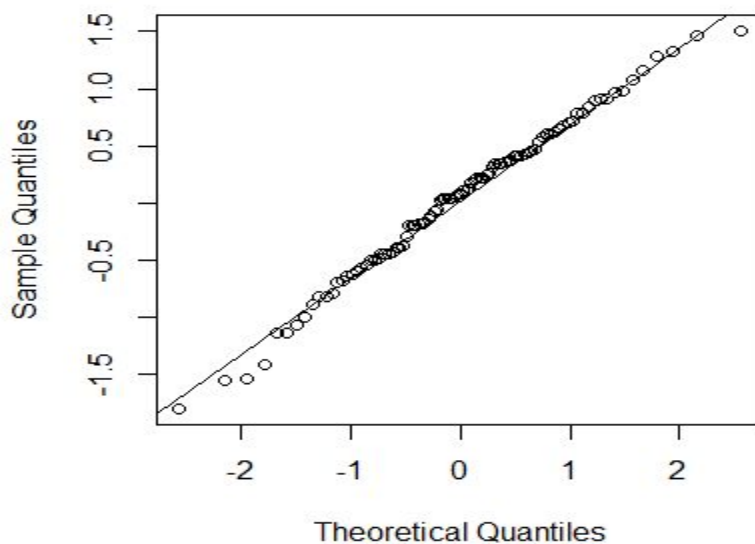
1	2	3	4	5	6	7
NA	NA	-1.80543625	-1.54829128	-1.13467907	-0.64205047	-0.89496903
8	9	10	11	12	13	14
-1.07778810	-0.68617283	-0.38641903	-0.17748992	-0.43125911	-0.69571635	-0.57757500
15	16	17	18	19	20	21
-0.48221985	-0.20342571	-0.50784984	-0.49821848	0.10644951	-0.60509240	0.05686458
22	23	24	25	26	27	28
-0.82178205	0.25211810	-0.39883298	-0.16399833	0.02735191	0.03604001	-0.44162079
29	30	31	32	33	34	35
-0.05864202	-0.07879595	-0.18108121	-0.44422176	0.12583545	0.57315513	0.64330683
36	37	38	39	40	41	42
-0.63254756	-0.11745732	0.45548522	-1.41142362	0.26221185	0.07018304	0.37210016
43	44	45	46	47	48	49
0.35041664	0.34845460	0.07836434	0.36950010	-1.55330434	0.20448726	0.78375910
50	51	52	53	54	55	56
0.67723662	0.78792142	0.21463211	0.53827745	0.03141377	-0.99484079	0.03733979
57	58	59	60	61	62	63
0.97176912	0.90074142	0.42588360	0.60953325	0.40875539	-0.38191084	-0.19539824
64	65	66	67	68	69	70
-0.54540734	0.98076929	0.60174819	0.31812190	-0.19367187	1.50934523	0.10163998
71	72	73	74	75	76	77
0.21158047	0.69858054	0.41519945	-0.29743202	-0.79838083	-0.13720459	0.03999679
78	79	80	81	82	83	84
-0.45409444	0.41659295	0.33656214	1.46788205	1.15985219	0.21380769	0.34562794
85	86	87	88	89	90	91
0.71395172	-0.82612960	1.32518529	0.91560218	0.17354679	0.62573193	0.90771722
92	93	94	95	96	97	
0.47357808	0.43895732	-1.13956679	1.28358142	1.07965595	0.84292575	

#We observe that the first 2 values have come out as NA, means they are the outlier values.

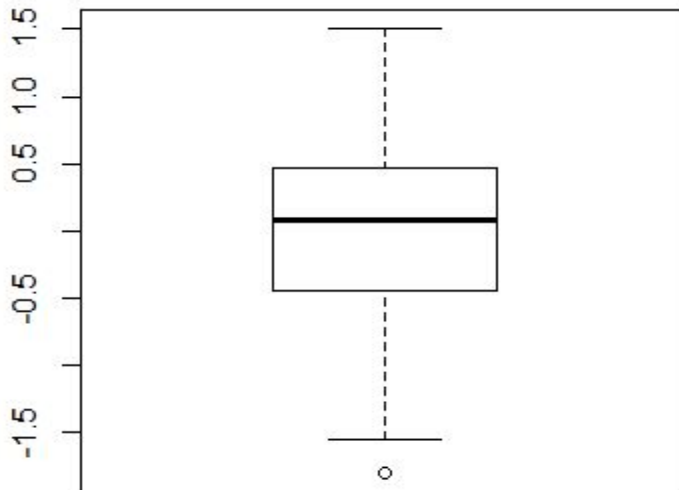
```
qqnorm(fit.out)
```

```
qqline(fit.out)
```

Normal Q-Q Plot



boxplot(fit.out)



#####now removing the outliers from the data(first 2 values)

prostate = tail(prostate1, -2)

##set y

y = log(prostate\$psa)

Note:

prostate1 was our initial dataset.

prostate is now the new dataset after the removal of residuals.

#####check for the best fit -manually

#full

fit.full <- lm(y ~ cancervol + weight + age + benpros + factor(vesinv) + capspen + factor(gleason), data = prostate)

> summary(fit.full)

#

Call:

lm(formula = y ~ cancervol + weight + age + benpros + factor(vesinv) +

capspen + factor(gleason), data = prostate)

#

Residuals:

Min 1Q Median 3Q Max

-1.83835 -0.47139 0.05644 0.45003 1.46694

#

Coefficients:

Estimate Std. Error t value Pr(>|t|)

(Intercept) 1.991266 0.692637 2.875 0.00509 **

cancervol 0.062206 0.014474 4.298 4.53e-05 ***

weight 0.001123 0.001734 0.647 0.51907

age -0.009134 0.011191 -0.816 0.41665

```
# benpros      0.082172  0.028201  2.914  0.00455 **
# factor(vesinv)1 0.791629  0.253819  3.119  0.00247 **
# capspen      -0.025558  0.030948 -0.826  0.41118
# factor(gleason)7 0.308906  0.179597  1.720  0.08903 .
# factor(gleason)8 0.768287  0.250041  3.073  0.00284 **
# ---
# Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
#
# Residual standard error: 0.7231 on 86 degrees of freedom
# Multiple R-squared:  0.5986,    Adjusted R-squared:  0.5613
# F-statistic: 16.03 on 8 and 86 DF,  p-value: 3.066e-14
```

#####try1---removing weight from fit.full as it has high p-value

```
fit.try1 <- lm(y ~ cancervol + age + benpros + factor(vesinv) + capspen + factor(gleason), data = prostate)
```

```
> summary(fit.try1)
```

Call:

```
lm(formula = y ~ cancervol + age + benpros + factor(vesinv) +
    capspen + factor(gleason), data = prostate)
```

Residuals:

```
    Min       1Q   Median       3Q      Max
-1.85269 -0.48381  0.05273  0.44920  1.51187
```

Coefficients:

```
            Estimate Std. Error t value Pr(>|t|)
(Intercept)  2.011339   0.689630   2.917  0.00450 **
cancervol     0.062319   0.014425   4.320 4.12e-05 ***
age          -0.008738   0.011137  -0.785  0.43480
benpros       0.087693   0.026791   3.273  0.00153 **
factor(vesinv)1 0.797177   0.252826   3.153  0.00222 **
capspen      -0.025681   0.030844  -0.833  0.40735
factor(gleason)7 0.290387   0.176711   1.643  0.10393
factor(gleason)8 0.760915   0.248946   3.057  0.00297 **
```

```
---
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Residual standard error: 0.7207 on 87 degrees of freedom
Multiple R-squared:  0.5966,    Adjusted R-squared:  0.5642
F-statistic: 18.38 on 7 and 87 DF,  p-value: 8.333e-15
```

```
anova(fit.try1,fit.full)
```

```
# Analysis of Variance Table
```

```
#
```

```
# Model 1: y ~ cancervol + age + benpros + factor(vesinv) + capspen + factor(gleason)
```

```
# Model 2: y ~ cancervol + weight + age + benpros + factor(vesinv) + capspen +
```

```
# factor(gleason)
```

```
# Res.Df  RSS Df Sum of Sq  F Pr(>F)
```

```
# 1    87 45.187
```

```
# 2    86 44.967  1  0.21918 0.4192 0.5191
```

concluding fit.try1 as a better pick because there is an increase in F-statistic and p-value is > 0.05, accept Null Hypothesis, ie. fit.try2 is as good as fit.try1, we find weight as insignificant predictor, so we proceed with fit.try2.

#####try2---removing age as it has the highest pvalue in fit.try1

```
fit.try2 <- lm(y ~ cancervol + benpros + factor(vesinv) + capspen + factor(gleason), data = prostate)
```

```
> summary(fit.try2)
```

Call:

```
lm(formula = y ~ cancervol + benpros + factor(vesinv) + capspen +  
    factor(gleason), data = prostate)
```

Residuals:

Min	1Q	Median	3Q	Max
-1.9566	-0.4941	0.0405	0.4521	1.4363

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	1.48341	0.15096	9.827	8.16e-16 ***
cancervol	0.06325	0.01434	4.409	2.92e-05 ***
benpros	0.08025	0.02500	3.210	0.00185 **
factor(vesinv)1	0.77965	0.25129	3.103	0.00258 **
capspen	-0.02613	0.03077	-0.849	0.39800
factor(gleason)7	0.27506	0.17524	1.570	0.12010
factor(gleason)8	0.72626	0.24446	2.971	0.00383 **

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.7191 on 88 degrees of freedom

Multiple R-squared: 0.5938, Adjusted R-squared: 0.5661

F-statistic: 21.44 on 6 and 88 DF, p-value: 2.294e-15

```
anova(fit.try2,fit.try1)
```

```
# > anova(fit.try2,fit.try1)
```

```
# Analysis of Variance Table
```

```
#
```

```
# Model 1: y ~ cancervol + benpros + factor(vesinv) + capspen + factor(gleason)
```

```
# Model 2: y ~ cancervol + age + benpros + factor(vesinv) + capspen + factor(gleason)
```

```
# Res.Df RSS Df Sum of Sq F Pr(>F)
```

```
# 1 88 45.506
```

```
# 2 87 45.187 1 0.31976 0.6157 0.4348
```

#concluding fit.try2 as better because p-value > 0.05 and F-statistic of fit.try2 has increased.

```
#####try3---removing capspen from try2
```

```
fit.try3 <- lm(y ~ cancervol + benpros + factor(vesinv) + factor(gleason), data = prostate)
```

```
> summary(fit.try3)
```

Call:

```
lm(formula = y ~ cancervol + benpros + factor(vesinv) + factor(gleason),  
    data = prostate)
```

Residuals:

Min	1Q	Median	3Q	Max
-1.95245	-0.51569	0.04707	0.46982	1.42865

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	1.49439	0.15017	9.951	4.03e-16 ***
cancervol	0.05797	0.01291	4.492	2.11e-05 ***

```
benpros      0.07964  0.02495  3.192 0.00196 **
factor(vesinv)1 0.68403  0.22431  3.050 0.00302 **
factor(gleason)7 0.26308  0.17440  1.508 0.13497
factor(gleason)8 0.70283  0.24252  2.898 0.00473 **
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Residual standard error: 0.718 on 89 degrees of freedom
Multiple R-squared:  0.5905,    Adjusted R-squared:  0.5674
F-statistic: 25.66 on 5 and 89 DF, p-value: 6.039e-16
```

```
anova(fit.try3,fit.try2)
#
# > anova(fit.try3,fit.try2)
# Analysis of Variance Table
#
# Model 1: y ~ cancervol + benpros + factor(vesinv) + factor(gleason)
# Model 2: y ~ cancervol + benpros + factor(vesinv) + capslen + factor(gleason)
# Res.Df  RSS Df Sum of Sq   F Pr(>F)
# 1    89 45.879
# 2    88 45.506  1  0.37304 0.7214 0.398
```

#Concluding fit.try3 as good because p-value > 0.05, we accept null hypothesis that try 2 and try 3 are same and F-statistic of fit.try2 has increased.

#####try4-----removing factor(gleason) from try3

```
fit.try4 <- lm(y ~ cancervol + benpros + factor(vesinv), data = prostate)
> summary(fit.try4)
```

```
Call:
lm(formula = y ~ cancervol + benpros + factor(vesinv), data = prostate)
```

```
Residuals:
    Min       1Q   Median       3Q      Max
-1.82574 -0.53983  0.07311  0.54884  1.43414
```

```
Coefficients:
            Estimate Std. Error t value Pr(>|t|)
(Intercept)  1.62256    0.12833  12.644 < 2e-16 ***
cancervol     0.07161    0.01193   6.001 3.94e-08 ***
benpros       0.08672    0.02553   3.397 0.001013 **
factor(vesinv)1 0.80978    0.22552   3.591 0.000534 ***
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Residual standard error: 0.743 on 91 degrees of freedom
Multiple R-squared:  0.5516,    Adjusted R-squared:  0.5368
F-statistic: 37.32 on 3 and 91 DF, p-value: 8.118e-16
```

```
anova(fit.try4,fit.try3)
#
# > anova(fit.try4,fit.try3)
# Analysis of Variance Table
#
# Model 1: y ~ cancervol + benpros + factor(vesinv)
# Model 2: y ~ cancervol + benpros + factor(vesinv) + factor(gleason)
# Res.Df  RSS Df Sum of Sq   F Pr(>F)
# 1    91 50.230
# 2    89 45.879  2  4.3505 4.2197 0.01775 *
```



```
# ---
# Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

#gleason is a significant variable and we can omit it. Moreover, $pval < 0.05$, reject null hyp, which means try4 and try3 are not same. So we proceed with fit.try3.

Variable Selection: (Other conventional methods)

There are many methods for selecting model variables. We will use the following ones:

- **Forward selection based on AIC**
- **Backward elimination based on AIC**
- **Stepwise Selection - Both forward/backward**

1. Forward selection based on AIC

1. Find the best possible one predictor model**
2. Keep that predictor and find the best possible two predictor model given the original predictor is selected.
3. Keep both those predictors and find the best possible three predictor model given the first two selected.
- ...
4. Keep going until adding another predictor no longer reduces AIC

****Assumption:**

assume an intercept is included and always retained

```
##### proceeding with AIC
##### Forward selection based on AIC
```

```
fit.full.forward <- step(lm(y ~ 1, data = prostate),
  scope = list(upper = ~cancervol + weight + age + benpros + factor(vesinv) + capspen +
    factor(gleason)),
  direction = "forward")
```

```
#
# Start: AIC=17.66
# y ~ 1
#
# Df Sum of Sq  RSS   AIC
# + cancerrvol    1  48.996 63.029 -34.977
# + factor(vesinv) 1   37.423 74.603 -18.962
# + factor(gleason) 2   36.524 75.502 -15.823
# + capspen       1   30.616 81.409 -10.667
# <none>                  112.025 17.661
# + benpros       1    1.705 110.321  18.204
# + age           1    1.358 110.667  18.502
# + weight        1    1.158 110.867  18.674
#
# Step: AIC=-34.98
# y ~ cancerrvol
#
# Df Sum of Sq  RSS   AIC
# + factor(gleason) 2    7.8988 55.130 -43.697
# + factor(vesinv) 1    6.4303 56.599 -43.200
# + benpros       1    5.6823 57.347 -41.952
# <none>                  63.029 -34.977
# + weight        1    1.2197 61.809 -34.833
# + age           1    1.1049 61.924 -34.657
# + capspen       1    0.9432 62.086 -34.409
```

```

#
# Step: AIC=-43.7
# y ~ cancervol + factor(gleason)
#
# Df Sum of Sq  RSS   AIC
# + benpros      1  4.4568 50.673 -49.705
# + factor(vesinv) 1  3.9998 51.130 -48.852
# + weight      1  1.4575 53.673 -44.242
# <none>                55.130 -43.697
# + age          1  0.1718 54.958 -41.994
# + capspen      1  0.1666 54.964 -41.985
#
# Step: AIC=-49.71
# y ~ cancervol + factor(gleason) + benpros
#
# Df Sum of Sq  RSS   AIC
# + factor(vesinv) 1  4.7940 45.879 -57.147
# <none>                50.673 -49.705
# + weight      1  0.2793 50.394 -48.230
# + capspen      1  0.1891 50.484 -48.060
# + age          1  0.1136 50.560 -47.918
#
# Step: AIC=-57.15
# y ~ cancervol + factor(gleason) + benpros + factor(vesinv)
#
# Df Sum of Sq  RSS   AIC
# <none>                45.879 -57.147
# + capspen  1  0.37304 45.506 -55.922
# + age      1  0.33275 45.547 -55.838
# + weight   1  0.19336 45.686 -55.548

```

Backward elimination based on AIC

```

fit.full.backward <- step(lm(y ~ cancervol + weight + age + benpros + factor(vesinv) + capspen +
factor(gleason), data = prostate),
scope = list(lower = ~1), direction = "backward")

```

```

#
# Start: AIC=-53.05
# y ~ cancervol + weight + age + benpros + factor(vesinv) + capspen +
# factor(gleason)
#
# Df Sum of Sq  RSS   AIC
# - weight      1  0.2192 45.187 -54.592
# - age         1  0.3483 45.316 -54.321
# - capspen     1  0.3566 45.324 -54.304
# <none>                44.967 -53.054
# - factor(gleason) 2  4.9839 49.951 -47.069
# - benpros      1  4.4394 49.407 -46.110
# - factor(vesinv) 1  5.0862 50.054 -44.874
# - cancervol    1  9.6576 54.625 -36.572
#
# Step: AIC=-54.59
# y ~ cancervol + age + benpros + factor(vesinv) + capspen + factor(gleason)
#
# Df Sum of Sq  RSS   AIC
# - age          1  0.3198 45.506 -55.922
# - capspen      1  0.3601 45.547 -55.838
# <none>                45.187 -54.592
# - factor(gleason) 2  4.8786 50.065 -48.852
# - factor(vesinv) 1  5.1637 50.350 -46.313

```

```
# - benpros      1  5.5647 50.751 -45.559
# - cancervol    1  9.6942 54.881 -38.128
#
# Step: AIC=-55.92
# y ~ cancervol + benpros + factor(vesinv) + capspen + factor(gleason)
#
# Df Sum of Sq  RSS   AIC
# - capspen      1  0.3730 45.879 -57.147
# <none>                45.506 -55.922
# - factor(gleason) 2  4.5887 50.095 -50.796
# - factor(vesinv)  1  4.9780 50.484 -48.060
# - benpros       1  5.3280 50.834 -47.404
# - cancervol     1 10.0544 55.561 -38.958
#
# Step: AIC=-57.15
# y ~ cancervol + benpros + factor(vesinv) + factor(gleason)
#
# Df Sum of Sq  RSS   AIC
# <none>                45.879 -57.147
# - factor(gleason) 2  4.3505 50.230 -52.540
# - factor(vesinv)  1  4.7940 50.673 -49.705
# - benpros       1  5.2510 51.130 -48.852
# - cancervol     1 10.4003 56.280 -39.737
```

Both forward/backward

```
fit.full.both <- step(lm(y ~ 1, data = prostate),
  scope = list(lower = ~1, upper = ~cancervol + weight + age + benpros + factor(vesinv) + capspen +
factor(gleason)),
  direction = "both")
#
# Start: AIC=17.66
# y ~ 1
#
# Df Sum of Sq  RSS   AIC
# + cancervol      1 48.996 63.029 -34.977
# + factor(vesinv)  1 37.423 74.603 -18.962
# + factor(gleason) 2 36.524 75.502 -15.823
# + capspen        1 30.616 81.409 -10.667
# <none>                112.025 17.661
# + benpros        1  1.705 110.321 18.204
# + age            1  1.358 110.667 18.502
# + weight         1  1.158 110.867 18.674
#
# Step: AIC=-34.98
# y ~ cancervol
#
# Df Sum of Sq  RSS   AIC
# + factor(gleason) 2  7.899 55.130 -43.697
# + factor(vesinv)  1  6.430 56.599 -43.200
# + benpros        1  5.682 57.347 -41.952
# <none>                63.029 -34.977
# + weight         1  1.220 61.809 -34.833
# + age            1  1.105 61.924 -34.657
# + capspen        1  0.943 62.086 -34.409
# - cancervol      1 48.996 112.025 17.661
#
# Step: AIC=-43.7
# y ~ cancervol + factor(gleason)
```

```

#
# Df Sum of Sq  RSS   AIC
# + benpros      1  4.4568 50.673 -49.705
# + factor(vesinv) 1  3.9998 51.130 -48.852
# + weight        1  1.4575 53.673 -44.242
# <none>                      55.130 -43.697
# + age           1  0.1718 54.958 -41.994
# + capspen       1  0.1666 54.964 -41.985
# - factor(gleason) 2  7.8988 63.029 -34.977
# - cancervol     1 20.3718 75.502 -15.823
#
# Step: AIC=-49.71
# y ~ cancervol + factor(gleason) + benpros
#
# Df Sum of Sq  RSS   AIC
# + factor(vesinv) 1  4.7940 45.879 -57.147
# <none>                      50.673 -49.705
# + weight        1  0.2793 50.394 -48.230
# + capspen       1  0.1891 50.484 -48.060
# + age           1  0.1136 50.560 -47.918
# - benpros       1  4.4568 55.130 -43.697
# - factor(gleason) 2  6.6734 57.347 -41.952
# - cancervol     1 22.7560 73.429 -16.467
#
# Step: AIC=-57.15
# y ~ cancervol + factor(gleason) + benpros + factor(vesinv)
#
# Df Sum of Sq  RSS   AIC
# <none>                      45.879 -57.147
# + capspen       1  0.3730 45.506 -55.922
# + age           1  0.3327 45.547 -55.838
# + weight        1  0.1934 45.686 -55.548
# - factor(gleason) 2  4.3505 50.230 -52.540
# - factor(vesinv) 1  4.7940 50.673 -49.705
# - benpros       1  5.2510 51.130 -48.852
# - cancervol     1 10.4003 56.280 -39.737

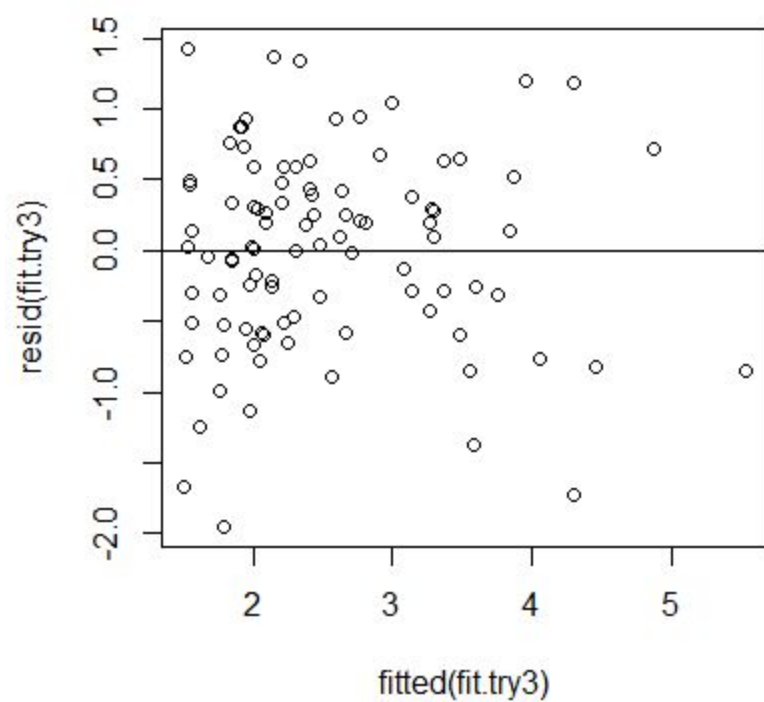
```

#####Here are the results for AIC
#for step forward-----y ~ cancervol + factor(gleason) + benpros + factor(vesinv)
#for step backward----y ~ cancervol + benpros + factor(vesinv) + factor(gleason)
#for step both----- y ~ cancervol + factor(gleason) + benpros + factor(vesinv)

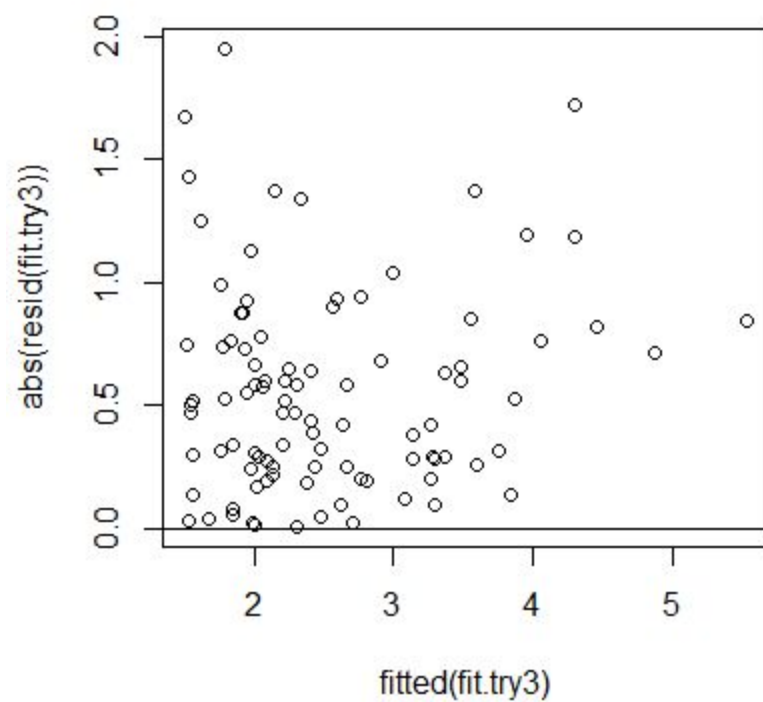
#In all the three AIC, we get the model with these four variables to be the best, which we also concluded for fit.try3. So we choose fit.try3 as the best fit.

#So our final model will contain: cancervol + factor(gleason) + benpros + factor(vesinv)

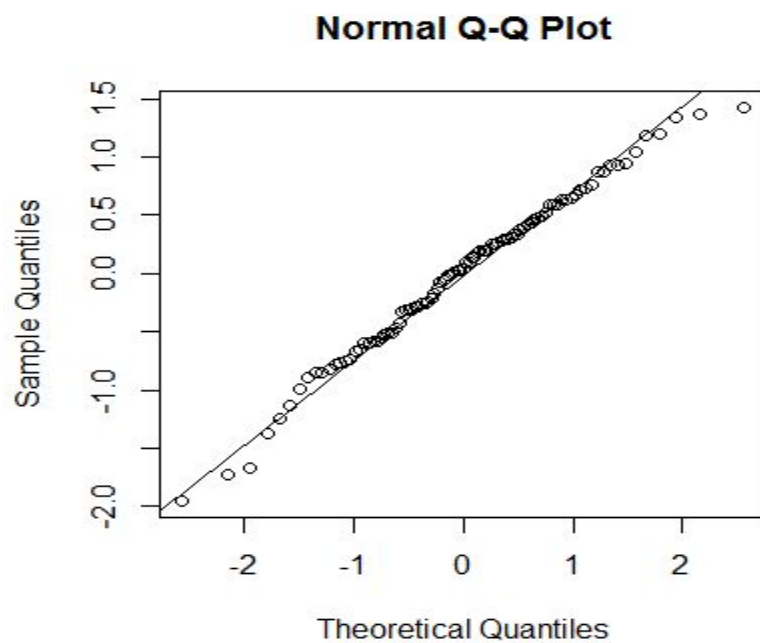
#residual plot
 plot(fitted(fit.try3), resid(fit.try3))
 abline(h = 0)



```
# plot of absolute residuals  
plot(fitted(fit.try3), abs(resid(fit.try3)))  
abline(h=0)
```



```
# normal QQ plot  
qqnorm(resid(fit.try3))  
qqline(resid(fit.try3))
```



#finding the modes of categorical variables vesinv and gleason

```
mode.vesinv = table(factor(vesinv))
new.vesinv = names(mode.vesinv)[mode.vesinv==max(mode.vesinv)]

mode.gleason = table(factor(gleason))
new.gleason = names(mode.gleason)[mode.gleason==max(mode.gleason)]
```

#we check the class type of each variable in prostate

```
sapply(prostate, class)
subject    psa    cancervol    weight    age    benpros    vesinv    capspen    gleason
"integer" "numeric" "numeric" "numeric" "integer" "numeric" "integer" "numeric" "integer"
```

#create x.new to use in predict function**#we use means for cancervol and benpros and****#modes for vesinv and gleason**

```
x.new=data.frame(cancervol=mean(cancervol),benpros=mean(benpros),vesinv=new.vesinv,gleason=new.gleason)
```

```
> x.new
  cancervol benpros vesinv gleason
1  7.136218 2.588087    0        7
```

#predict

```
> ans = predict(fit.try3,newdata=x.new)
> ans
      1
2.37726
```

#now, log(psa)=2.37726**#We need psa. So, $psa = e^{2.37726}$**

```
> exp(ans)
      1
10.77533
```
