

Indian Institute of Technology Kanpur



SESSION:-2014-15

Regression Analysis Project

Predicting the chance of recurrence of Breast Cancer and the number of days it will recur in

Submitted to:
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DATA SET:

Title: Wisconsin Prognostic Breast Cancer (WPBC)

Source: Source Information

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b) Donor: Nick Street

c) Date: December 1995

Number of instances: 198

Number of attributes: 34 (ID, outcome, 32 real-valued input features)

Attribute information

- 1) ID number
- 2) Outcome (R = recur, N = nonrecur)
- 3) Time (recurrence time if field 2 = R, disease-free time if field 2 = N)
The Recurrence Surface Approximation (RSA) method is a linear programming model which predicts Time To Recur using both recurrent and nonrecurrent cases.

4-33) Ten real-valued features are computed for each cell nucleus:

- a) radius (mean of distances from center to points on the perimeter)
- b) texture (standard deviation of gray-scale values)
- c) perimeter
- d) area
- e) smoothness (local variation in radius lengths)
- f) compactness ($\text{perimeter}^2 / \text{area} - 1.0$)

- g) concavity (severity of concave portions of the contour)
- h) concave points (number of concave portions of the contour)
- i) symmetry
- j) fractal dimension ("coastline approximation" - 1)

Several of the papers are listed on the site

<http://www.google.com/url?q=http%3A%2F%2Farchive.ics.uci.edu%2Fml%2Fdatasets%2FBreast%2BCancer%2BWisconsin%2B%2528Prognostic%2529&sa=D&sntz=1&usg=AFQjCNHrBM4U7mSnJME2VjrRiN1IVUbgqQ> which contain detailed description of how these features are computed.

The mean, standard error, and "worst" or largest (mean of the three largest values) of these features were computed for each image, resulting in 30 features. For instance, field 4 is Mean Radius, field 14 is Radius SE, field 24 is Worst Radius.

Values for features 4-33 are recorded with four significant digits.

- 34) Tumor size - diameter of the excised tumor in centimeters
- 35) Lymph node status - number of positive axillary lymph nodes observed at time of surgery

Missing attribute values:

Lymph node status is missing in 4 cases.

Class distribution: 151 non-recur, 47 recur

Acknowledgements

We would like to thank our instructor of the course Dr. Sharmishtha Mitra for providing constant guidance and motivation for this project, without which it would have been an impossible task to accomplish.

Next we would like to thank the effort of the UCI machine learning repository team to give access to the innumerable datasets to students like us. A well structured data have been an important facet in the project.

Last but not the least we appreciate each others' contribution and hard work for the fulfillment of the project.

ABSTRACT

In this project we have tried to predict the number of patients who are at a chance of having the disease again. For this, we have used a logistic regression model to classify the patients into those who shall have breast cancer again in the future and those that shall not. From among the patients in our observation set who show recurrent breast cancer, we have used multiple linear regression model to predict the time period in which the disease shall recur.

The columns in the data set have been coded in the following way:

- rad=radius
- tex=texture
- peri=perimeter
- smooth=smoothness
- compac=compactness
- conc=concavity
- cpts=concave points
- sym=symmetry
- fd=facial dimension
- tum_size=tumor size
- lymph=lymph node status
- m=mean
- se=standard error
- w="worst" or largest(mean of the three largest values)

So rad_m implies mean radius of the nuclei, rad_se implies standard error from the nuclei radii values, etc.

Logistic Regression model

We fit a logistic regression model on our training dataset with 156 observations and 33 regressors and obtain the prediction obtained from our training data set[80% of the data]

```
fit=glm(formula = outcome ~ time + rad_m + tex_m + peri_m + area_m +  
smooth_m + compac_m + conc_m + cpts_m + sym_m + fd_m + rad_se +  
tex_se + peri_se + area_se + smooth_se + compac_se + conc_se +  
cpts_se + sym_se + fd_se + rad_w + tex_w + peri_w + area_w +  
smooth_w + compac_w + conc_w + cpts_w + sym_w + fd_w + tum_size +  
lymph, family = "binomial", data = train)
```

The number of observations misclassified =12.

The misclassification percentage=7.69%

The test set consists of 38 data points. The prediction obtained is shown below where 1 denotes that the disease is likely to happen again and 0 specifies otherwise.

[R code:

```
predict(fit, testdata, type=response) ]
```

```

157 158 159 160 161 162 163 164 165 166 167 168 169 170 171 172 173 174
0 0 0 0 0 0 0 0 0 0 0 0 0 0 1 0 1 0
175 176 177 178 179 180 181 182 183 184 185 186 187 188 189 190 191 192
0 0 1 0 0 0 0 0 0 0 0 0 0 0 1 0 0 1
193 194
0 0

```

And the number of observations misclassified=14

The misclassification percentage=36.84%

However the VIF's of some regressors are quite high, indicating high multicollinearity, hence we need to remove multicollinearity from the model by variance decomposition method.

[R code:

```
>library(faraway)
```

```
>vif(fit)
```

```
]
```

```

time      rad_m      tex_m      peri_m      area_m      smooth_m
102.38364 63796.38497 217.29630 67398.32418 11982.17207 479.09867
compac_m   conc_m   cpts_m      sym_m      fd_m      rad_se
1401.49209 1107.55100 1022.00796 237.12490 441.04466 3762.40524
tex_se     peri_se     area_se     smooth_se  compac_se  conc_se
207.63171 3822.36763 1278.34208 249.07073 1169.82759 1509.19851
cpts_se    sym_se     fd_se      rad_w      tex_w      peri_w
244.12018 465.43306 799.22344 19129.92818 352.19082 7207.77312
area_w     smooth_w    compac_w    conc_w     cpts_w     sym_w
10248.26733 264.43651 1132.72258 1047.78158 279.53559 597.17596
fd_w      tum_size    lymph
1054.13322 30.96778 40.46549

```

The model after removing successively removing collinearity using variance decomposition method.

```

outcome ~ time + tex_m + compac_m + tex_se + peri_se +
  compac_se + conc_se + cpts_se + sym_se + fd_se + area_w +
  conc_w + tum_size + lymph

```

The VIF's of the regressors:

```

time      tex_m      compac_m      tex_se      peri_se      compac_se      conc_se
17.75276 10.28624 45.40091 17.02112 35.26033 64.50147 60.46200
cpts_se    sym_se     fd_se      area_w     conc_w     tum_size     lymph
35.40312 16.86470 40.21718 18.62755 37.72544 10.15669 10.31293

```

The number of observations misclassified in the training data set of the model after removing collinearity=27

The misclassification percentage=17.31%

The prediction on the test data is as below.

157	158	159	160	161	162	163	164	165	166	167	168	169	170	171	172	173	174
1	0	1	0	0	0	0	0	0	0	1	0	0	0	0	0	0	1
175	176	177	178	179	180	181	182	183	184	185	186	187	188	189	190	191	192
1	0	1	1	0	0	0	0	0	1	1	1	1	0	0	1	1	0
193	194																
1	1																

The number of observations misclassified in the test data set=11

The misclassification percentage=28.94%

We further apply variable selection to reduce the number of regressors.

Model after removing collinearity and variable selection

After successively removing collinearity and variable selection, we obtain the model as:

outcome ~ time + compac_m + peri_se + conc_se + fd_se + conc_w

And the VIF's of the regressors obtained

time	compac_m	peri_se	conc_se	fd_se	conc_w
15.28565	35.46256	13.19941	32.81442	23.31598	29.28844

The number of observations misclassified in the training data set of the model =29

The misclassification percentage=18.56%

The prediction on the test data is as below.

157	158	159	160	161	162	163	164	165	166	167	168	169	170	171	172	173	174
1	0	1	0	0	0	0	0	1	1	1	1	0	0	0	0	0	1
175	176	177	178	179	180	181	182	183	184	185	186	187	188	189	190	191	192
1	0	1	0	1	1	0	0	0	1	1	1	1	1	1	0	1	0
193	194																
0	1																

The number of observations misclassified in the test data set=15

The misclassification percentage=39.47%

MLR Model

For MLR model, the actual dataset included only 46 observations with the initially 33 regressors. 80% of the data (36 observations) were used for training and rest for testing.

Issues associated:

- High number of regressors i.e $n \sim p$
- Multicollinearity
- Non-linear model

Step 1: To handle the data more efficiently an initial variable selection was done so that adjusted r-squared was maximum. This was performed using `regsubsets()` command from “car” library in R.

As a result, 26 variables were selected accordingly and the summary of the fit is as follows:

```
lm(formula = time ~ rad_m + tex_m + peri_m + area_m + compac_m +  
  conc_m + cpts_m + sym_m + fd_m + rad_se + tex_se + peri_se +  
  smooth_se + compac_se + conc_se + cpts_se + sym_se + tex_w +  
  peri_w + area_w + smooth_w + compac_w + conc_w + cpts_w +  
  sym_w + fd_w + tum_size, data = train)
```

Residuals:

Min	1Q	Median	3Q	Max
-7.873	-3.005	-1.278	3.149	12.749

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	1.299e+02	1.153e+02	1.127	0.292548
rad_m	1.448e+02	4.045e+01	3.579	0.007197 **
tex_m	-4.974e+00	2.573e+00	-1.933	0.089321 .
peri_m	-2.728e+01	6.451e+00	-4.228	0.002884 **
area_m	5.172e-01	1.176e-01	4.399	0.002289 **
compac_m	1.940e+03	4.635e+02	4.185	0.003058 **
conc_m	2.249e+03	3.849e+02	5.844	0.000386 ***
cpts_m	-5.072e+03	7.581e+02	-6.691	0.000154 ***
sym_m	8.418e+02	3.717e+02	2.265	0.053323 .
fd_m	-5.343e+03	2.222e+03	-2.405	0.042872 *
rad_se	-4.875e+02	1.078e+02	-4.523	0.001942 **
tex_se	-4.005e+01	1.767e+01	-2.267	0.053176 .
peri_se	7.601e+01	1.818e+01	4.180	0.003078 **
smooth_se	-1.455e+04	4.182e+03	-3.480	0.008314 **

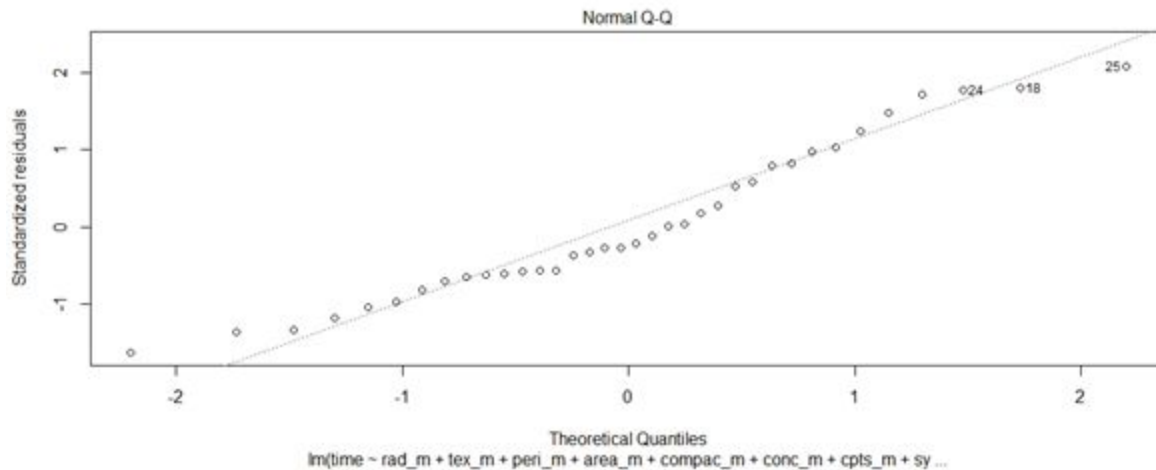
compac_se	-8.842e+02	6.444e+02	-1.372	0.207246
conc_se	-7.235e+03	1.228e+03	-5.891	0.000365 ***
cpts_se	2.503e+04	5.042e+03	4.965	0.001100 **
sym_se	2.847e+03	1.255e+03	2.268	0.053031 .
tex_w	5.667e+00	2.175e+00	2.606	0.031343 *
peri_w	-2.251e-01	1.666e+00	-0.135	0.895827
area_w	-4.229e-02	5.642e-02	-0.750	0.475014
smooth_w	2.259e+03	4.994e+02	4.523	0.001942 **
compac_w	-2.693e+02	9.563e+01	-2.816	0.022630 *
conc_w	3.180e+02	1.242e+02	2.561	0.033590 *
cpts_w	-1.582e+03	3.189e+02	-4.960	0.001107 **
sym_w	-5.330e+02	2.164e+02	-2.463	0.039144 *
fd_w	1.012e+03	7.664e+02	1.320	0.223309
tum_size	8.857e+00	1.821e+00	4.864	0.001250 **

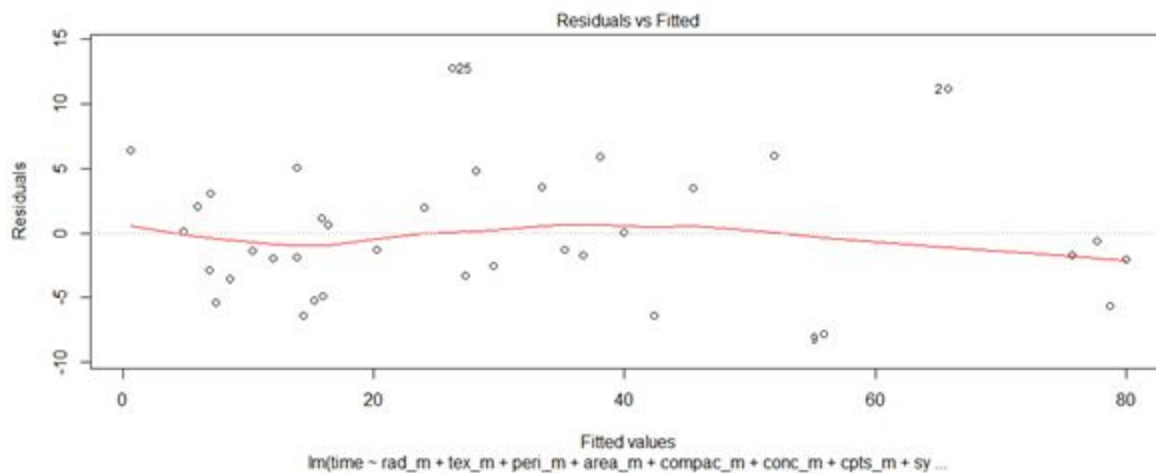
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 10.07 on 8 degrees of freedom

Multiple R-squared: 0.9588, Adjusted R-squared: 0.8196

F-statistic: 6.89 on 27 and 8 DF, p-value: 0.003978





The plots show a relatively better fitted model but the predicted values for the testing data did not give proper results.

[pred_init](#)

	37	38	39	40	41	42	43	44	45	46
	-0.695916	5.113816	35.780140	-23.043968	-5.652990	39.342355	99.413660	192.139541	62.405499	6.137621

Step2: Next we checked for multicollinearity using variance decomposition method, the same way we did in Logistic model.

Removal of variables was stopped when VIF's reached a value around 20 (max VIF=21.03), but variance decomposition gave non significant variance decomposition proportions for high VIF regressors.

The model obtained after removing multicollinearity is as follows:

Call:

```
lm(formula = time ~ tex_m + compac_m + conc_m + sym_m + tex_se +
    peri_se + smooth_se + compac_se + sym_se + area_w + smooth_w +
    compac_w + conc_w + cpts_w + fd_w + tum_size, data = train)
```

Residuals:

	Min	1Q	Median	3Q	Max
	-28.004	-14.492	1.234	12.045	31.476

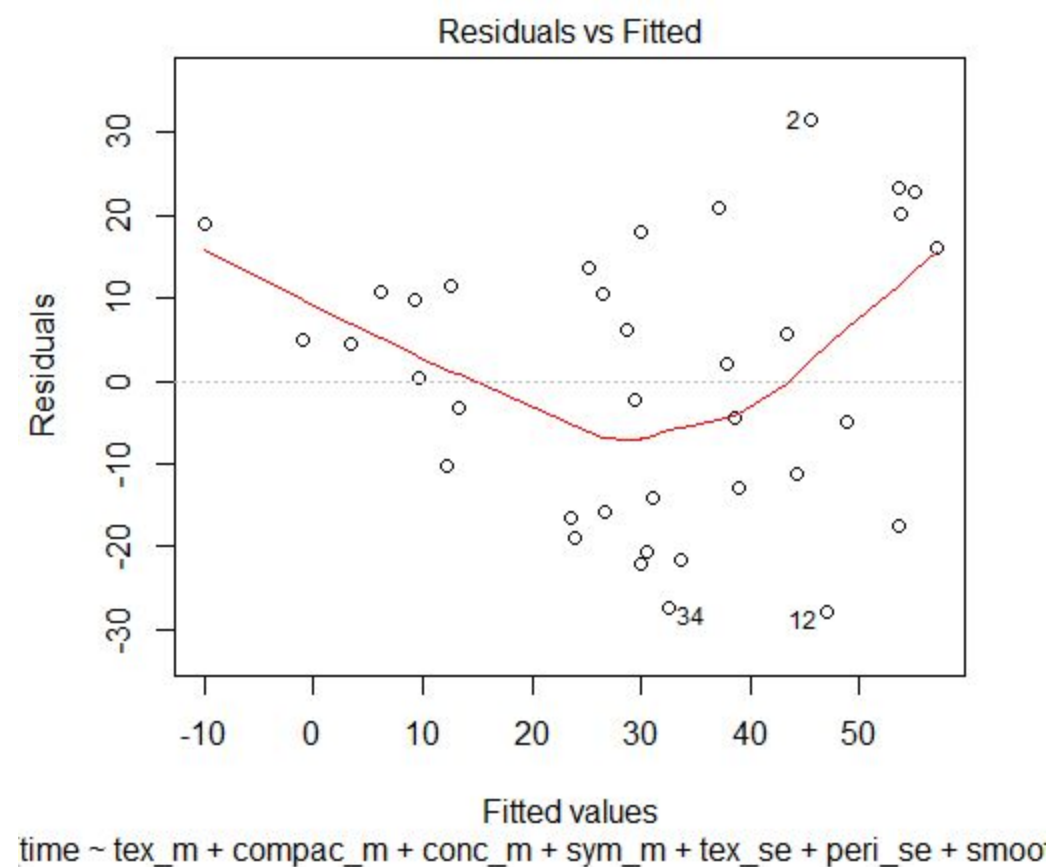
Coefficients:

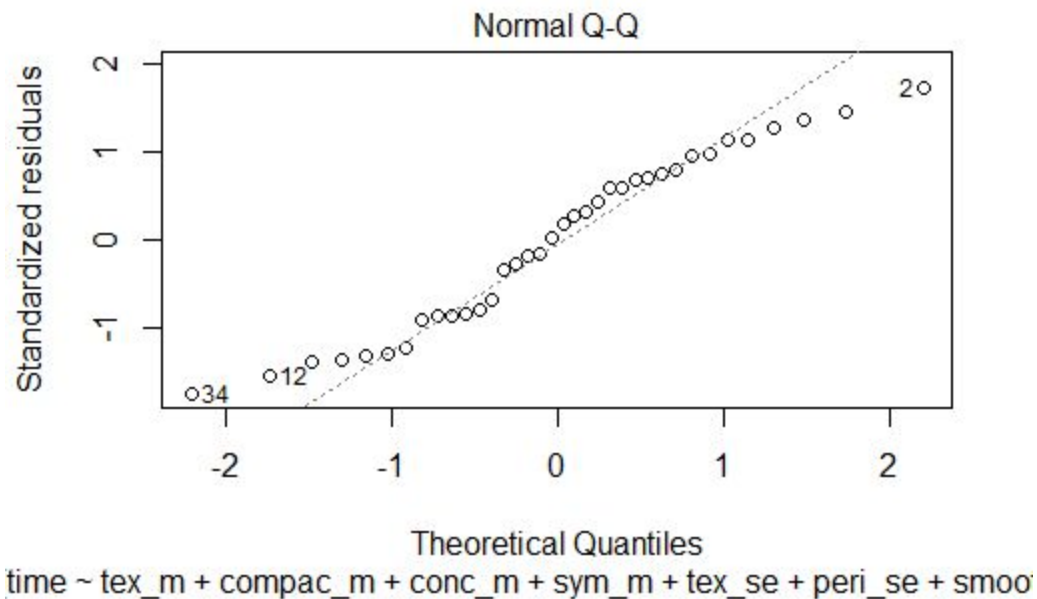
	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	4.496e+01	7.020e+01	0.640	0.5295
tex_m	-1.788e+00	1.530e+00	-1.169	0.2568
compac_m	-3.795e+02	3.842e+02	-0.988	0.3356
conc_m	2.692e+02	2.501e+02	1.076	0.2953
sym_m	3.575e+02	2.654e+02	1.347	0.1939

tex_se	2.713e+01	1.495e+01	1.815	0.0854
peri_se	6.189e+00	4.920e+00	1.258	0.2236
smooth_se	-2.825e+03	4.175e+03	-0.677	0.5068
compac_se	-5.289e+02	7.322e+02	-0.722	0.4789
sym_se	-1.047e+03	7.111e+02	-1.472	0.1574
area_w	-2.456e-02	1.681e-02	-1.461	0.1603
smooth_w	2.571e+02	5.271e+02	0.488	0.6313
compac_w	1.048e+02	1.279e+02	0.820	0.4227
conc_w	-4.978e+01	8.454e+01	-0.589	0.5629
cpts_w	-2.805e+02	2.382e+02	-1.177	0.2536
fd_w	-3.423e+01	6.591e+02	-0.052	0.9591
tum_size	4.182e-01	2.595e+00	0.161	0.8737

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

Residual standard error: 22.21 on 19 degrees of freedom
 Multiple R-squared: 0.5232, Adjusted R-squared: 0.1216
 F-statistic: 1.303 on 16 and 19 DF, p-value: 0.2882

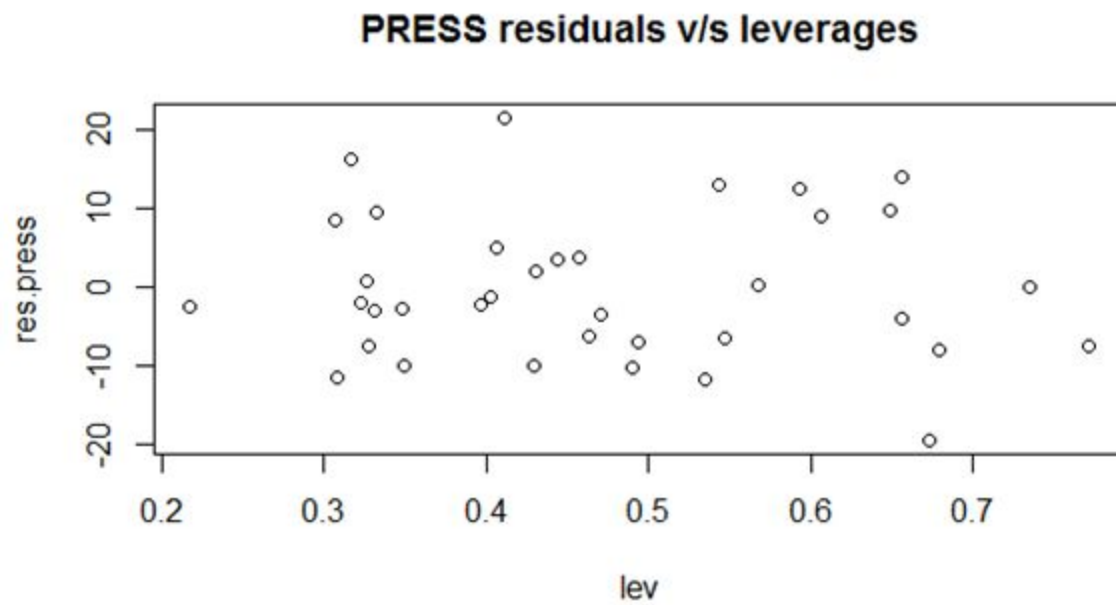
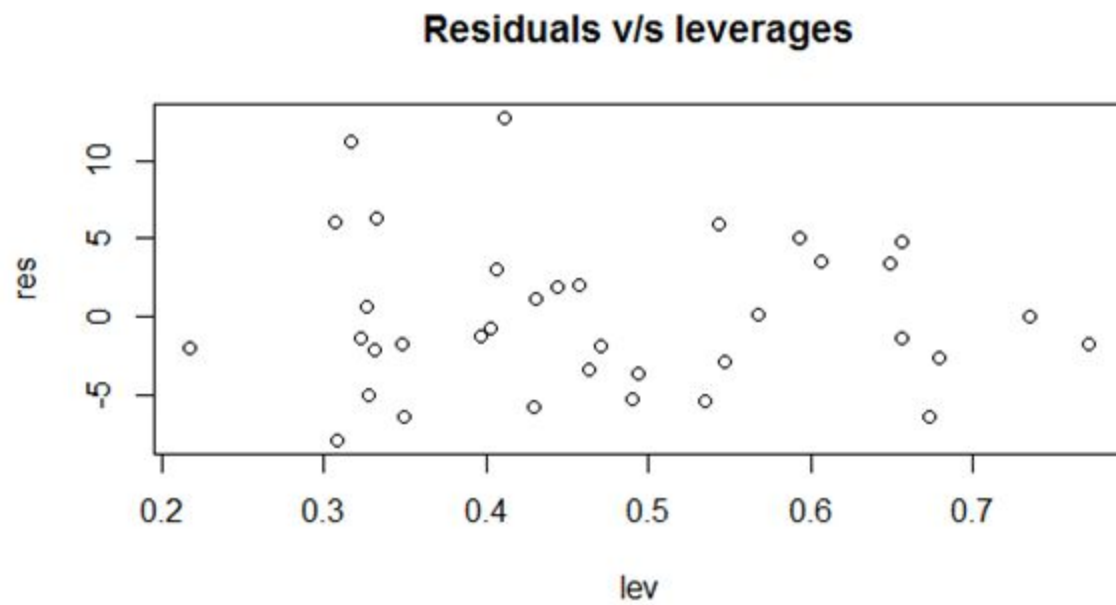




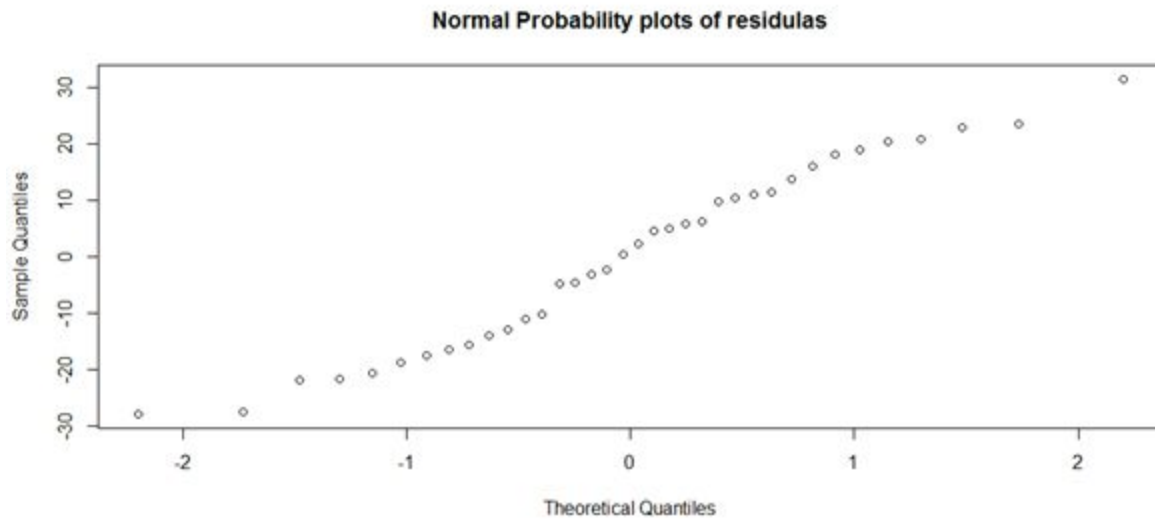
37 38 39 40 41 42 43 44 45
 1.694452 39.198678 12.546668 22.692806 42.760911 34.724093 29.558598 14.852857
 42.213418
 46
 9.471197

The model did not fit as well as the previous model but the prediction was better in this case. The residual plots and the q-q plots gave a clear idea of deviation from our assumptions of normality and presence of a linear model.

Step3: Residual analysis was done to check model inadequacies. PRESS residuals were calculated. Plots of leverage values (hii) and residuals were as follows. Also no leverage value came significantly different from others and were close to the mean value of 0.477



No influential observation was recorded as ordinary residuals and Press residuals had similar values.



Step4: The residual plot was assumed to be an opening funnel type and box-cox was applied to check for variance dependence. The lambda chosen was -1.9 by subsequent checking of SSRs for each iteration.

The final output showed that the assumption was wrong

Call:

```
lm(formula = ((time^lambda) - 1)/lambda ~ tex_m + compac_m +
    conc_m + sym_m + tex_se + peri_se + smooth_se + compac_se +
    sym_se + area_w + smooth_w + compac_w + conc_w + cpts_w +
    fd_w + tum_size, data = train)
```

Residuals:

Min	1Q	Median	3Q	Max
-0.060048	-0.007831	0.002109	0.008929	0.031181

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	4.569e-01	6.816e-02	6.704	2.08e-06 ***
tex_m	-2.214e-03	1.485e-03	-1.491	0.1524
compac_m	-7.574e-02	3.730e-01	-0.203	0.8412
conc_m	-2.598e-01	2.428e-01	-1.070	0.2981
sym_m	9.306e-02	2.577e-01	0.361	0.7220
tex_se	2.665e-02	1.451e-02	1.837	0.0820 .
peri_se	-4.819e-03	4.776e-03	-1.009	0.3257
smooth_se	2.089e+00	4.053e+00	0.515	0.6122
compac_se	4.048e-01	7.109e-01	0.569	0.5758
sym_se	7.445e-02	6.905e-01	0.108	0.9153
area_w	3.070e-05	1.632e-05	1.881	0.0753 .

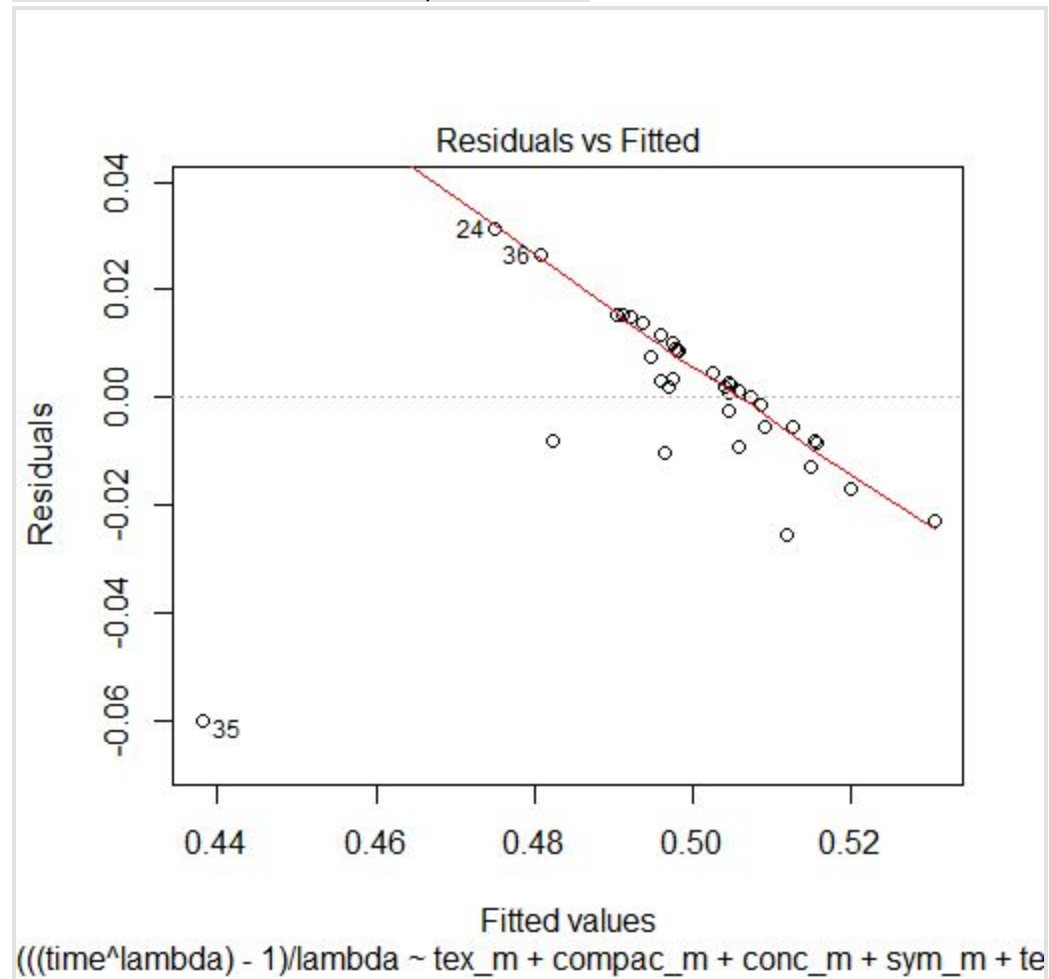
smooth_w	2.798e-01	5.118e-01	0.547	0.5909
compac_w	-5.431e-03	1.242e-01	-0.044	0.9656
conc_w	7.446e-02	8.208e-02	0.907	0.3757
cpts_w	-2.730e-01	2.313e-01	-1.180	0.2525
fd_w	1.297e-01	6.399e-01	0.203	0.8415
tum_size	1.793e-03	2.520e-03	0.712	0.4853

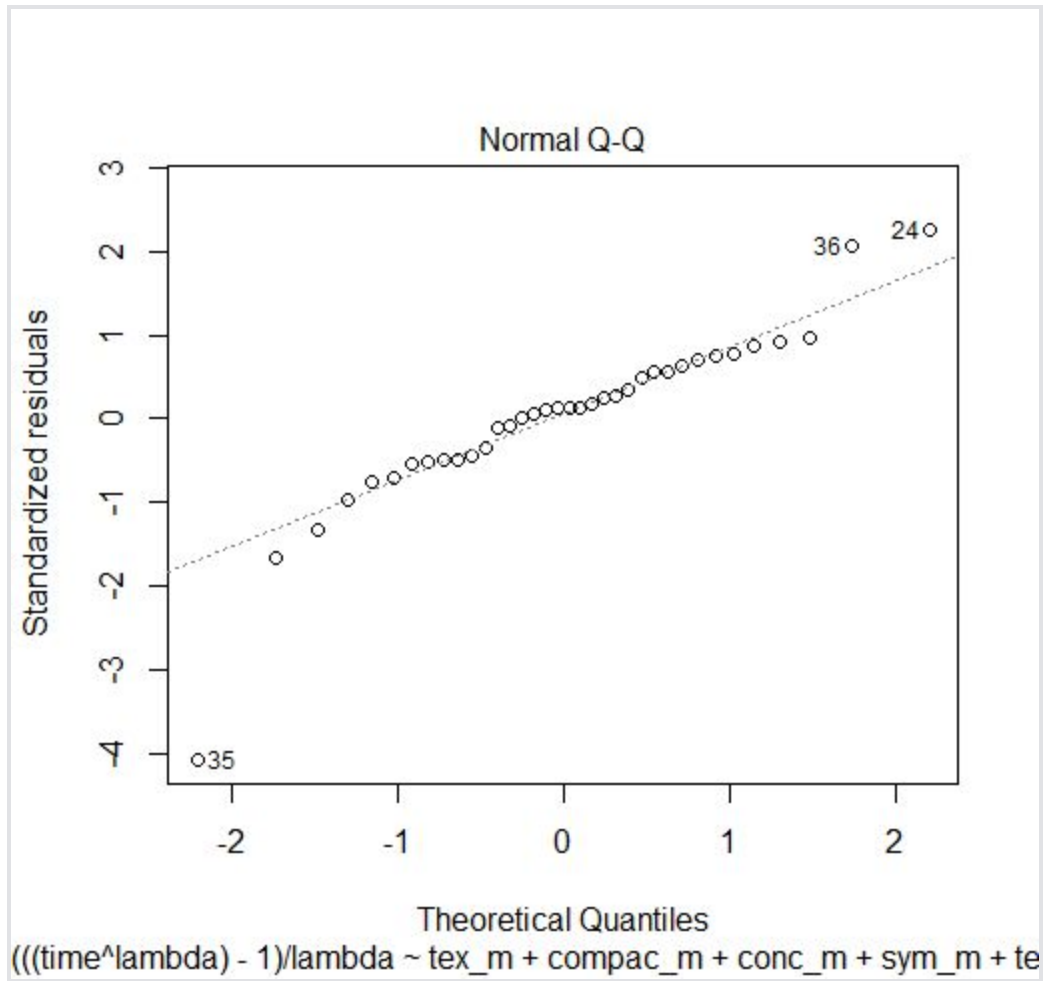
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.02157 on 19 degrees of freedom

Multiple R-squared: 0.4827, Adjusted R-squared: 0.04711

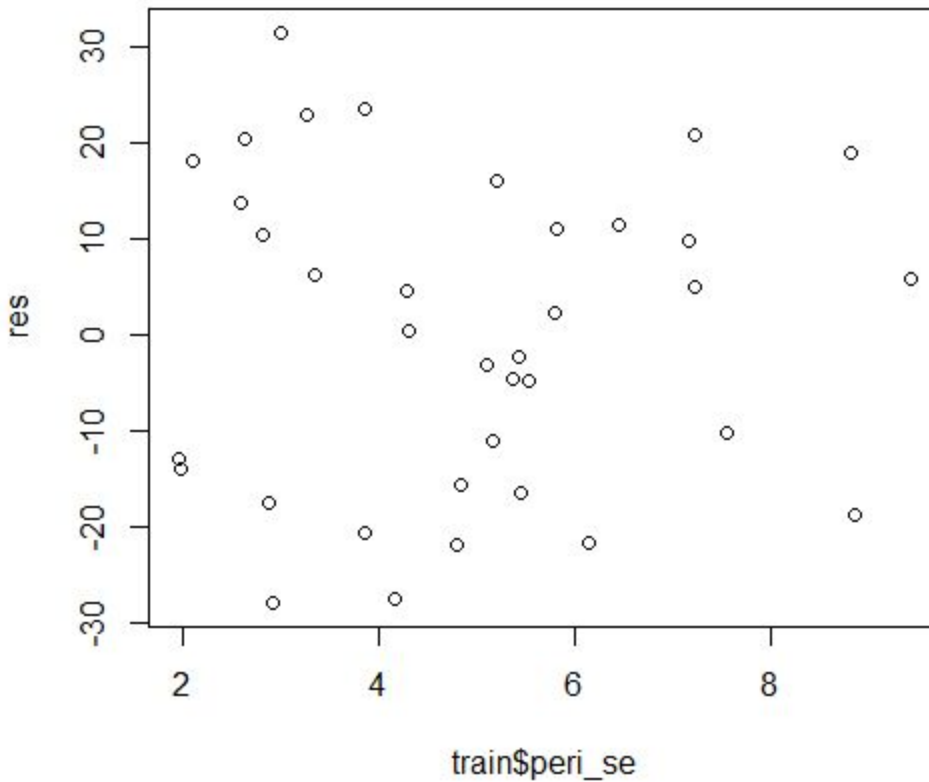
F-statistic: 1.108 on 16 and 19 DF, p-value: 0.411





Step5: Non-linearity was assumed finally. The fit of residuals v/s regressor for each regressor(in the model or not) was compared with residuals v/s fitted.

One such example:



Since the number of observations were less, we could not make a proper estimation of the regressor that could be creating the non-linearity.

Results and conclusions.

The final model that was considered was the one after removing multicollinearity as it gave the best predictions among all the models and our basic aim was prediction only. But still there were significant differences from the actual values.

```
model: model2<-lm(formula = time ~ tex_m + compac_m +  
  conc_m + sym_m + tex_se + peri_se +  
  smooth_se + compac_se + sym_se +  
  area_w + smooth_w + compac_w + conc_w + cpts_w +  
  fd_w + tum_size, data = train)
```

predictions:

predicted	actual
1.69	1
39.19	9
12.54	16
22.69	9
42.760	14
34.724	12
29.55	11
14.85	7
42.21	14
9.47	1

The data available is not a good dataset for prediction as it gives just 10% accuracy, but it is a good dataset to predict whether the cancer will recur in a particular patient in due time. Some other parameters would have been needed for making a better prediction.