2.1: Introduction to Simple Linear Regression

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See Handout 2.1.1 for information regarding the Toluca power company example.

1 Why Linear Regression?

Linear regression is good for:

- **Inference:** determine if there is a statistically significant linear relationship between two variables, while possibly accounting for the effect of additional variables.
 - Example: after accounting for the effects of square footage and age, are lot size and home sale price significantly linearly related?
- **Prediction:** use variables that are "easy" to measure to predict variables that are harder to measure.
 - Example: Use elevation (easy to measure) to predict annual snow accumulation (hard to measure).

Linear regression only works for variables that share a statistical relationship.

Terminology:

- Y response variable
- X_i predictor variables
- ϵ error (or difference) term
- β_i model parameters (true values are unknown and are estimated)

Linear Regression focuses on finding appropriate estimates of the model parameters (b_i) :

The idea is that we want to select paramater estimates that make the predicted values of Y (\hat{Y}) close to the actual values of Y.

2 Ordinary Least Squares (OLS) Regression

If assumptions regarding residuals are satisfied (more in Handout 2.2), then the OLS estimates of the model parameters are "best."

What does it mean to be "best"?

- unbiased given an infinite number of different samples of data, the average of my estimates will be equal to true (and unknown) value of the parameter.
 - In other words, my estimates are "centered" on the truth.
- minimum variance the variation in the estimate from sample to sample is the smallest of all possible estimation methods.

Applications - Toluca Example:

Let X represent the lot size and let Y represent the total work hours. Based on the initial scatterplot, we assume that the relationship between X and Y can be modeled as

$$Y = \beta_0 + \beta_1 X + \epsilon$$

OLS seeks to minimize:

$$Q = \sum_{i=1}^{n} \left(Y_i - \hat{Y}_i \right)^2,$$

which requires us to select estimates b_0 and b_1 that minimize

$$Q = \sum_{i=1}^{n} (Y_i - (b_0 + b_1 X_i))^2 = f(\mathbf{X}).$$

We can use multivariable calculus to find the minimum of Q by finding the critical points, i.e.

$$\nabla Q = \nabla f(\mathbf{X}) = 0.$$

The single critical point that minimizes Q is

$$b_{1} = \frac{\sum_{i} (X_{i} - \bar{X})(Y_{i} - \bar{Y})}{\sum_{i} (X_{i} - \bar{X})^{2}}$$
$$b_{0} = \bar{Y} - b_{1}\bar{X}.$$

Obtain OLS estimates automatically in SAS with:

```
proc reg data=toluca;
  model workhours = lotsize;
  title1 'Simple linear model';
run:
```

Equation Estimates:

$$b_0 = 62.37, b_1 = 3.57$$

Model Equation:

$$\hat{Y} = 62.37 + 3.57(lotSize)$$

The Critical Assumption

OLS least squares hinges on the assumption that

$$\epsilon \stackrel{iid}{\sim} N(0, \sigma^2)$$

- **independent:** Knowing the value of one of the model residuals tells you nothing about any of the others.
- identically distributed: All of the residuals come from the same distribution.
- Normal Distribution: The model residuals follow a normal (bell shaped) distribution.

- **zero mean:** The average of the residuals is zero (unbiased estimates).
- **constant variance:** The spread of the residuals about the line is the same across the range of X and the range of predicted values.

If the assumptions hold, then the simple linear regression can be visualized as in Figure 1.

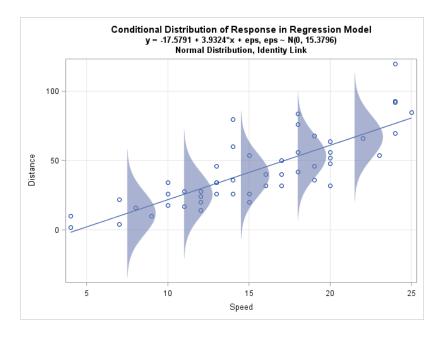


Figure 1: Sample visualization taken from Rick Wicklin on The DO Loop.

In other words, Y follows a normal distribution with a center that is conditional on X.

Estimating σ

Estimating the variance about the regression line:

- Allows us to get a measure of the model fit: lower relative MSE \rightarrow better model.
- All significance tests of model coefficients are based on our estimate of σ .

Estimation of ϵ in Theory

Suppose that $\epsilon_1, \epsilon_2, \dots, \epsilon_n$ were an observed sample from some population. (In practice, ϵ is estimated as the residuals of our OLS model, represented as e_i .)

We could then estimate $Var(\epsilon)$ as

$$\frac{1}{n-1} \sum_{i=1}^{n} \left(\epsilon_i - \bar{\epsilon} \right)^2$$

Note that the variance calculation requires the estimation of $\mu_{\epsilon} = \bar{\epsilon} = \frac{1}{n} \sum_{i} \epsilon_{i}$.

This calculation "constrains" one of the ϵ_i . This means that if we know epsilon and $\epsilon_1, \ldots \epsilon_{n-1}$, then we can know ϵ_n .

We call the number of unconstrained observations the "degrees of freedom" (DF).

Every time you estimate a parameter, you lose one degree of freedom.

Think of observations as currency. We spend money to estimate things and our degrees of freedom are the leftover cash.

Estimation of ϵ in Practice

Why is it that we can't directly obtain the values of ϵ ?

We don't know the true regression line, so we cannot know the true values of epsilon.

We can obtain estimates of the residuals e_i through the regression line:

$$e_i = Y_i - \hat{Y}_i = Y_i - (b_0 + b_1 X_i).$$

OLS, by design, makes $\sum_i e_i = 0 \rightarrow \bar{e} = 0$, meaning I don't have to spend any DF to obtain \bar{e} .

Variance of the Residuals

$$\hat{\sigma}^2 = s^2 = \frac{1}{df_E} \sum_{i=1}^n e_i^2 = \frac{1}{n-2} \sum_{i=1}^n (e_i - \bar{e})^2$$

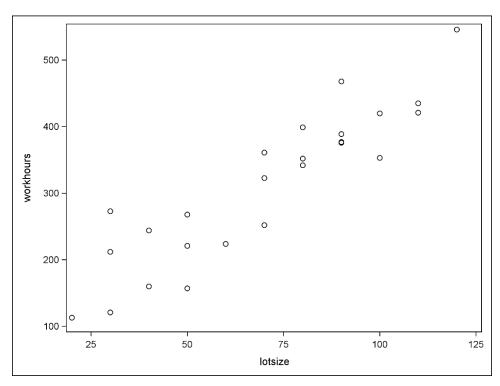
We call this estimate the "mean square error" or MSE.

2.1.1: SAS: Simple Linear Regression

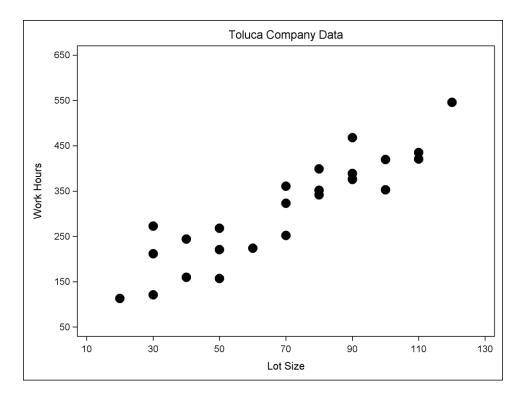
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Example: The Toluca Company makes replacement parts for refrigeration equipment. For a certain part, it takes some time to set up the production process, and then the production of a given lot size can begin. As part of a cost improvement program, the company wished to better understand the relationship between the lot size (X) and the total work hours (Y). Data were reported for 25 representative lots of varying size.

```
/* Input data */
data toluca; input lotsize workhours @@; cards;
  80
      399
           30
               121
                    50
                        221
                             90
                                 376
                                      70
                                          361
                                                   224
                                               60
 120
      546
           80
               352
                   100
                        353
                             50
                                 157
                                      40
                                          160
                                               70
                                                   252
      389
                   110
                            100
                                 420
                                          212
  90
           20
               113
                        435
                                      30
                                               50
                                                   268
  90
      377
          110
               421
                    30
                        273
                             90
                                 468
                                      40
                                          244
                                               80
                                                   342
  70
      323
  ;
run;
/* Make a scatterplot of Y=workhours and X=lotsize */
proc sgplot data=toluca;
  scatter x=lotsize y=workhours ;
run;
```



```
/* Be professional -- make it look nice */
proc sgplot data=toluca;
    scatter x=lotsize y=workhours /
        markerattrs=(symbol=CIRCLEFILLED size=3pt);
    title1 'Toluca Company Data';
    xaxis label='Lot Size' values=(10 to 130 by 20);
    yaxis label='Work Hours' values=(50 to 650 by 100);
run;
```



```
/* Look at correlation between these variables */
proc corr data=toluca;
    var workhours lotsize;
run;
```

Toluca Company Data

The CORR Procedure

2 Variables: workhours lotsize

Simple Statistics							
Variable	N	Mean	Std Dev	Sum	Minimum	Maximum	
workhours	25	312.28000	113.13764	7807	113.00000	546.00000	
lotsize	25	70.00000	28.72281	1750	20.00000	120.00000	

Pearson Correlation Coefficients, N = 25 Prob > r under H0: Rho=0					
	workhours lotsize				
workhours	1.00000	0.90638			
		<.0001			
lotsize	0.90638	1.00000			
	<.0001				

```
/* Now fit simple linear model with Y=workhours and
X=lotsize */
proc reg data=toluca;
  model workhours = lotsize;
  title1 'Simple linear model';
run;
```

Simple linear model

The REG Procedure Model: MODEL1

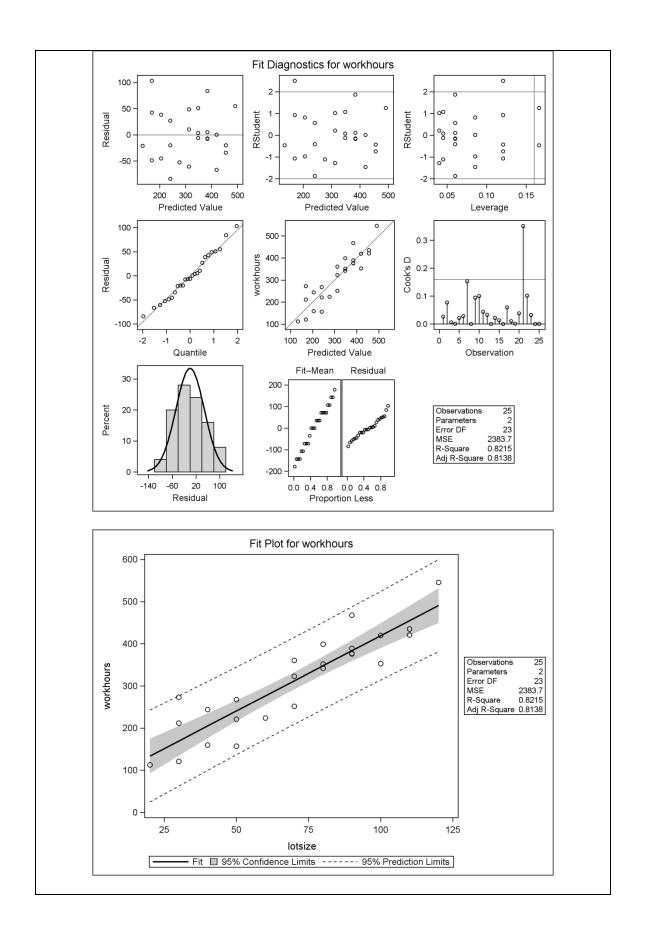
Dependent Variable: workhours

Number of Observations Read	25
Number of Observations Used	25

Analysis of Variance						
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F	
Model	1	252378	252378	105.88	<.0001	
Error	23	54825	2383.71562			
Corrected Total	24	307203				

Root MSE	48.82331	R-Square	0.8215
Dependent Mean	312.28000	Adj R-Sq	0.8138
Coeff Var	15.63447		

Parameter Estimates					
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t
Intercept	1	62.36586	26.17743	2.38	0.0259
lotsize	1	3.57020	0.34697	10.29	<.0001



```
/* See predicted values */
proc reg data=toluca noprint;
  model workhours = lotsize;
  output out=PredictedValues p=Predict;
proc print data=PredictedValues;
  title1 'Predicted Values';
run;
```

Predicted Values					
	Obs	lotsize	workhours	Predict	
	1	80	399	347.982	
	2	30	121	169.472	
	3	50	221	240.876	
	•••				
	24	80	342	347.982	
	25	70	323	312.280	

2.2: Diagnostics and Remedial Measures

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1 Why Diagnostics

Recall that the nice properties of the OLS coefficient estimates relied on the assumption that

$$\epsilon \stackrel{iid}{\sim} N(0, \sigma^2).$$

Model Assumptions in Linear Regression

- 1. X and Y share a linear relationship
 - X and Y can be related in a non-linear way, but OLS regression cannot be used in this case
- 2. model describes all observations
 - no outliers or influential points
- 3. additional predictor variables are unnecessary
 - there is no additional information to "extract" from ϵ
- 4. ϵ 's follow a normal distribution
 - Crucial for small sample sizes, not so critical for large (> 500) sample sizes due to central limit theorem.
- 5. ϵ 's have constant variance
- 6. ϵ 's are independent (possibly related to item #3)

We check assumptions using **diagnostics** and fix violated assumptions using **remedial measures**. Violations are most apparent in the **error terms** $(\epsilon_1, \ldots, \epsilon_n)$ so we focus on **residuals** (e_1, \ldots, e_n) .

There are both **graphical** and **numerical** checks of the assumptions regarding residuals, but the graphical assumptions are more informative.

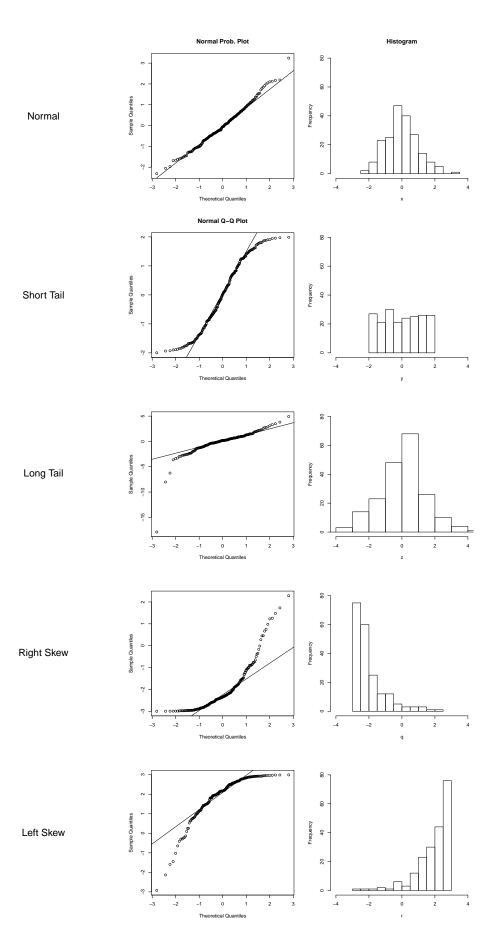


Figure 1: Example scatterplot and histograms for different data distributions.

2 Graphical Diagnostics

- Boxplot: quick way to check for the symmetry of residuals.
- **Histogram:** Way to check the shape of a distribution.
 - SAS will overlay a normal curve on the histogram of residuals to help check for normality.
- Normal Probability Plot: a qq-plot where the quantiles of the data are compared to the expected quantiles under a normal distribution
 - Expected values under normality have a mean of 0 and a SD = $\sqrt{\text{MSE}}$.
 - See page 111 in textbook for method for details about how to approximate expected observations under normality.
 - If data are approximately normal, the residuals in the Normal Probability Plot should closely follow a straight line.
- Sequence Plot: Line plot with residual values on the X axis and observation number on the X-axis.
 - Can "connect" the dots because there is only one Y value for every X value.
 - Look for patterns in the residuals across time/observation number.
 - * Patterns would suggest that the residuals are **not independent**.

• Residual Plot

- Plot e vs X or e vs \hat{Y}
- Look for non-linearity and or non-constant variance in these scatterplots.

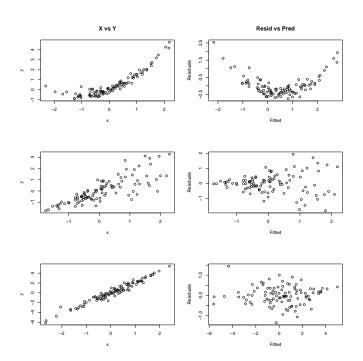


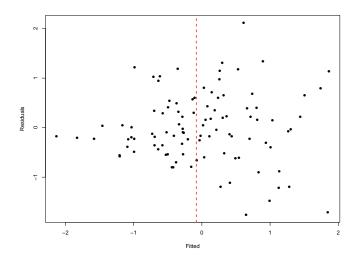
Figure 2: Plots showing 1) non-linearity, 2) non-constant variance, and 3) satisfied assumptions.

3 Numerical Diagnostics

Numerical diagnostics seek to determine if violations of model assumptions are statistically significant.

3.1 Some Numerical Tests

• Broth-Forsythe (BF) test of constant variance:



- Split the data into two groups based on the median predicted value.
- Calculate the median absolute deviation (MAD) $d_i = |e_i \tilde{e}|$, where \tilde{e} is the median within each group (lower and upper).
- Conduct a two-sample t-test of d's to determine if the average of d_i 's within each group are significantly different.
- Null Hypothesis: The variance of ϵ is constant. (Estimated with residuals e).

Toluca Example: BF p-value is .2, which suggests there is not significant evidence of non-constant variance.

- Correlation Test of Normality
 - Calculate correlation between observed e's and the "normal-expected" e's (e*). Similar to expected residuals in applot.
 - Null hypothesis: ϵ follows a normal distribution.
 - If the correlation isn't at least as big as the cirtical value for $\alpha = 0.05$ in Table B.6 for a given sample size n, then reject H_0 .
 - NOTE: The p-values provided in the SAS macro output mean *nothing* for the correlation test of normality.

Toluca Example: Correlation of 0.992 > 0.96, so there is not significant evidence of non-normality.

- F-test for lack of fit (test of linearity between X and Y)
 - See textbook 3.7 for details.
 - Requires multiple observations at one or more X-levels. (Hard to do in observational studies or studies with multiple X-variables).
 - Basically, the test compares the regression predictions to the empirical average of Y at X-levels with multiple observations.
 - Null hypothesis: The regression function is linear.

Toluca Example: p-value 0.69 > .05 suggests there is no significant evidence of non-linearity.

TABLE B.6
Critical Values
for Coefficient
of Correlation
between
Ordered
Residuals and
Expected
Values under
Normality
when
Distribution of
Error Terms
Is Normal.

		Level of Significance $lpha$						
n	.10	.05	.025	.01	.005			
5	.903	.880	.865	.826	.807			
6	.910	.888	.866	.838	.820			
7	.918	.898	.877	.850	.828			
8	.924	.906	.887	.861	.840			
9	.930	.912	894	.871	.854			
10	.934	.918	.901	.879	.862			
12	942	.928	.912	.892	.876			
14	.948	.935	.923	.905	.890			
16	.953	.941	.929	.913	.899			
18	.957	946	.935	.920	.908			
20	.960	.951	.940	.926	.916			
22	.963	.954	.945	.933	.923			
24	.965	.957	.949	,937	.927			
26	.967	.960	.952	,941	.932			
-28	.969	.962	.955	.944	.936			
30	.971	.964	957	.947	.939			
40	.977	.972	.966	.959	.953			
∄50	.981	.977	.972	.966	.961			
60	984	.980	.976	.971	.967			
70	.986	.983	.979	.975	.971			
80	.987	.985	.982	.978	.975			
90	.988	.986	.984	.980	.977			
100	.989	.987	.985	.982	.979			

Source: Reprinted, with permission, from S. W. Looney and T. R. Gulledge, Jr., "Use of the Correlation Coefficient with Normal Probability Plots," *The American Statistician* 39 (1985), pp. 75–79.

4 Remedial Measures

If assumptions are violated, your options are:

• Give up (at least on linear regression).

- Alternatives to OLS like Regression Trees, Quantile Regression etc. (more later in the semester).
- Depending on the X vs Y relationship, try non-linear regression (more later in the semester).
- For non-normality and heteroskedasticity: Variable Transformations on X or Y (not e).
 - Sometimes, a combination of transformations on both X and Y variables is needed.
 - NOTE: Removing "outlier" points from a model should be seen as a measure of last resort.

Box-Cox Approach

Great starting point to determine candidate transformations, but no "magical" solution.

• Define new response variable

$$Y' = \begin{cases} \operatorname{sign}(\lambda)Y^{\lambda} & \lambda \neq 0\\ \log(Y) & \lambda = 0 \end{cases}$$

(Note that $sign(\lambda)$ preserves the original ordering of the response variable).

• Consider the theoretical model

$$Y_i^{\lambda} = \beta_0 + \beta_1 X_i + \epsilon_i$$

- Consider a set of candidate lambda values, use maximum likelihood estimation to determine the "best value."
 - Maximum likelihood estimation: "which value of λ is the most likely, given the data that I have observed?
- When possible: pick an *interpretable* transformation that is close to the transformation recommended by SAS.

Ex: if $\lambda = .009$ is recommended, probably go with $\lambda = 0 \rightarrow \log(\lambda)$

In SAS:

Omitted Predictors

Think of regression as a form of data mining: we want to extract as much information as we can from our data.

If we failed to extract all the information from the data, this may show up as a trend in the plot the residuals (which SHOULD have a constant mean of 0).

We will discuss more about how to extract *time* related information in data at the end of the semester.

If you apply multiple methods to fix violations of assumptions, make sure to check that the final model actually fixed the violations of assumptions.

2.2.1: SAS - Residual Diagnostics

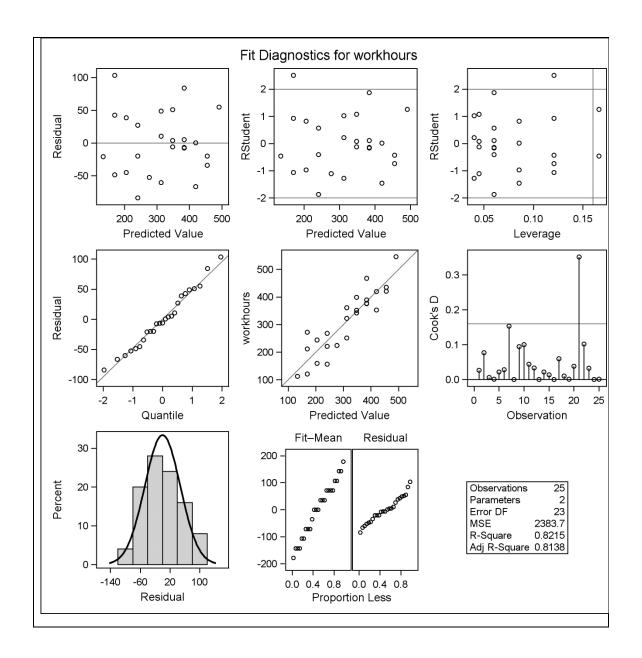
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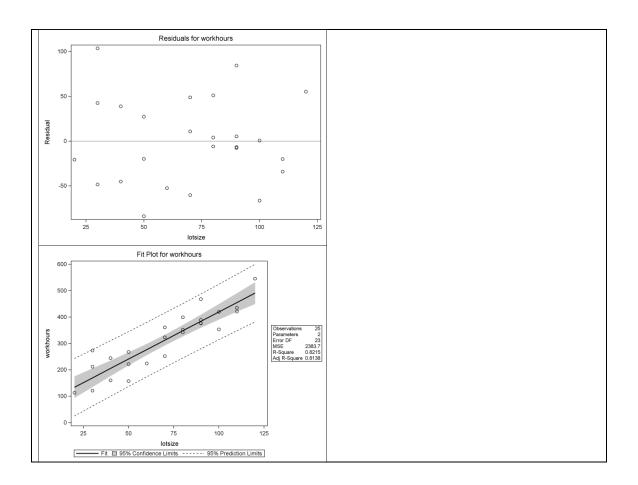
Example: (The Toluca Company data from Handout #2).

```
/* Input Toluca data (recall Ch. 1 example) */
data toluca; input lotsize workhours @@; cards;
             30
                           221
                                 90
                                                     60
                                                         224
   80
      399
                121
                       50
                                     376
                                           70 361
  120
       546
             80
                 352
                      100
                           353
                                 50
                                     157
                                           40 160
                                                     70 252
                113 110
                                                     50 268
   90
      389
                          435
                               100 420
             20
                                           30 212
                                           40 244
                                                     80 342
   90
      377 110 421
                       30 273
                                 90 468
   70
      323
run;
/* Now fit simple linear model with Y=workhours and X=lotsize,
  with residuals and predicted values saved in data set
   tolucaout */
proc reg data=toluca;
 model workhours = lotsize;
  output out=tolucaout r=resid p=pred;
  title1 'Simple linear model';
run;
```

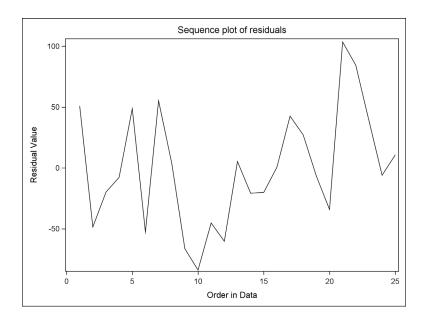
Analysis of Variance						
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F	
Model	1	252378	252378	105.88	<.0001	
Error	23	54825	2383.71562			
Corrected Total	24	307203				

Parameter Estimates						
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t	
Intercept	1	62.36586	26.17743	2.38	0.0259	
lotsize	1	3.57020	0.34697	10.29	<.0001	





```
/* Look at sequence plot */
data temp; set tolucaout;
  order = _n_;
proc sgplot data=temp;
  series x=order y=resid / lineattrs=(pattern=solid);
  xaxis label='Order in Data';
  yaxis label='Residual Value';
  title1 'Sequence plot of residuals';
run;
```



```
/*********** Numerical Diagnostics *********/
/* F-test for lack of fit */
proc rsreg data=toluca;
  model workhours = lotsize / lackfit covar=1 noopt;
  title1 'F-test for lack of fit';
run;
```

Residual	DF	Sum of Squares	Mean Square	F Value	Pr > F
Lack of Fit	9	17245	1916.069540	0.71	0.6893
Pure Error	14	37581	2684.345238		
Total Error	23	54825	2383.715617		

```
/** Brown-Forsythe and Correlation Test of Normality (shortcut)
**/
/* Two [unused] ways to access shortcut:
    filename macrourl "C:\[filepath]\resid num diag.sas";
     %include macrourl;
 */
%macro resid num diag(dataset,datavar,label= ...
/*
  This resid num diag.sas file provides a convenient shortcut
   to obtaining numerical checks of residuals from
   a fitted linear regression model.
   The macro takes five arguments:
    dataset is the name of the data set
    datavar is the name of the variable in the data set
             for which numerical diagnostics are desired
             (usually a residual)
    label is a character string for detail in output
    predvar is the name of the variable (usually predicted
          value) on which to sort for the Brown-Forsythe test
          (t-statistic and p-value reported)
    predlabel is the character string for detail in output
          related to the predvar variable
*/
```

```
/* Call the shortcut: */
%resid_num_diag(dataset=tolucaout, datavar=resid,
    label='residual', predvar=pred, predlabel='predicted');
```

P-value for Brown-Forsythe test of constant variance in residual vs. predicted

Obs	t_BF	BF_pvalue
1	1.31648	0.20098

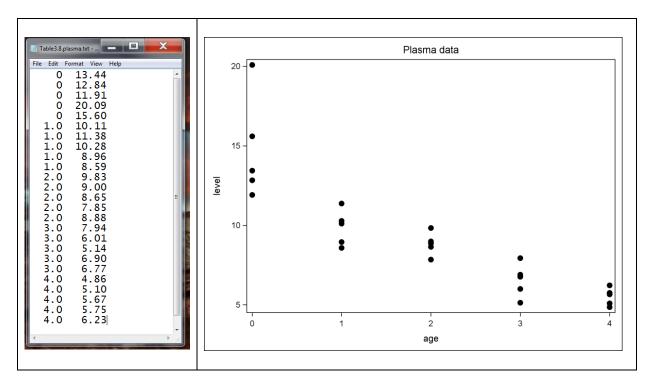
Output for correlation test of normality of residual (Check text Table B.6 for threshold)

Pearson Correlation Coefficients, N = 25 Prob > r under H0: Rho=0				
	resid	expectNorm		
resid residual	1.00000	0.99151 <.0001		
expectNorm	0.99151 <.0001	1.00000		

2.2.2: SAS - Linear Regression Remedial Measures

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<u>Example</u>: Age and plasma level for 25 healthy children in a study are reported. Of interest is how plasma level depends on age. (Text Table 3.8 – first column is age; second column is plasma level)



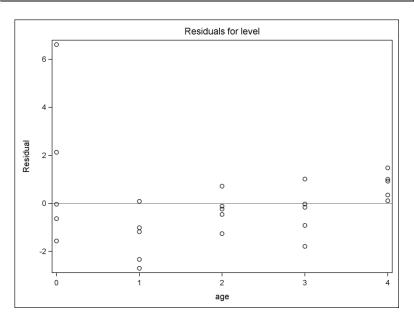
```
/*
data plasma;
   infile "[File Path]/Table3.8.plasma.txt";
   input age level;
run;
*/
data plasma; input age level @@; cards;
                                                           15.60
      13.44
                0
                   12.84
                             0
                                11.91
                                          0
                                              20.09
      10.11
              1.0
                                                            8.59
 1.0
                   11.38
                           1.0
                                10.28
                                        1.0
                                               8.96
                                                     1.0
 2.0
       9.83
              2.0
                    9.00
                           2.0
                                               7.85
                                                     2.0
                                                            8.88
                                  8.65
                                        2.0
 3.0
       7.94
                    6.01
                                  5.14
                                               6.90
                                                     3.0
                                                            6.77
              3.0
                           3.0
                                        3.0
 4.0
       4.86
              4.0
                    5.10
                          4.0
                                  5.67
                                        4.0
                                               5.75
                                                     4.0
                                                            6.23
  ;
/* Fit regression model and check assumptions */
proc reg data=plasma;
  model level = age;
  output out=out1 r=resid p=pred;
  title1 'Simple model for plasma data';
run;
```

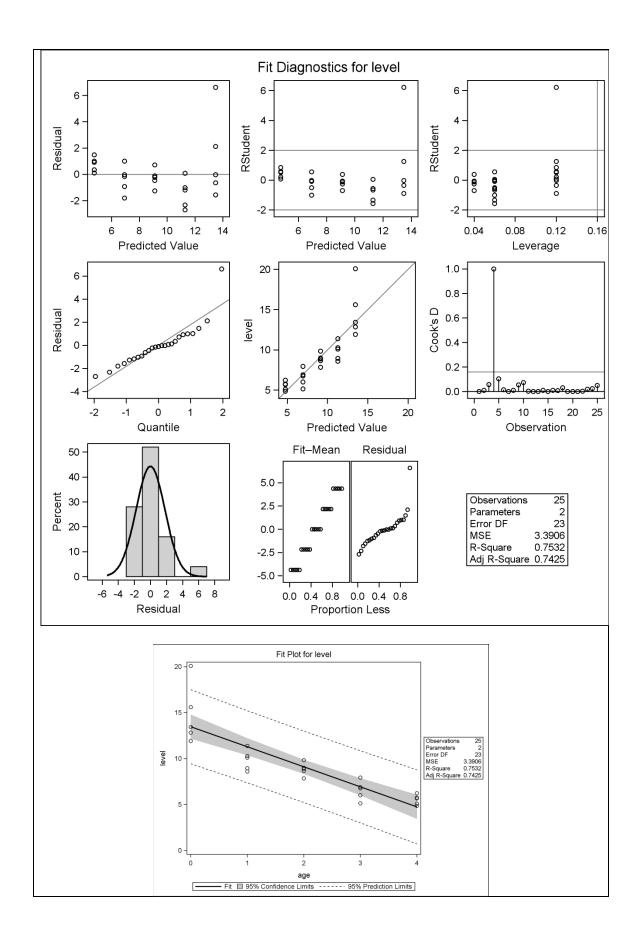
Simple model for plasma data

Analysis of Variance					
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F
Model	1	238.05620	238.05620	70.21	<.0001
Error	23	77.98306	3.39057		
Corrected Total	24	316.03926			

Root MSE	1.84135	R-Square	0.7532
Dependent Mean	9.11120	Adj R-Sq	0.7425
Coeff Var	20.20974		

Parameter Estimates					
Variable DF Parameter Estimate Standard Error t Value Pr >				Pr > t	
Intercept	1	13.47520	0.63786	21.13	<.0001
age	1	-2.18200	0.26041	-8.38	<.0001





```
/* Define shortcut macro, using line copied from
    [File Path]/resid_num_diag_lline.sas
    */
%macro resid_num_diag(dataset,...
/* Call shortcut macro */
%resid_num_diag(dataset=out1, datavar=resid, label='Residual',
    predvar=pred, predlabel='Predicted Value');
```

P-value for Brown-Forsythe test of constant variance in Residual vs. Predicted Value

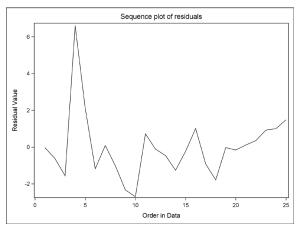
Obs	t_BF	BF_pvalue
1	1.50583	0.14572

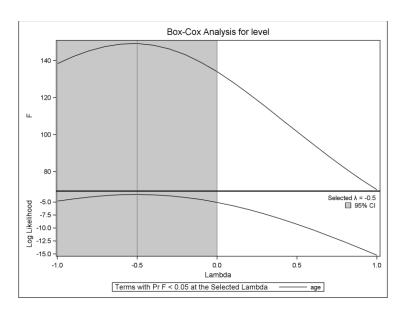
Output for correlation test of normality of Residual (Check text Table B.6 for threshold)

Pearson Correlation Coefficients, N = 25 Prob > r under H0: Rho=0				
	resid	expectNorm		
resid Residual	1.00000	0.90360 <.0001		
expectNorm	0.90360 <.0001	1.00000		

Residual	DF	Sum of Squares	Mean Square	F Value	Pr > F
Lack of Fit	3	22.748784	7.582928	2.75	0.0699
Pure Error	20	55.234280	2.761714		
Total Error	23	77.983064	3.390568		

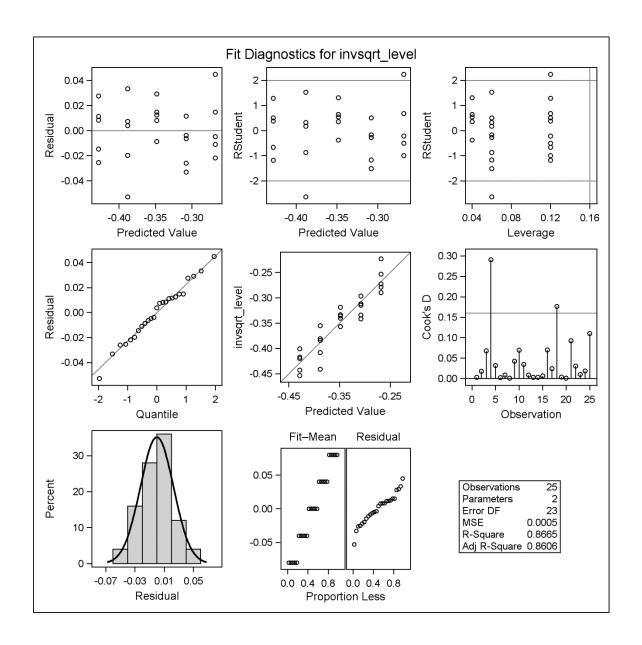
```
/* Look at sequence plot */
data temp; set out1;
  order = _n_;
proc sgplot data=temp;
  series x=order y=resid / lineattrs=(pattern=solid) ;
  xaxis label='Order in Data';
  yaxis label='Residual Value';
  title1 'Sequence plot of residuals';
run;
```





```
data plasma; set plasma;
  log_level = log(level);
  invsqrt_level = -1/sqrt(level);
run;

/* Inverse square root */
proc reg data=plasma;
  model invsqrt_level = age;
  output out=out2 r=resid p=pred;
  title1 'Simple model for negative inverse root plasma data';
run;
```



```
%resid_num_diag(dataset=out2, datavar=resid,
    label='Residual (neg. inverse root)',
    predvar=pred, predlabel='Predicted Value (neg. inverse
root)');
```

P-value for Brown-Forsythe test of constant variance in Residual (neg. inverse root) vs. Predicted Value (neg. inverse root)

Obs	t_BF	BF_pvalue
1	0.16654	0.86918

Output for correlation test of normality of Residual (neg. inverse root) (Check text Table B.6 for threshold)

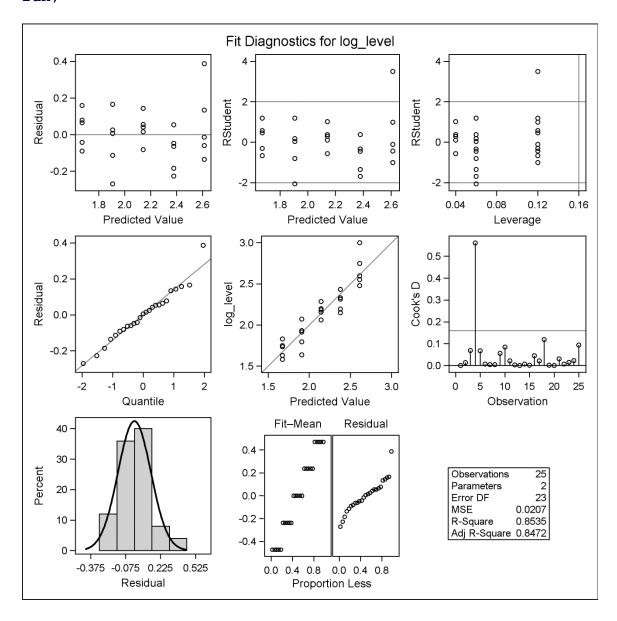
Pearson Correlation Coefficients, N = 25 Prob > r under H0: Rho=0				
resid expectNorm				
resid	1.00000	0.99188		
Residual (neg. inverse root)		<.0001		
expectNorm	0.99188	1.00000		
	<.0001			

```
proc rsreg data=plasma;
  model invsqrt_level = age / lackfit covar=1 noopt;
  title1 'F-test for lack of fit (neg. inverse root)';
run;
```

F-test for lack of fit (neg. inverse root)

Residual	DF	Sum of Squares	Mean Square	F Value	Pr > F
Lack of Fit	3	0.001556	0.000519	0.96	0.4312
Pure Error	20	0.010813	0.000541		
Total Error	23	0.012369	0.000538		

```
/* Log */
proc reg data=plasma;
  model log_level = age;
  output out=out3 r=resid p=pred;
  title1 'Simple model for log plasma data';
run;
```



```
%resid_num_diag(dataset=out3, datavar=resid,
label='Residual (log)',
    predvar=pred, predlabel='Predicted Value (log)');
```

P-value for Brown-Forsythe test of constant variance in Residual (log) vs. Predicted Value (log)

Obs	t_BF	BF_pvalue
1	0.95179	0.35110

Output for correlation test of normality of Residual (log) (Check text Table B.6 for threshold)

Pearson Correlation Coefficients, N = 25 Prob > r under H0: Rho=0				
	resid	expectNorm		
resid Residual (log)	1.00000	0.98071 <.0001		
expectNorm	0.98071 <.0001	1.00000		

```
proc rsreg data=plasma;
  model log_level = age / lackfit covar=1 noopt;
  title1 'F-test for lack of fit (log)';
run;
```

F-test for lack of fit (log)

Residual	DF	Sum of Squares	Mean Square	F Value	Pr > F
Lack of Fit	3	0.081944	0.027315	1.39	0.2758
Pure Error	20	0.394004	0.019700		
Total Error	23	0.475948	0.020693		

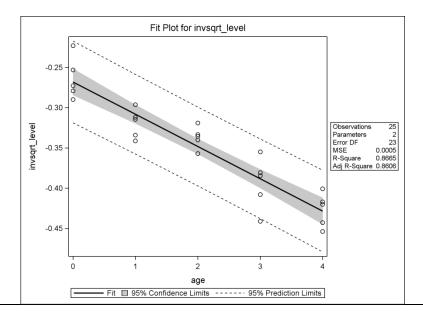
```
/* Probably go with inverse square root */
proc reg data=plasma;
  model invsqrt_level = age;
  title1 'Negative inverse root plasma data';
run;
```

Negative inverse root plasma data

Analysis of Variance						
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F	
Model	1	0.08025	0.08025	149.22	<.0001	
Error	23	0.01237	0.00053778			
Corrected Total	24	0.09262				

Root MSE	0.02319	R-Square	0.8665
----------	---------	----------	--------

Parameter Estimates						
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t	
Intercept	1	-0.26803	0.00803	-33.36	<.0001	
age	1	-0.04006	0.00328	-12.22	<.0001	



2.3: Simple Model Inference

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Recall the simple linear model

$$Y_i = \beta_0 + \beta_1 X_{i,1} + \epsilon_i.$$

Inference is the process by which we make a decision about whether an observed difference from an expectation was simply due to chance or not.

In other words, inference is the process of making conclusions given incomplete information.

1 Why Inference?

Hypothetical questions:

- Suppose you found out that there is not significant relationship between study time and final grades in Stat 5100, how would this effect your approach to this course?
- Suppose you have the flu and you find out from a clinical trial of a new flu drug that those who took the drug had slightly shorter flu durations than those who took the placebo, but that the difference was likely due to chance. How likely would you be to purchase this drug?

In the absence of complete information, inference is an efficient way to decide what associations are "real" and which are not.

2 Hypothesis Testing

Recall that hypothesis testing is the formal way by which we determine if an observed difference from an expectation was due to chance.

Process

- Define a null and alternative hypothesis.
 - $-H_0$: "no effect"
 - $-H_a$: "some effect"
- Define a test statistic:
 - Compares what we observed to what we expected if the null hypothesis was true.
- Determine the "sampling distribution"
 - Determines the natural variation in the test statistic that we would expect if we took many different samples from the same population.
 - In practice, we only ever take one sample. Statistical theory is what allows us to determine what the distribution would look like if we could take many samples.
 - The distribution often relies on model assumptions.
- Get p-value

- This is the probability of obtaining an observation as far, or farther, away from what we expected if the null was true.
- Make conclusion in context.
 - If the p-value is small ($< \alpha$), then it is unlikely that we would have obtained our observation if the null hypothesis is true. This provides evidence that the observed difference between our observation and expectation is REAL, and not simply due to chance.

2.1 Toluca Example:

If model assumptions are satisfied, then $b_1 \sim N(\beta_1, \sigma^2\{b_1\})$.

 \sim means "follows" while $\sigma^2\{b_1\}$ represents the true variance of b_1 , as estimated by $s\{b_1\}$.

Recall that, if the null hypothesis is true, then $\beta_1 = 0$. Thus, our test statistic becomes

$$t = \frac{b_1 - 0}{s\{b_1\}} \sim t_{df_E} = 10.29$$

with 25 - 2 = 23 degrees of freedom with a **p-value** < 0.0001.

where df_E is the degrees of freedom for the residuals, which is n-2 in the simple linear model case draw t-distribution and shade the area that represents the p-value

Since our p-value is lower than our level of significance (which is typically 0.05 and something we set beforehand), we would **reject** the null hypothesis **and conclude** that there is significant evidence that lot size and work hours are linearly related.

Where did $\alpha = 0.05$ come from?

Short answer: Sir Ronald Fisher, a prominent statistician, made it up:

It is a common practice to judge a result significant, if it is such a magnitude that it would have been produced by chance not more frequently than once in twenty trials. This is an arbitrary, but convenient, level of significance for the practical investigator...¹

However, $\alpha = 0.05$ has proven to be a good level of significance that balances the probability of Type I (claiming a difference when there isn't one) and Type II (claiming *no* difference when there is one).

Consider the following

You wish to determine if Aggie ice cream is more fattening than other ice cream shops in Logan. Suppose your null hypothesis is: "Aggie ice cream has the same number of calories per cup as Charlie's ice cream." You then conduct a test and obtain a p-value of 0.048, indicating that there is evidence that the average caloric counts are significantly different. You then realize that you forgot

¹As on p99 of "The Lady Tasting Tea" (2001) by David Salsburg. See http://jse.amstat.org/v16n2/velleman. pdf for more discussion about statistical theory.

to include five recorded observations in your study. When you include these additional observations, you obtain a p-value of 0.052, indicating no significant difference.

P-values should inform an analysis, rather than become the analysis.

Confidence Intervals

• General Form:

estimate \pm (critical value) \times (SE of estimate)

• For β_1 :

$$b_1 \pm t^* \times s\{b_1\}$$

- Interpretation:
 - We are 95% confident that the true value of β_1 is contained in this interval.
 - If we were to create 100 confidence intervals from 100 different samples, we would expect about 95 of them to contain the true β_1 .

Testing $H_0: \beta_1$ at level α is the same as checking whether 0 is inside the $(1-\alpha)100\%$ CI for β_1 .

Model Inference

All previous examples test whether an individual X variable has a significant linear relationship with Y. We will now look at some measures of model usefulness that apply when there is more than one X variable.

Ingredients of Model Inference

• Sum of Squares

–
$$SS_{total} = \sum_{i} (Y_i - \bar{Y})^2 \propto \text{variance of Y}$$

$$-SS_{error} = \sum_{i} (Y_i - \hat{Y}_i)^2 = \sum_{i} e_i^2 \propto \text{variance not explained by model}$$

$$-SS_{model} = SS_{total} - SS_{error}$$

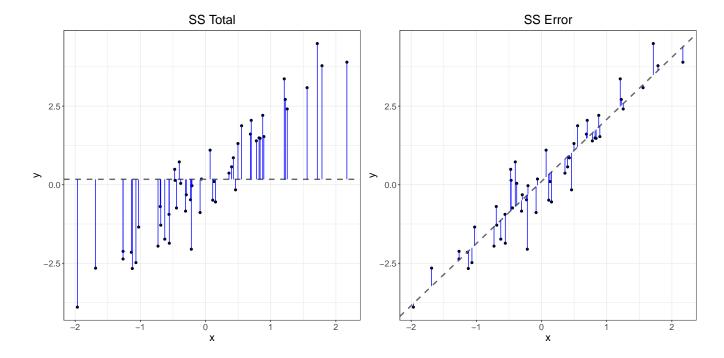


Figure 1: Illustration of SS_{total} and SS_{error} .

• Mean Square: $MS = \frac{SS}{df}$

•
$$F = \frac{MS_{model}}{MS_{error}}$$

•
$$R^2 = \frac{SS_{model}}{SS_{total}} = 1 - \frac{SS_{error}}{SS_{total}}$$

- Interpretation: the percent of the variation in Y that is explained by the model.

• MSE = Mean Square Error = $\hat{\sigma}^2$ = our best estimate of the error variance ($\epsilon \sim N(0, \sigma^2)$).

Toluca Example:

 $R^2=0.82$ (from Handout 2.1.1) which means that about 82% of the variation in work hours is explained by lot size.

Two other ways to look at $H_0: \beta_1 = 0:$

1. How much worse would the model fit be if we dropped the β_1 term?

Reduced Model (null hypothesis): $Y_i = \beta_0 + \epsilon_i$.

Full Model:
$$Y_i = \beta_0 + \beta_1 X_{i,1} + \epsilon_i$$
.

F-statistic looks at chance in SS_{error} between these two models.

Can be extended to consider removal of subsets of X variables (more later in the semester).

- 2. Let $\rho = Corr(X, Y) = \text{true}$, unknown correlation coefficient, which we estimate with the sample correlation (r).
 - When there is only one x-variable in the model it follows that

$$H_0: \beta_1 = 0 \equiv H_0: \rho = 0.$$

Inference on the Response Variable Y

We can create interval estimates for the response variable.

$$\hat{Y} \pm t_{df_E} (1 - \frac{\alpha}{2}) * SE{\{\hat{Y}\}}$$

Two Intervals:

• Confidence Interval: Interval estimate of mean (or expected) Y for population of all $X = X_h$.

$$SE\{\hat{Y}\} = s\{\hat{Y}_h\} = \hat{\sigma}^2 \left[\frac{1}{n} + \frac{(X_h - \bar{X})^2}{\sum_i (X_i - \bar{X})^2} \right]$$

• **Prediction Interval**: Interval estimate of **predicted** Y for a single [new] observation at $X = X_h$

$$SE\{\hat{Y}\} = s\{\hat{Y}_{h(new)}\} = \hat{\sigma}^2 \left[1 + \frac{1}{n} + \frac{(X_h - \bar{X})^2}{\sum_i (X_i - \bar{X})^2} \right]$$

Toluca Example (with $X_h = 10$)

- If we were to go to a new (single) lot of 10 acres, we are 95% confident that the work hours would be between
 - -13.6 (truncate at 0) and 209.7.
- If we were to consider all possible 10 acre sized lots, we are 95% confident that the mean work hours across all these lots would be between

50.5 and 145.6.

Note: Most models have more than one predictor variable, we will use the following common notation throughout the remainder of this course:

- n = sample size
- $p = \text{number of } \beta_j$'s in the model (including the intercept)
- $df_E = n p$

2.3.1: SAS - Simple Inference

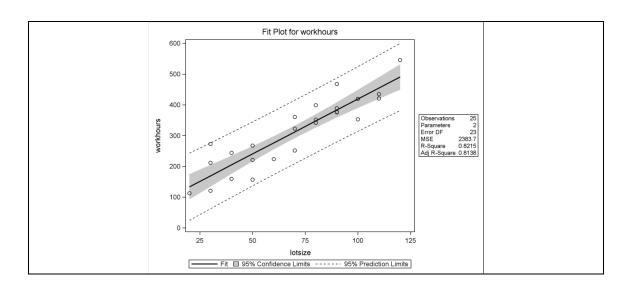
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<u>Example</u>: (The Toluca Company data from Chapter 1 & Chapter 3 Handouts) We really want to say something about how lotsize affects workhours – does it?

```
/* Input Toluca data (recall Ch. 1 example) */
data toluca; input lotsize workhours @@; cards;
  80
      399
           30
              121
                    50 221
                             90 376
                                      70 361
                                               60
                                                  224
 120
     546
           80
              352
                   100
                       353
                             50
                                157
                                      40 160
                                               70 252
  90
      389
           20
               113
                  110
                       435
                            100
                                420
                                      30 212
                                               50 268
  90
     377 110 421
                    30 273
                             90 468
                                      40 244
                                               80 342
  70
     323
run;
/* Now fit simple linear model with Y=workhours and
  X=lotsize, with residuals and predicted values saved
   in data set tolucaout */
proc reg data=toluca;
 model workhours = lotsize;
  output out=tolucaout r=resid p=pred;
  title1 'Simple linear model';
run;
/* Check assumptions */
/* Define shortcut macro, using line copied from
   www.stat.usu.edu/jrstevens/stat5100/resid num diag 1line.sas
 */
%macro resid num diag(dataset, ...
%resid num diag(dataset=out1, datavar=resid, label='Residual',
  predvar=pred, predlabel='Predicted Value');
/* See output from this on p.5 of Handout #4.
  Only when assumptions are met does inference make sense!
*/
```

```
/* Fit a simple linear model with Y=workhours and X=lotsize;
   output the 95% confidence intervals for the coefficients.
   Get predicted values (call them Predict here) and
   upper and lower 95% prediction and confidence intervals
   for each X value; put all this in a dataset called confidence.
  Also, include prediction for two X-levels not in original
   data set (X=10 \text{ and } X=130). */
data dummy; input lotsize @@; cards;
 10 130
data trick; set toluca dummy;
run;
proc reg data=trick;
 model workhours = lotsize / clb alpha=.05;
                                          /* 1-alpha is level */
  output out=confidence p=Predict
                     ucl=uPred /* upper and lower limits for */
                                /*
                                        individual prediction */
                     lcl=lPred
                     uclm=uConf /* upper and lower limits for */
                     lclm=lConf; /* group mean confidence */
  title1 'Regression with 95% interval estimation';
run;
```

	Regression with 95% interval estimation									
	Parameter Estimates									
Variable	D F	Paramete r Estimate	Standar d Error	t Valu e	Pr > t		onfidence mits			
Intercep t	1	62.36586	26.17743	2.38	0.0259	8.2137 1	116.5180 1			
lotsize	1	3.57020	0.34697	10.29	<.0001	2.8524 4	4.28797			



Predicted values and confidence and predicted intervals for lotsize < 50; these are 95% intervals.

O	bs	lotsize	workhours	Predict	lPred	uPred	lConf	uConf
	2	30	121	169.472	62.5464	276.397	134.367	204.577
1	11	40	160	205.174	99.9483	310.400	175.649	234.698
1	14	20	113	133.770	24.6977	242.842	92.587	174.952
	17	30	212	169.472	62.5464	276.397	134.367	204.577
2	21	30	273	169.472	62.5464	276.397	134.367	204.577
2	23	40	244	205.174	99.9483	310.400	175.649	234.698
2	26	10	•	98.068	-13.5719	209.708	50.500	145.636

/****************

Note: there are other ways to get the CI for Y in SAS, but they

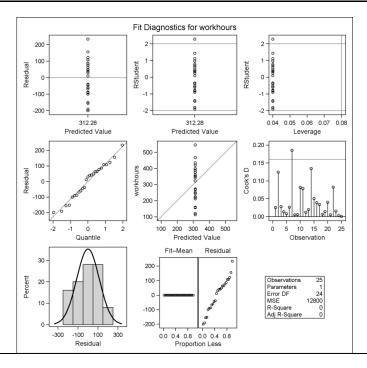
aren't included here; just know that if you needed to, you could get the SE for Yhat using the stdp and stdi options in proc reg.

```
/* Look at Reduced model */
proc reg data=toluca;
  model workhours = ;
  title1 'Reduced Model (dropped lotsize predictor)';
run;
```

Reduced Model (dropped lotsize predictor)

Analysis of Variance								
Source DF Sum of Square F Value Pr >								
Model	0	0						
Error	24	307203	12800					
Corrected Total	24	307203						

Parameter Estimates							
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t		
Intercept	1	312.28000	22.62753	13.80	<.0001		



2.4: Simultaneous Inference and Important Considerations

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Simultaneous inference is when we want to conduct multiple tests of significance at the same time.

1 Why Simultaneous Inference?

In handout 2.3, we conducted inference for parameters one at a time. We need to change our approach when looking at multiple parameters simultaneously.

How and why do we need to change our approach when conducting simultaneous inference?

(check out **this comic** for help).

If we conduct several tests at the same level of significance, the probability of getting one false positive result (a type I) error becomes much higher than α .

As a result, we need to adjust the level of significance to account for a "multiplicity" of testing.

2 Bonferroni Adjustment

Multiplicity:

- Let A_j = event that an individual $(1 \alpha)100\%$ CI does not contain the true value of β_j .
- $P(A_0) = P(A_1) = \alpha \rightarrow \text{Type I Error}$
 - $P(NOTA_j)$ = probability that an interval contains the true value of β_j .
- Bonferroni Inequality: $P(\text{NOT}A_0 \text{ AND NOT } A_1) \ge 1 P(A_0) P(A_1)$

This means that if we conduct g tests at a confidence level of $(1 - \frac{\alpha}{g})$, then we are guaranteed that overall level of confidence for all intervals *considered jointly* will be at least $(1 - \alpha)$, we call this the **Bonferroni adjustment**.

- Bonferroni Advantage: Can be applied in *any* situation that requires a multiplicity adjustment, including simultaneous intervals for \hat{Y} at multiple X_h levels.
- Bonferroni Disadvantage: Can be overly conservative, producing inefficient (unnecessarily wide) intervals.

Comparison of Simultaneous Intervals for \hat{Y}

- Confidence intervals (mean response)
 - Bonferroni

$$\hat{Y} \pm t_{n-p} (1 - \frac{\alpha}{2g}) * s{\hat{Y}_h}$$

- Working-Hotelling (WH)

$$\hat{Y} \pm W * s{\{\hat{Y}_h\}} \qquad \left(W = \sqrt{pF_{p,n-p}(1-\alpha)}\right)$$

Notice that the W-statistic does not consider g

- * WH provides a "confidence band" for the entire regression line (all possible X_h levels).
- * This means the WH interval at any individual X_h will be wider than the t-based confidence interval, but the WH intervals will eventually be narrower than Bonferroni confidence intervals if enough X_h are considered.
- Prediction intervals (new response)
 - Bonferroni

$$\hat{Y} \pm t_{n-p} (1 - \frac{\alpha}{2q}) * s{\{\hat{Y}_{h(new)}\}}$$

Scheffe (chef-eh)

$$\hat{Y} \pm S * s{\hat{Y}_{h(new)}}$$
 $\left(S = \sqrt{gF_{g,n-p}(1-\alpha)}\right)$

Rule of Thumb: Always pick the most efficient interval that guarantees your intended type I error (α) .

Table 1: Summary of Methods for Simultaneous Intervals

Simultaneous Interval on:	Methods
β 's	Bonferroni
Population means of Y at multiple X_h	Bonferroni or Working-Hotelling
Predictions for Y at multiple X_h	Bonferroni of Scheffe

3 Inverse Prediction

Problem: What is the value of X_h necessary to achieve a specific value of \hat{Y} .

Solution: solve for X.

$$\hat{Y} = b_0 + b_1 X_h$$

$$b_1 X_h = \hat{Y} - b_0$$

$$X_h = \frac{\hat{Y} - b_0}{b_1}$$

Problem: Use Y to predict values of X.

Solution: DO NOT solve for X.

Why?

- The least squares slope estimate of regression model that predicts Y using X: $\rho_{\overline{SD\{X\}}}^{SD\{Y\}}$.
- The least squares slope estimate of regression model that predicts X using Y: $\rho \frac{SD\{X\}}{SD\{Y\}}$.
- Notice that the slopes are NOT inverses of each other.

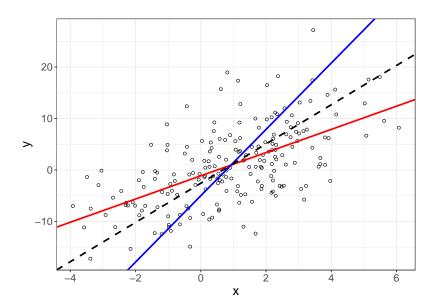


Figure 1: Scatterplot of points along with the regression line that uses X to predict Y (red), the regression line that uses Y to predict X (blue), the SD line (black).

4 Cautions for Linear Regression

- Remedial measures may not fix violations of assumptions
 - May need to abandon OLS regression altogether
- Interpretation: Sometimes the X vs Y relationship may look counterintuitive
 - May be the result of omitted predictors
- R^2 can be abused
 - Higher $R^2 \to \text{not always better model}$
 - Lower $\mathbb{R}^2 \to \text{does not mean there is no linear relationship}$

2.4.1: SAS - Simultaneous Inference and Regression Through Origin

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```
/* Input Toluca data (recall Ch. 1 example) */
data toluca; input lotsize workhours @@; cards;
80 399
        30 121
                50 221
                        90 376
                                          60 224
120 546
       80 352 100 353
                       50 157
                                 40
                                    160
                                         70 252
90 389 20 113 110 435 100 420
                                    212
                                          50 268
                                 30
90 377 110 421 30 273 90 468 40 244
                                         80 342
70 323
 ;
run;
/* Simultaneous 95% interval estimation of betas */
proc reg data=toluca;
 model workhours = lotsize / clb alpha=.025;
 title1 'Simultaneous 95% confidence intervals on betas';
run;
```

Simultaneous 95% confidence intervals on betas

Parameter Estimates										
Variable	D F	Paramete r Estimate	Standar d Error	t Valu e	Pr > t	97.5% Confidence Limits				
Intercep t	1	62.36586	26.17743	2.38	0.0259	0.40436	125.1360 7			
lotsize	1	3.57020	0.34697	10.29	<.0001	2.73821	4.40220			

```
/* Simultaneous 90% interval estimation of mean workhours
   at lotsize levels 30, 65, 100 (using Working-Hotelling
   and Bonferroni)
 */
data dummy; input lotsize check; cards;
  30 1
  65
     1
  100 1
data temp; set toluca dummy;
proc reg data=temp noprint;
  model workhours = lotsize;
  output out=out1 p=Yhat stdp=seYhat;
    /* KEY: stdp is SE of mean prediction */
data out1; set out1;
  alpha = 0.10; /* 1-alpha is simult. conf. level */
 p = 2;
                 /* # of beta's (including intercept) */
 n = 25;
                /* sample size */
                 /* number of simultaneous intervals */
  q = 3;
  W = sqrt(p*finv(1-alpha,p,n-p)); /* WH crit. val. */
                                   /* Bonf. crit. val. */
  t = tinv(1-alpha/(2*g), n-p);
 WH upper = Yhat + W*seYhat;
 WH lower = Yhat - W*seYhat;
 B upper = Yhat + t*seYhat;
  B lower = Yhat - t*seYhat;
proc print data=out1;
  where check = 1;
  var lotsize Yhat seYhat WH lower WH upper
      B lower B upper;
  title1
   'Simultaneous 90% interval estimation of mean response';
   'at three X-levels, using Working-Hotelling and
   Bonferroni';
run;
```

Simultaneous 90% interval estimation of mean response at three X-levels, using Working-Hotelling and Bonferroni

Ob s	lotsiz e	Yhat	seYhat	WH_lowe	WH_uppe	B_lowe r	B_uppe r
26	30	169.47 2	16.969 7	131.154	207.790	131.057	207.887

27	65	294.42 9	9.9176	272.035	316.823	271.978	316.880
28	100	419.38 6	14.272	387.159	451.613	387.077	451.695

```
/* Simultaneous 95% prediction limits on next two lots,
  with sizes 80 and 100 units (using Scheffe and
  Bonferroni)
 */
data dummy; input lotsize check; cards;
  80 1
 100 1
data temp; set toluca dummy;
proc reg data=temp noprint;
  model workhours = lotsize;
  output out=out1 p=Yhat stdi=seYhatnew;
    /* KEY: stdi is SE of individual prediction */
data out1; set out1;
  alpha = 0.05; /* 1-alpha is simult. pred. level */
                 /* # of beta's (including intercept) */
  n = 25;
                /* sample size */
  q = 2;
                 /* number of simultaneous intervals */
  S = sqrt(g*finv(1-alpha,g,n-p)); /* Scheffe crit val */
                                   /* Bonf. crit. val. */
  t = tinv(1-alpha/(2*q), n-p);
  S upper = Yhat + S*seYhatnew;
  S_lower = Yhat - S*seYhatnew;
  B upper = Yhat + t*seYhatnew;
  B lower = Yhat - t*seYhatnew;
proc print data=out1;
  where check = 1;
  var lotsize Yhat seYhatnew S lower S upper
      B lower B upper;
  title1 'Simultaneous 95% interval estimation of
          individual prediction';
  title2 'at two X-levels, using Scheffe and Bonferroni';
run;
```

Simultaneous 95% interval estimation of individual prediction at two X-levels, using Scheffe and Bonferroni

Obs	lotsize	Yhat	seYhatnew	S_lower	S_upper	B_lower	B_upper
26	80	347.982	49.9110	217.407	478.557	228.302	467.662
27	100	419.386	50.8666	286.311	552.461	297.414	541.358

```
/* Regression through origin example: plumbing supplies
  company looking at relationship between number of
  work units (X) and labor costs (Y) at its 12 warehouses
*/
data warehouse; input work cost @@; cards;
20 114
        196 921 115 560
                          50 245
                                  122 575
                                           100 475
33 138
        154 727 80 375
                                  182 828
                         147 670
                                           160 762
 0 .
proc reg data=warehouse;
 model cost = work / noint;
 output out=out1 p=pred;
 title1 'Regression through origin';
run;
```

Regression through origin

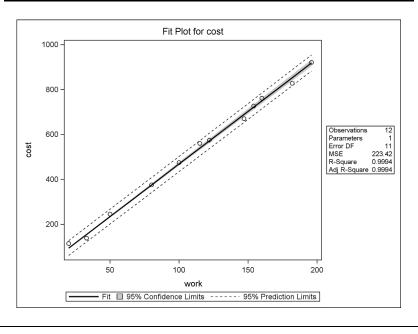
Number of Observations Read	13
Number of Observations Used	12
Number of Observations with Missing Values	1

Note: No intercept in model. R-Square is redefined.

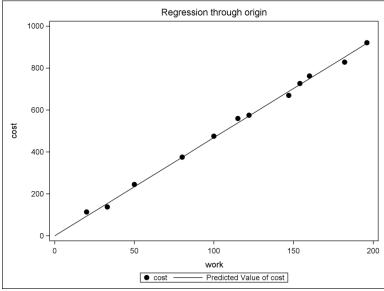
Analysis of Variance								
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F			
Model	1	4191980	4191980	18762.5	<.0001			
Error	11	2457.65933	223.42358					
Uncorrected Total	12	4194438						

Root MSE	14.94736	R-Square	0.9994
Dependent Mean	532.50000	Adj R-Sq	0.9994
Coeff Var	2.80702		

Parameter Estimates									
Variable DF Parameter Standard t Value Pr >									
work	1	4.68527	0.03421	136.98	<.0001				



run;
/* Note forced inclusion of
 work=0 dummy observation
 for graphical purposes */



2.5: Multiple Linear Regression

Dr. Bean - Stat 5100

1 Why Multiple Linear Regression?

- Models that use a single explanatory variable to predict a response are very limited in terms of its capability.
- We are often interested in determining the effect of an explanatory variable on the response variable *after* accounting for the effects due to other explanatory variables.
 - Example: Is there a difference in the pay based on gender after accounting for job type and hours worked?

2 What Changes from Simple Linear Regression?

1. Interpretation of coefficients

$$Y_i = \beta_0 + \beta_1 X_{i1} + \beta_2 X_{i2} + \ldots + \beta_{p-1} X_{i,p-1} + \varepsilon_i$$

$$\beta_0 = E[Y|X_1 = X_2 = \dots = X_{p-1} = 0]$$

 β_k = expected (or average) change in Y for every unit increase in predictor X_k , while holding all other predictors constant

Need all three elements for a correct interpretation.

 β_k sometimes called "partial regression coefficient" because it reflects partial effect of X_k on Y after accounting for effects of other predictors

- 2. ANOVA table
 - model df = p 1 = # of predictor variables
 - error df = n p– we have to "spend" more degrees of freedom to calculate the additional coefficients
 - model F-test more meaningful:

$$H_0$$
: $\beta_1 = \beta_2 = \dots = \beta_{p-1} = 0$
 H_a : $\beta_k \neq 0$ for at least one $k = 1, \dots, p-1$

• R^2 called coefficient of multiple determination (still interpret as % variance in Y explained by model); $\sqrt{R^2}$ called coeff. of multiple correlation

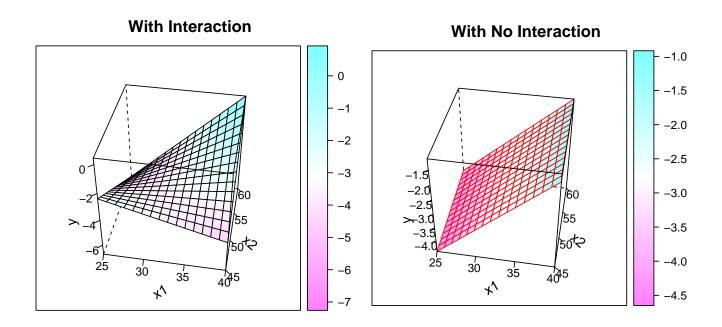


Figure 1: Regression surface using two X variables to predict Y. Harder to visualize when there are more than two predictor variables.

- 4. F-test for lack of fit less practical
 - requires multiple observations at one or more X profiles, which is hard to achieve when the number of X's is large.
 - "X-profile" or "covariate profile" refers to specific values for all predictors
- 5. More assumptions to check later regarding inter-related predictors
 - basically, if predictors are related to each other, the model becomes very hard to interpret
- 6. Other variable types can be included (interactions, qualitative, higher-order) (more in Module 3)

3 Matrix Approach to Multiple Linear Regression

When the number of X variables gets large, the matrix representation of linear regression models is easier to write and understand.

$$Y = (Y_1, \dots, Y_n)' = \text{vector of response variable}$$

$$\varepsilon = (\varepsilon_1, \dots, \varepsilon_n)' = \text{vector of error terms}$$

$$X_k = (X_{1k}, \dots, X_{nk})' = \text{vector of predictor variable } \#k \quad (k = 1, \dots, p - 1)$$

$$X = \begin{bmatrix} 1 & X_1 & \dots & X_{p-1} \end{bmatrix} = \text{matrix with } p \text{ columns and } n \text{ rows}$$

$$\beta = (\beta_0, \beta_1, \dots, \beta_{p-1})' = \text{vector of coefficients}$$

$$b = (b_0, b_1, \dots, b_{p-1})' = \text{vector of coefficient } \underbrace{\text{estimates}}$$

Then regression model is

$$Y = X\beta + \varepsilon$$

$$\varepsilon \sim N(0, \sigma^2 I) \quad I = \text{``identity'' matrix'}$$

 $b = (X'X)^{-1}X'Y$

Estimates:

truth:
$$Cov(b) = (X'X)^{-1}\sigma^2$$
 estimated: $s^2\{b\} = (X'X)^{-1} \cdot \text{MSE}$ $\sqrt{\text{diag. elements}}$ gives SE's of b_k 's

Matrices with variance on

diag., covariance on off-diag.

We'll come back to this, but for now, note that

$$\hat{Y} = Xb
= X(X'X)^{-1}X'Y
= HY$$

H projects Y down to column space of X:

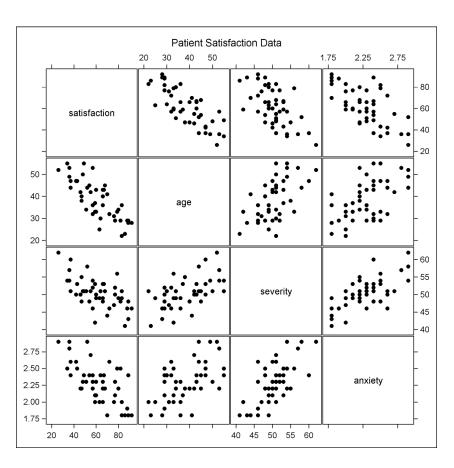
- Y = observed response values vector; is not a [perfect] linear combination of predictor variables
- \hat{Y} = predicted response values vector; <u>is</u> a [perfect] linear combination of predictor variables

2.5.1: SAS - Multiple Predictors

Dr. Bean – Stat 5100

<u>Example</u>: (Exercises 6.15-6.17) A hospital administrator is studying the relation between patient satisfaction (Y, an index) and patient's age (X1, in years), severity of illness (X2, an index), and anxiety level (X3, an index). Data are reported for 46 randomly selected patients. For all index variables, higher values indicate more (satisfaction, severity, anxiety).

```
/* Input data (see Exercises 6.15-6.17) */
data patient;
   input satisfaction age severity anxiety @@; cards;
  48
         50
                51
                     2.3
                             57
                                    36
                                           46
                                                 2.3
  66
         40
                48
                     2.2
                             70
                                    41
                                           44
                                                 1.8
  89
         28
                43
                     1.8
                             36
                                    49
                                           54
                                                 2.9
                     2.2
                                                 2.4
  46
         42
                50
                             54
                                    45
                                           48
                     2.9
                                                 2.1
  26
         52
                62
                             77
                                    29
                                           50
                                                 2.4
  89
         29
                48
                     2.4
                             67
                                    43
                                           53
  47
                     2.2
                                                 2.3
         38
                55
                             51
                                    34
                                           51
  57
         53
                54
                     2.2
                             66
                                    36
                                           49
                                                 2.0
                                                 1.9
  79
                     2.5
                                    29
         33
                56
                             88
                                           46
  60
         33
                49
                     2.1
                             49
                                    55
                                           51
                                                 2.4
  77
         29
                                                 2.9
                52
                     2.3
                             52
                                    44
                                           58
                                                 1.8
  60
         43
                50
                     2.3
                             86
                                    23
                                           41
                                                 2.5
  43
         47
                53
                     2.5
                             34
                                    55
                                           54
  63
         25
                49
                     2.0
                                    32
                                                 2.6
                             72
                                           46
  57
         32
                52
                     2.4
                                    42
                                           51
                                                 2.7
                             55
                     2.0
  59
         33
                42
                             83
                                    36
                                           49
                                                 1.8
  76
         31
                47
                     2.0
                             47
                                    40
                                           48
                                                 2.2
  36
         53
                57
                     2.8
                             80
                                    34
                                           49
                                                 2.2
                                                 2.4
         29
                     2.5
  82
                48
                             64
                                    30
                                           51
  37
         47
                     2.4
                                           50
                                                 2.6
                60
                             42
                                    47
                                                 2.0
  66
         43
                53
                     2.3
                             83
                                    22
                                           51
  37
         44
                51
                     2.6
                                           51
                                                 2.2
                             68
                                    45
  59
         37
                53
                     2.1
                             92
                                    28
                                           46
                                                 1.8
run;
/* Look at scatterplot matrix */
proc sgscatter data=patient;
  matrix satisfaction age severity anxiety /
       markerattrs=(symbol=CIRCLEFILLED size=2pt);
  title1 'Patient Satisfaction Data';
run;
```



```
/* Fit regression model */
proc reg data=patient;
  model satisfaction = age severity anxiety;
  output out=out1 r=resid p=pred;
  title1 'Patient Satisfaction Regression';
run;
```

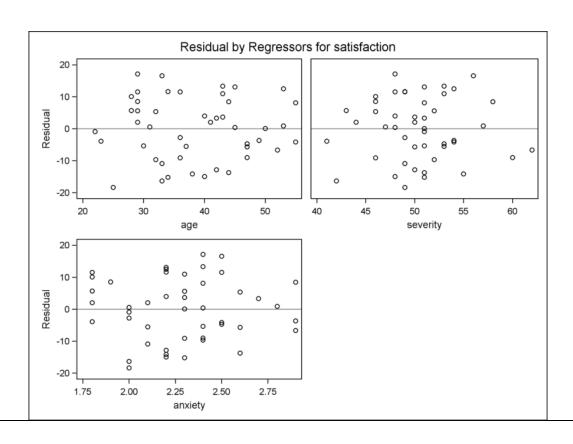
Patient Satisfaction Regression

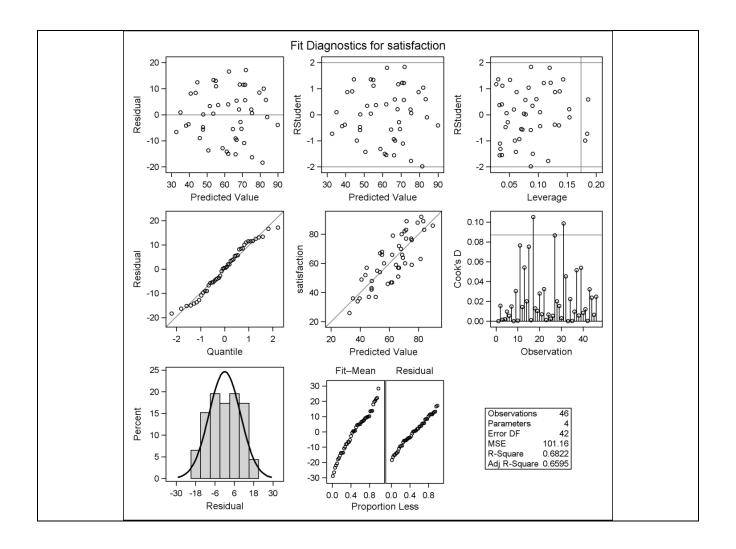
Number of Observations Used 46

Analysis of Variance											
Source	DF	Sum of Squares			Pr > F						
Model	3	9120.46367	3040.15456	30.05	<.0001						
Error	42	4248.84068	101.16287								
Corrected Total	45	13369									

Root MSE	10.05798	R-Square	0.6822
Dependent Mean	61.56522	Adj R-Sq	0.6595
Coeff Var	16.33711		

Parameter Estimates										
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t					
Intercept	1	158.49125	18.12589	8.74	<.0001					
age	1	-1.14161	0.21480	-5.31	<.0001					
severity	1	-0.44200	0.49197	-0.90	0.3741					
anxiety	1	-13.47016	7.09966	-1.90	0.0647					





```
/* Joint 90% intervals for beta1, beta2, and beta3 */
proc reg data=patient;
  model satisfaction = age severity anxiety /
      clb alpha=.0333;
  title1 'Simultaneous 90% intervals for three predictors
effects';
run;
```

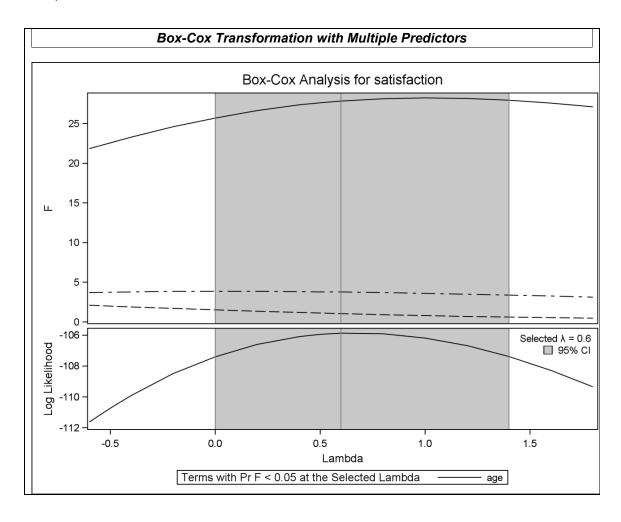
Simultaneous 90% intervals for three predictors effects

	Parameter Estimates											
Variable	DF	Parameter Estimate	Standard Error	t Value	alue Pr > t 96.67% Confidence Limits							
Intercept	1	158.49125	18.12589	8.74	<.0001	118.59967	198.38283					
age	1	-1.14161	0.21480	-5.31	<.0001	-1.61434	-0.66888					
severity	1	-0.44200	0.49197	-0.90	0.3741	-1.52473	0.64072					
anxiety	1	-13.47016	7.09966	-1.90	0.0647	-29.09514	2.15482					

```
/* Simultaneous 90% prediction limits on two new patients
   (using Scheffe and Bonferroni), with profiles
      age=35, severity=45, anxiety=2.2
  and
      age=42, severity=61, anxiety=1.8
 */
data dummy; input age severity anxiety check; cards;
  35 45 2.2 1
  42 61 1.8 1
data temp; set patient dummy;
proc reg data=temp noprint;
  model satisfaction = age severity anxiety;
  output out=out1 p=Yhat stdi=seYhatnew;
    /* KEY: stdi is SE of individual prediction */
data out1; set out1;
  alpha = 0.10; /* 1-alpha is simult. pred. level */
                /* # of beta's (including intercept) */
 p = 4;
                /* sample size */
  n = 46;
                /* number of simultaneous intervals */
  q = 2;
  S = sqrt(g*finv(1-alpha,g,n-p)); /* Scheffe crit val */
  t = tinv(1-alpha/(2*q), n-p);
                                   /* Bonf. crit. val. */
  S upper = Yhat + S*seYhatnew;
  S lower = Yhat - S*seYhatnew;
  B upper = Yhat + t*seYhatnew;
  B lower = Yhat - t*seYhatnew;
proc print data=out1;
  where check = 1;
  var age severity anxiety Yhat S_lower S_upper
      B lower B upper;
  title1 'Simultaneous 90% intervals of individual
prediction';
  title2 'at two X-profiles, using Scheffe and Bonferroni';
run;
```

Simultaneous 90% intervals of individual prediction at two X-profiles, using Scheffe and Bonferroni

	Obs	age	severity	anxiety	Yhat	S_lower	S_upper	B_lower	B_upper
	47	35	45	2.2	69.0103	46.0553	91.9652	48.0122	90.0083
Ī	48	42	61	1.8	59.3350	31.3797	87.2903	33.7629	84.9071



2.6: Multiple Inference and Multicollinearity

Dr. Bean - Stat 5100

1 Why Multiple Inference?

We already have tools to test the significance of model coefficients:

- Individual coefficients: t-tests $(H_0: \beta_k = 0)$
- All coefficients: model F-test $(H_0: \beta_1 = \beta_2 = \cdots = \beta_{p-1} = 0)$

What if we want to consider the significance of a subset of the X predictor variables? (More than one, but not all of them).

2 Subset Testing

Example: Bodyfat Dataset (Handout 2.6.1)

 $Y = \text{body}, X_1 = \text{triceps}, X_2 = \text{thigh}, X_3 = \text{midarm}$

$$Y = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \beta_3 X_3 + \epsilon$$

Consider $H_0: \beta_2 = \beta_3 = 0$.

How to test: See how much better the full model is (using tricep, thigh, and midarm) compared to the reduced one (using only triceps).

- Notation: $SSE(X_1, X_2, X_3) = SS_{error}$ when model has predictors X_1, X_2 , and X_3 represents amount variation in Y left unexplained by the full model
- Assuming H_0 : $\beta_2 = \beta_3 = 0$ is true, fit "reduced" model (only predictor X_1) and calculate $SSE(X_1)$
- Note that $SSE(X_1) > SSE(X_1, X_2, X_3)$
 - ALWAYS true, as a "worthless" X variable won't ever increase the SSE, but may reduce
 it slightly by chance.
 - $-\,$ NOT true of validation error (more discussion in Module 4).
 - then define "extra sum of squares"

$$SSR(X_2, X_3|X_1) = SSE(X_1) - SSE(X_1, X_2, X_3)$$

Note: this represents amount variation in Y accounted for by $X_2 \& X_3$ when X_1 already in model

• Define

$$MSR(X_2, X_3|X_1) = \frac{SSR(X_2, X_3|X_1)}{2}$$

- think of this as the mean square reduction

• Build test statistic for H_0 : $\beta_2 = \beta_3 = 0$

$$F^* = \frac{MSR(X_2, X_3|X_1)}{MSE(X_1, X_2, X_3)}$$
$$= \frac{SSR(X_2, X_3|X_1)/(2)}{SSE(X_1, X_2, X_3)/(16)}$$

• When H_0 : $\beta_2 = \beta_3 = 0$ is true, $F^* \sim F_{2,16}$

General test of any # of β_k 's:

$$Y = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \ldots + \beta_{p-1} X_{p-1} + \epsilon$$

$$H_0 : \beta_q = \beta_{q+1} = \ldots = \beta_{p-1} = 0$$

$$p = \# \text{ of } \beta \text{'s in full model (incl. intercept)}$$

$$q = \# \text{ of } \beta \text{'s in reduced model (incl. intercept)}$$

$$p - q = \# \text{ of } \beta \text{'s being tested in } H_0$$

$$F^* = \frac{[(\text{SSE in reduced model}) - (\text{SSE in full model})]/(p - q)}{[\text{SSE in full model}]/(n - p)}$$

Under H_0 , $F^* \sim F_{p-q,n-p}$

Recall the t-statistic from test of individual predictor $(H_0: \beta_k = 0)$?

$$t^* = \frac{b_k}{s\{b_k\}}$$

– if only have one predictor in model then $(t^*)^2 \sim F_{1,n-p}$

SSR also called sequential sums of squares or Type I SS; example in SAS:

- $SSR(X_1) \approx 352.27$
- $SSR(X_2|X_1) \approx 33.17$
- $SSR(X_3|X_1,X_2) \approx 11.55$

Related concept: "Coefficients of Partial Determination"

• what proportion of [previously unexplained] variation in Y can be explained by addition of predictor X_k to model

$$R_{Y3|12}^2 = \frac{SSR(X_3|X_1, X_2)}{SSE(X_1, X_2)}$$

- $SSR(X_3|X_1,X_2)$ - reduction in SSE that occurs when X_3 is added to the model when X_1 and X_2 are already in the model.

- $SSE(X_1, X_2)$ amount of unexplained variation in Y when X_1 and X_2 are in the model.
- example in SAS:
 - $-R_{V1}^2 \approx 0.711$
 - $-R_{Y2|1}^2 \approx 0.232$
 - $-R_{Y3|12}^2 \approx 0.105$

3 Multicollinearity

Textbook sections 7.6 and 10.5

The model F test says that the coefficients *collectively* are highly significant, but *none* of the individual variables are significant.

This is a symptom of **multicollinearity** (i.e. collinearity):

- Two X variables share a strong linear relationship with each other (independent of Y)
- One X variable is a near linear combination of two or more X variables

Problems with Multicollinearity:

- β_k hard to interpret as it no longer makes sense to "hold all other predictor variables constant."
- The variance of b_k will be very large (inflated) as our estimates are starting to become non-unique \rightarrow makes inference of β_k difficult if not impossible.
 - Could make estimate of b_k counter-intuitive (example: getting a negative estimate of b_k despite knowing that X and Y are positively correlated)).
- Contradictory results between individual t-tests and model F tests (or subset F tests).

NOT Problems with Multicollinearity:

• Multicollinearity does NOT affect a model's predictive ability.

3.1 Standardizing Variables

One way to better understand multicollinearity is by standardizing variables.

$$Y_i^* = \frac{1}{\sqrt{n-1}} \left(\frac{Y_i - \bar{Y}}{\text{SD of } Y} \right) \qquad , \qquad X_{ik}^* = \frac{1}{\sqrt{n-1}} \left(\frac{X_{ik} - \bar{X}_k}{\text{SD of } X_k} \right)$$

- sometimes called "correlation transformation" because

$$Corr(X_k, Y) = \sum_{i} X_{ik}^* Y_i^*$$

If all variables have been standardized, then consider matrix approach (with no Intercept column in matrix X^*):

$$Y^* = X^*\beta^* + \varepsilon$$

$$b^* = (X^{*\prime}X^*)^{-1}X^{*\prime}Y^*$$

$$Cov(b^*) = (X^{*\prime}X^*)^{-1}\sigma^2$$

There is no intercept column because, by construction, the intercept will be Y=0 as all points must past through $(\bar{X}, \bar{Y}) = (0, 0)$

To un-standardize regression coefficient estimates:

$$b_k = \left(\frac{\text{SD of } Y}{\text{SD of } X_k}\right) \cdot b_k^*$$

$$b_0 = \bar{Y} - \sum_{k=1}^{p-1} b_k \bar{X}_k$$

Relevance to multicollinearity:

- the correlation matrix among the [original] predictor variables is $X^{*'}X^*$
- the "closer" X_j and X_h are, the larger will be the j^{th} and h^{th} diagonal elements of $Cov(b^*)$, so the estimated variance is higher for b_j and b_h
- We can use the correlation matrix to obtain a set of **condition indices** as obtained from the **eigenvalues** of the matrix.

While standardizing helps to better mathematically understand the effect of multicollinearity, it is not necessary to standardize to detect multicollinearity.

3.2 Ways to Diagnose Multicollinearity

3.2.1 Condition Index/Principal Components

- Recall from linear algebra: λ is an **eigenvalue** of a symmetric, square matrix A iff there exists a vector x (the **eigenvector** for λ) such that $Ax = \lambda x$.
- Let $\lambda_1, \ldots, \lambda_k$ be the eigenvalues of $X^{*\prime}X^*$, and let

Condition Index_i =
$$\left(\frac{\lambda_{max}}{\lambda_i}\right)^{1/2}$$

- Each condition index is associated with a **principal component**
 - Each principal component is a linear combination of the original predictor variables. Each principal component shares no correlation with any other principal component (i.e. $cor(PC_1, PC_2) = 0$).

$$PC_1 = a_1 X_1^* + \dots + a_{p-1} X_{p-1}^*$$

$$PC_2 = c_1 X_1^* + \dots + c_{p-1} X_{p-1}^*$$

$$\vdots$$

• Each principal component explains some percentage of the variation in the original predictors.

IF the condition index is high (more than 10 or so) AND the associated principal component explains a high proportion of the variance (usually more than 50% variability) in two or more predictor variables, then we have potentially problematic multicollinearity.

3.2.2 Variance Inflation Factor (VIF)

- Let R_k^2 be the coefficient of multiple determination (the R^2 value) when predictor X_k^* is regressed on the other predictors
 - This is a measure of how much of the variance of X_k^* is explained by the other X variables.
- Define $VIF_k = (1 R_k^2)^{-1}$, for k = 1, ..., p 1 as the "Variance Inflation Factor" for b_k (the estimate of β_k)

IF the largest VIF is much more than 10 **OR** the average VIF is much more than 1, then we have evidence of potentially problematic multicollinearity.

We usually use a combination of the VIF and condition index to asses multicollinearity.

3.2.3 Important things to remember about standardization

- Relative magnitude of b_k^* estimates not meaningful if predictors are on different scales
- Standardization most common when predictors X_1, \ldots, X_{p-1} have very different scales
- β_k^* is expected change in Y for every <u>SD</u> (not unit) increase in predictor X_k , while all other predictors are held constant
- Standardizing has:
 - no effect on VIF
 - marginal effect on proportions of variance in Condition Index output
 - possibly substantial effect on magnitude of Condition Indexes
- Recommendations:
 - Standardize if either:
 - * desire common scale of b_k^* estimates
 - * need uncorrelated, higher-order predictors

3.3 Multicollinearity Summary

Three ways to diagnose multicollinearity:

- 1. combination of condition index <u>and</u> proportion of variation
- 2. variance inflation factors
- 3. model F-test vs. individual t-tests

Possible remedial measures for multicollinearity:

- Collect more data
- Choose a subset of predictor variables
- Ridge regression
- Latent root regression use Principal Components as predictors (may lack interpretability)

$$PC = a_1 X_1 + a_2 X_2 + \ldots + a_{p-1} X_{p-1}$$

Stat 5100 Handout 2.6.1 – SAS: Inference with Multiple Predictors

Example: (Table 7.1) Study seeks to relate (in females) amount of body fat (Y) to triceps skinfold thickness (X_1) , thigh circumference (X_2) , and midarm circumference (X_3) . Amount of body fat is expensive to measure, requiring immersion of person in water. This expense motivates the desire for a predictive model based on these inexpensive predictors.

- Q1: Do thigh and midarm both have no effect on body fat when triceps is in the model?
- Q2: Do the relationships among the predictors cause any problems in the fitted model?

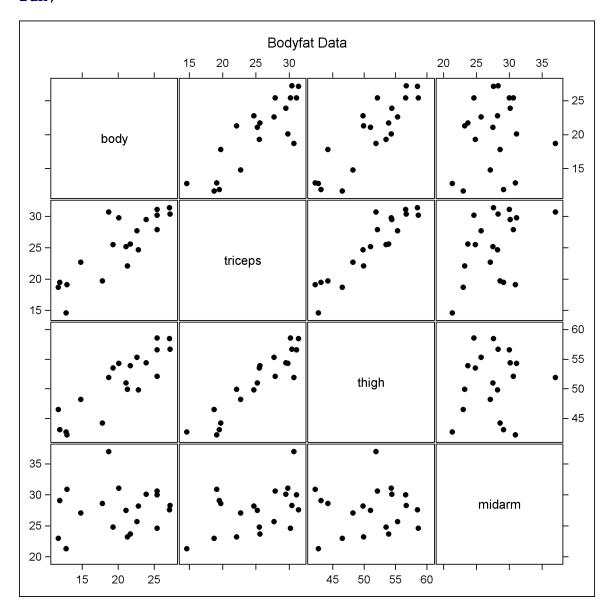
```
/* Input data */
data bodyfat;
   input triceps thigh midarm body @@; cards;
  19.5
        43.1
              29.1
                    11.9
                                     49.8
                                           28.2
                               24.7
                                                 22.8
  30.7
        51.9
              37.0
                    18.7
                               29.8
                                     54.3
                                           31.1
                                                 20.1
  19.1
        42.2
              30.9
                    12.9
                                           23.7
                               25.6
                                     53.9
                                                 21.7
  31.4
        58.5
              27.6
                    27.1
                               27.9
                                     52.1
                                           30.6
                                                 25.4
  22.1
        49.9
              23.2
                    21.3
                                           24.8
                               25.5
                                     53.5
                                                 19.3
  31.1
        56.6
              30.0
                    25.4
                                     56.7
                                           28.3
                                                 27.2
                               30.4
  18.7
        46.5
              23.0
                    11.7
                               19.7
                                     44.2
                                           28.6
                                                 17.8
                    12.8
  14.6
        42.7
                                     54.4
                                           30.1
              21.3
                               29.5
                                                 23.9
  27.7
        55.3
              25.7
                    22.6
                               30.2
                                     58.6
                                           24.6
                                                 25.4
  22.7
        48.2
              27.1
                    14.8
                               25.2
                                     51.0
                                           27.5
                                                 21.1
```

proc corr data=bodyfat;
 var body triceps
 thigh midarm;
 title1 'Correlation
matrix';
run;

run;

	Pearson Correlation Coefficients, N = 20 Prob > r under H0: Rho=0									
	body	body triceps thigh								
body	1.00000	0.84327 <.0001	0.87809 <.0001	0.14244 0.5491						
triceps	0.84327	1.00000	0.92384	0.45778						
thigh	0.87809	0.92384	1.00000	0.0424						
	<.0001	<.0001		0.7227						
midarm	0.14244 0.5491	0.45778 0.0424	0.08467 0.7227	1.00000						

```
proc sgscatter data=bodyfat;
  matrix body triceps thigh midarm/
     markerattrs=(symbol=CIRCLEFILLED size=2pt);
  title1 'Bodyfat Data';
run;
```



```
/* Q1: Test whether thigh and midarm BOTH have
   no effect on body when triceps is in the model */
proc reg data=bodyfat;
  model body = triceps thigh midarm;
  title1 'Bodyfat Regression';
  title2 '(full model)';
run;
```

Bodyfat Regression (full model)

Analysis of Variance										
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F					
Model	3	396.98461	132.32820	21.52	<.0001					
Error	16	98.40489	6.15031							
Corrected Total	19	495.38950								

```
proc reg data=bodyfat;
  model body = triceps;
  title1 'Bodyfat Regression';
  title2 '(reduced model)';
run;
```

Bodyfat Regression (reduced model)

Analysis of Variance										
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F					
Model	1	352.26980	352.26980	44.30	<.0001					
Error	18	143.11970	7.95109							
Corrected Total	19	495.38950								

```
data temp;
   F = ( (143.11970-98.40489)/2 ) / ( 6.15031 );
   p = 1-probf(F,2,16);
proc print data=temp;
   title1 'Subset F-test, by hand';
run;
```

Subset F-test, by hand							
Obs	F	p					
1	3.63517	0.049950					

```
/* Do this subset F-test, automatically.
   Also look at related quantities:
   See all sequential sums of squares and
   coefficients of partial determination */
proc reg data=bodyfat;
   model body = triceps thigh midarm / ss1 pcorr1;
   subsetcheck: test thigh=midarm=0;
   title1 'Subset F-test, automatically';
run;
```

Subset F-test, automatically

	Parameter Estimates											
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t	Type I SS	Squared Partial Corr Type I					
Intercept	1	117.08469	99.78240	1.17	0.2578	8156.76050	•					
triceps	1	4.33409	3.01551	1.44	0.1699	352.26980	0.71110					
thigh	1	-2.85685	2.58202	-1.11	0.2849	33.16891	0.23176					
midarm	1	-2.18606	1.59550	-1.37	0.1896	11.54590	0.10501					

Test subsetcheck Results for Dependent Variable body									
Source DF Mean F Value Pr > F									
Numerator	2	22.35741	3.64	0.0500					
Denominator	16	6.15031							

```
/* Q2: Investigate effect of relationships among
predictors. */

/* Standardizing all variables */
proc reg data=bodyfat;
  model body = triceps thigh midarm / stb;
  title1 'Standardized regression coefficients';
  title2 '(note extra column in output)';
run;
```

Standardized regression coefficients (note extra column in output)

Parameter Estimates							
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t	Standardized Estimate	
Intercept	1	117.08469	99.78240	1.17	0.2578	0	
triceps	1	4.33409	3.01551	1.44	0.1699	4.26370	
thigh	1	-2.85685	2.58202	-1.11	0.2849	-2.92870	
midarm	1	-2.18606	1.59550	-1.37	0.1896	-1.56142	

```
/* Test for multicollinearity */
proc reg data=bodyfat;
  model body = triceps thigh midarm / vif collin;
  title1 'Test for multicollinearity';
run;
```

Test for multicollinearity

Analysis of Variance							
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F		
Model	3	396.98461	132.32820	21.52	<.0001		
Error	16	98.40489	6.15031				
Corrected Total	19	495.38950					

Parameter Estimates								
Variable	DF	Parameter Estimate	Standard Error	t Value	Pr > t	Variance Inflation		
Intercept	1	117.08469	99.78240	1.17	0.2578	0		
triceps	1	4.33409	3.01551	1.44	0.1699	708.84291		
thigh	1	-2.85685	2.58202	-1.11	0.2849	564.34339		
midarm	1	-2.18606	1.59550	-1.37	0.1896	104.60601		

Collinearity Diagnostics								
Number	Eigenvalue	Condition Index	Proportion of Variation					
			Intercept	triceps	thigh	midarm		
1	3.96796	1.00000	0.00000195	0.00000320	0.00000110	0.00000980		
2	0.02052	13.90482	0.00037152	0.00132	0.00003262	0.00139		
3	0.01151	18.56570	0.00059915	0.00021875	0.00032550	0.00693		
4	0.00000865	677.37207	0.99903	0.99846	0.99964	0.99167		