Homework 3

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Problem 1

(a) $\Sigma_{j=0}^{p-1}c_j\hat{\beta}_j$ can be denoted as $c^T\hat{\beta}$,where c and $\hat{\beta}$ are vectors. In least squares estimate, $\hat{\beta}=(X^TX)^{-1}X^TY$, so

Mean:
$$E(\sum_{i=0}^{p-1} c_i \hat{\beta}_i) = c^T \beta$$

Variance:
$$Var(c^T\hat{\beta}) = c^T\sigma^2(X^TX)^{-1}c$$

Distribution:
$$c^T \hat{\beta} \sim N(c^T \beta, c^T \sigma^2 (X^T X)^{-1} c)$$

(b) Consider that $c^T\hat{\beta} \sim N(c^T\beta, c^T\sigma^2(X^TX)^{-1}c)$. Then we know that $c^T\hat{\beta} - c^T\beta \sim N(0, c^T\sigma^2(X^TX)^{-1}c)$.

Under H_0 , we use T test, and the test statistics is

$$T = \frac{c^T \hat{\beta} - h}{c^T \sigma^2 (X^T X)^{-1} c}$$

Under significant level α , the reject region is

$$|T| > t_{n-n,\alpha/2}$$

In this case, $\hat{\beta}$ is independent.

(c)

$$E(Y_{n+1} - \hat{Y}_{n+1}) = 0$$

$$Var(Y_{n+1} - \hat{Y}_{n+1}) = \sigma^2 (1 + Z^T (X^T X)^{-1} Z)$$

$$Y_{n+1} - \hat{Y}_{n+1} \sim N(0, \sigma^2 (1 + Z^T (X^T X)^{-1} Z))$$

(d) Proof:

$$MSE(\hat{Y}_{n+1}) = E(\hat{Y}_{n+1} - Y_{n+1})^2 = \sigma^2 (1 + Z^T (X^T X)^{-1} Z)$$
$$Z^T (X^T X)^{-1} Z = Z^T (X^T X)^{-1} X^T X (X^T X)^{-1} Z > 0$$
$$MSE(\hat{Y}_{n+1}) = \sigma^2 (1 + Z^T (X^T X)^{-1} Z) > \sigma^2$$

(e) We consider the T test.

$$T = \frac{Y_{n+1} - \hat{Y}_{n+1}}{\sqrt{\sigma^2 (1 + Z^T (X^T X)^{-1} Z)}} \sim t_{n-p}$$

So, the interval I

$$(\hat{Y}_{n+1} - t_{n-p,\alpha/2} \sqrt{\sigma^2 (1 + Z^T (X^T X)^{-1} Z)}, \hat{Y}_{n+1} + t_{n-p,\alpha/2} \sqrt{\sigma^2 (1 + Z^T (X^T X)^{-1} Z)}$$

Problem 2

(a)

The first model can be described as a single line because it is a SLR.

The second model can be described as a parallel lines because it contains indicator variables thus it can change the intercept when lmass=0, but cannot change the slope.

The third model can be described as a separate lines because it contains cross variables so it has both different intercepts and slopes.

(b)

F-test: whether there is a difference between the in-flight energy and body mass among birds, echolocating and non-echolocation bats. The second model is the reduced model and the first model is the full model.

Other test: whether there is a difference for birds, echolocating bats and nonecholocating bats after body mass is accounted for. The second model is the full model and the first model is the reduced model. (c)

> anova(lm1,lm2,lm3)

Analysis of Variance Table

Model 1: lenergy ~ lmass

Model 2: lenergy ~ lmass + Type

Model 3: lenergy ~ Type * Imass

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

1 18 0.58289

2 16 0.55332 2 0.029574 0.4100 0.6713

3 14 0.50487 2 0.048450 0.6718 0.5265

Comparing the first, the second and third model, the p-values are 0.6713, 0.5265 seperately. Hence, the first model is the best, wich means that the full model is reduced to

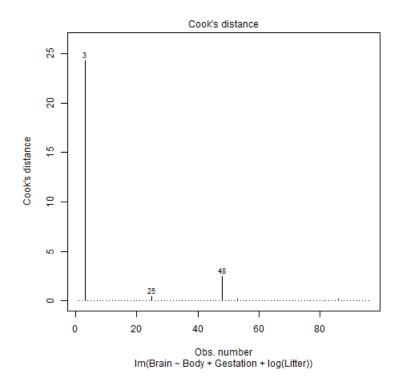
 $lenergy \sim lmass$

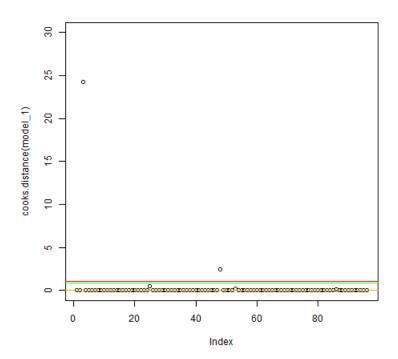
Problem 3

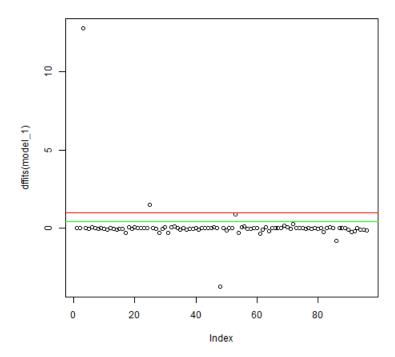
(a) By using R, we do the regression:

$$Y = -231.58 + 0.97X_1 + 1.93X_2 + 89ln(X_3)$$

To find influential points, we compute the cook's distance and fiffits value and the graphs are shown below:





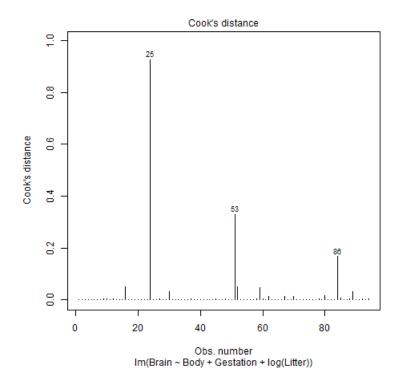


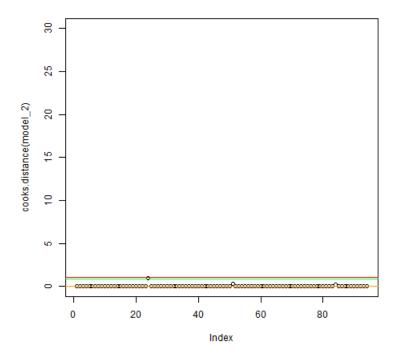
Based on the graphs, we can see that there are two influential points: 3, 25, 48, 53 and 86, which corresponding to African elephant, dolpin, hippopotamus, human being and Tapir respectively.

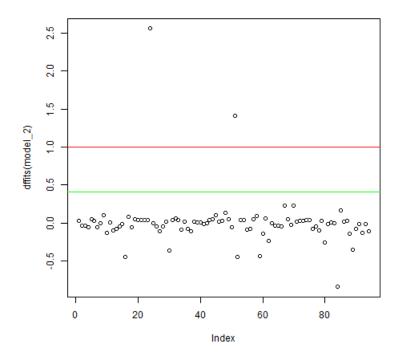
(b) After deleting the influential points, we do the regression again and get

$$Y = -135.44 + 0.47X_1 + 1.972X_2 + 39.38ln(X_3)$$

To find influential points, we compute the cook's distance and diffits value and the graphs are shown below:







Based on the graphs, there are still two influential points; 25 and 53.

Main differences: The coefficient of body weight and litter size and interception changed, F-value and t-value are decreasing.

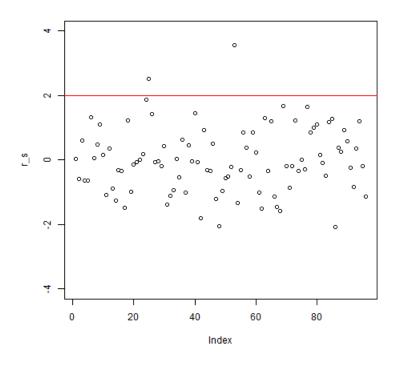
(c) Consider all the data and fit the regression of log brain weight on log body weight, log gestation, and log litter size:

$$ln(Y) = 0.85482 + 0.57507 ln(X_1) + 0.41794 ln(X_2) - 0.31007 ln(X_3)$$

Then, we can calculate the studentized residuals:

> r_s							
1	2	3	4	5	6		8
0.03743244	-0.59640636	0.60328414	-0.62769966	-0.63113474	1.32746311	0.04998926	0.47261422
9	10	11	12	13	14	15	16
1.09307100	0.16232902						-0.33613386
17	18	19	20	21	22	23	24
	1.11.30330						1.87603881
25	26	27	28	29	30	31	32
CONTRACTOR STANDARDS	1.42291877						-1.10691827
33		35	36		38	39	40
-0.94276157				-1.00847048			1.43773970
41	42	43	44	45		47	48
10000	-1.81623410						
49	50	51	52		54	55	56
	-0.55595462						
57		59	60	61	62	63	64
and a more and a second property	-0.51348066						
65	66	67	68	69	70	71	72
	-1.14859836						
73	74	75	76	77	78	79	80
	-0.33484868						
81	82	83	84	85	86	87	88
The second secon	-0.10334766						0.26424150
89	90	91	92		94	95	96
0.92300908	0.57397277	-0.23412/28	-0.8280693/	0.35813822	1.194588/0	-0.1884//18	-1.14395/81

(d) To find influential points, we compute the cook's distance and fiffits value and the graphs are shown below:



we can conclude that at the points 25 and 53, which corresponding to dolphin and human, have substantially larger brain weights than were predicted by the model.

(e) According to the graphics shown in (d), the conclusion is — there is no mammals have substantially smaller brain weights than were predicted by the model.

(f)

- 1. After log transformation, F-value increased, which means the model fits better.
- 2. Removing influential points and refit the regression may not be a good way to make thing right, even make it worse.
- 3. The assumption of normality can hold by log transformation.

Problem 4

(a) SSTO = 8100 and $\hat{\sigma^2} = \frac{SSE}{df}$. The estimates of σ^2 are shown below:

Model variables	$\hat{\sigma}^2$
None	300
Α	240
В	230
С	260
AB	220
AC	210
ВС	230
ABC	215

(b) Adjusted $R^2=1-\frac{\frac{SSE_p}{n-p}}{\frac{SSTO}{N-1}}.$ The adjusted R^2 are shown below:

Model variables	Adjusted \mathbb{R}^2	
None	0	
Α	0.2	
В	0.233	
С	0.133	
AB	0.267	
AC	0.3	
ВС	0.233	
ABC	0.283	

(c) $C_p(M) = \frac{SSE(M)}{\hat{\sigma}^2} - n + 2p(M)$, the C_p statistics are shown below:

Model variables	C_p
None	11.674
Α	5.023
В	3.814
С	7.442
AB	3.851
AC	2.419
ВС	4.744
ABC	4

(d) $BIC_p = nln(SSE_p) - nln(n) + pln(n)$. The BIC are shown below:

Model variables	BIC_p
None	162.0198
Α	158.0473
В	156.8556
С	160.2885
AB	157.8450
AC	156.5424
ВС	159.0896
ABC	159.3905

(e)

(i)The last model with variables A, B and C has the smallest $\hat{\sigma}^2$, 198.46.

(ii)The model with variables A and C has the largest adjusted $\ensuremath{\mathit{R}}^2$, 0.3.

(iii)The model with variables A and C has the smallest C_p statistic, 2.419.

(iv)The model with variables A and C has the smallest $\it BIC$, 156.54.

(f)

Starting with none variable, we get the model contains Bhas the smallest SSE. Then we perform an extra-sum-of-squares F -test to see whether that variable is significant.

Under $\alpha = 0.05$, F = 7.75. Thus, B is significant.

Then, we get AB has the smaller SSE. Samely, we perform an extra-sum-of-squares F -test to see whether the additional variable is significant.

Under $\alpha = 0.05$, $F = 2.182 < 4.24 = F_{1,n-4,0.05}$. So Ais insignificant.

Finally, by forward selecion, we have the model form:

$$Y\beta_0 + \beta_2 B + \epsilon$$

(g) To find the posterior probability for these 8 models, we first subtract the smallest BIC from the other ones, and then calculate the difference of each model as below:

$$BIC_{j}^{*} = BIC_{j} - BIC_{min}$$

Then, we have the posterior probability:

$$p(M_j) = \frac{e^{-\frac{1}{2}BIC_j^*}}{\sum_{j=1}^k e^{-\frac{1}{2}BIC_j^*}}$$

The posterior distributions are shown below:

Model variables	Posterioir distribution	
None	0.01802701	
Α	0.13138647	
В	0.23840663	
С	0.04284313	
AB	0.14537115	
AC	0.27882110	
ВС	0.07802002	
ABC	0.06712450	

A Appendix: Problem 2

```
bats = read.csv( file = "bats.csv" , header = T)
bats = transform(bats , lenergy = log(Energy))
bats = transform(bats , lmass = log(Mass))
lm1 = lm(lenergy ~ lmass , data = bats)
lm2 = lm(lenergy ~ lmass + Type, data = bats)
lm3 = lm(lenergy ~ Type*lmass , data = bats)
anova(lm1,lm2,lm3)
```

B Appendix: Problem 3

```
data = read.csv( "brain.csv")
model_1=Im(Brain ~ Body+Gestation+log(Litter) ,data = data)
summary (model_1)
cooks.distance(model_1)
n = dim(model.matrix(model_1))[1]
p = dim(model.matrix(model_1))[2]
png("3_a_1.png")
plot(model_1, which = 4)
dev. off ()
png( "3_a_2.png")
plot(cooks.distance(model_1) , ylim = c (0,30))
abline(h=1, col="red")
abline (h=qf (0.5 , p, n-p), col="green")
abline(h=4/n, col="blue")
abline (h=4/(n-p-1-1), col="orange")
dev. off ()
dffits (model_1)
png("3_a_3.png")
plot( dffits (model_1) )
abline(h=1, col="red")
abline(h=2*sqrt(p/n), col="green")
```

```
dev. off ()
summary(influence.measures(model 1))
data_new = data[-48,][-3,]
model 2=Im(Brain ~ Body+Gestation+log(Litter), data = data new)
summary (model 2)
cooks.distance(model 2)
n = dim(model.matrix(model 2))[1]
p = dim(model.matrix(model 2))[2]
png("3_b_1.png")
plot(model_2, which = 4)
dev. off ()
png( "3_b_2.png")
plot(cooks.distance(model_2), ylim = c (0,30))
abline(h=1, col="red")
abline (h=qf (0.5, p, n-p), col="green")
abline(h=4/n, col="blue")
abline (h=4/(n-p-1-1), col="orange")
dev. off ()
dffits (model_2)
png("3_b_3.png")
plot( dffits (model_2) )
abline(h=1, col="red")
abline(h=2*sqrt(p/n), col="green")
dev. off ()
summary(influence.measures(model_2))
```