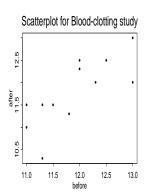
## Homework #8, ST518

1. A person's blood-clotting ability is typically expressed in terms of a "prothrombin time," which is defined to be the interval between the initiation of the prothrombin-thrombin (two proteins) reaction and the formation of the final clot. Does *aspirin* affect this function? Measurements made before administration of two tablets and three hours after.

Prothrombin Times (seconds)				
Subject	Before Aspirin $(Y_1)$	After Aspirin $(Y_2)$	Difference $(D)$	
1	12.3	12.0	0.3	
2	12.0	12.3	-0.3	
3	12.0	12.5	-0.5	
4	13.0	12.0	1.0	
5	13.0	13.0	0.0	
6	12.5	12.5	0.0	
7	11.3	10.3	1.0	
8	11.8	11.3	0.5	
9	11.5	11.5	0	
10	11.0	11.5	-0.5	
11	11.0	11.0	0	
12	11.3	11.5	-0.2	



- (a) Carry out a paired t-test of the hypothesis that prothrombin time is unaffected by aspirin.
- (b) Carry out an F-test of the same hypothesis treating subjects as blocks in an analysis for a RCBD.
- (c) (not graded) Show that, in general, the paired t-test is equivalent to the F-test for the RCBD with block size equal to 2. One approach is to write squared differences involving  $Y_{ij}, \overline{Y}_{i\cdot}, \overline{Y}_{\cdot j}, \overline{Y}_{\cdot i}$  in terms of differences,  $D_j, \overline{D}$ .
- (d) Consider the mixed model

$$Y_{ij} = \mu + \tau_i + B_j + E_{ij}$$

where  $B_j \stackrel{iid}{\sim} N(0, \sigma_B^2)$  and  $E_{ij} \stackrel{iid}{\sim} N(0, \sigma^2)$  with  $B \perp E$  for i = 1, ..., a and j = 1, ..., b.

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- i. Show that  $E[MS(block)] = \sigma^2 + a\sigma_B^2$
- ii. Use this result to estimate the variance component for subject effects in a mixed model for the prothrombin data.
- iii. Report an estimate of the intra-subject correlation. Is the scatterplot above consistent with this estimate?

2. (taken from Ott and Longnecker 15.10, p. 889) Fuel efficiency of four blends of gasoline is measured in MPG. There is considerable variability due to driver. Another source of variability is model of car. An experiment randomizes four models of car and gasoline blends (A,B,C,D) to drivers according to the design below

	Model			
Driver	1	2	3	4
1	15.5(A)	33.8(B)	13.7(C)	29.2(D)
2	16.3(B)	26.4(C)	19.1(D)	22.5(A)
3	10.5(C)	31.5(D)	17.5(A)	30.1(B)
4	14.0(D)	34.5(A)	19.7(B)	21.6(C)

- (a) Assuming normally distributed data, propose a model in which the effects of model, driver and blend are additive on the mean.
- (b) Obtain an ANOVA table for this model.
- (c) Report the average fuel efficiency for each blend and the lowest level of significance  $\alpha$  at which these averages can be said to differ significantly.
- 3. For each of several species, an Ecology researcher ran an exposure assay in which groups of n=25 ants are measured for mortality after exposure to one of three different bacteria. For each species, ants are randomized to the three bacteria treatments within each of 15 colonies. That is, a randomized complete block design is used for each species, with colonies serving as complete blocks. Find the results for the DB species on moodle as "DBdat.mtx".
  - (a) Watch the video on the hiddenf package. What is the name of the function in R that will create a directory system containing all of the help files that a developer can complete to create documentation for a package? \_\_\_\_\_\_\_skeleton.
  - (b) Obtain an interaction plot with proportion of ants dying out of n = 50 on the vertical axis, bacteria treatment on the horizontal axis and lines connecting mortality rates from the same colony.
  - (c) Either by using the hiddenf package in R, or by assigning colonies to groups, obtain an ANOVA table with the following sources of variability: treatments, groups, group-by-treatment interaction, colony within group.

In SAS use can use this code:

```
data DB;
   set DB;
   group=2-(colony in (1,2,4,6,7));
run;
```

```
proc glm;
  *by species; *there were other species but HW8 only uses species=DB;
  class treatment group colony;
  model y=treatment|group colony(group);
  lsmeans treatment|group;
run;
```

- (d) Obtain the p-value for the test of group-by-treatment interaction after and report it after multiplying by  $2^{15-1} 1$ . Is there evidence that the resistance to bacteria varies across colonies?
- (e) Obtain another plot of survival versus bacteria with different lines for colonies. Color the colony lines according to group.

- 4. An experiment investigates the growth of oysters. Four bags with ten oysters each are randomly placed at four underwater stations next to a power plant:
  - Trt1: At the bottom of a discharge canal
  - Trt2: At the top of a discharge canal
  - Trt3: At the bottom of an intake canal
  - Trt4: At the top of an intake canal

(See "oysters.sas" and "oysters.dat" on moodle.) Average initial weight x and final weight y are measured for each of the 16 bags. (Bags serve as the experimental units.) Let  $z = x - \bar{x}$  denote the difference from average of the initial weights. SAS code and output to fit an ANCOVA model appear at the end of the exam.

- (a) Obtain the F-ratio for a test of equal final weights in a one-way ANOVA where initial weight z is ignored.
- (b) Obtain the F-ratio for a test of equal final weights in a one-way ANOVA after controlling for initial weight z.
- (c) Use the output to report the unadjusted means for treatments 1 and 4. (Use the  $2^{nd}$  column in the table below.)

	Unadjusted	Adjusted	Std.
Treatment	Mean	Mean	Error
1			
_	_	_	
4			

(d) For bag i, let  $x_{i1}, \ldots, x_{i4}$  denote indicator variables for treatments 1-4, respectively. Consider the analysis of covariance model:

$$y_i = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i3} + \beta_4 x_{i4} + \beta_5 z_i + E_i$$

- i. Use the fitted model to report the mean final weight, after adjustment to the average initial weight  $\bar{x}$ , for treatments 1 and 4. Fill in the appropriate column in the table above. Show work.
- ii. Report the standard error for the adjusted means for locations 1 and 4. (Fill in the table, writing "NP" if it is not possible to give a number based on the provided output.)
- (e) Consider the difference between mean final weights under treatments 1 and 4. Estimate this difference after controlling for initial weight. Report a standard error and a p-value for a test of no difference.

```
proc glm; /* These data taken from Freund, Littell and Spector */
   class trt;
   model final=trt z /solution;
   means trt;
run;
```

output given next page

		he SAS System e GLM Procedu		1
	111	e dim Flocedu	16	
	Class	Levels	Values	
	trt	4	1 2 3 4	
	2.5	Sum of	у с п	
Source	DF	Squares	Mean Square F 36.1888402	Value Pr > F
Model	4 11	144.7553608		113.59 <.0001
Error			0.3186036	
Corrected Total	15	148.2600000		
R-Squ	are Coeff	Var Root	MSE final Me	an
0.976			4450 32.300	
Source	DE	Tuno I cc	Moon Square E	Volue Pr > E
trt	3 JU	19pe 1 55	Mean Square F 9.6816667	20 20 < 0001
z	1	115 7103609	115.7103608	363 18 < 0001
<sup>2</sup>	1	115.7103606	115.7103606	363.16 \.0001
Source	DF	Type III SS	Mean Square F 2.5580345	Value Pr > F
trt				
z	1	115.7103608	115.7103608	363.18 <.0001
		S+a	ndard	
Parameter	Estimate		Error t Value	Pr >  t
Intercept	32.82685402	B 0.283	98639 115.59	<.0001
	-1.23028630			0.0172
trt 2	-1.36002698	B 0.401	24637 -3.39	0.0060
trt 3	-1.36002698 0.48289720	B 0.410	86022 1.18	0.2647
trt 4				
z	1.04670265		92404 19.06	<.0001
NOTE: The X'X matrix has been found to be singular, and a generalized				
11			quations. Terms	
estimates a	re followed by	the letter '	B' are not unique	ely estimable.
Level of -		L	z	
trt N	Mean	Std Dev		Std Dev
1 4	34.4750000 31.6500000	3.18891309	2.75000000	3.20572405
2 4	31.6500000	1.53731367	2.75000000 0.17500000	0.96046864
3 4	30.8500000	2.95578529	-2.35000000	2.75862284
		4.29757684	-0.57500000	4.04917687