

### STAT 512 Homework Assignment 3: Due in class, Friday September 14, 2011

1. Cline et al. (1997) investigated the effect of the level of vitamin A on bone density in dogs using a complete block design. From their abstract:

*“Thirty-two random-source dogs were assigned to four treatments in a randomized complete block design. The diets contained 15,000, 50,000, 116,000, or 225,000 IU vitamin A/1,000 kcal ME. Diets were fed up to 1 yr. Computed tomography was used to determine bone density of the right tibia at ... 12 mo. ... There was no difference ( $p > .10$ ) in tibia bone or marrow density in any of the dogs fed the various concentrations of vitamin A.”*

Assume the investigators used blocks of 4 similar dogs, and assigned one dog from each block to each treatment.

- (a) Suppose that the diets actually *do* influence bone density, i.e. that the investigators made a type II error in this study, testing at the 0.10 level. What would the value of the non-centrality parameter for this test have to be in order for the probability of this error to be 0.5? (Determine this to 2 significant places numerically by trial and error.)
  - (b) Now suppose a larger follow-up experiment of the same kind (CBD in blocks of size 4) might be performed with the hope of establishing differences among the 4 diets. If the non-centrality parameter for this first study actually was the value you calculated above, how many blocks would be needed in the follow-up study to result in a power of 0.9?
2. Consider a block experiment with 3 blocks of 4 units each ( $N = 12$ ). Using the notation developed in class:
    - (a) What is  $\mathbf{H}_1$ ?
    - (b) Suppose  $t = 4$ , and one unit is assigned to each treatment in each block (i.e. our definition of a standard CBD). What is  $\mathbf{X}_{2|1}$ ? Show why this design is or is not “Condition E-equivalent” to a CRD with  $n_1 = n_2 = n_3 = n_4 = 3$ .
    - (c) Suppose  $t = 3$ , and in each block one unit is assigned to each of treatments 1 and 2 and two units are assigned to treatment 3. What is  $\mathbf{X}_{2|1}$ ? Show why this design is or is not “Condition E-equivalent” to a CRD with  $n_1 = n_2 = 3$  and  $n_3 = 6$ .
    - (d) Suppose  $t = 3$ , and in block  $i$  ( $i = 1, 2, 3$ ), two units are assigned to treatment  $i$  and one unit is assigned to each of the other two treatments. What is  $\mathbf{X}_{2|1}$ ? Show why this design is or is not “Condition E-equivalent” to a CRD with  $n_1 = n_2 = n_3 = n_4 = 3$ .
    - (e) Continuing with the design specified in part (d) of this problem, derive the reduced normal equations for  $\hat{\tau}$ . (Note that you will have to be careful about the way you define notation for data here. I suggest using  $y_{i,j}$  for the datum from block  $i$  and treatment  $j$  when  $i \neq j$ , and  $\bar{y}_{i,i}$  for the average of the two responses with common treatment in each block.)

**Reference:** Cline, J.L., G.L.Czarnecki-Maulden, J.M. Losonsky, C.R. Sipe, and R.A. Easter (1997). “Effect of increasing dietary vitamin A on bone density in adult dogs,” *J. Anim. Sci.* **75**, 2980-5.