GLM/GAM Online Course Exercises: Day 1

Linear Modeling

- 1) The R data frame warpbreaks gives the number of breaks per fixed length of wool during weaving, for two different wool types, and 3 different weaving tensions. Using a linear model, establish whether there is evidence that the effect of tension on break rate is dependent on the type of wool. If there is, use interaction.plot() function to examine the nature of the dependence.
- 2) The **R** data frame **cars** contains data about the stopping distance and speed of cars when the driver was signaled to stop. It takes a fixed reaction time for drivers to apply their brakes, so the car will travel a distance directly proportional to its speed before beginning to slow. However, an automobile's kinetic energy is proportional to the square of its speed, but the brakes can only dissipate that energy, and slow the car, at a constant rate per unit distance traveled.

Fit three different linear models to this data. Report the results.:

- a) dist $\sim \beta_0 + \beta_1$ (speed) + β_2 (speed²) + e
- b) dist ~ β_1 (speed) + β_2 (speed²) + e
- c) dist $\sim \beta_1$ (speed) + e

Which model seems to fit better? Why?

Using the second model in the above list, estimate the average time that it takes a driver to apply the brakes (there are 5280 feet in a mile).

Generalized Linear Modeling

3) The following table shows numbers of occasions when inhibition (i.e., no flow of current across a membrane) occurred within 120 s, for different concentrations of the protein peptide-C. The outcome *yes* implies that inhibition has occurred. Use logistic regression to model the probability of inhibition as a function of protein concentration. Report and plot your results fully. Interpret your results.

conc	0.1	0.5	1	10	20	30	50	70	80	100	150
no	7	1	10	9	2	9	13	1	1	4	3
yes	0	0	3	4	0	6	7	0	0	1	7

4) The R data frame ACF1 in the package DAAG consists of two columns: count and endtime. The first column contains the counts of simple aberrant foci (ACFs). These are aberrant aggregations of tube-like structures in the rectal end of 22 rat colons after administration of a dose of the carcinogen azoxymethane. Each rat was sacrificed after 6, 12 or 18 weeks. Create a scatterplot of count by (~) endtime.

Run two glm models. The first specifies count (as the response variable) as predicted by endtime (the explanatory variable) and uses a poisson family for the distribution. Plot the results. Interpret the results. Then run a second model adding an endtime^2 term to the right hand side to accommodate a possible quadratic effect. Plot the results. Interpret the results. Compare the two models with an anova table. Which model 'fits' better? Why?