

# Problem set 8

## Analysis for Neuroscientists

1. In the next class, we will encounter the *general linear model*, in which we relate dependent variables  $\mathbf{Y}$  to independent variables  $\mathbf{X}$  according to the equation:

$$\mathbf{Y} = \mathbf{X}\boldsymbol{\beta} + \mathbf{U}.$$

Assume

$$\mathbf{Y} = \begin{pmatrix} Y_{11} \\ Y_{21} \\ Y_{31} \end{pmatrix}, \mathbf{X} = \begin{pmatrix} X_{11} & X_{12} \\ X_{21} & X_{22} \\ X_{31} & X_{32} \end{pmatrix}, \boldsymbol{\beta} = \begin{pmatrix} \beta_{11} \\ \beta_{21} \end{pmatrix}, \mathbf{U} = \begin{pmatrix} U_{11} \\ U_{21} \\ U_{31} \end{pmatrix}.$$

Write down the expression for  $Y_{11}$ ,  $Y_{21}$ , and  $Y_{31}$  in terms of the other variables using the rules of matrix multiplication and addition.

2. In `prob2.mat` there is a variable `spikes`. `spikes{c}` contains a  $500 \times 100$  array of 0s and 1s representing the spikes of neuron  $c$  (3 neurons total) during 100 trials of 500 ms each. We're going to compute cross-correlograms (CCGs) for different pairs of neurons to see which ones are connected.

The function `calcccg(s1,s2,Nlags,dt)` will compute the cross-correlogram given `s1` and `s2`, which are vectors of 0s and 1s representing spikes. `Nlags` is the number of bins to the left and right of 0 to be included. For this problem, you can use `Nlags=20`. A peak to the right of 0 means `s1` tends to fire after `s2`.

- (a) Compute the average (over all 100 trials) cross-correlogram for a pair of neurons by using `calcccg` and averaging the result over the trials. You can base your code on this snippet:

```
load('prob2.mat');

Nlags = 20;
t = dt*(-Nlags:Nlags);

ccg = zeros(length(t),1);

%fill in code here
```

```
plot(1000*t,ccg); %factor of 1000 to convert seconds to milliseconds
xlabel('Lag (ms)');
ylabel('Cross-correlation (Hz)');
```

- (b) Do this for all pairs of cells. Which pairs seem to have a synaptic connection?
- (c) For one of the connections you identified in the previous question, remake the CCG but by shuffling the trials of one of the neurons. You can use `randperm(Ntrials)` to get a randomly reordered vector of trial numbers. Plot the shuffled CCG on top of the true CCG.

## References