## POLS0012 Causal Analysis: Tutorial Exercise 4

In this question, we'll use a simulated dataset to illustrate randomisation inference, based on Table 3.3 in the Gerber and Green textbook, an example of an experiment created with block randomisation and 14 observations. We'll code the exact p-values by hand first, then using R's built-in code from the ri package. We'll also illustrate it with and without accounting for blocked randomisation. The dataset is called "a" and is in the file "t4\_data.Rda". It contains the following variables for 14 observations:

- Y: Outcome variable
- block: Two blocks used for randomisation, 1 or 2
- tr: =1 if in the treatment group, 0 if in the control group

You first need to install and load R's ri package using the code install.packages("ri") and library("ri")

- a) Calculate the ATE and its p-value from the experiment, ignoring the blocking (suppose we didn't realise this was a block-randomised experiment). Store the ATE as an object
- b) Using the genperms() function, create a matrix of all possible permutations of treatment assignment, ignoring the blocking. How many possible permutations are there?
- c) Now, create the null distribution of ATEs using the permutations matrix:
  - i) Create an empty vector called "ates" using ates <- c().
  - ii) Using a for() loop, fill in this vector with the ATE under each possible permutation of treatment

Code Hint: The columns of the matrix from (b) give you each possible permutation of treatment

- d) Calculate the exact two-tailed p-value using the estimated ATE and the null distribution. Is it higher or lower than the p-value calculated the traditional way in (a)?
  - **Hint:** You need to calculate the proportion of the null distribution that is at least as extreme as the estimated ATE. It can help to start with a drawing
- e) To visualise what is happening, plot the null distribution as a density or histogram, adding vertical lines for the estimated ATE and the critical values for the exact hypothesis test Code Hint: you can add vertical lines to a plot using abline(v=) and you can use the quantile() function to find the critical values

- f) Now we'll carry out the inference automatically using R's built-in functions from the ri library. The results should be identical to your hand-coded estimate in (d):
  - i) Declare the potential outcomes under the sharp null using the genouts() function. It takes three arguments: the observed outcome variable, the treatment variable, and the null hypothesis specified as ate=
  - ii) Generate the null distribution using the gendist() function. It takes two arguments: the potential outcomes from step (i) and the matrix of permutations from question (b)
  - iii) Calculate the p-value and plot the results using the dispdist() function. It takes two arguments: the distribution from step (ii) and the estimated ATE
- g) Finally, let's re-estimate everything, this time accounting for the blocking. First, calculate the ATE and its p-value from the experiment, accounting for the blocking. Is it different to (a)?

**Hint:** Remember that you need to control for block membership

h) Using the genperms() function, create a matrix of all possible permutations of treatment assignment, accounting for the blocking. How many possible permutations are there, compared to part (b)?

Code Hint: You need to add the argument blockvar= to the genperms() function

i) Repeat (f), this time using the permutations matrix from question (h). Comment on how the null distribution has changed, compared to (f)