# Buffalo First Analysis

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```
read_data <- function(i){
   filename <- paste("../data/nsim_1000/angle_sigma1_results_",i,"_10_1000", sep = "")
   dat <- read.csv(filename) %>%
      mutate(rho = i/10)
   return(dat)
}
dat <- lapply(1:10, read_data) %>% bind_rows
```

### Set Up

We compare the performance of propensity score matching, Mahalanobis distance matching, and Buffalo Matching (described in the previous section) on simulated data, varying the dimensionality of the problem, the fixed treatment to control ratio during matching, and the correlation between the true propensity and prognostic score. The generative model for all of our simulations is the following:

$$\begin{split} X_i \sim_{iid} \text{Normal}(0, I_p), \\ T_i \sim_{iid} \text{Bernoulli}\left(\frac{1}{1 + \exp(-\phi(X_i))}\right), \\ Y_i = \tau T_i + \Psi(X_i) + \epsilon_i, \\ \epsilon_i \sim_{iid} N(0, \sigma^2), \end{split}$$

where the true propensity and prognositic scores are given by the linear combinations

$$\phi(X_i) = X_{i1}/3 - 4,$$
  

$$\Psi(X_i) = \rho X_{i1} + \sqrt{(1 - \rho^2)} X_{i2},$$

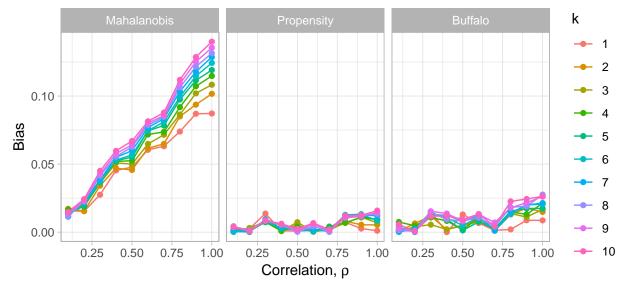
so that  $\operatorname{Cor}(\phi(X_i), \Psi(X_i)) \propto \rho$ . The propensity score formula was chosen such that there were approximately 100 treated observations in each dataset (But there aren't actually 100 treated in each dataset???). We consider  $p=10, \, \rho=0,0.1,\ldots,0.9,1.0$ , and  $k=1,\ldots,10$ . Each simulation consisted of a dataset of size n=2000 and was repeated N=1000 times. We fix the treatment effect to be constant with  $\tau=1$  and the noise to be  $\sigma=1$ . For a given matching, we estimate ATT and design sensitivity  $\tilde{\Gamma}$  using the permutation t-statistic from the package sensitivtymv

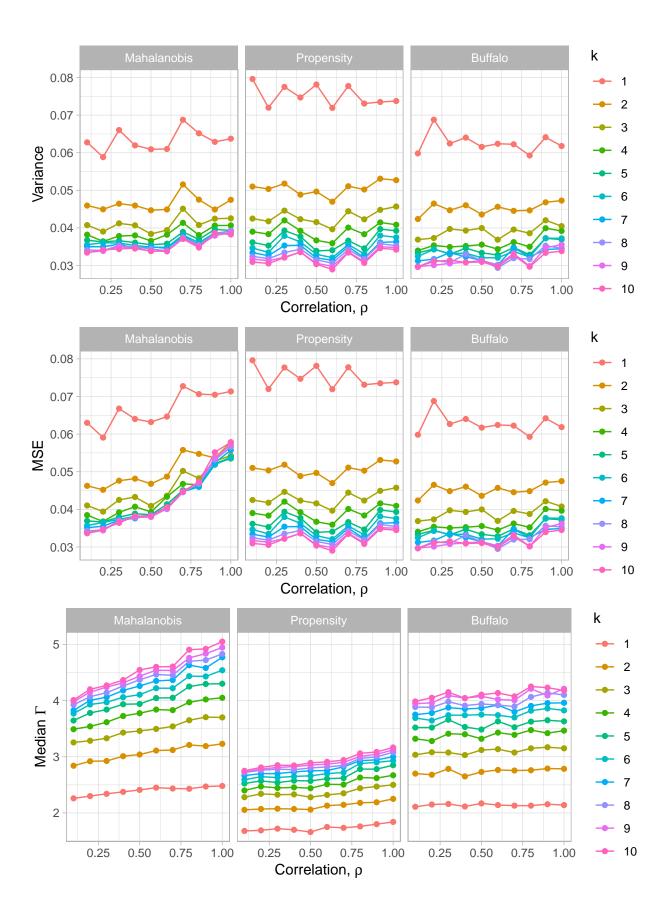
## Replication of Dylan's Plots

#### **Basic Visualization**

Here, I assume propensity and prognostic information are uncorrelated  $\rho = 0$ 

# Performance for 1:1 to 1:10 matching as correllation of propensity and prognostic score is modified





## Rocky Versions

