Homework1

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```
library(tidyverse)
library(knitr)
```

Part 1

Question 1

```
Y_obs <- c(8.62,1.48,8.93,9.57,2.65,7.3,.06,1.72,2.19,7.32,7.53,7.62)
Z <- c(rep(0,6), rep(1,6))
```

a)

```
Y_obs <- c(8.62,1.48,8.93,9.57,2.65,7.3,.06,1.72,2.19,7.32,7.53,7.62)
Z <- c(rep(0,6), rep(1,6))

tstat_obs <- mean(Y_obs[Z == 1]) - mean(Y_obs[Z == 0])

ind_combos <- combn(1:12,6)

tstats <- vector(mode = "double",length = ncol(ind_combos))

for (i in 1:ncol(ind_combos)) {
    Zperm <- rep(0,12)
    Zperm[ind_combos[,i]] = 1
    tstats[i] = mean(Y_obs[Zperm==1]) - mean(Y_obs[Zperm==0])
}

pval <- mean(abs(tstats) >= abs(tstat_obs))
```

The two-tailed p-value is 0.2706.

b)

```
set.seed(2929)
sampled_tstats <- sample(tstats, size = 1000, replace = TRUE)
pval_1000 <- mean(abs(sampled_tstats) >= abs(tstat_obs))
```

The two-tailed p-value from 1000 samples from the distribution under the Sharp Null Hypothesis is 0.27.

c)

```
ttest_pval <- t.test(Y_obs[Z==1], Y_obs[Z==0],var.equal = TRUE)$p.value
```

The p-value using a t-test is 0.3368.

d)

- (b)'s approximation of (a) is part of the assignment mechanism component of the potential outcome framework as it draws from the distribution of all possible treatment assignments.
- (c)'s approximation of (a) falls under the probabilistic model component of the potential outcome framework as it assumes the data in both groups is normally distributed with equal variance.

Question 2

a)

```
pval <- mean(abs(tstats) >= abs(tstat_obs))
```

The p-value from randomizing within pairs is 0.375.

b)

```
set.seed(2121)
sampled_tstats <- sample(tstats, size = 1000, replace = TRUE)
pval_1000 <- mean(abs(sampled_tstats) >= abs(tstat_obs))
```

The p-value from sampling is 0.398.

c)

```
ttest_pval <- t.test(Y_obs[Z==1], Y_obs[Z==0], var.equal = TRUE, paired = TRUE)</pre>
```

Using a paired t-test the p-value is 0.3652.

d)

Part (2b) is a part of the assignment mechanism as it makes sure $Z \perp X$ through randomization

Part (2c) is a part of the probabilistic model just like question 1.

Question 3

$$\begin{split} Y_i^{obs} &= Z_i Y_i(1) + (1 - Z_i) Y_i(0) \\ \hat{\tau} &= \frac{1}{n_1} \sum_{i=1}^n Z_i Y_i^{obs} - \frac{1}{n_0} \sum_{i=1}^n (1 - Z_i) Y_i^{obs} \\ \hat{\tau} &= \frac{1}{n_1} \sum_{i=1}^n Z_i Y_i(1) - \frac{1}{n_0} \sum_{i=1}^n (1 - Z_i) Y_i(0) \\ \text{Under CRE } Z_i \perp Y_i(0), Y_i(1) \text{ and } E[Y(z)] &= Y(z) \\ E[\hat{\tau}] &= \frac{1}{n_1} \sum_{i=1}^n E[Z_i] Y_i(1) - \frac{1}{n_0} \sum_{i=1}^n E(1 - Z_i) Y_i(0) \\ &= \frac{1}{n_1} \sum_{i=1}^n \frac{n_1}{n} Y_i(1) - \frac{1}{n_0} \sum_{i=1}^n \frac{n_0}{n} Y_i(0) \\ &= \frac{1}{n_1} \sum_{i=1}^n Y_i(1) - \frac{1}{n_0} \sum_{i=1}^n Y_i(0) = \tau \end{split}$$

Question 4

Yes, it is possible for there to be evidence of a additive treatment effect under randomization. If this is the case we can change the Null Hypothesis from $Y_i(0) = Y_i(1)$ to $Y_i(0) - Y_i(1) = \tau$ and set τ equal to some fixed value. To impliment this with the data above we can do a two-sample t-test but set μ equal to some fixed value.

Part 2

```
indexes <- sample_ind[,i]</pre>
    sample <- pot_outcomes[indexes,]</pre>
    for (j in 1:ncol(assign_ind)) {
       assignment <- assign_ind[,j]</pre>
       assignment <- assign_ind[,sample(1:6,1)]</pre>
       YO_obs <- sample[assignment, "YO"]
       Y1_obs <- sample[-assignment, "Y1"]
       diff <- mean(Y1_obs) - mean(Y0_obs)</pre>
      mean_diffs[i,j] = diff
    }
  }
From Simulation: 232.3602032
  var1 <- sum((pot_outcomes[,"Y1"]-mean(pot_outcomes[,"Y1"]))^2)/11</pre>
  var0 <- sum((pot_outcomes[,"Y0"]-mean(pot_outcomes[,"Y0"]))^2)/11</pre>
  diff_means <- mean(pot_outcomes[,"Y1"]) - mean(pot_outcomes[,"Y0"])</pre>
  var01 <- sum((pot_outcomes[,"Y1"] - pot_outcomes[,"Y0"] - diff_means)^2)/11</pre>
  form <- var0/6 + var1/6 - var01/12
```

From Formula: 75.4261364

```
sigma <- var(Y0_obs)/2 + var(Y1_obs)/2

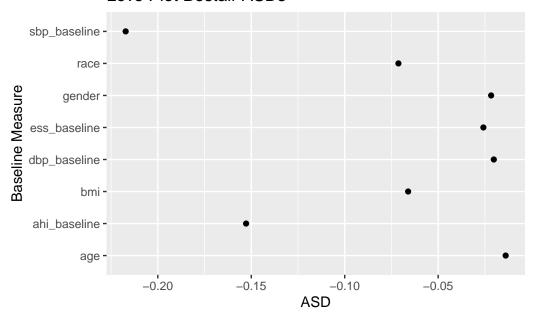
variances[i,j] = sigma
}
</pre>
```

Neyman Estimator of Variance: 230.2083333

Part 3

```
baselines <- c("gender", "age", "bmi",</pre>
                 "race", "sbp_baseline", "dbp_baseline", "ahi_baseline", "ess_baseline")
ASDs = matrix(NA, nrow = 1, ncol = 8)
colnames(ASDs) <- baselines</pre>
Z <- bestair$treatment arm</pre>
for (bl in baselines) {
  X <- pull(bestair[,bl])</pre>
  s1 <- var(X[Z==1])
  s0 \leftarrow var(X[Z==0])
  diff_sum \leftarrow sum(X*Z)/sum(Z)-sum(X*(1-Z))/sum(1-Z)
  asd <- diff_sum/sqrt(s1+s0)</pre>
  ASDs[,bl] <- asd
}
#love plot
asd_dat <- tibble(</pre>
 bls = baselines,
  asd = ASDs[1,]
```

Love Plot Bestair ASDs



```
Z <- bestair$treatment_arm
Y <- bestair$sbp_6mo
estimate_unadj <- mean(Y[Z==1]) - mean(Y[Z==0])
se_unadj <- summary(lm(Y~Z))$coefficients[2,"Std. Error"]
hw_unadj <- sqrt(car::hccm(lm(Y ~ Z), type = "hc2")[2, 2])
unadj_res <- c(estimate_unadj,se_unadj,hw_unadj)

bestair_centered <- bestair|>
mutate(across(gender:ess_baseline, ~ .x-mean(.x)))
ancova1 <- lm(formula = sbp_6mo~.,data = bestair_centered)
estimate_anc1 <- ancova1$coefficients[["treatment_arm"]]</pre>
```

	Unadjusted ATE	ANCOVA1	ANCOVA2
Estimate	-4.907	-2.540	-2.078
s.e.	2.230	1.683	2.092
Huber-White s.e.	2.266	1.696	NA

The Estimates decrease as we use a more flexible model and add interaction terms.

```
bestair_hyperten <- bestair_centered |>
   mutate(resist_hyperten = if_else(sbp_6mo>=130,1,0)) |>
   select(treatment_arm:ess_baseline,resist_hyperten)
```

```
Z <- bestair$treatment_arm
Y <- bestair_hyperten$resist_hyperten
bin_unadj_est <- mean(Y[Z==1]) - mean(Y[Z==0])
bin_ols <- lm(resist_hyperten~., data = bestair_hyperten)
bin_est <- bin_ols$coefficients["treatment_arm"]</pre>
```

```
bin_ols_inter <- lm(resist_hyperten~.^2, data = bestair_hyperten)
bin_est_inter <- bin_ols_inter$coefficients["treatment_arm"]

results <- cbind(bin_unadj_est,bin_est,bin_est_inter)
colnames(results) <- c("Unadjusted ATE","ANCOVA1","ANCOVA2")
rownames(results) <- "Binary Estimates"
results |> round(3) |> kable()
```

	Unadjusted ATE	ANCOVA1	ANCOVA2
Binary Estimates	-0.208	-0.128	-0.107

b)

```
logist_unadj <- glm(resist_hyperten~treatment_arm, family = binomial(link = "logit"), data
logist_anc1 <- glm(resist_hyperten~., family = binomial(link = "logit"), data = bestair_hy
logist_anc2 <- glm(resist_hyperten~.^2, family = binomial(link = "logit"), data = bestair_
unadj_ate <- logist_unadj$coefficients[[2]]
anc1 <- logist_anc1$coefficients[["treatment_arm"]]
anc2 <- logist_anc2$coefficients[["treatment_arm"]]
results <- cbind(unadj_ate,anc1,anc2)
colnames(results) <- c("Unadjusted ATE","ANCOVA1","ANCOVA2")
rownames(results) <- "Logistic Estimates"
results |> round(3) |> kable()
```

	Unadjusted ATE	ANCOVA1	ANCOVA2
Logistic Estimates	-0.985	-0.818	-5.34

The linear model seems to perform better than logistic regression.