# Field Experiments: Design, Analysis and Interpretation Solutions for Chapter 3 Exercises

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## Question 1

Important concepts: [10 points]

a) What is a standard error? What is the difference between a standard error and a standard deviation?

Answer:

The standard error is a measure of the statistical uncertainty surrounding a parameter estimate. The standard error is a measure of dispersion in a sampling distribution; the standard deviation is the measure of dispersion of any distribution but is most often used to describe the dispersion in an observed variable. The standard error is the standard deviation of the sampling distribution, or the set of estimates that could have arisen under all possible random assignments.

b) How is randomization inference used to test the sharp null hypothesis of no effect for any subject? Answer:

The sharp null hypothesis of no effect is a case in which  $Y_i(1) = Y_i(0)$ ; under this assumption, all potential outcomes are observed because treated and untreated potential outcomes are identical. In order to form the sampling distribution under the sharp null hypothesis of no effect, we simulate a random assignment and calculate the test statistic (for example, the difference-inmeans between the assigned treatment and control groups). This simulation is repeated a large number of times in order to form the sampling distribution under the null hypothesis. The p-value of the test statistic that is observed in the actual experiment is calculated by finding its location in the sampling distribution under the null hypothesis. For example, if the observed test statistic is as large or larger than 9,000 of 10,000 simulated experiments, the one-tailed p-value is 0.10.

c) What is a 95% confidence interval?

Answer:

A confidence interval consists of two estimates, a lower number and an upper number, that are intended to bracket the true parameter of interest with a specified probability. An estimated confidence interval is a random variable that varies from one experiment to the next due to random variability in how units are allocated to treatment and control. A 95% interval is designed to bracket the true parameter with a 0.95 probability across hypothetical replications of a given experiment. In other words, across hypothetical replications, 95% of the estimated 95% confidence intervals will bracket the true parameter.

<sup>\*</sup>Solutions prepared by Peter M. Aronow and revised by Alexander Coppock

- d) How does complete random assignment differ from block random assignment and clustered random assignment? Answer:
  - Under complete random assignment, each subject is assigned separately to treatment or control groups such that m of N subjects end up in the treatment condition. Under block random assignment, complete random assignment occurs within each block or subgroup. Under clustered assignment, groups of subjects are assigned jointly to treatment or control; the assignment procedure requires that if one member of the group is assigned to the treatment group, all others in the same group are also assigned to treatment.
- e) Experiments that assign the same number of subjects to the treatment group and control group are said to have a "balanced design." What are some desirable statistical properties of balanced designs?

Answer:

One desirable property of a balanced design is that under certain conditions, it generates less sampling variability than unbalanced designs; this property of balanced designs holds when the variance of  $Y_i(0)$  is approximately the same as the variance of  $Y_i(1)$ . Another attractive property is that estimated confidence intervals are, on average, conservative (they tend to overestimate the true amount of sampling variability) under balanced designs. (A final attractive property, which comes up in Chapter 4, is that regression is less prone to bias under balanced designs.)

## Question 2

Rewrite equation (3.4) substituting for  $Y_i(1)$  using the equation  $Y_i(1) = Y_i(0) + \tau_i$ . Assume that N = 2m, and interpret the implications of the resulting formula for experimental design. [5 points] Answer:

Substituting N = 2m and  $Y_i(1) = Y_i(0) + \tau_i$  gives

$$\begin{split} SE(\widehat{ATE}) &= \sqrt{\frac{1}{(N-1)} \left\{ \frac{mVar(Y_i(0))}{2m-m} \frac{mVar(Y_i(0)+\tau_i)}{2m-m} \right\} + 2cov(Y_i(0),Y_i(0)+\tau_i)} \\ &= \sqrt{\frac{1}{(N-1)} \left\{ Var(Y_i(0)) + Var(Y_i(0)+\tau_i) + 2Var(Y_i(0)) + 2cov(Y_i(0),\tau_i) \right\}} \\ &= \sqrt{\frac{1}{(N-1)} \left\{ 3Var(Y_i(0)) + [Var(Y_i(0)) + Var(\tau_i) + 2cov(Y_i(0),\tau_i)] + 2cov(Y_i(0),\tau_i) \right\}} \\ &= \sqrt{\frac{1}{(2m-1)} \left\{ 4Var(Y_i(0)) + Var(\tau_i) + 4cov(Y_i(0),\tau_i) \right\}} \end{split}$$

All else being equal, the true standard error is smaller when the variance of the treatment effect is smaller, the variance of  $Y_i(0)$  is smaller, and the covariance of the treatment effect and  $Y_i(0)$  is smaller.

## Question 3

Using the equation  $Y_i(1) = Y_i(0) + \tau_i$ , show that when we assume that treatment effects are the same for all subjects,  $Var(Y_i(0)) = Var(Y_i(1))$  and the correlation between  $Y_i(0)$  and  $Y_i(1)$  is 1.0.[5 points]

Under constant treatment effects,  $Var(Y_i(1) = Var(Y_i(0) + \tau) = Var(Y_i(0))$ , and the correlation between  $Y_i(1)$  and  $Y_i(0)$  is:

$$cor(Y_{i}(1), Y_{i}(0)) = \frac{Cov(Y_{i}(1), Y_{i}(0))}{\sqrt{Var(Y_{i}(1)) * Var(Y_{i}(0))}}$$

$$= \frac{Cov(Y_{i}(0) + \tau, Y_{i}(0))}{\sqrt{Var(Y_{i}(0)) * Var(Y_{i}(0))}}$$

$$= \frac{Var(Y_{i}(0))}{Var(Y_{i}(0))}$$

$$= 1$$

## Question 4

Consider the schedule of outcomes in the table below. If treatment A is administered, the potential outcome is  $Y_i(A)$ , and if treatment B is administered, the potential outcome is  $Y_i(B)$ . If no treatment is administered, the potential outcome is  $Y_i(0)$ . The treatment effects are defined as  $Y_i(A) - Y_i(0)$  or  $Y_i(B) - Y_i(0)$ . [5 points]

Table 1: Question 4 Table

Subject		A A	<b>%</b>
Miriam	,C	2	3
Benjamin	2	3	3
Helen	)3	4	3
Eya	4	5	3
Billie	5	6	3

Suppose a researcher plans to assign two observations to the control group and the remaining three observations to just one of the two treatment conditions. The researcher is unsure which treatment to use.

a) Applying equation (3.4), determine which treatment, A or B, will generate a sampling distribution with a smaller standard error.

Answer:

First, notice that  $Y_i(A) = Y_i(0) + 1$ . Then using the results developed in the previous exercise:

$$SE(\widehat{ATE_A}) = \sqrt{\frac{1}{5-1} \left\{ \frac{3*2}{2} + \frac{2*2}{3} + 2*2 \right\}}$$

$$= 1.44$$

$$SE(\widehat{ATE_B}) = \sqrt{\frac{1}{5-1} \left\{ \frac{3*2}{2} + \frac{2*0}{3} + 2*0 \right\}}$$

$$= 0.86$$

The standard error for the B vs. control comparison is smaller than the standard error for the A vs. control comparison. Thus, administering treatment B gives rise to a narrower sampling distribution.

b) What does the result in part (a) imply about the feasibility of studying interventions that attempt to close an existing "achievement gap"?

#### Answer:

When treatment B is administered, the achievement gap between the best and worst student narrows, leaving no variance in  $Y_i(B)$ . Two of the three terms in equation (3.4) therefore become zero, and the resulting standard error is much lower than it would be under treatment A, which has a constant effect across all subjects. The basic principle here is that it helps to study treatments that reduce the covariance between untreated and treated potential outcomes.

#### Question 5

Using Table 2.1, imagine that your experiment allocates one village to treatment. [10 points]

a) Calculate the estimated difference-in-means for all seven possible randomizations. Answer:

There are 7 subjects, 1 of which is assigned to treatment, and thus the number of randomizations is  $\frac{7!}{1!(7-1)!} = 7$ . Now let's define  $\widehat{ATE_i}$  as the difference in means constructed when assuming village i is assigned to treatment.

 $ATE_i$ Village  $Y_i(0)$  $\tau_i$ 1 5 10 2 0 15 3 20 10 20 15 -5 4 5 10 20 10 6 15 15 0 15 7 15 39 5 Mean 15 20  $4(15-20)^2+2(30-20)^2$ SD

=6.755

Table 2: Question 5 Table

b) Show that the average of these estimates is the true ATE.

100

Answer:

The table shows that the average across all randomizations is 5, which is the true ATE.

c) Show that the standard deviation of the seven estimates is identical to the standard error implied by equation (3.4).

Beginning with Equation 3.4:

$$\begin{split} SE(\widehat{ATE}) &= \sqrt{\frac{1}{(N-1)} \left\{ \frac{mVar(Y_i(0))}{N-m} + \frac{(N-m)*Var(Y_i(1))}{m} + 2cov(Y_i(0), Y_i(1)) \right\}} \\ &= \sqrt{\frac{1}{6} \left\{ \frac{Var(Y_i(0))}{6} + 6Var(Y_i(1)) + 2cov(Y_i(0), Y_i(1)) \right\}} \\ &cov(Y_i(0), Y_i(1)) = \frac{(10-15)(15-20) + (20-15)(30-20) + (20-15)(15-20)}{7} = \frac{50}{7} \\ &= \sqrt{\frac{1}{6} \left\{ \frac{100}{7} + 6\frac{300}{7} + 2\frac{50}{7} \right\}} \\ &= 6.755 \end{split}$$

This is identical to the standard deviation calculated in the table above.

d) Referring to equation (3.4), explain why this experimental design has more sampling variability than the design in which two villages out of seven are assigned to treatment.

Answer:

The covariance term is unaffected, but the first two variance terms are multiplied by different numbers. The first term is multiplied by 1/6 in this example as opposed to 2/5 in the 2-of-7 example. The second term is multiplied by 6/1 in this example as opposed to 5/2 in the 2-of-7 example. Because the second variance term is larger than the first, allocating more sample to the treatment group reduces sampling variance.

$$SE(\widehat{ATE}) = \sqrt{\frac{1}{(N-1)} \left\{ \frac{mVar(Y_i(0))}{N-m} + \frac{(N-m)*Var(Y_i(1))}{m} + 2cov(Y_i(0), Y_i(1)) \right\}}$$

$$= \sqrt{\frac{1}{6} \left\{ \frac{1}{6} \frac{100}{7} + \frac{6}{1} \frac{300}{7} + 2\frac{50}{7} \right\}} = 6.755, \text{ if } m = 1$$

$$= \sqrt{\frac{1}{6} \left\{ \frac{2}{5} \frac{100}{7} + \frac{5}{2} \frac{300}{7} + 2\frac{50}{7} \right\}} = 4.603, \text{ if } m = 2$$

e) Explain why, in this example, a design in which one of seven observations is assigned to treatment has more<sup>1</sup> sampling variability than a design in which six villages out of seven are assigned to treatment.

<sup>&</sup>lt;sup>1</sup>Text mistakenly printed "less"

$$SE(\widehat{ATE}) = \sqrt{\frac{1}{(N-1)} \left\{ \frac{mVar(Y_i(0))}{N-m} + \frac{(N-m)*Var(Y_i(1))}{m} + 2cov(Y_i(0), Y_i(1)) \right\}}$$

$$= \sqrt{\frac{1}{6} \left\{ \frac{1}{6} \frac{100}{7} + \frac{6}{6} \frac{300}{7} + 2\frac{50}{7} \right\}} = 6.755, \text{ if } m = 1$$

$$= \sqrt{\frac{1}{6} \left\{ \frac{6}{1} \frac{100}{7} + \frac{1}{6} \frac{300}{7} + 2\frac{50}{7} \right\}} = 4.23, \text{ if } m = 6$$

By the same logic as above – allocating more units to the condition in which potential outcomes are more variable can reduce sampling variability.

#### Question 6

The Clingingsmith, Khwaja, and Kremer study discussed in section 3.5 may be used to test the sharp null hypothesis that winning the visa lottery for the pilgrimage to Mecca had no effect on the views of Pakistani Muslims toward people from other countries. Assume that the visa authorities conducted a complete random assignment; generate 10,000 shaulated random assignments under the sharp null hypothesis. How many of the simulated random assignments generate an estimated ATE that is at least as large as the actual estimate of the ATE? What is the implied one-tailed p-value? How many of the simulated random assignments generate an estimated ATE that is at least as large in absolute value as the actual estimate of the ATE? What is the implied two-tailed p-value? [10 points]

```
. // download data from: http://hdl.handle.net/10079/6hdr852
. // copy and paste the url to your web browser
. import delim "Clingingsmith_et_al_QJE_2009dta.csv",clear
(8 vars, 958 obs)
. set seed 1234567
 rename success D
  rename views Y
//findit tsrtest
. //package name: st0158.pkg install
. cap program drop ate
. program define ate, rclass
      args Y D
       sum `Y' if `D'==1, meanonly
       local Y_treat=r(mean)
       sum `Y' if `D'==0, meanonly
       local Y_con=r(mean)
        return scalar ate_avg = `Y_treat'-`Y_con'
  7. end
. // ssc install tsrtest
. tsrtest D r(ate_avg) using 3_6_resam.dta, overwrite: ate Y D
Two-sample randomization test for theta=r(ate_avg) of ate Y D by D
```

```
Combinations: 8.4503047638e+285 = (958 \text{ choose } 448)
Assuming null=0
Observed theta: .4748
Minimum time needed for exact test (h:m:s): 4.2e+278:00:00
Reverting to Monte Carlo simulation.
Mode: simulation (10000 repetitions)
progress: |.....|
 p=0.00190 [one-tailed test of Ho: theta(D==0) <= theta(D==1)]
p=0.99830 [one-tailed test of Ho:
                                   theta(D==0)>=theta(D==1)]
p=0.00360 [two-tailed test of Ho:
                                   theta(D==0) == theta(D==1)]
Saving log file to 3_6_resam.dta...done.
. preserve
. use "3_6_resam.dta", clear
. global ate = theta[1]
. di $ate
.4748337
. drop if _n==1
(1 observation deleted)
. count if theta >= $ate
 19
. scalar p_onesided = r(N)/N
. count if abs(theta) >= $ate
  36
. scalar p_twosided = r(N)/N
. di "p.value.onesided = "p_onesided
p.value.onesided = .0019
. di "p.value.twosided = "p_twosided
p.value.twosided = .0036
. restore
```

The estimated ATE is 0.4748337. The number of simulated ATEs under the sharp null hypothesis of no effect that were as large was 15, corresponding to a p-value of 0.0017. The number of simulated ATEs under the sharp null hypothesis of no effect that were as large in absolute value was was 32, corresponding to a p-value of 0.0034.

## Question 7

A diet and exercise program advertises that it causes everyone who is currently dieting to lose at least seven pounds more than they otherwise would have during the first two weeks. Use randomization inference (the procedure described in section 3.4) to test the hypothesis that  $\tau_i = 7$  for all i. The treatment group's weight losses after two weeks are (2, 11, 14, 0, 3) and the control

group's weight losses are (1, 0, 0, 4, 3). In order to test the hypothesis  $\tau_i = 7$  for all i using the randomization inference methods discussed in this chapter, subtract 7 from each outcome in the treatment group so that the exercise turns into the more familiar test of the sharp null hypothesis that  $\tau_i = 0$  for all i. When describing your results, remember to state the null hypothesis clearly, and explain why you chose to use a one-sided or two-sided test. [10 points]

Table 3: Question 7 Table

Subject	$Y_i(0)$	$Y_i(1)$	$Y_i(1) - 7$
1	?	2	-5
2	?	11	4
3	?	14	7
4	?	0	-7
5	?	3	-4
6	1	?	?
7	0	?	?
8	0	?	?
9	4	?	? * * *
10	3	?	

```
. clear
. set seed 1234567
. set obs 10
number of observations (_N) was 0, now 10
. input D Y
                       Y
            D
 1. 0 1
 2. 0 0
 3. 0 0
 4.04
 5. 0 3
 6. 1 2
 7. 1 11
 8. 1 14
 9.10
 10.13
 gen Y_star = Y+D*(-7)
. cap program drop ate
. program define ate, rclass
           args Y D
 1.
 2.
            sum `Y' if `D'==1, meanonly
 3.
            local Y_treat=r(mean)
           sum `Y' if `D'==0, meanonly
           local Y_con=r(mean)
            return scalar ate_avg = `Y_treat'-`Y_con'
 6.
 7. end
 // findit tsrtest (to install the package)
. tsrtest D r(ate_avg): ate Y_star D
```

```
Two-sample randomization test for theta=r(ate_avg) of ate Y_star D by D
Combinations:
                252 = (10 \text{ choose } 5)
Assuming null=0
Observed theta: -2.6
Minimum time needed for exact test (h:m:s): 0:00:00
Mode: exact
 p=0.83730 [one-tailed test of Ho: theta(D==0)<=theta(D==1)]
p=0.20635 [one-tailed test of Ho: theta(D==0)>=theta(D==1)]
 p=0.41270 [two-tailed test of Ho: theta(D==0)==theta(D==1)]
. // ate
. di r(obsvStat)
-2.6
. // p.value.onesided
. di r(lowertail)
.20634921
```

There are 10 subjects, 5 of which are assigned to treatment, and thus the number of randomizations is  $\frac{10!}{5!5!} = 252$ . The null hypothesis is that the true ATE is a 7 pound loss; the alternative hypothesis is that the weight loss ATE is less than 7 pounds. A one-sided hypothesis test is used because we only want to reject the weight loss program's claims if the observed weight loss is less than what they claimed; if they understated the degree of weight loss, their program would be even more effective than claimed, and one would hardly fault them for that. Using the code for randomization inference posted on the website, we find that the observed difference in weight loss between the treatment and control groups (6 - 1.6 = 4.4) is smaller than 79% of all simulated experiments under the null hypothesis of a 7-pound effect for everyone. Thus, the p-value is 0.21, meaning we cannot reject the null hypothesis of a 7-pound effect at the conventional 0.05 significance threshold.

## 1 Question 8

Natural experiments sometimes involve what is, in effect, block random assignment. For example, Titiunik studies the effect of lotteries that determine whether state senators in Texas and Arkansas serve two-year or four-year terms in the aftermath of decennial redistricting.<sup>2</sup> These lotteries are conducted within each state, and so there are effectively two distinct experiments on the effects of term length. An interesting outcome variable is the number of bills (legislative proposals) that each senator introduces during a legislative session. The table below lists the number of bills introduced by senators in each state during 2003. [10 points]

a) For each state, estimate of the effect of having a two-year term on the number of bills introduced.

```
// download data from : http://hdl.handle.net/10079/s1rn910
. // copy and paste the url to your web browser
. use "Titiunik_WorkingPaper_2010.csv.dta",clear
.
. set seed 1234567
```

 $<sup>^2</sup>$ Titiunik 2010.

Table 4: Question 8 Table

Texas		Arkansas		
Term Length: 0 = four-year term; 1 = two-year term	# of bills introduced	Term Length: 0 = four-year term; 1 = two-year term	# of bills introduced	
0	18	0	11	
0	29	0	15	
0	41	0	17	
0	53	0	23	
0	60	0	24	
0	67	0	25	
0	75	0	26	
0	79	0	28	
0	79		31	
0	88	0.	33	
0	93		34	
0	101		35	
0	103	0	35	
0	106	0	36	
0	107	0	38	
0	131	0	52	
1	29	0	59	
1	37	1	9	
1	42	1	10	
1	45	1	14	
1	45	1	15	
1	54	1	15	
1	54	1	17	
1	58	1	18	
1	61	1	19	
1	64	1	19	
1	69	1	20	
1	73	1	21	
1	75	1	23	
1	92	1	23	
1	104	1	24	
		1	28	
		1	30	
		1	32	
		1	34	

```
rename term2year D
rename bills_introduced Y
rename texas0_arkansas1 block
qui tabstat Y if block ==0, by(D) stat(mean) save
scalar ate_texas = el(r(Stat2),1,1) - el(r(Stat1),1,1)
qui tabstat Y if block ==1, by(D) stat(mean) save
scalar ate_ark = el(r(Stat2),1,1) - el(r(Stat1),1,1)

di "ate_texas="%18.5f ate_texas
ate_texas= -16.74167

di "ate_arkansas="%18.5f ate_ark
ate_arkansas= -10.09477
```

The estimated ATE in Texas is -16.742. In Arkansas, the estimated ATE is -10.095.

b) For each state, estimate the standard error of the estimated ATE.

The estimated se in Texas is 9.346. In Arkansas, the estimated se is 3.396.

c) Use equation (3.10) to estimate the overall ATE for both states combined.

The overall ATE, -13.217 is the weighted average of the two separate ATEs, where the weights are the shares of overall N in each state.

d) Explain why, in this study, simply pooling the data for the two states and comparing the average number of bills introduced by two-year senators to the average number of bills introduced by four-year senators leads to biased estimates of the overall ATE.

#### Answer:

The two states differ in terms of the probability that a given legislator will be assigned to the treatment. Therefore, we cannot pool the data without introducing a correlation between treatment assignment and the potential outcomes associated with the two states. In this study, the experiments take place within each state, and the analyst should pool the state-level results in order to obtain an overall result.

e) Insert the estimated standard errors into equation (3.12) to estimate the standard error for the overall ATE.

The overall standard error is (4.745).

f) Use randomization inference to test the sharp null hypothesis that the treatment effect is zero for senators in both states.

```
// calculate probs under block assignment
. bysort block: egen probs=mean(D).
. cap program drop ate_block
. program define ate_block, rclass
 1. args Y D probs
 2. tempvar ipw
 3. gen `ipw' = .
 4. // calculate inverse probability weight under block assignment
. replace `ipw' = `D'/`probs' + (1-`D')/(1-`probs')
 5. qui reg `Y' `D' [iw=`ipw']
 6. return scalar ate=_b[`D']
. // ssc install ritest (to install ritest package)
. ritest D r(ate), strata(block) reps(10000) nodots: ///
> ate_block Y D probs
(66 missing values generated)
(66 real changes made)
     command: ate_block Y D probs
       _pm_1: r(ate)
 res. var(s): D
  Resampling: Permuting D
Clust. var(s): __000000
    Clusters: 66
Strata var(s): block
      Strata: 2
T | T(obs) c n p=c/n SE(p) [95% Conf. Interval]
```

Here, we use a two-tailed test because it is not clear theoretically whether longer or shorter terms should make legislators more responsive. Comparing the observed difference-in-means to the distribution of 10,000 simulated randomizations under the sharp null hypothesis reveals a two-tailed p-value of 0.0071, leading us to reject the null hypothesis.

## Question 9

Camerer reports the results of an experiment in which he tests whether large, early bets placed at horse tracks affect the betting behavior of other bettors. Selecting pairs of long-shot horses running in the same race whose betting odds were approximately the same when betting opened, he placed two \$500 bets on one of the two horses approximately 15 minutes before the start of the race. Because odds are determined based on the proportion of total bets placed on each horse, this intervention causes the betting odds for the treatment horse to decline and the betting odds of the control horse to rise. Because Cameror's bets were placed early, when the total betting pool was small, his bets caused marked changes in the odds presented to other bettors. (A few minutes before each race started, Camerer canceled his bets.) While the experimental bets were still "live," were other bettors attracted to the treatment horse (because other bettors seemed to believe in the horse) or repelled by it (because the diminished odds meant a lower return for each wager)? Seventeen pairs of horses in this study are listed below. The outcome measure is the number of dollars that were placed on each horse (not counting Camerer's own wagers on the treatment horses) during the test period, which begins 16 minutes before each race (roughly 2 minutes before Camerer began placing his bets) and ends 5 minutes before each race (roughly 2 minutes before Camerer withdrew his bets). [10 points]

<sup>&</sup>lt;sup>3</sup>Camerer 1998. This example draws on the second of Camerer's studies and restricts the sample to cases in which a treatment horse is compared to a single control horse.

Table 5: Question 9 Table

Pair 1 533 1503 Pair 2 376 1186 Pair 3 576 1366 Pair 4 1135 1666 Pair 5 158 367 Pair 6 282 542 Pair 7 909 1597 Pair 8 566 933 Pair 9 0 555 Pair 10 330 786 Pair 11 74 959 Pair 12 138 319 Pair 13 347 812		Pair	Contr	ol Horse in P	air		
			Change	Total bets $T-16 \text{ min}$	Total bets $T-5 \min$	Change	Difference in changes
Pair 1	533	1503	970	587	2617	2030	-1060
Pair 2	376	1186	810	345	1106	761	49
Pair 3	576	1366	790	653	2413	1760	-970
Pair 4	1135	1666	531	1296	2260	964	-433
Pair 5	158	367	209	201	574	373	-164
Pair 6	282	542	260	269	489	220	40
Pair 7	909	1597	688	775	1825	1050	-362
Pair 8	566	933	367	629	1178	549	-182
Pair 9	0	555	555	0	355	355	200
Pair 10	330	786	456	233	842	609	-153
Pair 11	74	959	885	130	256	126	759
Pair 12	138	319	181	179	356	177	4
Pair 13	347	812	465	382	604	222	243
Pair 14	169	329	160	165 A	355	190	-30
Pair 15	41	297	256	33)	75	42	214
Pair 16	37	71	34	33	121	88	-54
Pair 17	261	485	224	282	480	198	26

a) One interesting feature of this study is that each pair of horses ran in the same race. Does this design feature violate the non-interference assumption, or can potential outcomes be defined so that the non-interference assumption is satisfied?

Answer:

This design feature violates non-interference if the estimand is defined as the difference between the following two potential outcomes: total bets on a given horse when experimental bets are placed on that horse versus no experimental bets on any horse in the race. One could avoid violating non-interference by redefining the estimand as the difference between the following two potential outcomes: total bets on a horse when experimental bets are placed on that horse versus experimental bets are placed on a competing horse in the same race.

b) A researcher interested in conducting a randomization check might assess whether, as expected, treatment and control horses attract similarly sized bets prior to the experimental intervention. Use randomization inference to test the sharp null hypothesis that the bets had no effect prior to being placed.

```
// download data from : http://hdl.handle.net/10079/1g1jx43
. // copy and paste the url to your web browser
.
. use "Camerer_JPEsubset_1998.dta.dta", clear
.
. set seed 1234567
. rename treatment D
. rename pair block
```

```
rename preexperimentbets covs
        // calculate probs under block assignment
        bysort block: egen probs=mean(D)
        // permuation to calculate F stat and one-side P value
        ritest D e(F), strata(block) reps(10000) right nodots: ///
        regress D covs
    Source | SS
                          df MS
                                          Number of obs
                                                               34
_____
                                          F(1, 32) =
                                                             0.02
  0.8914
                                                       =
                           32 .265467988 R-squared
                                                           0.0006
                                          Adj R-squared =
                                                          -0.0306
    Total | 8.5 33 .257575758
                                          Root MSE
       D | Coef. Std. Err. t P>|t| [95% Conf. Interval]
  ______

    covs | -.0000386
    .0002809
    -0.14
    0.891
    -.0006109
    .0005336

    _cons | .5137818
    .1335793
    3.85
    0.001
    .2416896
    .785874

     command: regress D covs
      _pm_1: e(F)
 res. var(s): D
  Resampling: Permuting D
Clust. var(s): __000000
   Clusters: 34
Strata var(s): block
     Strata: 17
     | T(obs) c n p=c/n SE(p) [95% Conf. Interval]
    _pm_1 | .0189265 3736 10000 0.3736 0.0048 .3641064 .3831672
Note: Confidence interval is with respect to p=c/n.
Note: c = \#\{T >= T(obs)\}
        // p.value
        di el(r(p),1,1)
.3736
```

We conducted 10,000 random assignments, and for each we calculated the F-statistic of a regression of treatment assignment on pre-experimental bets (controlling for blocks). The observed F-statistic for the actual experiment is larger than 3696 of the simulated experiments, implying a p-value of 0.37.

c) Calculate the average increase in bets during the experimental period for treatment horses and control horses. Compare treatment and control means, and interpret the estimated ATE.

```
. rename experimentbets change
```

The average treatment group change was \$461.24, as opposed to an average change of \$571.41 in the control group. Therefore, the estimated ATE is \$-110.18.

d) Show that the estimated ATE is the same when you subtract the control group outcome from the treatment group outcome for each pair and calculate the average difference for the 17 pairs. Answer:

The average difference between treatment and control outcomes for each pair is also 110.18.

e) Use randomization inference to test the sharp null hypothesis of no treatment effect for any subject. When setting up the test, remember to construct the simulation to account for the fact that random assignment takes place within each pair. Interpret the results of your hypothesis test and explain why a two-tailed test is appropriate in this application.

```
cap program drop ate_block

program define ate_block, rclass
1. args Y D probs
2. tempvar ipw
3. gen `ipw' = .
```

```
4. // calculate inverse probability weight under block assignment
. replace `ipw' = `D'/`probs' + (1-`D')/(1-`probs')
 5. qui reg `Y' `D' [iw=`ipw']
 6. return scalar ate=_b[`D']
. ritest D r(ate), strata(block) reps(10000) nodots: ///
> ate_block change D probs
(34 missing values generated)
(34 real changes made)
    command: ate_block change D probs
     _pm_1: r(ate)
 res. var(s): D
  Resampling: Permuting D
Clust. var(s): __000000
   Clusters: 34
Strata var(s): block
     Strata: 17
    T(obs) c n p=c/n SE(p) [95% Conf. Interval]
______
Note: Confidence interval is with respect to p=c/n.
Note: c = \#\{|T| >= |T(obs)|\}.
. // ate
. di el(r(b), 1, 1)
-110.17647
. // p.value.twosided
. di el(r(p), 1, 1)
.317
```

A two-tailed test generates a p-value of 0.3092, indicating that one cannot reject the sharp null of no effect for any unit. A two-tailed test is appropriate because some theories predict a positive effect while others predict a negative effect: "were other bettors attracted to the treatment horse (because other bettors seemed to believe in the horse) or repelled by it (because the diminished odds meant a lower return for each wager)?" The appropriate null hypothesis in this case is no effect, which would be rejected if we observed either strongly positive or strongly negative differences between treatment and control horses.

## Question 10

Suppose that 800 individual students were randomly assigned to classrooms of 25 students apiece, and these classrooms were then randomly assigned as clusters to treatment and control. Assume the non-interference assumption holds. Use equations (3.4) and (3.22) to explain why this clustered design has the same standard error as complete random assignment of individual students to treatment and control. [10 points] Answer:

The equation for the standard error under individual assignment:

$$SE(\widehat{ATE}) = \sqrt{\frac{1}{(N-1)} \left\{ \frac{mVar(Y_i(0))}{N-m} + \frac{mVar(Y_i(1))}{N-m} + 2cov(Y_i(0), Y_i(1)) \right\}}$$

The equation for the standard error under clustered assignment with equal-size clusters:

$$SE(\widehat{ATE}) = \sqrt{\frac{1}{(k-1)} \left\{ \frac{mVar(\bar{Y}_j(0))}{N-m} + \frac{mVar(\bar{Y}_j(1))}{N-m} + 2cov(\bar{Y}_j(0), \bar{Y}_j(1)) \right\}}$$

When the clusters are formed randomly (i.e., individuals are randomly allocated to clusters prior to assignment), the two formulas give approximately the same answer. In order to see the correspondence, notice that the variance of the average treated outcome from random draw of 25 students is  $Var(\bar{Y}_j(0)) = \frac{Var(Y_i(0))}{25}$ , and similarly,  $Var(\bar{Y}_j(1)) = \frac{Var(Y_i(1))}{25}$ , and  $cov(\bar{Y}_j(0), \bar{Y}_j(1)) = \frac{cov(Y_i(0), Y_i(1))}{25}$ . Thus, the quantity inside the braces in both equations differs by a factor of 25, which is approximately  $\frac{N-1}{k-1}$ .

## Question 11

Use the data in Table 3.3 to simulate cluster randomized assignment. [10 points]

a) Suppose that clusters are formed by grouping observations  $\{1,2\}, \{3,4\}, \{5,6\} \dots \{13,14\}$ . Use equation (3.22) to calculate the standard error assuming half of the clusters are randomly assigned to the treatment.

```
. clear
 set seed 1234567
. set obs 14
number of observations (_N) was 0, now 14
. input YO
            Υ0
 1. 0
 7.6
 8.9
 9.14
 10.15
 11. 16
 12. 16
 13. 17
 14. 18
 end
. input Y1
```

```
Y 1
                 1. 0
                 2. 0
                 3. 1
                 4. 2
                5. 0
                6.0
              7. 2
           8.3
          9. 12
       10. 9
       11. 8
      12. 15
      13. 5
     14. 17
  . end
  . gen int cluster = (n+1)/2
  . //ssc install tabstatmat (install the package)
   . // save tabstat summary result to matrix % \left( 1\right) =\left( 1\right) \left( 
  . tabstat YO, by(cluster) stat(mean) save
  Summary for variables: YO
          by categories of: cluster
      cluster | mean
                                          1 | .5
2 | 3
3 | 5
                                                                                                                         7.5
                                                    4 |
                                                                                                                  14.5
                                                    5 I
                                                      6 I
                                                                                                                                       16
                                                      7 | 17.5
   -----
                 Total | 9.142857
   ______
  . tabstatmat Ybar0, nototal
 Ybar0[7,1]
                                                                  ΥO
                                                          . 5
3
5
 1:mean
 2:mean
3:mean
                                                           7.5
 4:mean
 5:mean 14.5
 6:mean 16
7:mean 17.5
  . mat colnames Ybar0=Ybar0
 . tabstat Y1, by(cluster) stat(mean) save
```

```
Summary for variables: Y1
   by categories of: cluster
cluster | mean
     1 | 0
     3 |
               0
             2.5
      4 |
      5 | 10.5
6 | 11.5
7 | 11
_____
  Total | 5.285714
______
. tabstatmat Ybar1, nototal
Ybar1[7,1]
       Y1
        0
1:mean
2:mean 1.5
3:mean
        0
4:mean 2.5
5:mean 10.5
6:mean 11.5
7:mean 11
. mat colnames Ybar1=Ybar1
. // function to calculate population variance
. cap program drop var_pop
. program define var_pop, rclass
 1. args varname
          tempvar x_dev
 2.
          qui sum `varname'
 3.
 4.
           local avg = r(mean)
 5.
          local length = r(N)
          gen `x_dev' = (`varname'-`avg')^2/`length'
          qui tabstat `x_dev', stat(sum) save
 7.
 8.
          return scalar variance_pop = el(r(StatTotal),1,1)
 9. end
. // function to calculate population covariance
. cap program drop cor_pop
. program define cor_pop, rclass
 1. args x y
 2.
           tempvar xy_dev
 3.
          qui sum `x'
 4.
           local avg_x = r(mean)
 5.
          local length = r(N)
 6.
       qui sum `y'
7. local avg_y = r(mean)
```

```
8.
          gen `xy_dev' = (`x'-`avg_x')*(`y'-`avg_y')
 9.
            qui tabstat `xy_dev', stat(sum) save
 10.
             return scalar cor_pop = el(r(StatTotal),1,1)/`length'
 11. end
. preserve
. clear
. set obs 7
number of observations (_N) was 0, now 7
. svmat Ybar0, names(col)
number of observations will be reset to 7
Press any key to continue, or Break to abort
number of observations (_N) was 0, now 7
. svmat Ybar1, names(col)
. // var_Ybar0
. var_pop Ybar0
. scalar var_Ybar0=r(variance_pop)
. // var_Ybar1
. var_pop Ybar1
. scalar var_Ybar1=r(variance_pop)
. // cov_Ybar0
. cor_pop Ybar0 Ybar1
. scalar cov_Ybar0=r(cor_pop)
. scalar se_ate = sqrt((1/6)*((4/3)*var_Ybar0+(3/4)*var_Ybar1+2*cov_Ybar0))
. di %8.6f se_ate
4.706192
```

Assuming that 4 out of 7 clusters are assigned to treatment, the standard error of the ATE will be 4.71.

b) Suppose that clusters are instead formed by grouping observations  $\{1, 14\}, \{2, 13\}, \{3, 12\} \dots \{7, 8\}$ . Use equation (3.22) to calculate the standard error assuming half of the clusters are randomly assigned to the treatment.

```
replace cluster = _n
(7 real changes made)
. replace cluster = 15-cluster if (cluster>7)
(7 real changes made)
.
```

```
. clear matrix
. // Ybar0
. tabstat YO, by(cluster) stat(mean) save
Summary for variables: YO
 by categories of: cluster
cluster | mean
  1 | 9
2 | 9
3 | 9
            9
10
9.5
     4 |
     5 |
     6 l
              10
     7 |
             7.5
 Total | 9.142857
_____
. tabstatmat YbarO, nototal
Ybar0[7,1]
     YO
1:mean 9
2:mean 9
3:mean 9
4:mean 10
5:mean 9.5
6:mean 10
7:mean 7.5
. mat colnames Ybar0=Ybar0
. // Ybar1
. tabstat Y1, by(cluster) stat(mean) save
Summary for variables: Y1
 by categories of: cluster
cluster | mean
   1 | 8.5
     2 |
             2.5
     3 |
             8
5
     4 |
     5 I
             4.5
     6 |
     7 | 2.5
  Total | 5.285714
. tabstatmat Ybar1, nototal
Ybar1[7,1]
```

```
Y 1
1:mean 8.5
2:mean 2.5
3:mean 8
4:mean 5
5:mean 4.5
6:mean 6
7:mean 2.5
. mat colnames Ybar1=Ybar1
. preserve
. clear
. set obs 7
number of observations (_N) was 0, now 7
. svmat Ybar0, names(col)
number of observations will be reset to 7
Press any key to continue, or Break to abort
number of observations (_{\rm N}) was 0, now 7
. svmat Ybar1, names(col)
. // var_Ybar0 <- var.pop(Ybar0)
. var_pop Ybar0
. scalar var_Ybar0=r(variance_pop)
. // var_Ybar1 <- var.pop(Ybar1)
. var_pop Ybar1
. scalar var_Ybar1=r(variance_pop)
. // cov_Ybar0 <- cov.pop(Ybar0,Ybar1)
. cor_pop Ybar0 Ybar1
. scalar cov_Ybar0=r(cor_pop)
. // se_ate
. scalar se_ate = sqrt((1/6)*((4/3)*var_Ybar0+(3/4)*var_Ybar1+2*cov_Ybar0))
. di %8.7f se_ate
0.9766259
. restore
```

Assuming that 4 out of 7 clusters are assigned to treatment, the standard error of the ATE will be 0.98.

c) Why do the two methods of forming clusters lead to different standard errors? What are the implications for the design of cluster randomized experiments?

Answer:

The first method clusters the most similar villages together, and the second method clusters the most dissimilar villages together. As a result, the variances of the average within-cluster potential outcomes are much larger in the first method and smaller in the second. As a result, the second method produces a much narrower standard error of the ATE estimate. The implication for clustered design is that the more similar the observations with a cluster, the less precise the estimates we can produce. When possible, cluster heterogeneous observations together.

#### Question 12

Below is a schedule of potential outcomes for six classrooms, which are located in three schools. Using a cluster randomized design, researchers will assign one of the three schools (and all the classrooms it contains) to the treatment group. [5 points]

School	Classroom	$Y_i(0)$	$Y_i(1)$	^
A	A-1	0	0	~
В	B-1	0	1	
В	B-2	0	Oi	
$\mathbf{C}$	C-1	0	2	
$\mathbf{C}$	C-2	0	2	
$\mathbf{C}$	C-3	0	2	

Table 6: Question 12 Table

a) What is the average treatment effect among the six classrooms?

$$\frac{2+2+2+1+1+0}{6} = 1.333$$

b) There are three possible randomizations. Is the difference-in-means estimator unbiased? Answer:

The estimated ATE is 0 if school A is assigned to treatment, 1 if school B is assigned to treatment, and 2 if school C is assigned to treatment. So if we take the average of three estimates the ATE is  $\frac{0+1+2}{3}=1\neq 1.33$  and is therefore biased. When potential outcomes are related to cluster size, cluster randomization is prone to bias in small samples, as in this case. This condition holds in this case: the biggest cluster, Cluster C, has larger than average Y(1) values.

c) In general, cluster random assignment generates biased results when (i) clusters vary in size, (ii) potential outcomes vary by cluster, and (iii) the number of clusters is too small to ensure that m of N units are placed into the treatment condition in each randomization. Show what happens in this example when School A and School B are combined for purposes of random assignment, so that there is a 0.5 probability that either School C is placed in treatment or Schools A and B are placed in treatment. Does this design yield unbiased estimates? What are the implications of this exercise for the design of cluster randomized experiments?

Answer:

If A and B are combined and put into treatment, the estimated ATE is 2/3; if C is treated, the estimated ATE is 2. Therefore, the average estimated ATE is  $\frac{2/3+2}{2} = 1.33$ , which is the

true ATE. Therefore, combining clusters to make cluster size constant eliminates bias. The implication is that bias can be avoided by constructing clusters of equal size.

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