# Completely randomized designs

#### Course website

I'll put slides and labs here:

http://vicpena.github.io/doe

#### Lab groups

We have made two lab groups

- ► Group A: Meets from 3pm to 5pm at PC2
- ► Group B: Meets from 5pm to 7pm **at PC1** (different classroom)

You can find your group on my website

#### Last time

We reviewed the model

$$Y_j = \mu + \varepsilon_j, \qquad \varepsilon_j \stackrel{\text{iid}}{\sim} N(0, \sigma^2)$$

- $\blacktriangleright$   $\mu$  is a deterministic component, the expectation of Y
- $ightharpoonup arepsilon_j$  is a random "noise" term, centered at zero
- We know how to estimate  $\mu$  and  $\sigma^2$ , find confidence intervals for them, etc.

#### **Today**

We work with the model

$$Y_{ij} = \mu_i + \varepsilon_{ij}, \qquad \varepsilon_{ij} \stackrel{\text{iid}}{\sim} N(0, \sigma^2),$$

- i indexes group membership, j indexes observations within a group
- There are k groups, each with its own expected value  $\mu_1, \mu_2, \dots, \mu_k$ ; that is,  $\mathbb{E}(Y_{ij}) = \mu_i$
- Within each group, there are  $n_i$  observations, if  $n_1 = \cdots = n_k$ , we say that the design is balanced
- ▶ The total sample size is  $N = n_1 + n_2 + ... + n_k$
- The variance of the data within groups is the same; that is,  $Var(Y_{ij}) = \sigma^2$  for all i and j

#### Example

- ▶ A pharmaceutical wants to compare the effectiveness of 3 treatments (T1, T2, and T3).
- ► The treatments are **randomly** assigned to 24 patients. Large values of the outcome are associated with high effectiveness.

T1	T2	Т3	
4	7	9	
2	6	12	
6	5	6	
6	7	11	
5	6	10	
6	4	11	
2	7	9	
6	5	10	

## Model assumptions

The model comes with assumptions:

- ► The data are normally distributed
- The data are independent (there is no time dependence, for example)
- ▶ The variance is the **same** for all groups:  $Var(y_{ii}) = \sigma^2$

## Designed experiments vs observational studies

▶ The model

$$Y_{ij} = \mu_i + \varepsilon_{ij}, \qquad \varepsilon_{ij} \stackrel{\text{iid}}{\sim} N(0, \sigma^2),$$

can be used for both designed experiments and observational studies

- In this course, we will assume that the group assignment is controlled by us and made at random. In observational studies, we cannot assign observations to groups ourselves
- ➤ As we saw last time, randomizing can protect us from the effect of confounding variables
- In our context, this model is called a completely randomized design or one-way ANOVA (both names refer to the same thing)

## Inferential goals

#### We want to

- estimate the parameters  $\mu_1, \mu_2, \dots, \mu_k$  and  $\sigma^2$
- find confidence intervals for  $\mu_1, \mu_2, \dots, \mu_k$
- know whether the  $\mu_i$  are all equal or not, and to see which ones are different, if any
- check that the model assumptions are satisfied

#### Point estimation

- $\blacktriangleright$  The data come from groups with different means but equal variance  $\sigma^2$
- It won't surprise you that we'll estimate  $\mu_i$  with the sample mean of the observations coming from group i

$$\widehat{\mu}_i = \overline{Y}_i = \frac{1}{n_i} \sum_{i=1}^{n_i} Y_{ij}$$

▶ How to estimate  $\sigma^2$ ? Let the sample variance for group *i* be

$$S_i^2 = \frac{\sum_{j=1}^{n_i} (Y_{ij} - \overline{Y}_i)^2}{n_i - 1}$$

Then, we estimate  $\sigma^2$  with a weighted average of the  $S_i^2$ :

$$\widehat{\sigma}^2 = \frac{\sum_{i=1}^k (n_i - 1) S_i^2}{\sum_{i=1}^k (n_i - 1)}$$

# Confidence intervals for $\mu_i$

▶ A  $100 \cdot (1 - \alpha)$ % confidence interval for  $\mu_i$  is

$$\mathrm{CI}_{1-lpha}(\mu_i) = \overline{Y}_i \pm \mathrm{qt}(N-k,1-lpha/2) \frac{\widehat{\sigma}}{\sqrt{n_i}}$$

▶ In practice, we can use the confint function in R

# Back to example: Reading in data

Reading in and formatting the data

```
library(tidyverse)
wide = matrix(c(4, 7, 9,
                   2, 6, 12,
                   6, 5, 6,
                   6, 7, 11,
                   5, 6, 10,
                   6. 4. 11.
                   2. 7. 9.
                   6, 5, 10), byrow = T, ncol = 3)
wide = as.data.frame(wide)
colnames(wide) = c("T1", "T2", "T3")
df = wide %>% pivot_longer(cols = c(T1, T2, T3),
                               names_to = "treat",
                               values_to = "outcome")
df$treat = factor(df$treat)
```

#### Example: coefficients and confidence intervals

```
By default, confint does 95% confidence intervals
mod = aov(outcome ~ treat - 1, data = df)
dummy.coef(mod)
## Full coefficients are
##
## treat:
            T1 T2
                            Т3
            4.625 5.875 9.750
##
confint(mod)
##
              2.5 % 97.5 %
## treatT1 3.443244 5.806756
## treatT2 4.693244 7.056756
## treatT3 8.568244 10.931756
```

#### Alternative parametrization: sum-to-zero

The model

$$Y_{ij} = \mu_i + \varepsilon_{ij}, \qquad \varepsilon_{ij} \stackrel{\text{iid}}{\sim} N(0, \sigma^2),$$

can be reparametrized as

$$Y_{ij} = \mu + \tau_i + \varepsilon_{ij}, \qquad \varepsilon_{ij} \stackrel{\text{iid}}{\sim} N(0, \sigma^2), \qquad \sum_{i=1}^k \tau_i = 0$$

In this parametrization,

- $\triangleright$   $\mathbb{E}(Y_{ij}) = \mu + \tau_i$
- $\blacktriangleright$   $\mu$  is a "grand mean" common to all groups and  $\tau_i$  is a group-specific effect
- ▶ The models are equivalent: given  $\mu_1, \mu_2, ..., \mu_k$ , we can find  $\mu$  and  $\tau_1, \tau_2, ..., \tau_k$  (and viceversa)

$$\mu = \frac{\sum_{i=1}^{k} \mu_i}{k}, \qquad \tau_i = \mu_i - \frac{\sum_{i=1}^{k} \mu_i}{k}$$

Sum-to-zero: estimation

Point estimates

$$\widehat{\mu} = \frac{1}{k} \sum_{i=1}^{k} \overline{Y}_{i}, \qquad \tau_{i} = \overline{Y}_{i} - \widehat{\mu}$$

As before, we can find point estimates and intervals with R

# Sum-to-zero parametrization in R

```
Only CIs for \mu, \tau_1, and \tau_2 because \tau_3 = -\tau_1 - \tau_2
options(contrasts = c("contr.sum", "contr.poly"))
mod = aov(outcome ~ treat, data = df)
dummy.coef(mod)
## Full coefficients are
##
                6.75
## (Intercept):
                        T1 T2
                                       Т3
## treat:
##
                    -2.125 -0.875 3.000
confint(mod)
                    2.5 % 97.5 %
##
## (Intercept) 6.067713 7.43228732
```

## treat1 -3.089900 -1.16010001 ## treat2 -1.839900 0.08989999

## Hypothesis tests

We'll see two types of hypothesis tests

► Global test:

$$H_0: \mu_1 = \mu_2 = \ldots = \mu_k \qquad H_1: \mu_i \neq \mu_j \text{ for some } i, j,$$

which, in the sum-to-zero parametrization, is equivalent to

$$H_0: \tau_1 = \tau_2 = \dots = \tau_k = 0$$
  $H_1: \tau_i \neq 0$  for some  $i$ 

▶ Pairwise tests: for all pairs  $\{i,j\} \subset \{1,...,k\}$ , test

$$H_{0,ij}: \mu_i = \mu_j \qquad H_{1,ij}: \mu_i \neq \mu_j$$

#### Global test

$$H_0: \mu_1 = \mu_2 = \dots = \mu_k = \mu$$
  $H_1: \mu_i \neq \mu_j$  for some  $i, j, j \neq j$ 

#### How we do the test:

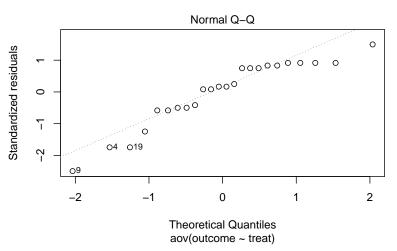
- Check that model assumptions are satisfied; if not, do not perform test
- 2. Set a significance level  $\alpha$
- 3. Given the data, compute a test statistic with a known distribution under  $H_0$
- 4. Find p-value: probability of finding a t-statistic as extreme or more extreme than the one we observed assuming that  $H_0$  is true; if the p-value is less than  $\alpha$ , reject  $H_0$ ; otherwise, do not reject  $H_0$

## Checking model assumptions

#### Model assumptions and how to check them:

- 1. **Normality:** qq-plot of model residuals
- 2. **Equality of variance:** visually, inspecting that the data from the groups have similar variance; there are formal tests as well, like bartlett.test in R
- Independence: Harder to check... If we have recorded the order in which the experiments have been run, we can check for time dependence

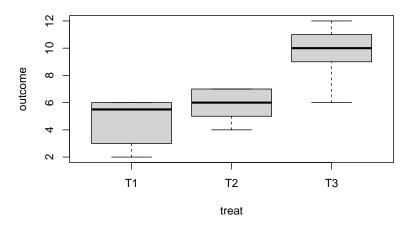
# Example: Checking normality



If the assumption is met, most points should be near the dashed line

## Example: Checking equality of variance

```
boxplot(outcome ~ treat, data = df)
```



If the assumption is met, the groups should have similar variability

# Finding a test statistic

Let

$$\overline{Y}_i = \frac{1}{n_i} \sum_{j=1}^{n_i} Y_{ij}, \qquad \overline{Y} = \frac{1}{N} \sum_{i=1}^k \sum_{j=1}^{n_i} Y_{ij}.$$

Then, the total sum of squares (SS total) as

$$\sum_{i=1}^{k} \sum_{j=1}^{n_i} (Y_{ij} - \overline{Y})^2 = \sum_{i=1}^{k} n_i (\overline{Y}_i - \overline{Y})^2 + \sum_{i=1}^{k} \sum_{j=1}^{n_i} (Y_{ij} - \overline{Y}_i)^2$$

that is,

SS total = SS between groups + SS within groups

## Finding a test statistic

We are testing:

$$H_0: \mu_1 = \mu_2 = \dots = \mu_k$$
  $H_1: \mu_i \neq \mu_i$  for some  $i, j,$ 

Intuitively...

- ▶ If  $H_0$  is true  $\overline{Y}_1 \approx \cdots \approx \overline{Y}_k \approx \overline{Y}$ , so we expect to have more variability within groups than between groups
- ▶ If  $H_0$  is not true, we expect to see more variability between groups

The test statistic will take that into account

#### The test statistic

Define the between groups (SSB), and within groups (SSW) sum of squares as below:

$$SSB = \sum_{i=1}^{k} n_i (\overline{Y}_i - \overline{Y})^2, \qquad SSW = \sum_{i=1}^{k} \sum_{j=1}^{n_i} (Y_{ij} - \overline{Y}_i)^2$$

And then define the "mean squares"

$$MSB = \frac{SSB}{k-1}, \qquad MSW = \frac{SSW}{N-k}$$

If  $H_0$  is true,

$$F = \frac{\text{MSB}}{\text{MSW}} \stackrel{H_0}{\sim} F_{k-1,N-k},$$

where  $F_{k-1,N-k}$  is the F-distribution with k-1 and N-k degrees of freedom.

## Finding the *p*-value

The *p*-value is one-sided: given data  $y_{ij}$ , we can compute the observed *F*-statistic  $f_{\rm obs}$  and compute the *p*-value as

$$P(F_{k-1,N-k} > f_{\rm obs})$$

Why is it one-sided? Recall that the F-statistic is

$$F = \frac{\text{MSB}}{\text{MSW}},$$

- ▶ If  $F \approx 0$ , we shouldn't be rejecting  $H_0$  because those are cases where the variability between groups is small, which is consistent  $H_0$ .
- ▶ We should only reject  $H_0$  for large values of F

# ANOVA (Analysis of Variance) table

df				<i>p</i> -value
Between $k$ - Within $N$	- 1 SSB - <i>k</i> SSW	$MSB = \frac{SSB}{k-1}$ $MSW = \frac{SSW}{N-k}$	$f_{\mathrm obs} = rac{\mathrm{MSB}}{\mathrm{MSW}}$	р

where the p-value is

$$p = P(F_{k-1,N-k} > f_{\rm obs})$$

## A note on terminology

- Some authors refer to SSB as SST (sum of squares treatment) and SSW as SSE (sum of squares error)
- R will use the term Residuals to refer to SSW

## Example: ANOVA table

##

#### Suppose our significance level is $\alpha = 0.05$

```
Df Sum Sq Mean Sq F value Pr(>F)
## treat 2 114.25 57.12 22.11 6.79e-06 ***
## Residuals 21 54.25 2.58
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.3
p-value is 6.79 \cdot 10^{-6} < \alpha, so we reject H_0
```

#### Pairwise comparisons

- ► The global test can detect if there are at least two group means that are different, but it doesn't tell us which ones are different
- If we're interested in seeing where the differences between groups lie, we can perform all  $\binom{k}{2}$  pairwise comparisons

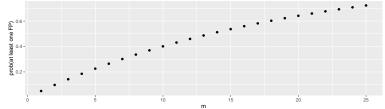
$$H_{0,ij}: \mu_i = \mu_j, \qquad H_{1,ij}: \mu_i \neq \mu_j$$

## Probability of a false positive

- We have to be careful when we perform such tests
- ▶ If we perform m independent tests at significance level  $\gamma$  and the null hypothesis is true for all of them, the probability of getting at least one false positive is

$$P(\text{at least one FP}) = 1 - P(\text{no FP}) = 1 - (1 - \gamma)^m$$

If  $\gamma=0.05$ , here's a graph of the probability



## Controlling false positive rate

There are many methods to perform the pairwise tests we're interested in while controlling that

$$P(\text{at least one FP}) = \alpha$$

- One such method is the Tukey Honestly Significant Difference test: TukeyHSD in R
- ▶ TukeyHSD reports point estimates for all pairwise differences, along with adjusted confidence intervals and p-values. If we reject an adjusted p-value if it is less than  $\alpha$ , the probability of at least one false positive is  $\alpha$ .
- TukeyHSD is based on the distribution of the Studentized range  $q = \sqrt{N} \, (\overline{Y}_{\rm max} \overline{Y}_{\rm min}) / \widehat{\sigma}$  under the same assumptions we've been making

#### Example: TukeyHSD

```
The default confidence level (1 - \alpha) is 0.95
```

```
TukeyHSD(mod, conf.level = 0.95)
##
     Tukey multiple comparisons of means
       95% family-wise confidence level
##
##
## Fit: aov(formula = outcome ~ treat, data = df)
##
## $treat
##
          diff
                      lwr
                               upr padj
## T2-T1 1.250 -0.7756249 3.275625 0.2865063
## T3-T1 5.125 3.0993751 7.150625 0.0000073
## T3-T2 3.875 1.8493751 5.900625 0.0002592
Significant differences between T3 and T1 and T3 and T2 at
\alpha = 0.05
```