

MEDB 5501, Module12

2024-11-05

Topics to be covered

- What you will learn
 - Two factor analysis of variance
 - Relationship to linear regression
 - Checking assumptions
 - R code for two factor analysis of variance
 - Interactions
 - R code for interactions
 - Your homework

Two factor analysis of variance

- Continuous outcome
- Two categorical predictors
- Example
 - Hearing test (decibels at high frequency)
 - Age group (Old or Young)
 - Gender (Female or Male)

Speaker notes

Two factor analysis of variance uses two categorical variables to predict a continuous outcome. We will focus today on the balanced case. In the balanced case, each combination of category levels has the same number of observations.

Balanced data

- Proportional number in each category level combination group
 - 3 old females, 3 old males, 3 young females, 3 young males
 - 6 old females, 6 old males, 2 young females, 2 young males

Speaker notes

Two factor analysis of variance uses two categorical variables to predict a continuous outcome. We will focus today on the balanced case. In the balanced case, each combination of category levels has the same number of observations.

Unbalanced data

- Unequal numbers in some category combinations
 - 3 old females, 3 old males, 3 young females, 2 young males
- Extreme case: empty category combinations
 - 3 old females, 3 old males, 3 young females, 0 young males

Speaker notes

In the unbalanced case the observations are not equal or proportional. These cases are quite complex from several perspectives.

Mathematical model, 1

- Y_{ijk}
 - i = which level of first category
 - j = which level of second category
 - k = which patient within a category combination

Speaker notes

With two levels, you need three subscripts (i, j, k) to keep track of the observations.

Mathematical model, 2

<i>Age</i>	<i>Gender</i>	<i>Outcome</i>
<i>Old</i>	<i>Female</i>	Y_{111}
<i>Old</i>	<i>Female</i>	Y_{112}
<i>Old</i>	<i>Female</i>	Y_{113}
<i>Old</i>	<i>Male</i>	Y_{121}
<i>Old</i>	<i>Male</i>	Y_{122}
<i>Old</i>	<i>Male</i>	Y_{123}
<i>Young</i>	<i>Female</i>	Y_{211}
<i>Young</i>	<i>Female</i>	Y_{212}
<i>Young</i>	<i>Female</i>	Y_{213}
<i>Young</i>	<i>Male</i>	Y_{221}
<i>Young</i>	<i>Male</i>	Y_{222}
<i>Young</i>	<i>Male</i>	Y_{223}

Speaker notes

Here's a simple example where the first categorical predictor and the second categorical predictor have two levels and there are three observation in each combination.

Mathematical model, 3

- $Y_{ijk} = \mu + \alpha_i + \beta_j + \epsilon_{ijk}$
 - $i = 1, \dots, a, j = 1, \dots, b$
 - $\sum \alpha_i = 0, \sum \beta_j = 0$
 - ϵ_{ijk} is $N(0, \sigma)$
- $\bar{Y}_{i.}$ is the average for the i th level of first factor
- $\bar{Y}_{.j}$ is the average for the j th level of second factor
- $\bar{Y}_{..}$ is the average for all of the data

Speaker notes

The mathematical model includes an overall mean (μ), a deviation from the overall mean associated with the different levels of the first factor (α_i), a deviation from the overall mean associated with the different levels of the second factor (β_j) and an error term (ϵ_{ijk}).

Because the alphas and betas are deviations, they have to sum to zero. The error term is assumed to be normally distributed and the standard deviation is the same for all the data.

You will need to compute averages for the first factor, $\bar{Y}_{i..}$, the second factor, $\bar{Y}_{.j.}$, and an overall mean, $\bar{Y}_{...}$ which is an average across all of the data.

Mathematical model, 4

- $SS(Total) = \sum_i \sum_j \sum_k (Y_{ijk} - \bar{Y}_{...})^2$
 - $df=abn-1$
- $SS(A) = \sum_i bn(\bar{Y}_{i..} - \bar{Y}_{...})^2$
 - $df=a-1$
- $SS(B) = \sum_j an(\bar{Y}_{.j.} - \bar{Y}_{...})^2$
 - $df=b-1$
- $SS(Error) = SS(Total) - SS(A) - SS(B)$
 - $df=(abn-1)-(a-1)-(b-1)$

Speaker notes

To assess the impact of the two categorical predictors, you compute sums of squares. $SS(\text{Total})$ represents the deviation of the individual values from the overall mean. $SS(A)$ represents deviations of the first category means from the overall mean. $SS(B)$ represents deviations of the second category means from the overall mean. Whatever is left over is $SS(\text{Error})$.

Artificial data

```
# A tibble: 12 × 5
```

	id	age	gender	code	db
	<int>	<chr>	<chr>	<chr>	<dbl>
1	1	old	female	of	45
2	2	old	female	of	60
3	3	old	female	of	60
4	4	old	male	om	65
5	5	old	male	om	60
6	6	old	male	om	70
7	7	young	female	yf	20
8	8	young	female	yf	20
9	9	young	female	yf	5
10	10	young	male	ym	25
11	11	young	male	ym	20
12	12	young	male	ym	30

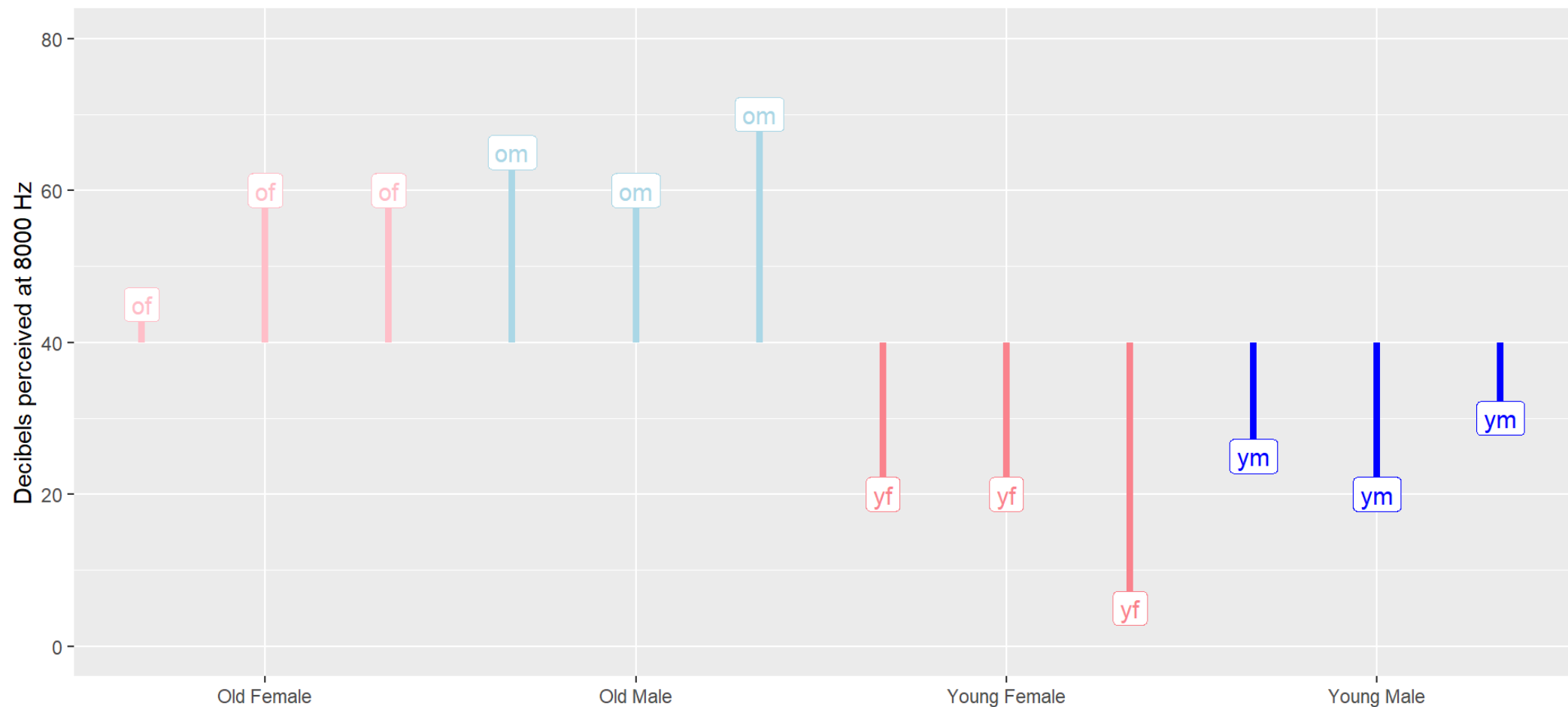
Artificial data with means

```
# A tibble: 12 × 8
```

	id	age	gender	code	db	age_mean	gender_mean	overall_mean
	<int>	<chr>	<chr>	<chr>	<dbl>	<dbl>	<dbl>	<dbl>
1	1	old	female	of	45	60	35	40
2	2	old	female	of	60	60	35	40
3	3	old	female	of	60	60	35	40
4	4	old	male	om	65	60	45	40
5	5	old	male	om	60	60	45	40
6	6	old	male	om	70	60	45	40
7	7	young	female	yf	20	20	35	40
8	8	young	female	yf	20	20	35	40
9	9	young	female	yf	5	20	35	40
10	10	young	male	ym	25	20	45	40
11	11	young	male	ym	20	20	45	40
12	12	young	male	ym	30	20	45	40

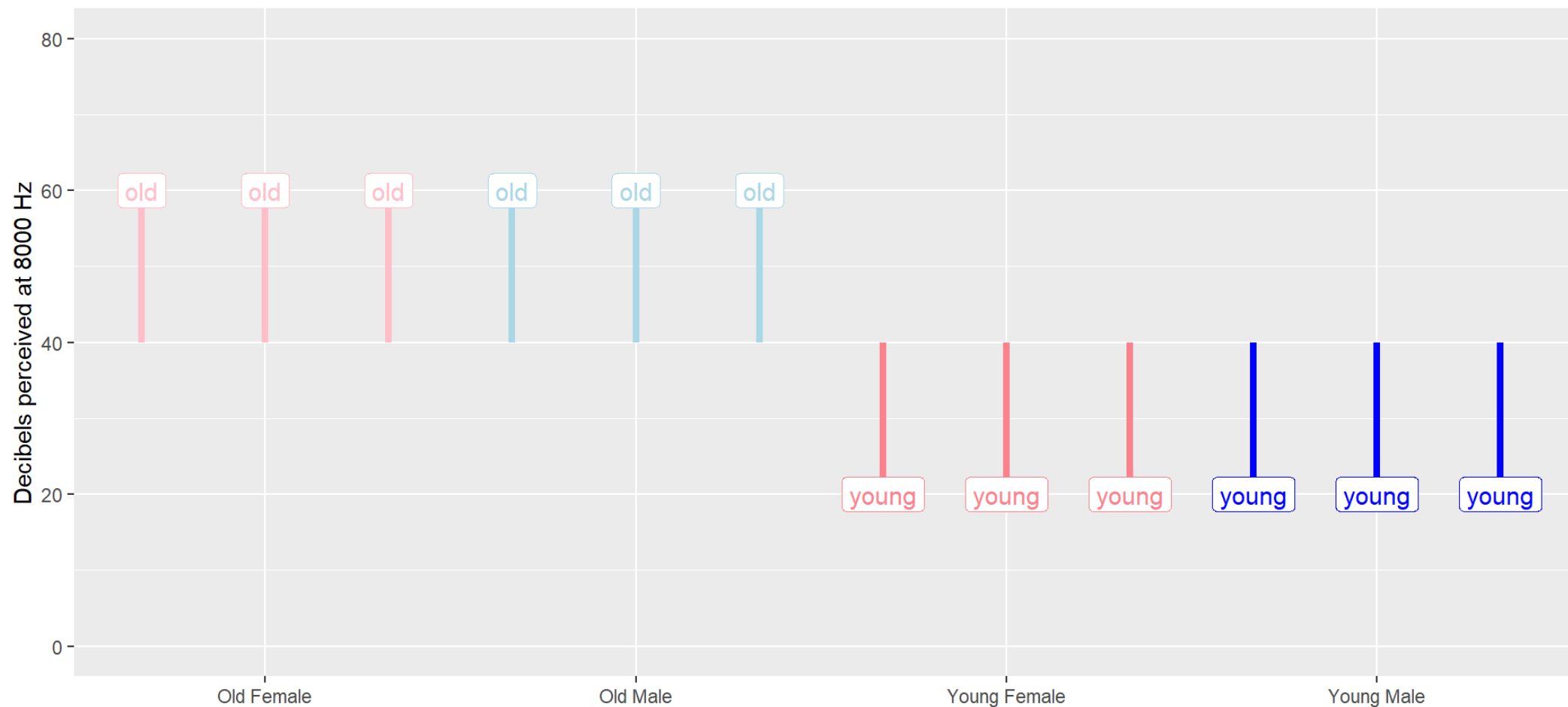
SS(Total)

Graph drawn by Steve Simon on 2024-11-03



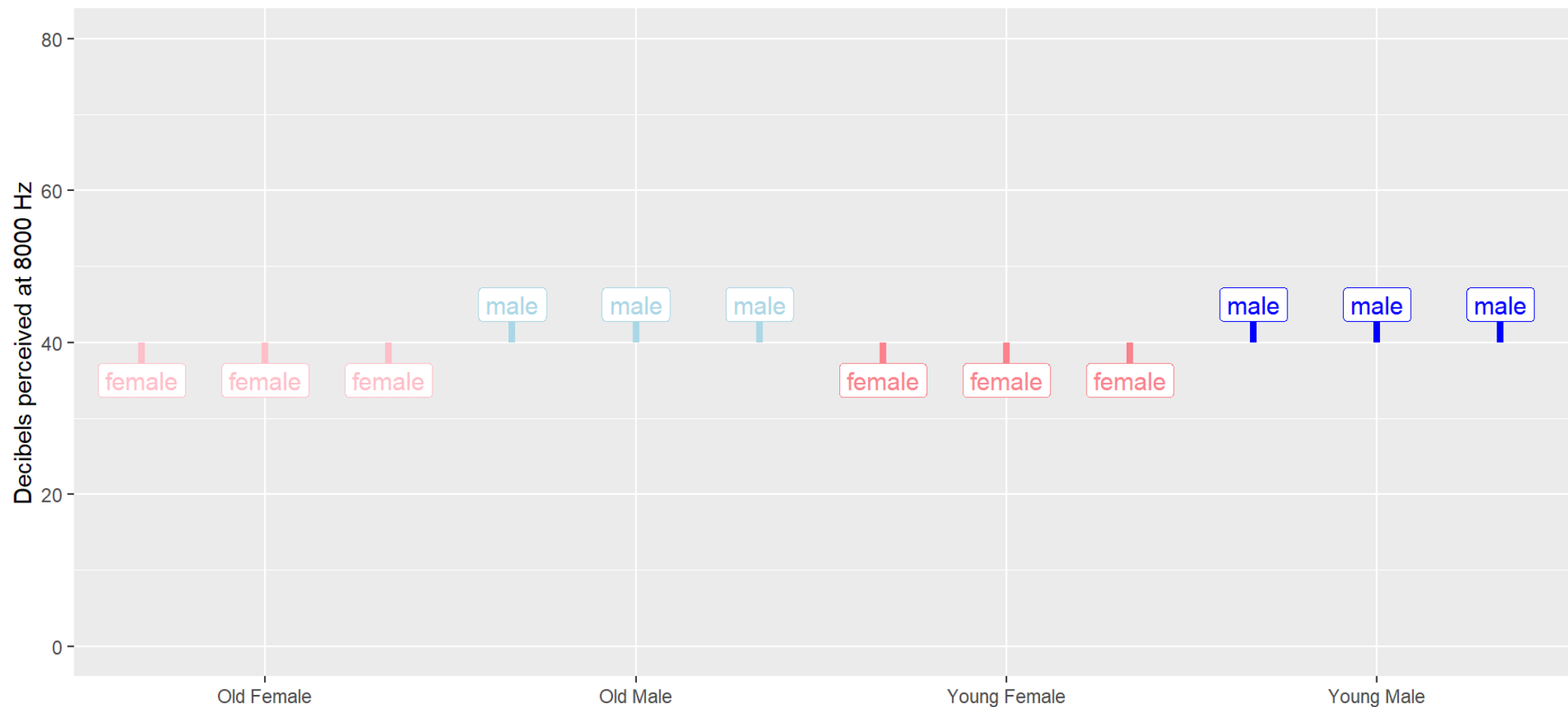
SS(Age)

Graph drawn by Steve Simon on 2024-11-03



SS(Gender)

Graph drawn by Steve Simon on 2024-11-03



Analysis of variance table

Analysis of Variance Table

Response: db

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
age	1	4800	4800.0	108.00	2.595e-06	***
gender	1	300	300.0	6.75	0.02883	*
Residuals	9	400	44.4			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Break #1

- What you have learned
 - Two factor analysis of variance
- What's coming next
 - Relationship to linear regression

Create indicator variables

```
# A tibble: 12 × 6
  age    gender code  i_young i_male    db
<chr> <chr>  <chr>    <dbl>  <dbl>  <dbl>
1 old    female of      0      0     45
2 old    female of      0      0     60
3 old    female of      0      0     60
4 old    male   om      0      1     65
5 old    male   om      0      1     60
6 old    male   om      0      1     70
7 young female yf      1      0     20
8 young female yf      1      0     20
9 young female yf      1      0      5
10 young male   ym      1      1     25
11 young male   ym      1      1     20
12 young male   ym      1      1     30
```

Speaker notes

You need $k-1$ indicators for a categorical predictor that has k levels. In this simple example, that just means one indicator for age and one indicator for gender.

Two factor analysis of variance using aov

```
1 m1 <- aov(db ~ age + gender, data=hearing)
2 anova(m1)
```

Analysis of Variance Table

Response: db

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
age	1	4800	4800.0	108.00	2.595e-06	***
gender	1	300	300.0	6.75	0.02883	*
Residuals	9	400	44.4			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Speaker notes

Here is a repeat of the analysis of variance table using aov.

Two factor analysis of variance using linear regression, 1

```
1 m2 <- lm(db ~ age + gender, data=hearing)
2 anova(m2)
```

Analysis of Variance Table

Response: db

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
age	1	4800	4800.0	108.00	2.595e-06	***
gender	1	300	300.0	6.75	0.02883	*
Residuals	9	400	44.4			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Speaker notes

The analysis of variance table is identical when you use `lm` in place of `aov`.

Two factor analysis of variance using linear regression, 2

```
1 tidy(m2)
```

```
# A tibble: 3 × 5
```

	term <chr>	estimate <dbl>	std.error <dbl>	statistic <dbl>	p.value <dbl>
1	(Intercept)	55	3.33	16.5	0.0000000492
2	ageyoung	-40	3.85	-10.4	0.00000260
3	gendermale	10.0	3.85	2.60	0.0288

Speaker notes

The intercept is the estimated db when both indicator variables equal zero. The first indicator, i_age , is the estimated average change in db when moving from female to male. The second indicator, i_gender , is the estimated average change in db when moving from old to young.

Break #2

- What you have learned
 - Relationship to linear regression
- What's coming next
 - Checking assumptions

Checking assumptions

- Normality (Non-normality)
- Homogeneity (Heterogeneity)
- Independence (Lack of independence)

Speaker notes

The assumptions for two factor analysis of variance is no different than for one factor analysis of variance or the two-sample t-test.

I'm a bit inconsistent in how I present this material. The assumptions are satisfied if you have normality, homogeneity, and independence. Equivalently, you could state that the assumptions are violated if you have non-normality or heterogeneity or lack of independence.

Use the boxplot to check assumptions

- Non-normality if boxplot shows skewness and/or outliers
- Heterogeneity if boxplot shows large change in variation
- Draw clustered boxplot to examine every combination of categories
 - Use $a \times b$ boxplots
- Independence is checked qualitatively

Alternatives if assumptions violated

- There is no analog to Kruskal-Wallis or Mann-Whitney-Wilcoxon
- Consider a log transformation
 - All values greater than 0
 - Groups with larger means have larger variation
 - Data is skewed right and outliers only on the high end

Break #3

- What you have learned
 - Checking assumptions
- What's coming next
 - R code for two factor analysis of variance

Listing of full-moon-er-admissions.yaml, 1

data_dictionary: full-moon.txt

description: >

The data give the admission rates to the emergency room of a Virginia mental health clinic before, during and after the 12 full moons from August 1971 to July 1972.

additional_description:

<https://gksmyth.github.io/ozdasl/general/fullmoon.html>

Listing of full-moon-er-admissions.yaml, 2

source:

- Blackman, S., and Catalina, D. (1973). The moon and the emergency room. Perceptual and Motor Skills 37, 624-626.
- Ovlin, J. F. (1943). Moonlight and nervous disorders. American Journal of Psychiatry 99, 578-584.
- Larsen, R.J., and Marx, M.L., (1986). An Introduction to Mathematical Statistics and Its Applications 2nd Edition. Prentice-Hall, Englewood Cliffs, New Jersey. Case Study 1.2.3.

copyright:

unknown

format:

tab-delimited

Listing of full-moon-er-admissions.yaml, 3

```
varnames:  
  first row of data  
  
size:  
  rows: 36  
  columns: 3  
  
vars:  
  Month:  
    label: Month of year  
    scale: Aug, Sep, ..., Jul
```

Listing of full-moon-er-admissions.yaml, 4

Moon:

label: time relative to full moon

values:

- Before
- During
- After

Admission:

label: Admission rate

Unit: patients/day

Listing of simon-5501-12-moon.qmd, 1

```
---  
title: "Analysis of full-moon-er-admissions data"  
format:  
  html:  
    embed-resources: true  
---
```

This program reads data on admissions to an emergency room before, during, and after a full moon. Find more information in the [data dictionary][dd].

[dd]: <https://github.com/pmean/data/blob/master/files/full-moon-er-admissions.yaml>

This code was written by Steve Simon on 2024-11-02 and is placed in the public domain.

Listing of simon-5501-12-moon.qmd,

2

```
## Load the tidyverse library
```

```
```{r setup}  
#| message: false
#| warning: false
library(broom)
library(tidyverse)
```
```

For most of your programs, you should load the tidyverse library. The broom library converts your output to a nicely arranged dataframe. The messages and warnings are suppressed.

Listing of simon-5501-12-moon.qmd,

3

```
## Read the data and view a brief summary

```{r read}
er <- read_tsv(
 "../data/full-moon-er-admissions.txt",
 col_types="ccn",
 col_names=TRUE)
names(er) <- tolower(names(er))
glimpse(er)
```
```

This is a tab delimited file. I changed all the variable names to all lowercase.

That's one less thing opportunity for inconsistency errors.

Listing of simon-5501-12-moon.qmd, 4

```
## Reorder the category levels, 1

```{r reorder-moon}
er$moon1 <- factor(
 er$moon,
 levels=c(
 "Before",
 "During",
 "After"))
````
```

If you let R order the categories, it will do it alphabetically. The factor function will place them in a more logical order.

Listing of simon-5501-12-moon.qmd, 5

```
## Reorder the category levels, 2

```{r reorder-month}
er$month1 <- factor(
 er$month,
 levels=c(
 "Aug", "Sep", "Oct",
 "Nov", "Dec", "Jan",
 "Feb", "Mar", "Apr",
 "May", "Jun", "Jul"))
````
```

Since the study started in August, it makes sense to order the months starting then. An alphabetical ordering (April first then August?) makes no sense here.

Listing of simon-5501-12-moon.qmd,

6

```
## Draw boxplot of admission versus moon

```{r boxplots-admission-vs-moon}
#| fig.width: 6
#| fig.height: 2.0
er |>
 ggplot(aes(moon1, admission)) +
 geom_boxplot() +
 xlab("Moon phase") +
 ylab("ER admissions per day") +
 ggtitle("Graphs drawn by Steve Simon on 2024-11-02") +
 coord_flip()
```,
```

Although there are a few outliers, I would not worry too much about the

Listing of simon-5501-12-moon.qmd, 7

```
## Draw boxplot of admission versus month

```{r boxplots-admission-vs-month}
#| fig.width: 6
#| fig.height: 6.0
er |>
 ggplot(aes(month1, admission)) +
 geom_boxplot() +
 xlab("Month") +
 ylab("ER admissions per day") +
 ggtitle("Graphs drawn by Steve Simon on 2024-11-02") +
 coord_flip()
````
```

Although there are some unusual patterns here, keep in mind that there are

Listing of simon-5501-12-moon.qmd, 8

```
## Calculate average ER admissions versus moon

```{r by-moon}
er |>
 group_by(moon1) |>
 summarize(
 admission_mn=mean(admission),
 admission_sd=sd(admission),
 n=n())
```
```

There is a slightly higher average admission rate and slightly more vaiation during a full moon.

Listing of simon-5501-12-moon.qmd,

9

```
## Calculate average ER admissions versus month

```{r by-month}
er |>
 group_by(month1) |>
 summarize(
 admission_mn=mean(admission),
 admission_sd=sd(admission),
 n=n())
```
```

Er admissions are trending upward over time and there is some evidence of heterogeneity. Again, treat this with caution as there are only three observations per month.

Listing of simon-5501-12-moon.qmd, 10

```
## Analysis of variance

```{r anova}
m1 <- aov(admission ~ moon1 + month1, data=er)
anova(m1)
```
```

There is a borderline, but statistically significant difference in average admission rates during different phases of the moon. There is a statistically significant difference in average admission rates between some of the months.

Listing of simon-5501-12-moon.qmd, 11

```
## Using lm, 1

```{r lm-1}
m2 <- lm(admission ~ moon1 + month1, data=er)
anova(m2)
```
```

The analysis of variance table using lm is the same as the analysis of variance table using aov.

Listing of simon-5501-12-moon.qmd, 12

```
## Using lm, 2
```

```
```{r lm-2}  
tidy(m2)
```
```

The intercept represents the estimated average admission rate before a full moon

and during the month of August. The admission rate is 2.5 patients per day higher during a full moon compared to a full moon. This is small but probably still important. The difference in average admission rates is 0.54 patients per

day higher after a full moon compared to before a full moon. The remaining estimates compare each month back to the first month (August).

Listing of simon-5501-12-moon.qmd, 13

```
## Using lm, 3

```{r lm-3}
er |>
 group_by(moon1) |>
 summarize(admission_mn=mean(admission)) |>
 mutate(admission_before=first(admission_mn)) |>
 mutate(difference=admission_mn-admission_before)
```
```

This table shows how to calculate the regression coefficients associated with phases of the moon. It is not really needed for the analysis, but helps illustrate how these coefficients are computed.

Listing of simon-5501-12-moon.qmd, 14

```
## Using lm, 4

```{r lm-4}
er |>
 group_by(month1) |>
 summarize(admission_mn=mean(admission)) |>
 mutate(admission_aug=first(admission_mn)) |>
 mutate(difference=admission_mn-admission_aug)
```
```

This table shows how to calculate the regression coefficients associated with months of the year.

Listing of simon-5501-12-moon.qmd, 15

```
## Save important files for later use

```{r save}
save(
 er,
 file="../data/full-moon-er-admissions.RData")
```
```

Break #4

- What you have learned
 - R code for two factor analysis of variance
- What's coming next
 - Interactions

What is an interaction

- Impact of one variable is influenced by a second variable
- Example, influence of alcohol on sleeping pills
- Three types of interactions
 - Between two categorical predictors
 - Between a categorical and a continuous predictor
 - Between two continuous predictors
- Interactions greatly complicate interpretation

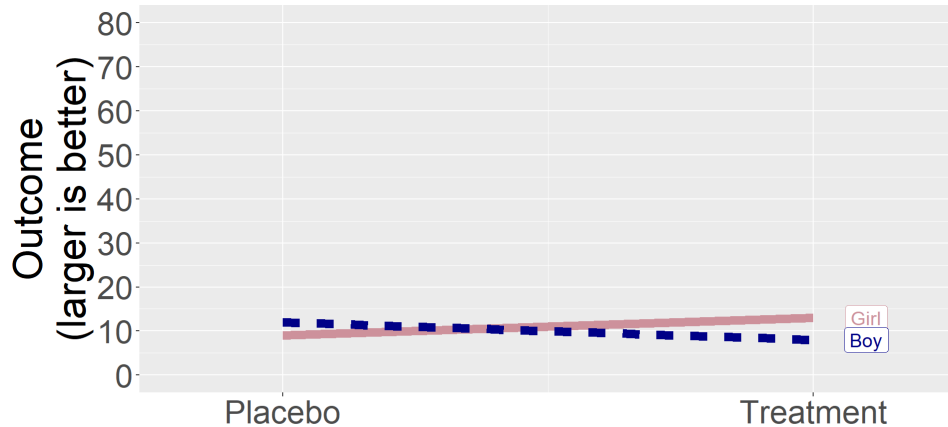
Speaker notes

Interactions are important to look for, but if you find one, don't rejoice. Interactions are a headache. They tell you that a simple interpretation of your research won't work. That's important to know, of course, but it also means that you will have to spend more time explaining your results in a paper or presentation.

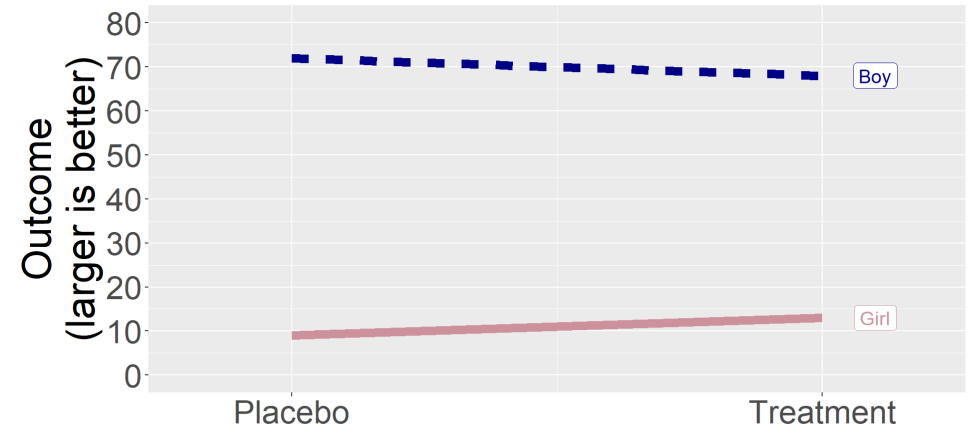
Interaction plot

- X axis, first categorical variable
- Separate lines for second categorical variable
- Y axis, average outcome

Hypothetical interaction plots, 1



- No interaction
- Ineffective treatment
- Boys/girls similar



- No interaction
- Ineffective treatment
- Boys fare better than girls

Speaker notes

An interaction plot shows the mean values for each of the two categories. In this example, there is a placebo and a treatment. The outcome is unspecified, but a larger value is presumed to represent a better outcome. This is a pediatric example and the data is subdivided into two populations, boys and girls.

The flatness or steepness of the lines indicates whether patients given the treatment fare better than patients given the placebo.

The separation (if there is any) between the lines measures whether boys fare better or worse than girls.

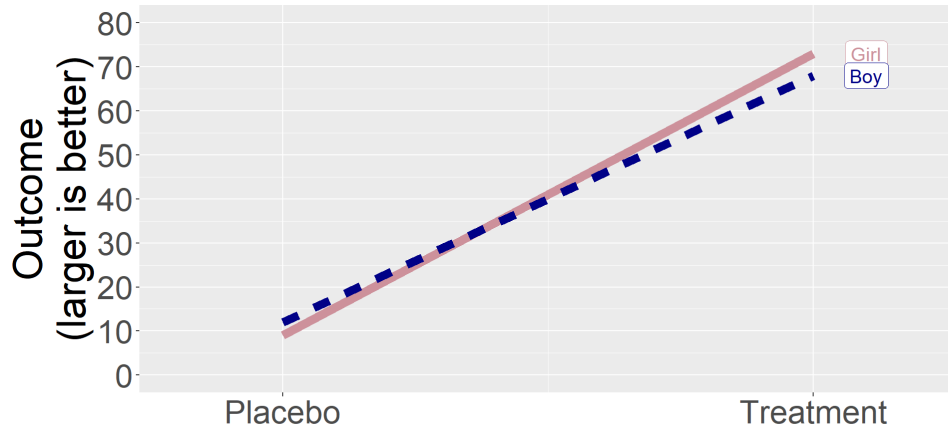
If the lines have roughly the same slope (both are flat or both are steep), then there is no interaction.

In the plot on the left, the two lines are flat, indicating that the treatment is ineffective. The outcome is not changed from the placebo.

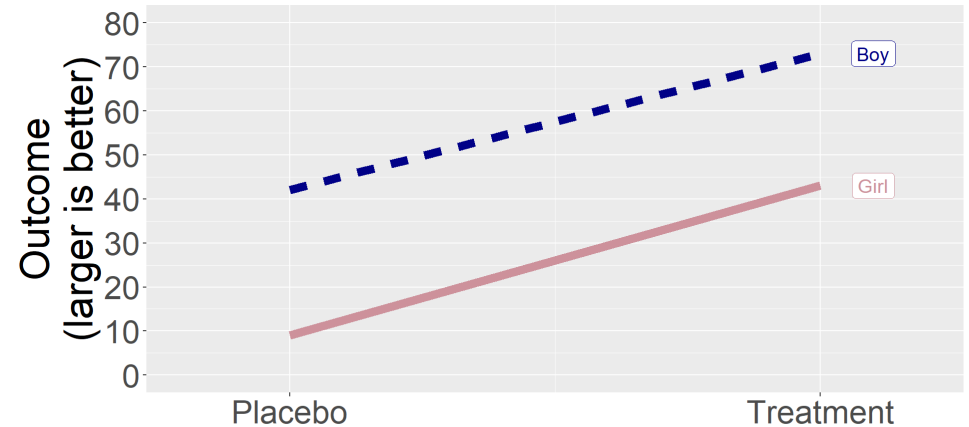
The two lines lie more or less on top of one another. This indicates that there is no difference in average outcome between boys and girls.

In the plot on the right, the two lines are flat. The treatment is ineffective. There is, however, a difference. The average outcome for boys is a lot better both in the placebo group and the treatment group. The lines are roughly parallel, indicating no interaction.

Hypothetical interaction plots, 2



- No interaction
- Effective treatment
- Boys/girls similar



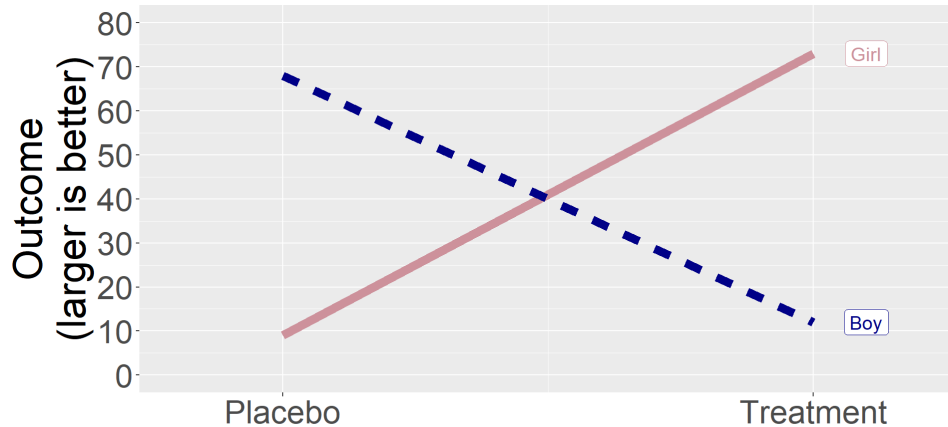
- No interaction
- Effective treatment
- Boys fare better than girls

Speaker notes

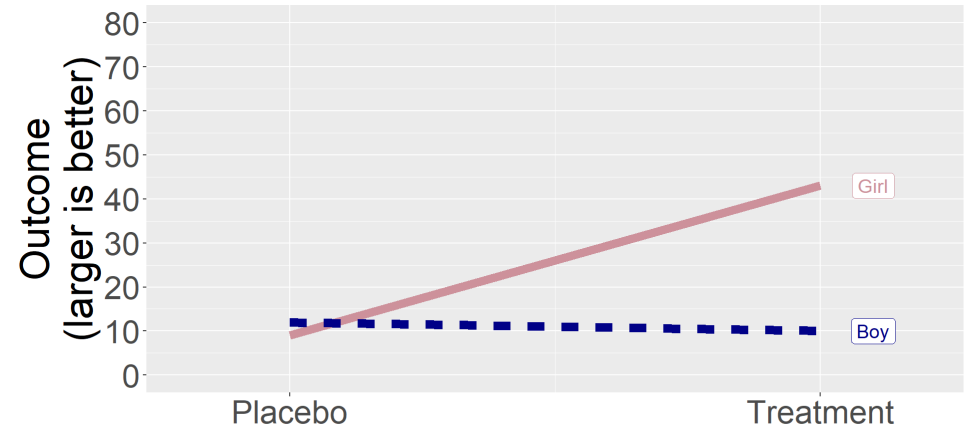
In the plot on the left, there is a steep slope for both boys and girls. The treatment is effective. There is no separation in the lines. Boys do not fare any better or worse on average than girls.

In the plot on the left, there is a steep slope and a separation between the lines. Boys fare better than girls on average. Both lines have a steep slope. The treatment. The lines are parallel, so there is no interaction.

Hypothetical interaction plots, 3



- Significant interaction
- Harmful treatment in boys
- Effective treatment in girls



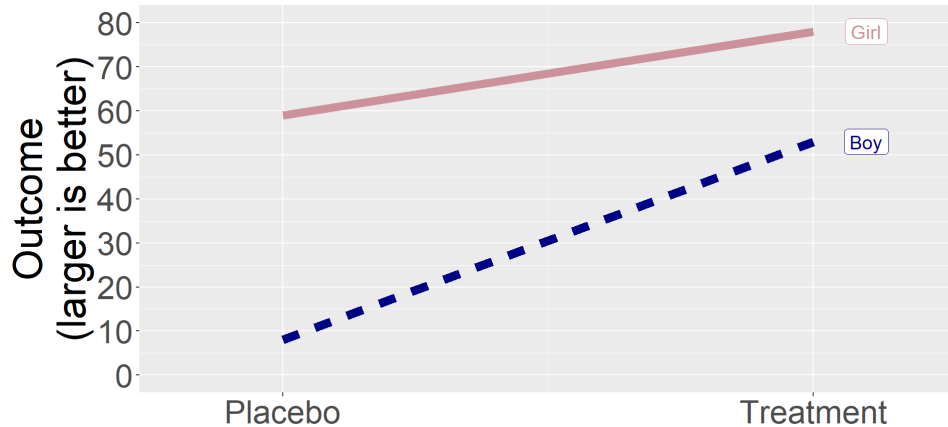
- Significant interaction
- Ineffective treatment in boys
- Effective treatment in girls

Speaker notes

In the plot on the left, the lines are not parallel, so this is evidence of an interaction. In fact, the two lines cross. This is an extreme interaction. Boys fare better on the treatment and girls fare better on the placebo.

In the plot on the right, the lines are not parallel, so this is also evidence of an interaction, but a different sort of interaction. The line for boys is flat and the line for girls is steep. The treatment is worthless for boys, but quite helpful for girls.

Hypothetical interaction plots, 4



- Significant interaction
- Girls fare better overall
- Effective treatment
- Much more effective in boys

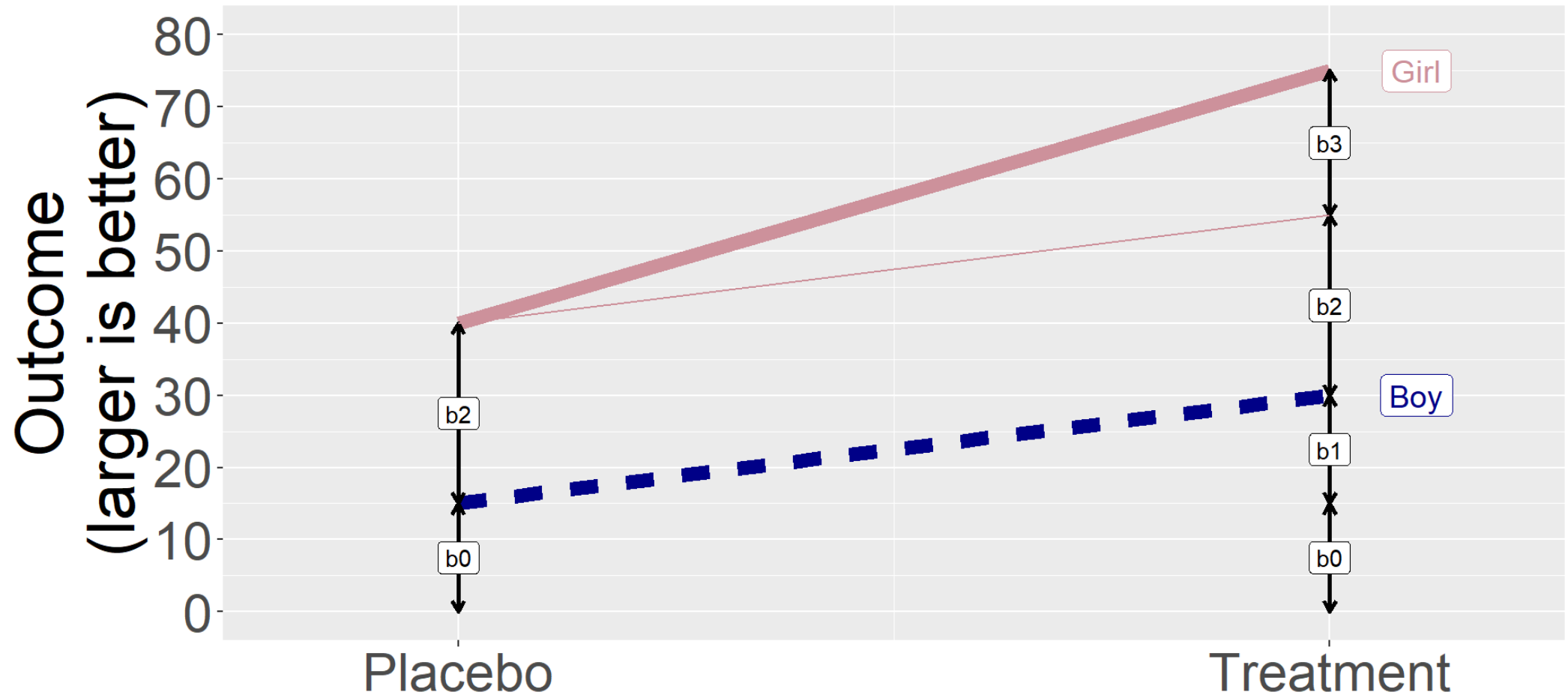
Speaker notes

In this final plot, the lines are not parallel, indicating a third type of interaction. The slope is much steeper for boys. Girls see a moderate improvement on average, but boys see a really large improvement.

Indicator variable for an interaction

```
# A tibble: 12 × 7
  age    gender code  i_young i_male i_m_by_y    db
  <chr> <chr>  <chr>    <dbl>  <dbl>  <dbl>  <dbl>
1 old   female of      0      0      0     45
2 old   female of      0      0      0     60
3 old   female of      0      0      0     60
4 old   male   om      0      1      0     65
5 old   male   om      0      1      0     60
6 old   male   om      0      1      0     70
7 young female yf      1      0      0     20
8 young female yf      1      0      0     20
9 young female yf      1      0      0      5
10 young male   ym      1      1      1     25
11 young male   ym      1      1      1     20
12 young male   ym      1      1      1     30
```

Interpretation of intercept and slopes



When you can't estimate an interaction

- Special case, $n=1$
 - Only one observation for categorical combination

Speaker notes

There is a special case where you have two categorical independent variables and you cannot estimate an interaction. If you have $n=1$, exactly one observation for each combination of your two categorical variables, then you don't have enough degrees of freedom to estimate an interaction and still be able to test whether that interaction is statistically significant.

It's sort of like that old joke I told about married life (it's okay but you lose a degree of freedom). Interactions cause an even bigger loss of degrees of freedom and in the case with only one observation per combination of categories, you lose enough degrees of freedom that it is not marriage, it being in prison.

Example, full moon study, 1 of 2

```
# A tibble: 36 × 3
  month1 moon1      n
  <fct>   <fct>   <int>
1 Aug     Before     1
2 Aug     During     1
3 Aug     After      1
4 Sep     Before     1
5 Sep     During     1
6 Sep     After      1
7 Oct     Before     1
8 Oct     During     1
9 Oct     After      1
10 Nov    Before     1
# i 26 more rows
```

Speaker notes

Here is an example where you only have one observation for each combination of categories.

Example, full moon study, 2 of 2

```
1 m1 <- aov(admission ~ month*moon, data=er)
2 anova(m1)
```

Analysis of Variance Table

Response: admission

| | Df | Sum Sq | Mean Sq | F value | Pr(>F) |
|------------|----|--------|---------|---------|--------|
| month | 11 | 455.58 | 41.417 | NaN | NaN |
| moon | 2 | 41.51 | 20.757 | NaN | NaN |
| month:moon | 22 | 127.82 | 5.810 | NaN | NaN |
| Residuals | 0 | 0.00 | NaN | | |

Speaker notes

You lose two degrees of freedom for moon (3 phases: before, during, and after). You lose 11 degrees of freedom for month (12 months -1). The interaction has 2 times 11 or 22 degrees of freedom. You only started with 35 degrees of freedom. Subtract 2, 11, and 22, and you are left with zero degrees of freedom for error.

If you find yourself in this situation, just state that no test for interaction was possible in your methods section and highlight this as a weakness of your study in the discussion section.

Break #5

- What you have learned
 - Interactions
- What's coming next
 - R code for interactions

Listing of fruitfly.yaml, 1

data_dictionary: fruitfly.dat.txt

copyright: >

This dataset is copyrighted by the authors of the Journal of Statistics Education article, but should be available for individual educational uses under the Fair Use provisions of copyright law.

description: >

Does access to mating affect the lifespan of fruitflies? This data shows the longevity of male fruitflies in the presence or absence of female fruitflies to mate with. Male fruitflies were housed with 0, 1, or 8 females. In some groups, the females were pregnant and thus not available for mating. There are

two covariates, length of the thorax and percentage of time sleeping, that might also influence longevity.

Listing of fruitfly.yaml, 2

source:

- Partridge, L., and Farquhar, M. (1981), "Sexual Activity and the Lifespan of Male Fruitflies," Nature, 294, 580-581.
- James A. Hanley & Stanley H. Shapiro. Sexual Activity and the Lifespan of Male Fruitflies: A Dataset That Gets Attention. Journal of Statistics Education v.2, n.1 (1994)

additional_description:

<https://jse.amstat.org/datasets/fruitfly.txt>

download_url:

<https://jse.amstat.org/datasets/fruitfly.dat.txt>

Listing of fruitfly.yaml, 3

```
format:
  fixed width

varnames:
  not included

missing_value_code:
  not needed

size:
  rows: 125
  columns: 6
```

Listing of fruitfly.yaml, 4

```
vars:  
  id:  
    columns: 1-2  
  
partners:  
  columns: 4  
  label: Number of female partners  
  values: 0, 1, or 8
```

Listing of fruitfly.yaml, 5

```
type:
  columns: 6
  label: Type of female fruitfly
  values:
    0: newly pregnant female
    1: virgin female
    9: not applicable (when partners=0)

longevity:
  columns: 8-9
  label: Lifespan
  unit: days
  range: positive
```

Listing of fruitfly.yaml, 6

```
thorax:
  columns: 11-14
  label: Length of thorax
  unit: mm
  range: positive

sleep:
  columns: 16-17
  label: Percentage of each day sleeping
  range: 0 to 100
---
```

Listing of simon-5501-12-fruitfly.qmd, 1

```
---  
title: "Analysis of fruitfly data"  
format:  
  html:  
    embed-resources: true  
---
```

This program reads data on fruit fly longevity. Find more information in the [data dictionary][dd].

[dd]: <https://github.com/pmean/data/blob/master/files/fruitfly.yaml>

This code was written by Steve Simon on 2024-10-29 and is placed in the public domain.

Listing of simon-5501-12-fruitfly.qmd,

2

```
## Load the tidyverse library
```

```
```{r setup}  
#| message: false
#| warning: false
library(broom)
library(tidyverse)
```
```

```
#### Comments on the code
```

For most of your programs, you should load the tidyverse library. The broom library converts your output to a nicely arranged dataframe. The messages and warnings are suppressed.

Listing of simon-5501-12-fruitfly.qmd,

3

```
## List the variable names
```

```
```{r variable-list}  
vlist <- c(
 "id",
 "partners",
 "type",
 "longevity",
 "thorax",
 "sleep")
```
```

Listing of simon-5501-12-fruitfly.qmd, 4

```
#### Comments on the code
```

```
When a dataset does not have variables on the first line, you need to specify  
them in the code.
```


Listing of simon-5501-12-fruitfly.qmd, 5

```
## Read the data and view a brief summary

```{r read}
fly <- read_fwf(
 "../data/fruitfly.txt",
 col_types="nnnnnn",
 fwf_widths(
 widths=c(2, 2, 2, 3, 5, 3),
 col_names=vlist))
glimpse(fly)
```
```

Listing of simon-5501-12-fruitfly.qmd, 6

```
#### Comments on the code
```

The fruitfly dataset has a fixed width format (fwf). You need to specify the columns that each variable uses.

Notice that the two categorical variables, *partners* and *type*, are actually numbers rather than strings. To avoid having R treat these variables as if they were continuous, use the `factor` function in some of the code below.

Listing of simon-5501-12-fruitfly.qmd, 7

```
## Create a subset of fruitfly dataset

```{r subset}
fly |>
 filter(type != 9) -> fly_subset
```
```

Listing of simon-5501-12-fruitfly.qmd, 8

```
#### Comments on the code
```

```
If you exclude the pure control group (No females), you can analyze the two  
factors, partners and type individually. Partners has two category levels, 1  
for
```

```
when one female was included in the cage and 8 for when eight females were  
included in the cage. Type also has two category levels, 0 for pregnant female  
fly/flies and 1 for virgin fly/flies. A male fly will not mate with a pregnant  
females, so you can think of this as a second level of controls. The two  
factors
```

```
are crossed, meaning that every possible combination of partners and type has  
outcomes measured.
```

Listing of simon-5501-12-fruitfly.qmd,

9

```
## Draw boxplot of longevity by partners and type

```{r boxplots-partners-type}
#| fig.width: 6
#| fig.height: 2.5
fly_subset |>
 ggplot(aes(factor(partners), longevity, fill=factor(type))) +
 geom_boxplot() +
 xlab("Number of partners") +
 ylab("Lifespan in days") +
 ggtitle("Graphs drawn by Steve Simon on 2024-10-28") +
 labs(fill="Female fly type") +
 scale_fill_discrete(labels=c("Pregnant", "Virgin")) +
 coord_flip()
````
```

Listing of simon-5501-12-fruitfly.qmd, 10

```
#### Interpretation of the output
```

```
The lifespans tend to be quite a bit shorter in cages with 8 virgin females.  
There are no problems with the homogeneity and normality assumptions.
```

Listing of simon-5501-12-fruitfly.qmd, 11

```
## Calculate average longevity by partners and type

```{r by-partners-and-type}
#| message: false
fly_subset |>
 group_by(type, partners) |>
 summarize(
 longevity_mn=mean(longevity),
 longevity_sd=sd(longevity),
 n=n()) -> fly_means
fly_means
```
```

Listing of simon-5501-12-fruitfly.qmd, 12

```
#### Interpretation of the output
```

```
The means show the same pattern as noted above. The standard deviations are  
all  
roughly the same.
```


Listing of simon-5501-12-fruitfly.qmd, 13

```
## Draw line graph

```{r line-graph}
fly_means |>
 ggplot(aes(
 factor(partners),
 longevity_mn,
 group=factor(type),
 color=factor(type))) +
 geom_line(linewidth=2) +
 expand_limits(y=range(fly_subset$longevity)) +
 xlab("Number of partners") +
 ylab("Lifespan in days") +
 labs(color="Female fly type") +
 scale_color_discrete(labels=c("Pregnant", "Virgin"))
````
```

Listing of simon-5501-12-fruitfly.qmd, 14

```
#### Interpretation of the output
```

This plot shows strong evidence of an interaction.

Male fruitflies do appear to have shorter average lifespans when sharing a cage

with virgin females compared to pregnant females, but this effect is far stronger when the number of female fruitflies is eight rather than one.

The line for pregnant (sharing a cage with pregnant females) is close to flat, indicating that there is little impact of the number of pregnant females on average longevity.

Listing of simon-5501-12-fruitfly.qmd, 15

```
## Analysis of variance including the interaction
```

```
```{r anova-with-interaction}  
m1 <- aov(longevity ~ factor(partners)*factor(type), data=fly_subset)
anova(m1)
```
```

```
#### Interpretation of the output
```

There is a statistically significant interaction between the number of female partners and the type of partners (pregnant or virgin). When you have an interaction, you do not normally interpret the individual categorical predictors.

Listing of simon-5501-12-fruitfly.qmd, 16

```
## Using lm, 1
```

```
```{r lm-with-interactions-1}  
m2 <- lm(longevity ~ factor(partners)*factor(type), data=fly_subset)
anova(m2)
```
```

```
#### Interpretation of the output
```

The analysis of variance table is identical when using lm instead of aov.

Listing of simon-5501-12-fruitfly.qmd, 17

```
## Using lm, 2
```

```
```{r lm-with-interactions-2}  
tidy(m2)
```
```

```
#### Interpretation of the output
```

The lifespan of fruitflies in cages with 8 virgin females is 17 days shorter than what you would expect. If the effect of type and partners were independent, you'd expect to see a 9.4 (= 8.0 + 1.4) day decline on average. The interaction shows that this cage is actually 16.6 days worse than that on average.

Listing of simon-5501-12-fruitfly.qmd, 18

```
## Using lm, 3

```{r add-coefficients}
b <- coef(m2)
fly_means |>
 mutate(b0=b[1]) |>
 mutate(b1=b[2]) |>
 mutate(b2=b[3]) |>
 mutate(b3=b[4]) |>
 select(-longevity_sd, -n) -> fly_coefficients
fly_coefficients[1, 5:7] <- 0
fly_coefficients[2, 6:7] <- 0
fly_coefficients[3, c(5,7)] <- 0
fly_coefficients
```
```

Listing of simon-5501-12-fruitfly.qmd, 19

```
#### Comments on the code
```

The `coef` function takes the intercept and slopes from a linear regression model and stores them in a vector. The `mutate` command adds the individual intercept and slopes to separate columns. Finally, place zeros in places where the various slopes are not needed.

Listing of simon-5501-12-fruitfly.qmd, 20

```
#### Interpretation of the output
```

This table is not needed for the data analysis, but is included to illustrate how the regression coefficients are computed in a two factor analysis of variance with an interaction.

The intercept is the estimated average with both categories each to the "zero" or reference level. In this case, the average lifespan of flies in a cage with one pregnant female.

The first slope coefficient (b1) represents the estimated average difference in lifespan between eight pregnant females and one pregnant females.

The second slope coefficient (b2) represents the estimated average difference

Listing of simon-5501-12-fruitfly.qmd, 21

```
## Save important files for later use
```

```
```{r save}  
save(
 fly,
 fly_subset,
 file=" ../data/fruitfly.RData")
```
```

```
#### Comments on the code
```

It is always good form to save the important pieces of your data analysis that might be re-used sometime later.

Break #6

- What you have learned
 - R code for interactions
- What's coming next
 - Your homework

Listing of simon-5501-12- directions.md, 1

```
---  
title: "Directions for 5501-12 programming assignment"  
---
```

This programming assignment was written by Steve Simon on 2024-10-08 and is placed in the public domain.

Listing of simon-5501-12- directions.md, 2

`## Program`

- Download the [program][tem]
 - Store it in your src folder
- Modify the file name
 - Use your last name instead of "simon"
- Modify the documentation header
 - Add your name to the author field
 - Optional: change the copyright statement

[tem]: <https://github.com/pmean/classes/blob/master/general/simon-5501-12-fruitfly.md>

Listing of simon-5501-12- directions.md, 3

`## Data`

- Download the `[data][dat]` file
 - Store it in your data folder
- Refer to the `[data dictionary][dic]`, if needed.

`[dat]: https://github.com/pmean/data/blob/main/files/fruitfly.txt`

`[dic]: https://github.com/pmean/data/blob/main/files/fruitfly.yaml`

Listing of simon-5501-12- directions.md, 4

`## Question 1`

Create a subset of the fruitfly data by removing the age where type equals 9. Draw a clustered boxplot with sleep as the outcome and partners and type as the categorical predictors. Interpret this graph. Is there evidence of non-normality?

`## Question 2`

Calculate descriptive statistics for sleep (mean, standard deviation, and sample size) by the combination of the two categorical predictors, partners and type. Is there evidence of heterogeneity?

Listing of simon-5501-12- directions.md, 5

`## Question 3`

`Draw a line graph for the mean sleep levels compared by type and partners. Is there evidence of an interaction?`

`## Question 4`

`Analyze the sleep variable using a two factor analysis of variance with an interaction. Present and interpret the analysis of variance table.`

Listing of simon-5501-12- directions.md, 6

`## Question 5`

`What factors might make you consider using a log transformation for the sleep variable? Do not run such an analysis but tell us whether you think the data would warrant such a transformation?`

Listing of simon-5501-12- directions.md, 7

`## Your submission`

- `- Save the output in html format`
- `- Convert it to pdf format.`
- `- Make sure that the pdf file includes`
 - `- Your last name`
 - `- The number of this course`
 - `- The number of this module`
- `- Upload the file`
- `- Please note the [policy on late submissions and rework][sim3].`

`[sim3]: https://github.com/pmean/classes/blob/master/general/policy-on-extensions-and-rework.md`

Listing of simon-5501-12- directions.md, 8

```
## If it doesn't work
```

```
Please review the [suggestions if you encounter an error page][sim4].
```

```
[sim4]: https://github.com/pmean/classes/blob/master/general/suggestions-if-you-encounter-an-error.md
```

Summary

- What you have learned
 - Two factor analysis of variance
 - Relationship to linear regression
 - Checking assumptions
 - R code for two factor analysis of variance
 - Interactions
 - R code for interactions
 - Your homework