# ANOVA for two factor experiments

#### **Session 6**

MATH 80667A: Experimental Design and Statistical Methods for Quantitative Research in Management HEC Montréal

#### Outline

Factorial designs and interactions

**Model formulation** 

Effect size, contrasts and power

# Factorial designs and interactions

## Example from the OSC psychology replication

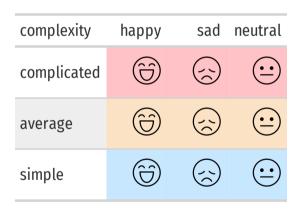
People can be influenced by the prior consideration of a numerical anchor when forming numerical judgments. [...] The anchor provides an initial starting point from which estimates are adjusted, and a large body of research demonstrates that adjustment is usually insufficient, leading estimates to be biased towards the initial anchor.

Replication of Study 4a of Janiszewski & Uy (2008, Psychological Science) by J. Chandler

#### Motivating example

Consider a study on the retention of information to children age 4 to which we read a story two hours after the reading.

We expect the ending (happy/sad/neutral) and the complexity (easy/average/hard) to impact their retention.



#### Why factorial designs?

To study the impact of story complexity and ending, we could run a series of one-way ANOVA.

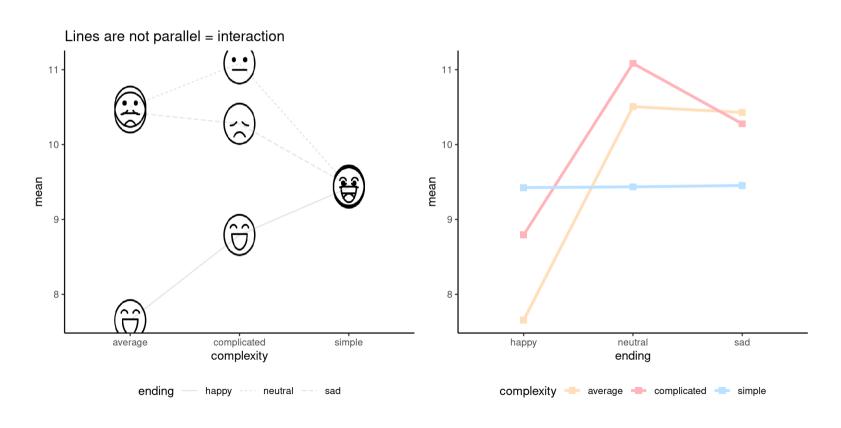
Factorial design is more efficient: can study the impact of multiple variables.

#### Estimates

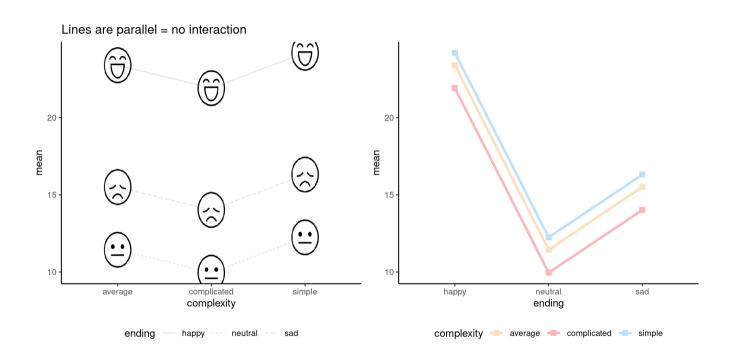
- Factorial design: study with multiple factors (subgroups)
- **simple effects**: difference between levels of one in a fixed combination of others (change in difficulty for happy ending)
- **main effects**: differences relative to average for each condition of a factor (happy vs neutral vs sad ending)
- **interaction effects**: when simple effects differ depending on levels of another factor

#### Interaction

An interaction is present when the effect of one factor depends on the levels of another factor.



#### Lack of interaction



In practice, the sample average are uncertain!

ullet Plot averages with confidence intervals or  $\pm 1$  standard error.

# Model formulation

#### Formulation of the two-way ANOVA

Two factors: A (complexity) and B (ending) with a and b levels.

Write the average response  $Y_{ijk}$  of the kth measurement in the group  $(A_i,B_j)$  as

$$Y_{ijk} = \mu_{ij} + arepsilon_{ijk}$$

#### where

- ullet  $Y_{ijk}$  is the kth replicate for ith level of factor A and jth level of factor B
- $\varepsilon_{ijk}$  are independent error terms with mean zero and variance  $\sigma^2$ .

Two-way ANOVA model with interaction: one average for each subgroup

#### Hypothesis tests

#### Interaction between factors $\boldsymbol{A}$ and $\boldsymbol{B}$

 $\mathscr{H}_0$ : no interaction between factors A and B vs  $\mathscr{H}_a$ : there is an interaction

Main effect of factor A

 $\mathscr{H}_0$ :  $\mu_{1.}=\cdots=\mu_{a.}$  vs  $\mathscr{H}_a$ : at least two marginal means of A are different

Main effect of factor B

 $\mathscr{H}_0$ :  $\mu_{.1} = \cdots = \mu_{.b}$  vs  $\mathscr{H}_a$ : at least two marginal means of B are different

#### Reparametrization

• Mean of  $A_i$  (average of row i):

$$\mu_{i.} = rac{\mu_{i1} + \cdots + \mu_{ib}}{b}$$

• Mean of  $B_j$  (average of column j):

$$\mu_{.j} = rac{\mu_{1j} + \cdots + \mu_{aj}}{a}$$

• Overall average:

$$\mu = rac{\sum_{i=1}^a \sum_{j=1}^b \mu_{ij}}{ab}$$

#### Formulation of the two-way ANOVA

Write the model for a response variable Y in terms of two factors  $A_i, B_j$ .

$$Y_{ijk} = \mu + \alpha_i + \beta_j + (\alpha \beta)_{ij} + \varepsilon_{ijk}$$

with the parameters in the sum-to-zero constraints

- $\alpha_i = \mu_{i.} \mu_{i.}$ 
  - $\circ$  mean of level  $A_i$  minus overall mean.
- $\beta_j = \mu_{.j} \mu_{.j}$ 
  - $\circ$  mean of level  $B_i$  minus overall mean.
- $\bullet \ (\alpha\beta)_{ij} = \mu_{ij} \mu_{i.} \mu_{.j} + \mu$ 
  - $\circ$  the interaction term for  $A_i$  and  $B_j$ .

## Breaking down the variability

For a **balanced** design, we can decompose the variability around sample means:

$$\begin{aligned} \mathsf{SS}_{ ext{total}} &= \mathsf{SS}_{ ext{model}} + \mathsf{SS}_{ ext{res}} \ &= \mathsf{SS}_A + \mathsf{SS}_B + \mathsf{SS}_{AB} + \mathsf{SS}_{ ext{res}} \end{aligned}$$

**Board work** 

## Analysis of variance table

term	degrees of freedom	mean square	F
A	a-1	$MS_A = SS_A/(a-1)$	$MS_A/MS_{\mathrm{res}}$
B	b-1	$MS_B = SS_B/(b-1)$	$MS_B/MS_{\mathrm{res}}$
AB	(a-1)(b-1)	$MS_{AB} = SS_{AB}/\{(a-1)(b-1)\}$	$MS_{AB}/MS_{\mathrm{res}}$
residuals	n-ab	$MS_{\mathrm{res}} = SS_{\mathrm{res}}/(n-ab)$	
total	n-1		

Factors are crossed, replicates are nested within AB groups

#### Example: Fiber strength

We consider a 5 imes 3 factorial design (no interaction term).

Consider five levels of application of potash

•  $T_1=36$ ,  $T_2=54$ ,  $T_3=72$ ,  $T_4=108$  and  $T_5=144$  lb K $_2$ O per acre, applied to a cotton crop.

The response is a measure of single-fiber strength, an average of a number of tests on the cotton from each plot.

There were three blocks each containing five plots.

# Data on fiber strength

#### Reordered data

block	<b>T1</b>	<b>T2</b>	<b>T3</b>	<b>T4</b>	<b>T5</b>
block 1	7.62	8.14	7.76	7.17	7.46
block 2	8.00	8.15	7.73	7.57	7.68
block 3	7.93	7.87	7.74	7.80	7.21

#### Residuals

block	<b>T1</b>	<b>T2</b>	<b>T3</b>	T4	<b>T5</b>
block 1	-0.14	0.18	0.11	-0.25	0.10
block 2	0.05	-0.01	-0.12	-0.05	0.13
block 3	0.09	-0.17	0.01	0.30	-0.23

# Analysis of variance table

	sum of squares	df	F	p-value
treatment	0.732	4	4.192	0.040
block	0.097	2	1.112	0.375
Residuals	0.349	8		

## Some pending questions

- Intuition behind degrees of freedom for the residuals?
- No interaction term (why?)
- Why blocking?

$A \setminus B$	$b_1$	$b_2$	$b_3$	$b_4$	$b_5$	sum
$a_1$	$AB_{11}$	$AB_{12}$	$AB_{13}$	$AB_{14}$	X	$A_1$
$a_2$	$AB_{21}$	$AB_{22}$	$AB_{23}$	$AB_{24}$	X	$A_2$
$a_3$	X	X	X	X	X	X
sum	$B_1$	$B_2$	$B_3$	$B_4$	X	total

Terms with X are fully determined by row/column/total averages

## Example from Keppel and Wichern (table)

Consider errors by monkeys under three drug conditions (  ${\cal A}$  ) and two degrees of food deprivation (  ${\cal B}$  )

Data for the  $3 \times 2$  factorial design

$A \setminus B$	<b>1h deprivation</b>	<b>24h deprivation</b>
Control	1, 4, 0, 7	15, 6, 10, 13
Drug 1	13, 5, 7, 15	6, 18, 9, 15
Drug 2	9, 16, 18, 13	14, 7, 6, 13

**R** Demonstration

# Effect size, contrasts and power

#### Noncentrality parameters

Consider a balanced design with n=a imes b imes k observations

For the mean squared errors, the expected values are

$$\mathsf{E}(\mathsf{MS}_A) = \sigma^2 + rac{bn}{a-1} \sum_{i=1}^a lpha_i^2$$

$$\mathsf{E}(\mathsf{MS}_{AB}) = \sigma^2 + rac{n}{(a-1)(b-1)} \sum_{j=1}^b \sum_{i=1}^a (lpha eta)_{ij}^2$$

Under the null hypothesis of no mean effect / interaction, these are thus unbiased estimators of the error variance.

#### Effect size

- We can report an estimate of the effect size for either of the main effects, for the interaction or overall.
- For a power calculation, do the calculations with each effect (whose size is of scientific interest and select the largest sample size per group.
- Given F statistics and degrees of freedom, we can find different measures: Cohen's f, (partial)  $\eta^2$  and  $\omega^2$ .
- Check the book for formulae and effectsize for estimates in R.
- Estimators of variability are noisy and biased

## Omega-squared and partial version

The proportion of variance explained by the effect T is

$$\omega_{ ext{effect}}^2 = rac{\sigma_{ ext{effect}}^2}{\sigma_{ ext{total}}^2}.$$

where 
$$\sigma_{ ext{total}}^2 = \sigma_A^2 + \sigma_B^2 + \sigma_{AB}^2 + \sigma_{ ext{error}}^2$$

The partial  $\omega^2$  for an effect is

$$\omega_{
m \langle effect
angle}^2 = rac{\sigma_{
m effect}^2}{\sigma_{
m effect}^2 + \sigma_{
m resid}^2}.$$

These get replaced by estimates based on F statistics and degrees of freedom (see Keppel & Wickens, p. 233).

The R package effectsize reports estimates with confidence intervals

Reference: Steiger (2004), Psychological Methods

#### One way for the two-way

We can cast the two-way model with an interaction as a one-way ANOVA with ab levels.

- Sometimes useful for using custom contrasts
- Used for some procedures that do not support two-way designs (unequal variance model) or Levene's test

#### Contrasts for the main effects

In the interaction model, we cast the main effect in terms of parameters Say the order of the coefficients is drug ( A ) and deprivation ( B ).

test	$\mu_{11}$	$\mu_{21}$	$\mu_{31}$	$\mu_{21}$	$\mu_{22}$	$\mu_{23}$
main effect $A$ (1 vs 2)	1	-1	0	1	-1	0
main effect $A$ (1 vs 3)	1	0	-1	1	0	-1
main effect $B$ (1 vs 2)	1	1	1	-1	-1	-1
interaction $AB$ (1 vs 2)	-1	2	-1	1	-2	1

## Testing hypothesis of interest

We only tests hypothesis that are of interest

- If there is a significant interaction, the marginal means are not of interest
- Rather, compute the simple effects.
- What is the number of hypothesis of interest? Often, this is pairwise comparisons within each level of the other factor
  - $\circ$  much less than  $\binom{ab}{2}$  pairwise comparisons
- Scheffé's method for all custom contrasts still applicable, but may be conservative
- Tukey's method also continues to hold (or generalization thereof)
- Omnibus procedures for controlling the FWER (Holm-Bonferroni) may be more powerful.