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

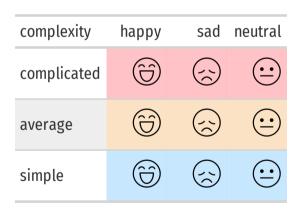
Effect size, contrasts and power

Factorial designs and interactions

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.

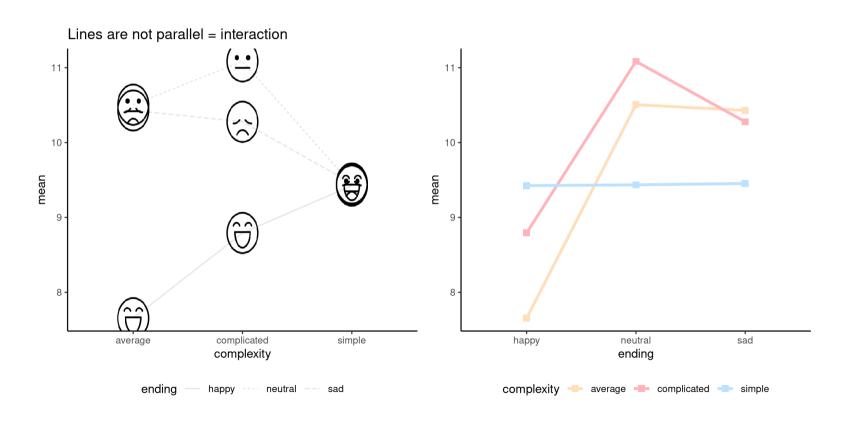
Factorial designs are more efficient: can study the impact of multiple variables simultaneously with **fewer overall observations**.

Estimates

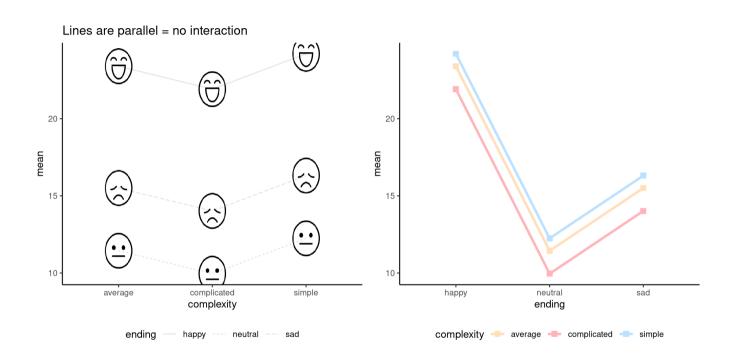
- Factorial design: study with multiple factors (subgroups)
- **simple effects**: difference between levels of one in a fixed combination of others (change in difficulty for happy ending)
- **main effects**: differences relative to average for each condition of a factor (happy vs neutral vs sad ending)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

• Plot averages with confidence intervals or ± 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

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

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

Hypothesis tests

Interaction between factors A and B

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

Main effect of factor A

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

Main effect of factor B

 \mathcal{H}_0 : $\mu_{.1} = \cdots = \mu_{.b}$ vs \mathcal{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):

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

• 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 .

$$Y_{ijk} = \mu + lpha_i + eta_j + (lphaeta)_{ij} + arepsilon_{ijk}$$

with the parameters in the sum-to-zero constraints

- ullet $lpha_i = \mu_{i.} \mu_{i.}$
 - \circ mean of level A_i minus overall mean.
- $\bullet \quad \beta_j = \mu_{.j} \mu$
 - \circ mean of level B_j minus overall mean.
- $ullet \qquad (lphaeta)_{ij} = \mu_{ij} \mu_{i.} \mu_{.j} + \mu_{.j}$
 - \circ the interaction term for A_i and B_j .

Sum-to-zero parametrization

The model in terms of α , β and $(\alpha\beta)$ is overparametrized.

For the sum-to-zero constraint, impose that

$$\sum_{i=1}^a lpha_i = 0, \quad \sum_{j=1}^b eta_j = 0, \quad \sum_{j=1}^b (lphaeta)_{ij} = 0, \quad \sum_{i=1}^a (lphaeta)_{ij} = 0.$$

which imposes 1 + a + b constraints.

Analysis of variance table

term	degrees of freedom	mean square	F
A	a-1	$MS_A = SS_A/(a-1)$	MS_A/MS_{res}
B	b-1	$MS_B = SS_B/(b-1)$	MS_B/MS_{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 5×3 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 K₂O 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	T1	T2	T3	T4	T5
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

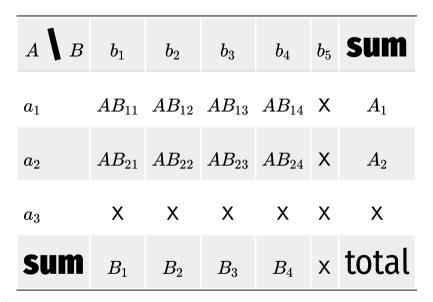
block	T1	T2	T3	T4	T5
block 1	-0.14	0.18	0.11	-0.25	0.10
block 2	0.05	-0.01	-0.12	-0.05	0.13
block 3	0.09	-0.17	0.01	0.30	-0.23

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?)



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

Effect size, contrasts and power

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.

Breaking down the variability

We can express the overall variability of the response around the global mean as

$$\sigma_{
m total}^2 = \sigma_A^2 + \sigma_B^2 + \sigma_{AB}^2 + \sigma_{
m resid}^2.$$

where $\sigma_A^2 = a^{-1} \sum_{i=1}^a \alpha_i^2$, $\sigma_{AB}^2 = (ab)^{-1} \sum_{i=1}^a \sum_{j=1}^b (\alpha \beta)_{ij}^2$, etc.

Partial Cohen's f

The **population** version of Cohen's partial *f* measures the proportion of variability that is explained by a main effect or an interaction, e.g.,

$$f_{\langle A
angle} = rac{\sigma_A^2}{\sigma_{
m resid}^2}, \qquad f_{\langle AB
angle} = rac{\sigma_{AB}^2}{\sigma_{
m resid}^2}.$$

These variance quantities are **unknown**, so need to be estimated somehow.

Partial effect size (variance)

Effect size are often reported in terms of variability via the ratio

$$rac{\sigma_{ ext{effect}}^2}{\sigma_{ ext{effect}}^2 + \sigma_{ ext{resid}}^2}.$$

- Both $\hat{\eta}_{\langle \text{effect} \rangle}^2$ and $\hat{\omega}_{\langle \text{effect} \rangle}^2$ are estimators of this quantity and obtained from the $_F$ statistic and degrees of freedom of the effect.
- $\widehat{\omega}^2_{\langle \mathrm{effect} \rangle}$ is less biased than $\eta^2_{\langle \mathrm{effect} \rangle}$.

Estimation of partial ω^2

$$\widehat{\omega}_{\langle ext{effect}
angle}^2 = rac{ ext{df}_{ ext{effect}}(F_{ ext{effect}}-1)}{ ext{df}_{ ext{effect}}(F_{ ext{effect}}-1)+n},$$

where *n* is the overall sample size.

In **R**, effectsize::omega_squared reports these estimates with one-sided confidence intervals.

Reference for confidence intervals: Steiger (2004), Psychological Methods

Converting ω^2 to Cohen's f

Given the estimated $\widehat{\omega}^2_{\langle \text{effect} \rangle}$, convert it into an estimate of Cohen's partial $f^2_{\langle \text{effect} \rangle}$ via

$${\widehat f}_{\, \langle {
m effect}
angle}^2 = rac{{\widehat \omega}_{\langle {
m effect}}^2
angle}{1 - {\widehat \omega}_{\langle {
m effect}}^2
angle}.$$

Note that effectsize::cohens_f returns $ilde{f}^2=n^{-1}F_{ ext{effect}} ext{df}_{ ext{effect}}$, a transformation of $\hat{\eta}^2_{\langle ext{effect} \rangle}$.

Power for factorial experiments

- G^*Power and **R** packages takee Cohen's f (or f^2) as input.
- Calculation based on F distribution with
 - \circ $\nu_1 = \mathrm{df}_{\mathrm{effect}}$ degrees of freedom
 - \circ $\nu_2 = n n_g$, where n_g is the number of mean parameters estimated.
 - \circ noncentrality parameter $\phi = Nf_{\langle ext{effect}
 angle}^2$

In **R**, if a and b denote the number of levels of each factor and omega. sq is $\omega_{\langle AB \rangle}^2$, then the sample size needed to detect this effect with power of 80% is

```
fhat <- sqrt(omega.sq/(1-omega.sq))
WebPower::wp.kanova(power = 0.8, f = fhat, ndf = (a-1)*(b-1), ng = ab)
```

Illustration of the bias

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 ($_A$, 3 levels) and deprivation ($_B$, 2 levels).

test	μ_{11}	μ_{12}	μ_{21}	μ_{22}	μ_{31}	μ_{32}
main effect A (1 vs 2)	1	1	-1	-1	0	0
main effect A (1 vs 3)	1	1	0	0	-1	-1
main effect $_B$ (1 vs 2)		-1	1	-1	1	-1
interaction $_{AB}$ (1 vs 2, 1 vs 2)	1	-1	-1	1	0	0
interaction AB (1 vs 3, 1 vs 2)	1	-1	0	0	-1	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
 - much less than (ab) 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.