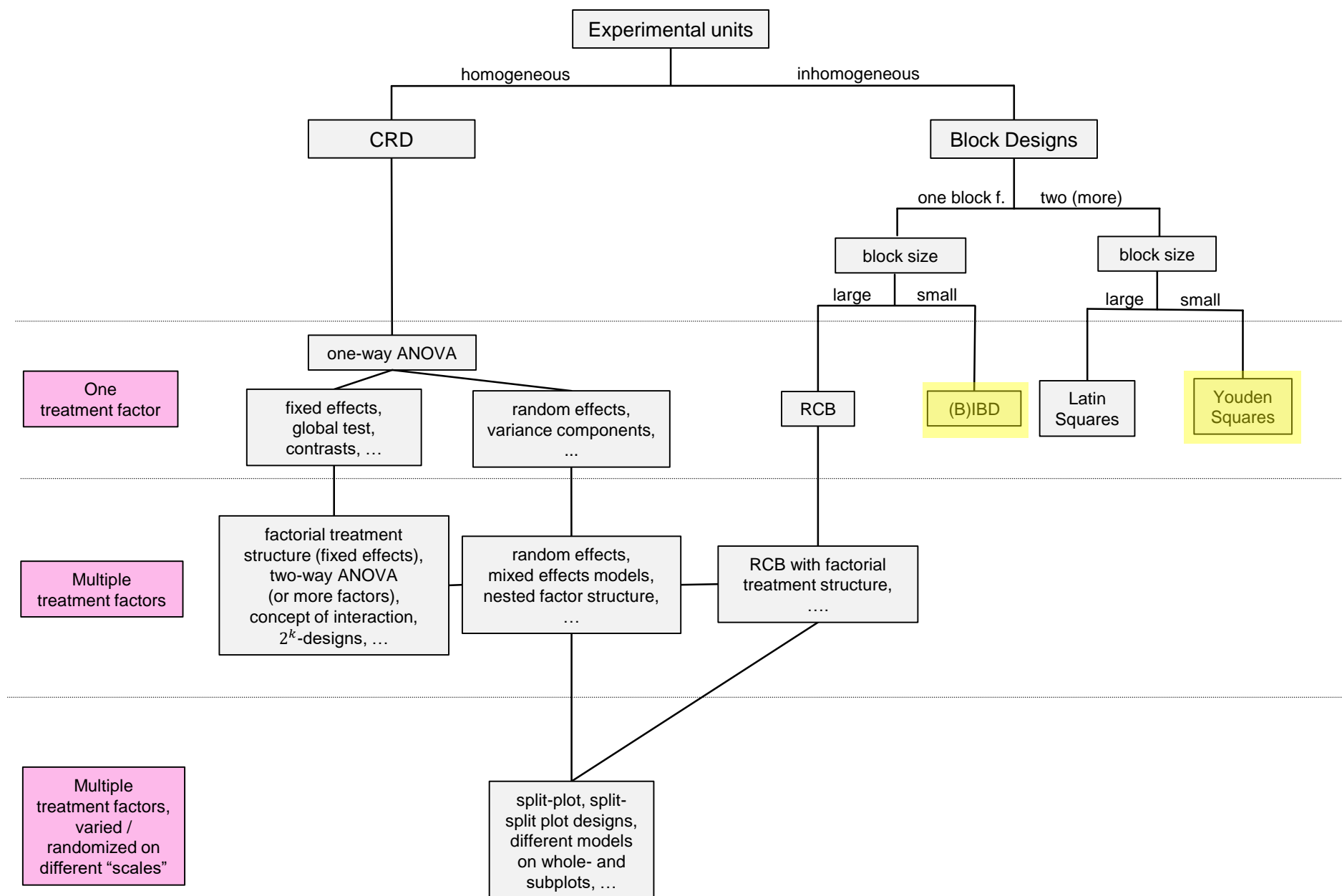


11



Incomplete Block Designs



Incomplete Block Designs

- Up to now we only considered **complete** block designs.
- This means we see **all** treatments in **each** block.
- In some situations this is **not** possible because
 - (physical) block size too small
 - too expensive
 - not advisable (think of raters having to rate 7 or more champagne brands each)
- Remember the eye drop example? What if we wanted to test **3** different eye drop types?
- It is still a good idea to **block** on subjects, but obviously it is **not** possible to have **complete** blocks in this example!



Example: Eye Drops (Oehlert, 2010)

- Suppose we have 3 subjects getting the treatments A, B, C :

| Subject 1 | Subject 2 | Subject 3 |
|-----------|-----------|-----------|
| A | A | B |
| B | C | C |

- This is a so-called **incomplete** block design.
- If we want to estimate the difference between A and B , we can use:
 - Subject 1: The estimate has variance $2\sigma^2$.
 - Combine subject 2 and subject 3:

$$A - B = (A - C) - (B - C)$$

This difference of differences has variance $2\sigma^2 + 2\sigma^2 = 4\sigma^2$.

- In a **complete** block design, we could estimate the difference in **each** block with the **same precision**.

Incomplete Block Designs

- We have to be careful about which treatment pairs we put together in the **same** block.
- We call a design **disconnected** if we can build two groups of treatments such that it **never** happens that we see members of both groups together in the **same block**.

| 1 | 2 | 3 | 4 | 5 | 6 |
|----------|----------|----------|----------|----------|----------|
| <i>A</i> | <i>A</i> | <i>B</i> | <i>D</i> | <i>D</i> | <i>E</i> |
| <i>B</i> | <i>C</i> | <i>C</i> | <i>E</i> | <i>F</i> | <i>F</i> |

- In a **disconnected** design, it is **not** possible to estimate all treatment differences!
- If the design is **not** disconnected, we call it **connected**.



Balanced Incomplete Block Designs (BIBDs)

- We call an incomplete block design **balanced (BIBD)** if all treatments **pairs** occur together in the same block **equally often** (we denote this number by λ).
- What is the benefit of the “balancedness” property?
- The **precision** (variance) of the estimated treatment differences $\alpha_i - \alpha_j$ is the **same** no matter what combination of i and j we are considering.
- This means that we can estimate all treatment differences with the **same accuracy**.
- Let us first give an overview of the different numbers involved in such a problem.

Balanced Incomplete Block Designs (BIBDs)

- We use the following notation:
 - g : number of treatments
 - b : number of blocks
 - k : number of units per block with $k < g$
 - r : number of replicates per treatment
 - N : total number of units
- In the eye drop example we had
 - $g = 3$ treatments (the different eye drops: A, B, C)
 - $b = 3$ blocks (the 3 subjects)
 - $k = 2$ units per block (the 2 eyes per subject)
 - $r = 2$ replicates per treatment
 - $N = 6$
- Of course it must hold that $N = b \cdot k = g \cdot r$.

Unreduced BIBDs

- We can always find a BIBD for every setting of $k < g$.
- How? Simply use **all** possible combinations.
- The number of combinations is $\binom{g}{k}$ (= binomial coefficient: $\frac{g!}{k!(g-k)!}$).
- E.g., for $g = 7$ and $k = 3$ we have $\binom{7}{3} = 35$.
- In R, have a look at the functions `choose` and `combn`.
- We call such a design an **unreduced** balanced incomplete block design.
- In practice, it is often **not** possible to have so many blocks.
- The big question: What number of blocks is “doable”?

Balanced Incomplete Block Designs (BIBDs)

- A treatment occurs in r blocks.
- There are $k - 1$ other “available units” in each of these blocks which makes a total of $r \cdot (k - 1)$ “available units”.
- The remaining $g - 1$ treatments must be divided **evenly** among them, otherwise the design is **not** balanced.
- Hence, $\frac{r \cdot (k - 1)}{g - 1}$ **must** be a **whole number** ($= \lambda$) for a BIBD to exist.
- This condition is only **necessary, not sufficient**.
- This means: Even if the condition is fulfilled, it might be the case that you **cannot** find a BIBD!

Example: Champagne (Roth, 2013)



- 14 raters, 7 champagne types, every rater rated 3 of them.

| 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 |
|---|---|---|---|---|---|---|---|---|----|----|----|----|----|
| 2 | 1 | 2 | 2 | 1 | 3 | 1 | 1 | 3 | 3 | 1 | 1 | 4 | 2 |
| 6 | 3 | 6 | 4 | 2 | 5 | 4 | 2 | 4 | 5 | 4 | 5 | 5 | 3 |
| 7 | 6 | 7 | 5 | 3 | 7 | 7 | 5 | 6 | 7 | 7 | 6 | 6 | 4 |

- This is a BIBD. We see every treatment pair combination exactly **twice** in the same block.
- In more detail, we have
 - $g = 7$ treatments
 - $b = 14$ blocks
 - $k = 3$ units per block
 - $r = 6$ replicates per treatment.
- Hence, $\lambda = \frac{r \cdot (k-1)}{g-1} = \frac{6 \cdot 2}{6} = 2$.

BIBD: Finding a Design

- First, make sure that the necessary condition is fulfilled.
- Old way: Check Appendix C.2 of the book with a list of BIBDs.
- Use R, e.g. function `find.BIB` in package `crossdes` or function `(b) ibd` in package `ibd` (among many others).
- See R-File for an example.

(B)IBD: Randomization

- How can we randomize a given (B)IBD?
- Randomize blocks to the groups of treatment letters.
- Within each block: Randomize assignment of treatment letters to physical units.
- Randomize assignment of treatment letters to actual treatments.
- How can we analyze an incomplete block design?

(B)IBD: Analysis

- The **model** for a (balanced) incomplete block design is the standard model, i.e.

$$Y_{ij} = \mu + \alpha_i + \beta_j + \varepsilon_{ij}$$

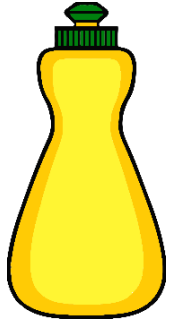
The diagram shows the equation $Y_{ij} = \mu + \alpha_i + \beta_j + \varepsilon_{ij}$. Below the equation, there are two boxes. The left box is labeled "effect of treatment" and has an arrow pointing to α_i . The right box is labeled "effect of block" and has an arrow pointing to β_j .

- However, as we don't observe all treatment \times block combinations, the “usual” estimates are **not** working and we need the computer to find the least squares estimates (which is no problem).
- We are using Type III sum of squares to test treatment effects **adjusted for block effects**.
- In other words: We analyze the treatment effects while **controlling for the block effects**.

Intra- and Interblock Analysis

- This is a so called **intrablock analysis** of the (B)IBD.
- It is also possible to recover some information by comparing different blocks.
- This would be called an **interblock analysis**.
- Information from both approaches can be suitably combined.
- This looks complicated in the book, but it is nothing else than the analysis when treating the block factor as **random**.
- We will **not** discuss this any further here.

Example: Dish Detergent (Oehlert, 2010, Ex. 14.2)



- Want to compare **9 different dishwashing solutions**.

| <i>Treatment</i> | <i>A</i> | <i>B</i> | <i>C</i> | <i>D</i> | <i>E</i> | <i>F</i> | <i>G</i> | <i>H</i> | <i>J</i> |
|-----------------------|----------|----------|----------|----------|-----------|-----------|-----------|-----------|----------|
| <i>Base detergent</i> | <i>I</i> | <i>I</i> | <i>I</i> | <i>I</i> | <i>II</i> | <i>II</i> | <i>II</i> | <i>II</i> | control |
| <i>Additive</i> | 3 | 2 | 1 | 0 | 3 | 2 | 1 | 0 | control |

- Available resources:
 - 3 washing basins
 - 1 operator for each basin (= 3 operators)
- The 3 operators wash at the same speed during each session, but the **speed might vary from session to session**.
- Response: **Number of plates washed** when foam disappears.

Example: Dish Detergent (Oehlert, 2010, Ex. 14.2)

- If we have 12 sessions, we can find a BIBD.
- The design was as follows:

| 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 |
|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|
| <i>A</i> | <i>D</i> | <i>G</i> | <i>A</i> | <i>B</i> | <i>C</i> | <i>A</i> | <i>B</i> | <i>C</i> | <i>A</i> | <i>B</i> | <i>C</i> |
| <i>B</i> | <i>E</i> | <i>H</i> | <i>D</i> | <i>E</i> | <i>F</i> | <i>E</i> | <i>F</i> | <i>D</i> | <i>F</i> | <i>D</i> | <i>E</i> |
| <i>C</i> | <i>F</i> | <i>J</i> | <i>G</i> | <i>H</i> | <i>J</i> | <i>J</i> | <i>G</i> | <i>H</i> | <i>H</i> | <i>J</i> | <i>G</i> |

- Analysis in R:

```
> fit <- aov(dishes ~ session + detergent, data = dish)
```

```
> drop1(fit, test = "F")
```

Single term deletions

Model:

dishes ~ session + detergent

| | Df | Sum of Sq | RSS | AIC | F value | Pr(>F) |
|-----------|----|-----------|---------|---------|----------|---------------|
| <none> | | | 13.19 | 3.841 | | |
| session | 11 | 10.06 | 23.25 | 2.260 | 1.1103 | 0.4127 |
| detergent | 8 | 1086.81 | 1100.00 | 147.104 | 164.8539 | 6.809e-14 *** |

Example: Dish Detergent (Oehlert, 2010, Ex. 14.2)

- If we call `summary.lm`, we get

```
> summary.lm(fit)
```

```
Call:
```

```
aov(formula = dishes ~ session + detergent, data = dis
```

```
Residuals:
```

```
      Min       1Q   Median       3Q      Max 
-1.1482 -0.5556  0.1111  0.4630  1.0000
```

```
Coefficients:
```

| | Estimate | Std. Error | t value | Pr(> t) | |
|-------------|----------|------------|---------|----------|-----|
| (Intercept) | 18.7037 | 0.6766 | 27.643 | 6.21e-15 | *** |
| session10 | 1.4074 | 0.8194 | 1.718 | 0.105170 | |
| session11 | 0.6296 | 0.8194 | 0.768 | 0.453458 | |
| session12 | 0.8519 | 0.8194 | 1.040 | 0.313998 | |
| session2 | 1.1111 | 0.8559 | 1.298 | 0.212612 | |
| session3 | 0.4444 | 0.8559 | 0.519 | 0.610667 | |
| session4 | 0.9259 | 0.8194 | 1.130 | 0.275148 | |
| session5 | 1.1481 | 0.8194 | 1.401 | 0.180266 | |
| session6 | 2.1481 | 0.8194 | 2.622 | 0.018513 | * |
| session7 | 1.8519 | 0.8194 | 2.260 | 0.038127 | * |
| session8 | 0.6296 | 0.8194 | 0.768 | 0.453458 | |
| session9 | 1.4074 | 0.8194 | 1.718 | 0.105170 | |
| detergent2 | -2.5556 | 0.7412 | -3.448 | 0.003309 | ** |
| detergent3 | -6.5556 | 0.7412 | -8.844 | 1.47e-07 | *** |
| detergent4 | -13.2222 | 0.7412 | -17.839 | 5.54e-12 | *** |
| detergent5 | 5.5556 | 0.7412 | 7.495 | 1.28e-06 | *** |
| detergent6 | 3.2222 | 0.7412 | 4.347 | 0.000499 | *** |
| detergent7 | 1.3333 | 0.7412 | 1.799 | 0.090928 | . |
| detergent8 | -0.5556 | 0.7412 | -0.750 | 0.464416 | |
| detergent9 | 9.7778 | 0.7412 | 13.192 | 5.16e-10 | *** |

Here we used `contr.treatment`. The coefficients are therefore comparisons to the **reference treatment** (= detergent 1). Note that the standard error is the same for all effects which is a property of the balanced design.

Partially Balanced Incomplete Block Designs

- It might very well be the case that we are in a situation where there is **no** BIBD available.
- In that case we could use a **partially balanced incomplete block design**, where some treatment pairs occur together more often than other pairs.
- Example (Kuehl, 2000, Display 9.3)

| <i>Block 1</i> | <i>Block 2</i> | <i>Block 3</i> |
|----------------|----------------|----------------|
| 1 | 2 | 3 |
| 4 | 5 | 6 |
| 2 | 3 | 1 |
| 5 | 6 | 4 |

- (1,4), (2,5), (3,6) are observed **twice**, remaining pairs only **once** together in the same block.
- The analysis is the same as for a BIBD!

Row-Column Incomplete Block Designs

- As we have seen with RCBs, we are sometimes facing the situation where we have **more than one** block factor (remember Latin Squares?).
- Latin Squares are often impractical due to their very strict constraint on the design.
- A **row-column incomplete block design** is a design where we block on rows **and** columns and one or both of them are **incomplete blocks**.

Example: Car Tires (Kuehl, 2000)

- Suppose we want to evaluate 7 treatments instead of 4.
- Assume that we have 7 cars and the following design:

| <i>Tire position</i> |  |  |  |  |  |  |  |
|-------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------|-----------------------------------------------------------------------------------|------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------|
|  | 3 | 4 | 5 | 6 | 7 | 1 | 2 |
|  | 5 | 6 | 7 | 1 | 2 | 3 | 4 |
|  | 6 | 7 | 1 | 2 | 3 | 4 | 5 |
|  | 7 | 1 | 2 | 3 | 4 | 5 | 6 |

- The tire positions are **complete blocks**, the columns form a **BIBD**. This is a so called **row-orthogonal design**.

Youden Squares

- A **Youden Square** is **rectangular** (!) such that
 - columns (rows) form a **BIBD**
 - rows (columns): each treatment appears **equally often** in each row (column)
- Hence, columns form a **BIBD**, rows an **RCB**.
- The model is as before:

$$Y_{ijk} = \mu + \alpha_i + \beta_j + \gamma_k + \epsilon_{ijk}$$

Diagram illustrating the model components and their corresponding experimental factors:

- α_i corresponds to **treatment**
- β_j corresponds to **Block factor 1 (rows)**
- γ_k corresponds to **Block factor 2 (columns)**

- Analysis in R “as usual”, just make sure to use `drop1` to ensure that the correct sum of squares is being used.

Example: Lithium in Blood (Oehlert, 2010, Ex. 14.5)



- Study was performed to measure **blood concentration of lithium** 12 hours after administering lithium carbonate using
 - *A*: 300mg capsule
 - *B*: 250mg capsule
 - *C*: 450mg time delay capsule
 - *D*: 300mg solution
- 12 subjects, each will be measured and treated **twice**, one week apart:

| Week | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 |
|------|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|----------|
| 1 | <i>A</i> | <i>D</i> | <i>C</i> | <i>B</i> | <i>D</i> | <i>D</i> | <i>B</i> | <i>B</i> | <i>C</i> | <i>A</i> | <i>A</i> | <i>C</i> |
| 2 | <i>B</i> | <i>C</i> | <i>A</i> | <i>C</i> | <i>A</i> | <i>B</i> | <i>A</i> | <i>D</i> | <i>D</i> | <i>D</i> | <i>C</i> | <i>B</i> |

- **Response:** Serum lithium level.

Example: Lithium in Blood (Oehlert, 2010, Ex. 14.5)

- We block on **both** rows (weeks) and columns (subjects).
- Every treatment appears 3 times in each week.
- The columns form a BIBD.

- Analysis in R:

```
> fit <- aov(hour.12 ~ subject + period + treatment, data = lith)
> drop1(fit, test = "F")
single term deletions
```

Model:

```
hour.12 ~ subject + period + treatment
```

| | Df | Sum of Sq | RSS | AIC | F value | Pr(>F) |
|-----------|----|-----------|----------|---------|---------|-----------|
| <none> | | | 0.016203 | -143.22 | | |
| subject | 11 | 0.029946 | 0.046149 | -140.09 | 1.3442 | 0.3449 |
| period | 1 | 0.031974 | 0.048177 | -119.06 | 15.7871 | 0.0041 ** |
| treatment | 3 | 0.005603 | 0.021806 | -142.09 | 0.9222 | 0.4728 |

- Unfortunately, we **cannot** detect any treatment effect here.



Summary

- Balancedness properties etc. ensure that we are performing the experiment as **efficient as possible**.
- If a design is **not** balanced anymore, we lose efficiency but **we can typically still analyze the data**.
- Exceptions are (e.g.) cases where a disconnected design has been used and the focus was on comparing all treatments.
- Package overview: <https://cran.r-project.org/web/views/ExperimentalDesign.html>

