LGN 5822 - Biometrical Genetics

### L08 – Randomized Complete Block Design

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"We shall need to judge of the magnitude of the differences introduced by testing our treatments upon the **different plots** by the discrepancies between the performances of the same treatment in **different blocks**"

> R.A Fisher, 1935 The Design of Experiments, Section 26



 If the researcher finds that any factor disturbs the homogeneity of the experimental units or environmental conditions



Controlled environmental conditions

How can we assure, then, that an observed agronomic difference is the result of a specific treatment, rather than the result of the experimental units to which it was allocated?

In other words, how do we prevent our treatment results from being confounded with our experimental units?



The **heterogeneity** of experimental units presents a problem:

Difficulty recognizing differences between experimental units!

 This could lead us to conclude that the differences in our variables are the result of the treatments applied, when in fact they were caused by the pre existing condition of the experimental units

### Randomized Complete Block Design (RCBD)

How to control undesirable variation between experimental units?



Minimize Field Differences

#### Randomized Complete Block Design (RCBD)

 The first step in using the RCBD is to recognize the source(s) of potential heterogeneity among plots (experimental units)



"Production gradient"

### How to block

- In blocking, we generally place an equal-sized block on every map unit. Each block, in this case, contains four experimental units (plots)
- Each treatment is applied to one experimental unit within the block



Blocking of randomized treatments to account for a known yield potential gradient

#### Randomized Complete Block Design (RCBD)



### Randomized Complete Block Design (RCBD)

 RCBD is the standard design for agricultural experiments where similar experimental units are grouped into blocks or replicates

• A **RCBD** is the most basic blocking design

RCDB uses the basic principles of repetition, randomization and local control

#### Randomized Complete Block Design (RCBD)

 It is used to control for variation in an experiment not predicted by the researcher (random error)

• e.g. variation in fertility or drainage differences in a field

#### Randomized Complete Block Design (RCBD)

• Assume we have r blocks containing g units each



Here, r = 3 blocks with g = 4 units

In every of the r blocks we randomly assign the g treatments to the g units, **independently** of the other blocks

#### Description of the Design: RCBD

- Probably the most used and useful of the experimental designs
- Takes advantage of grouping similar experimental units into blocks or replicates
- The blocks of experimental units should be as uniform as possible
- The purpose of grouping experimental units is to have the units in a block as uniform as possible so that the observed differences between treatments will be largely due to "true" differences between treatments
- Each block gets its "own" randomization

Description of the Design: RCBD

- We call a blocking design **complete** if every treatment is used in every block
- Each treatment must appear at least once per replicate
- In general we observe every treatment (only) once in every block, hence we have a total of r (the number of blocks) observations per treatment

### Example

 Researchers wanted to evaluate the effect of several different fertilization (nitrogen) timing schedules on leaf tissue of maize

- Treatment: Six different nitrogen application timing and rate schedules (including a control treatment of no nitrogen)
- Response: Leaf tissue nitrogen amount

### Example

 Experiment design: irrigated field with a water gradient along one direction, see next slide

 We already know: Available moisture in the soil (humidity) will have an influence on the response

### Example

Layout of Experimental Design



### Example

- Layout of Experimental Design
  - The differences in plant responses caused by the water gradient will be associated with blocks

• We also say: we **control for the water gradient** 

#### Advantages of the RCBD

- Generally more precise than the CRD
- No restriction on the number of treatments or replicates
- Some treatments may be replicated more times than others
- Missing plots are easily estimated

### Disadvantages of the RCBD

 In experiments with a large number of treatments, a RCDB may become inefficient because each block must contain all treatments. This can significantly increase the size of the experiment, require more resources, and make the analysis more complex

 If the variability between blocks is too large, this can negatively affect the ability of the RCBD to detect differences between treatments

Data table

Consider an experiment installed at the RCDB with *i* treatments and *j* replicates (blocks)

Treatments						
Blocks	1	2			Total	
1	Y <sub>11</sub>	Y <sub>21</sub>		Y <sub>I1</sub>	B <sub>1</sub>	
2	Y <sub>12</sub>	Y <sub>22</sub>		Y <sub>I2</sub>	B <sub>2</sub>	
J	Y <sub>1J</sub>	$Y_{2J}$		$Y_{IJ}$	BJ	
Total	T <sub>1</sub>	$T_2$		Т	G	

#### Data table

• Number of experimental units:  $N = I \ge J$ 

i

• Total for treatment *i*: 
$$T_i = \sum_{j=1}^{j} Y_{ij} = Y_{i.}$$

• Total for block 
$$j$$
:  $B_i = \sum_{i=1}^{r} Y_{ij} = Y_{j}$ 

• Mean for treatment *i*: 
$$\widehat{m_i} = \frac{T_i}{I}$$

• Mean for block j: 
$$\widehat{m}_j = \frac{B_j}{I}$$

• General mean of the experiment:  $\widehat{m} = \frac{G}{IJ}$ 

Statistical model

Model

• Data from the RCBD can be described with the following model:

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

where  $\mu$  is the intercept (an overall mean),  $\alpha_i$  is the effect of treatment *i*,  $\beta_j$  is the effect of block *j* and  $\varepsilon_{ij}$  is the associated random error term

#### Anova table

Source	Sum of	Degrees of	Mean	F-stat
	Squares	Freedom	Square	
Treatment	SST	t-1	MST	F*=MST/MSE
Block	SSB	b-1	MSB	
Error	SSE	(t-1)(b-1)	MSE	
Total	TSS	tb-1		

- Remember about the sum of squares!
- Typically, we are **not** making inference about blocks (we already know that blocks are different!)
- It is interesting to evaluate whether there is a difference between the treatments, which can be verified using the F test for treatments

### Hypotheses

Treatments

 $H_0: t_i = 0, i = 1, 2, \dots I$  $H_1: at \ least \ one \ value \ of \ t_k \neq 0$ 

Blocks

 $H_0: b_j = 0, j = 1, 2, \dots J$  $H_1: at \ least \ one \ value \ of \ b_k \neq 0$ 

In cases where the variation between blocks is doubtful, the researcher can perform the F test for blocks, to serve as guidance for setting up future experiments! #Load the "agridat" package



#Choose the dataset "federer.tobacco" from the agridat package

- Data: RCB of tobacco, height plants exposed to radiation: 56 observations
- Evaluation of plant growth in different doses of radiation

#Perform an analysis of variance (ANOVA) for the RCDB

#Tukey test for multiple comparisons of treatments

## Let's Practice!



<pre>&gt; head(federer.tobacco)</pre>							
	row	block	dose	height			
1	1	1	2500	1299.2			
2	1	2	250	1369.2			
3	1	3	0	1169.5			
4	1	4	2500	1219.1			
5	1	5	2500	1120.0			
6	1	6	5000	1031.5			
>							

### Let's Practice!



> summary(model\_anova)

```
Error: block

Df Sum Sq Mean Sq F value Pr(>F)

Residuals 1 278088 278088

Error: Within

Df Sum Sq Mean Sq F value Pr(>F)

dose 1 185289 185289 6.688 0.0125 *

Residuals 53 1468400 27706

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

## References

- Chapter 9 Analysis of Variance II: The Randomized Complete Block Design<sup>1</sup> (for a more classical view)
- Chapter 3 Complete Block Designs<sup>2</sup>

1. Steel, R. G. & Torrie, J. H. Principles and Procedures of Statistics: A Biometrical Approach. 2nd Edition. (1980).

2. Casella, G. Statistical Design. (2008).