

Incomplete Block Designs I

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Winter 2025



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Recall: Blocking

- **Blocking** is an error control technique that allows for heterogeneous experimental units to be studied to arrive at general conclusions.
- In a **randomized block design**, the heterogeneous experimental units are grouped into homogeneous sub-groups (blocks) before randomization.
- Treatment factors are then randomly assigned to the units within the smaller homogeneous blocks.

Introduction

- In randomized block designs, the number of EUs in the block must be greater than or equal to the number of factor levels.
 - RCB: Number of EUs per block = t .
 - GCB: Number of EUs per block = tr .
- When larger block sizes are needed, the blocks become less homogeneous.
- One solution is to construct block designs where each block only takes a subset of the possible factor levels.
 - *Incomplete block designs*

Road Map

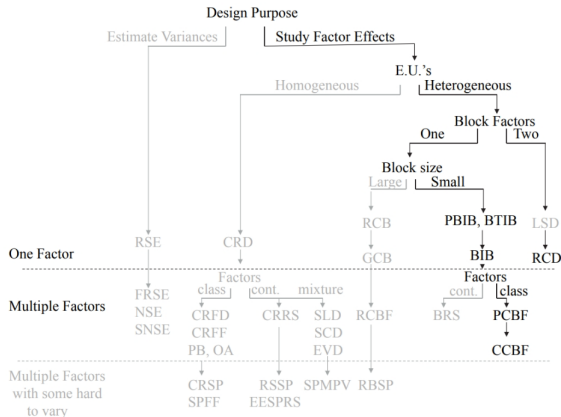


Figure: Source: (1)

Balanced Incomplete Block (BIB) Designs

- **Balanced incomplete block (BIB)** designs: Each treatment level is equally replicated and appears within a block with every other treatment level an equal number of times.
- All pairwise differences have the same standard error and the power of detection will be the same.
- *If one treatment factor level is left out of every block, it cannot be compared to the other treatment levels.*
- *If different treatment levels are unequally represented, some comparisons of factor levels will have more precision than others.*

Illustrative Example 1

- In food science, taste panels are used to test palatability of new recipes and food products.
- In this situation, the different food products represent the different factor levels.
- A random sample of subjects (EUs) is used to represent the consumer market.
 - Individual tastes are different therefore subjects represent the different blocks.
- It is not reasonable to make the subjects taste and rate every single recipe (lose their ability to discriminate).
- **Solution:** Each subject (block) only tastes a subset of the recipes.

Constructing Balanced Incomplete Block Designs I

- The simplest way to construct a BIB ($k < t$) is to form all possible subsets of k treatment levels chosen from t .
- For Example:
 - $t = 6$ recipes to be tested.
 - Each subject can taste at most $k = 3$ recipes without losing discriminatory power
 - $\binom{6}{3} = 20$ subjects (EUs) are required.
 - *All possible combinations are needed.*
 - Each recipe is (factor level) is replicated $r = 10$ times.
 - Each pair of treatment levels occurs in the same block $\lambda = 4$ times.
- This is only one way to set this experiment up, the number of blocks and replicates can change depending on the required practical difference.

Constructing Balanced Incomplete Block Designs

- The following requirements must be met in order for the design to be BIB:
 - 1 $b \geq t$
 - 2 $tr = bk$
 - 3 $\lambda(t - 1) = r(k - 1)$
 - r is the number of times each treatment level is replicated.
 - λ is the number of times each treatment level occurs with every other treatment level in the same block.
 - t is the total number of levels of the treatment factor.
 - k is the number of EUs **in a block**.
 - b is the number of blocks.
- These inequalities can be used to determine the minimum number of blocks required for a BIB design.

Constructing Balanced Incomplete Block Designs in R I

- R can be used to suggest values of λ and r to satisfy the inequalities:
 - `library(daewr)`
 - `BIBsize(t,k)`
- *Even if the inequalities are satisfied, a corresponding BIB design may not exist.*

Example 1

- Use the `BIBsize()` function to identify a possible design for the food science example ($t = 6, k = 3$).

Constructing Balanced Incomplete Block Designs in R II

- The following R code can be used to find BIB designs:
 - `library(AlgDesign)`
 - `BIB <- optBlock(~., withinData = factor(1:t),
 blocksizes = (k,b))`
 - `des <- BIB$rows`
 - `dim(des) <- NULL`
 - `des <- matrix(des, nrow = b, ncol = k, byrow = TRUE,
 dimnames = list(c("Block1", ..., "Blockb"), c("unit1",
 ..., "unitk")))`
 - `des`
- Once a BIB design is found, the levels of the treatment factor should be randomized within each block (like was done with RCB).

Example 2

- Use the `optBlock()` function to find a BIB design for the food science example ($t = 6$, $k = 3$, $b = 10$).
- What are the values of r and λ ?

Incomplete Block Design Model

- The model for the incomplete block design:

$$y_{ij} = \mu + b_i + \tau_j + \epsilon_{ij} \quad (1)$$

- It is the same model used for the randomized complete block design.
- The analysis is slightly different due to the missing observations.
 - May bias the estimates.

Analysis in R I

- May use the same methods used for analyzing data from factorial designs with an unequal number of replicates per cell (type III sum of squares).

OR

- If the differences in blocks are not of interest can use the following method in R:
 - `model <- aov(response ~ block + factor, data = data)`
 - `summary(model)`

Analysis in R II

- Can use the adjusted sum of squares to compare the adjusted means
 - `library(lsmeans)`
 - `lsmeans(model, pairwise ~ factor, adjust = ("tukey"))`
- Directly compares the adjusted means and performs hypothesis tests.

Example 3

- 1 Load the *taste* dataset from *daewr* package and take a moment to familiarize yourself with the data.
- 2 Construct an ANOVA table to examine the significance of the blocks and the recipe treatment.
- 3 Using the appropriate method, compare the adjusted means to determine if any of the recipes are *better* than any of the others.
- 4 Comment on the results.
- 5 *Do you notice anything about the standard errors of the differences?*

Determining the Number of Replicates

- Rough estimate for the number of replicates of each treatment level:

$$r = \frac{bk}{t}$$
- Rough estimate for the number of blocks: $b = \frac{tr}{k}$
- If an estimate of the variance of the heterogeneous EUs (σ^2) and the practical difference (Δ) is known:
 - Number of replicates needed (small blocks): $r = r_{crd}/RE$
 - Number of blocks of size k in an incomplete block design: $b = tr/k$
- *RE is the relative efficiency (how much blocking is expected to reduce the variance)*

Example 4

- Suppose:
 - $\hat{\sigma}_{crd}^2 = 0.040865$.
 - The number of replicates of each treatment required for a CRD design to achieve adequate power for detecting a practical difference in means was $r_{crd} = 20$.
 - Blocking is expected to reduce the variance by 90% ($\hat{\sigma}_{rcb}^2 = 10 \cdot \hat{\sigma}_{crd}^2$ or $RE = 10.0$)
- ❶ Determine the number of replicates (r) needed for for the small blocks to achieve the same power.
- ❷ How many blocks (b) are needed if the block size is $k = 2$ and there are $t = 4$ treatment levels.

References & Resources

- ① Lawson, J. (2014). *Design and Analysis of Experiments with R (Vol. 115)*. CRC press.
- BIBsize()
 - optBlock()