

Experiments in rectangular areas: restricted randomization or row-column designs?

R. A. Bailey

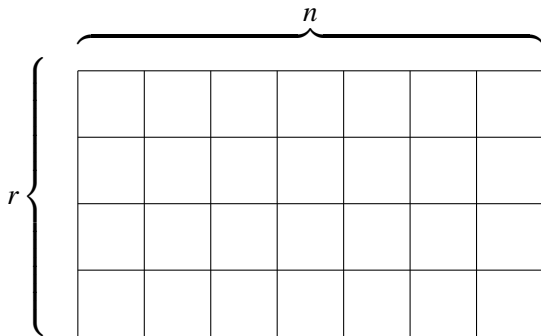


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The problem

An agricultural experiment to compare n treatments.
The experimental area has r rows and n columns.



Use a randomized complete-block design with rows as blocks.
(In each row, choose one of the $n!$ orders with equal probability.)

What should we do if the randomization produces a plan with one treatment always at one side of the rectangle?

Example

Federer (1955 book): guayule trees

<i>B</i>	<i>D</i>	<i>G</i>	<i>A</i>	<i>F</i>	<i>C</i>	<i>E</i>
<i>A</i>	<i>G</i>	<i>C</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>E</i>
<i>G</i>	<i>E</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>A</i>
<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
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<i>G</i>	<i>E</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>A</i>
<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
<i>G</i>	<i>B</i>	<i>F</i>	<i>C</i>	<i>D</i>	<i>A</i>	<i>E</i>

Proposed courses of action

Solution (Fisher): Continue to randomize and analyse as usual

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Solution: Use a Latinized design, but analyse as usual

Deliberately construct a design in which no treatment occurs more than once in any column.

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Solution: Use a carefully chosen Latinized design;
REML/ANOVA estimates of variance components

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- ▶ Simple to randomize.
- ▶ Simple to analyse.
- ▶ Some treatment comparisons in some experiments will have a specially low or specially high variance, but the estimated variance is unbiased when averaged over all comparisons and all possible randomized plans.

Assumed model

Y_α is the response on plot α .

$E(Y_\alpha) = \theta_i$ where i is the treatment on α .

$$\begin{aligned}\text{Var}(Y_\alpha) &= \sigma^2 && \text{for all } \alpha \\ \text{Cov}(Y_\alpha, Y_\beta) &= \begin{cases} \rho\sigma^2 & \text{if } \alpha \neq \beta \text{ in same row} \\ \tau\sigma^2 & \text{if } \alpha \neq \beta \text{ in same column} \\ 0 & \text{if } \alpha \neq \beta \text{ otherwise} \end{cases}\end{aligned}$$

with $0 \leq \rho \leq 1$ and $0 \leq \tau \leq 1$.

Concurrence

λ_{ij} = number of pairs of plots in the same column getting treatments i and j .

<i>B</i>	<i>D</i>	<i>G</i>	<i>A</i>	<i>F</i>	<i>C</i>	<i>E</i>
<i>A</i>	<i>G</i>	<i>C</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>E</i>
<i>G</i>	<i>E</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>A</i>
<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
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<i>G</i>	<i>E</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>A</i>
<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
<i>G</i>	<i>B</i>	<i>F</i>	<i>C</i>	<i>D</i>	<i>A</i>	<i>E</i>

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<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
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$$\lambda_{AD} = 0 + 1 + 0 + 1 + 0 + 0 + 1 = 3$$

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$$\lambda_{AA} = 1 + 1 + 0 + 1 + 0 + 1 + 1 = 5$$

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<i>G</i>	<i>E</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>A</i>
<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
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$$\lambda_{BB} = 4 + 1 + 0 + 0 + 1 + 1 + 0 = 7$$

Pairwise variance

$$\begin{aligned}\text{Var}(Y_\alpha) &= \sigma^2 \quad \text{for all } \alpha \\ \text{Cov}(Y_\alpha, Y_\beta) &= \begin{cases} \rho\sigma^2 & \text{if } \alpha \neq \beta \text{ in same row} \\ \tau\sigma^2 & \text{if } \alpha \neq \beta \text{ in same column} \\ 0 & \text{if } \alpha \neq \beta \text{ otherwise} \end{cases}\end{aligned}$$

$$\begin{aligned}V_{ij} &= \text{variance of the estimator of } \theta_i - \theta_j \\ &= \frac{\sigma^2}{r^2} [2r - 2r\rho + (\lambda_{ii} - r)\tau + (\lambda_{jj} - r)\tau - 2\lambda_{ij}\tau]\end{aligned}$$

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Pairwise variance in the example

<i>B</i>	<i>D</i>	<i>G</i>	<i>A</i>	<i>F</i>	<i>C</i>	<i>E</i>
<i>A</i>	<i>G</i>	<i>C</i>	<i>D</i>	<i>F</i>	<i>B</i>	<i>E</i>
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From
$$V_{BG} = \frac{2\sigma^2}{5} \left[1 - \rho - \frac{4}{5}\tau \right]$$

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From $V_{BG} = \frac{2\sigma^2}{5} \left[1 - \rho - \frac{4}{5}\tau \right]$ to $V_{EF} = \frac{2\sigma^2}{5} [1 - \rho + \tau]$

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<i>B</i>	<i>A</i>	<i>C</i>	<i>F</i>	<i>G</i>	<i>E</i>	<i>D</i>
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with average $V = \frac{2\sigma^2}{5} \left[1 - \rho - \frac{1}{15}\tau \right].$

Continue to randomize and analyse as usual: summary

- ▶ Simple to construct.
- ▶ Simple to randomize.
- ▶ Simple to analyse.
- ▶ Some treatment comparisons in some experiments will have a specially low or specially high variance, but the estimated variance is unbiased when averaged over all comparisons and all possible randomized plans.

Simple restricted randomization

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- ▶ Inefficient to produce plans: many will have to be rejected.
For the 5×7 rectangle, the proportion of plans with no repeat in any column is only 0.000006.
- ▶ The **actual** variance of treatment comparisons is **lower**, but the **estimate** of that variance is **higher**.

Calculations

λ_{ij} = number of pairs of plots in the same column getting treatments i and j

Note that $\sum_{j=1}^n \lambda_{ij} = r^2$ for each i .

We know that $V_{ij} = \frac{\sigma^2}{r^2} [2r(1 - \rho) + (\lambda_{ii} + \lambda_{jj} - 2\lambda_{ij} - 2r)\tau]$

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Put $V = \frac{1}{n(n-1)} \sum_{i=1}^n \sum_{j \neq i}^n V_{ij}$ and put $D = \sum_{i=1}^n \lambda_{ii}$.

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Calculations give $V = \frac{2\sigma^2}{r^2} \left[r(1 - \rho) + \left(\frac{D - r^2}{n-1} - r \right) \tau \right]$

Spectral form of covariance matrix

$$\text{Cov}(\mathbf{Y}) = \sigma^2[\mathbf{I} + \rho(\mathbf{R} - \mathbf{I}) + \tau(\mathbf{C} - \mathbf{I})]$$

where \mathbf{I} is the identity matrix,

\mathbf{R} is the matrix whose (α, β) -entry is equal to 1 if plots α and β are in the same row and to 0 otherwise,

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$$\text{So } \text{Cov}(\mathbf{Y}) = \xi_0 \mathbf{S}_0 + \xi_1 \mathbf{S}_1 + \xi_2 \mathbf{S}_2 + \xi_3 \mathbf{S}_3, \quad \text{where}$$

$$\xi_0 = \sigma^2(1 + (n-1)\rho + (r-1)\tau)$$

$$\xi_1 = \sigma^2(1 - \tau + (n-1)\rho)$$

$$\xi_2 = \sigma^2(1 - \rho + (r-1)\tau)$$

$$\xi_3 = \sigma^2(1 - \rho - \tau)$$

$$\text{rows stratum} \quad \mathbf{S}_1 = \frac{1}{n}\mathbf{R} - \frac{1}{rn}\mathbf{J} \quad \mathbf{S}_2 = \frac{1}{r}\mathbf{C} - \frac{1}{rn}\mathbf{J} \quad \text{columns stratum}$$

$$\text{grand mean} \quad \mathbf{S}_0 = \frac{1}{rn}\mathbf{J} \quad \mathbf{S}_3 = \mathbf{I} - \frac{1}{n}\mathbf{R} - \frac{1}{r}\mathbf{C} + \frac{1}{rn}\mathbf{J} \quad \text{plots stratum}$$

Strata for analysis of variance

stratum	df	variance
mean	1	ξ_0
rows	$r - 1$	ξ_1
columns	$n - 1$	ξ_2
plots	$(r - 1)(n - 1)$	ξ_3

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so $E(\text{SS for treatments}) = Q + r(n - 1)V/2$, where

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Q is a positive-definite quadratic form in the treatment effects.

Treatments are orthogonal to rows, so

$$\begin{aligned} E(\text{SS for contrasts } \perp \text{ to rows}) &= Q + (n - 1)\xi_2 + (n - 1)(r - 1)\xi_3 \\ &= \end{aligned}$$

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$$\begin{aligned} E(\text{SS for contrasts } \perp \text{ to rows}) &= Q + (n - 1)\xi_2 + (n - 1)(r - 1)\xi_3 \\ &= Q + (n - 1)(y + r - 1)\xi_3. \quad (y = \xi_2/\xi_3) \end{aligned}$$

Strata for analysis of variance

stratum	df	variance	
mean	1	ξ_0	
rows	$r - 1$	ξ_1	
columns	$n - 1$	ξ_2	} treatments
plots	$(r - 1)(n - 1)$	ξ_3	
			} residual

$$E(\text{SS for anything}) = \text{SS}(E(\text{anything})) + \text{variance term}$$

$$\text{so } E(\text{SS for treatments}) = Q + r(n-1)V/2, \quad \text{where}$$

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$$\text{So } E(\text{MS residual}) = E\left(\frac{\text{SS residual}}{(n-1)(r-1)}\right) = \frac{1}{r-1} \left[(y + r - 1)\xi_3 - \frac{rV}{2} \right].$$

Overestimation of variance

The estimator of V is $\hat{V} = \frac{2M}{r}$, where $M = \text{MS residual}$.

We have shown that $E\left(\frac{2M}{r}\right) = \frac{2}{r-1} \left[\frac{(y+r-1)}{r} \xi_3 - \frac{V}{2} \right],$

so smaller $V \implies$ larger \hat{V} .

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$$\begin{aligned} V &= \frac{2\xi_3}{r} \left[1 + \frac{(y-1)(n-r)}{r(n-1)} \right] \\ \text{and } \frac{2E(M)}{r} &= \frac{2\xi_3}{r} \left[1 + \frac{(y-1)n}{r(n-1)} \right] \end{aligned}$$

which over-estimates V by $2(y-1)\xi_3/[r(n-1)]$.

Simple restricted randomization: summary

Keep re-randomizing until you get a plan with no treatment more than once in any column. Analyse as usual.

- ▶ Inefficient to produce plans: many will have to be rejected.
- ▶ Variance is overestimated:

$$V = \frac{2\xi_3}{r} \left[1 + \frac{(y-1)(n-r)}{r(n-1)} \right]$$

and

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- ▶ Genuine treatment differences may not be detected.

Use a Latinized design, but analyse as usual

Deliberately construct a design in which no treatment occurs more than once in any column.

Easy to do this directly, eg

<i>A</i>						
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<i>G</i>	<i>A</i>	<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>
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<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>
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Randomize rows, columns, treatments.

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<i>G</i>	<i>A</i>	<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>
<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>
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Same bias in estimator of variance as for simple restricted randomization.

Super-valid restricted randomization

- ▶ Needs tables of designs.

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- ▶ The estimator of variance is unbiased when averaged over all comparisons **in this one experiment**.

Condition for unbiased estimator of variance

We have
$$V = \frac{2\xi_3}{r} \left[1 + \frac{(y-1)(D-r^2)}{r^2(n-1)} \right]$$

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$$\text{So } V = E(\hat{V}) \iff D = r(r+n-1)$$

If each pair of rows has one column with the same treatment but no treatment occurs more than twice in any column then $D = r(r+n-1)$ and $V = E(\hat{V})$.

A design from the tables

<i>A</i>	<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>
<i>D</i>	<i>E</i>	<i>F</i>	<i>C</i>	<i>A</i>	<i>B</i>	<i>G</i>
<i>A</i>	<i>G</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>E</i>	<i>D</i>
<i>D</i>	<i>B</i>	<i>G</i>	<i>F</i>	<i>C</i>	<i>A</i>	<i>E</i>
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1. In every pair of rows, there is exactly one column in which the two treatments are the same.

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<i>A</i>	<i>G</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>E</i>	<i>D</i>
<i>D</i>	<i>B</i>	<i>G</i>	<i>F</i>	<i>C</i>	<i>A</i>	<i>E</i>
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3. If m_i = the number of columns in which treatment i occurs twice, then $m_i - m_j \in \{-1, 0, 1\}$ for all other treatments j .
4. Subject to conditions (1)–(3), the spread of the variances of the estimators of simple treatment differences is as small as possible.

Pairwise variances in the example

<i>A</i>	<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>
<i>D</i>	<i>E</i>	<i>F</i>	<i>C</i>	<i>A</i>	<i>B</i>	<i>G</i>
<i>A</i>	<i>G</i>	<i>F</i>	<i>B</i>	<i>C</i>	<i>E</i>	<i>D</i>
<i>D</i>	<i>B</i>	<i>G</i>	<i>F</i>	<i>C</i>	<i>A</i>	<i>E</i>
<i>G</i>	<i>E</i>	<i>C</i>	<i>B</i>	<i>D</i>	<i>A</i>	<i>F</i>

Minimum $V_{AD} = \frac{2\sigma^2}{5} \left[1 - \rho - \frac{2}{5}\tau \right]$

Maximum $V_{AB} = \frac{2\sigma^2}{5} \left[1 - \rho + \frac{2}{5}\tau \right]$

Average $V = \frac{2\sigma^2}{5} (1 - \rho)$

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$$\text{Minimum} \quad V_{AD} = \frac{2\sigma^2}{5} \left[1 - \rho - \frac{2}{5}\tau \right] \quad \dots - \frac{4}{5}\tau$$

$$\text{Maximum} \quad V_{AB} = \frac{2\sigma^2}{5} \left[1 - \rho + \frac{2}{5}\tau \right] \quad \dots + \tau$$

$$\text{Average} \quad V = \frac{2\sigma^2}{5} (1 - \rho) \quad \dots - \frac{1}{15}\tau$$

one layout,
normal
method

Pairwise variances in the example

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Average $V = \frac{2\sigma^2}{5}(1 - \rho) \quad \dots - \frac{1}{15}\tau \quad \dots - \frac{2}{3}\tau$

one layout,
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simple
restricted
randomization

Super-valid restricted randomization: summary

- ▶ Needs tables of designs.
- ▶ Randomize rows, columns and treatments.
- ▶ Analyse as usual.
- ▶ Same average variance as in randomized complete-block design, but with smaller range.
- ▶ The estimator of variance is unbiased when averaged over all comparisons **in this one experiment**.

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- ▶ Same average variance as in randomized complete-block design, but with smaller range.
- ▶ The estimator of variance is unbiased when averaged over all comparisons **in this one experiment**.
- ▶ There is no separate estimate of ρ or τ (or y), so treatments must be randomized and a single standard error given for all differences.

Efficient row-column designs

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- ▶ More complicated analysis (should be available in software).
- ▶ Average variance may be less than, or more than, the average variance in randomized complete-block design, depending on the size of the correlations.
- ▶ Unbiased estimator of the variance of every treatment contrast.

Columns form an Incomplete-block design (IBD)

Given an incomplete-block design for n treatments in n blocks of size r ,
define the number A ($0 < A < 1$), depending on the design, by

$$A = \frac{2\sigma^2}{rV}$$

if the analysis uses information orthogonal to blocks.

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Randomize rows and columns.

Analyse by fitting rows, columns and treatments.

$$E(\text{MS residual}) = \xi_3$$

$$V_{ij} = \frac{2\xi_3}{rA_{ij}} \quad \text{where } A_{ij} \text{ is known}$$

$$V = \frac{2\xi_3}{rA} = \frac{2\sigma^2}{rA}(1 - \rho - \tau)$$

Example of a row-column design

<i>A</i>	<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>
<i>B</i>	<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>
<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>	<i>B</i>
<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>	<i>B</i>	<i>C</i>
<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>	<i>B</i>	<i>C</i>	<i>D</i>

$$V_{AB} = 1.044 \times \frac{2}{5} \xi_3$$

$$V_{AC} = 1.089 \times \frac{2}{5} \xi_3$$

$$V_{AD} = 1.091 \times \frac{2}{5} \xi_3$$

$$V = 1.075 \times \frac{2}{5} \xi_3$$

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<i>C</i>	<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>	<i>B</i>
<i>D</i>	<i>E</i>	<i>F</i>	<i>G</i>	<i>A</i>	<i>B</i>	<i>C</i>
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$$V_{AB} = 1.044 \times \frac{2}{5} \xi_3 \quad \text{normal method}$$

$$V_{AC} = 1.089 \times \frac{2}{5} \xi_3 \quad V = \frac{2}{5} \left(\frac{y+4}{5} \right) \xi_3$$

$$V_{AD} = 1.091 \times \frac{2}{5} \xi_3 \quad \text{averaged over randomizations}$$

$$V = 1.075 \times \frac{2}{5} \xi_3 \quad \text{N.B. } y = \xi_2 / \xi_3 \geq 1$$

Efficient row-column designs: summary

- ▶ Needs tables of designs.
- ▶ Randomize rows and columns.
- ▶ More complicated analysis (should be available in software).
- ▶ Average variance may be less than, or more than, the average variance in randomized complete-block design, depending on the size of the correlations.
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- ▶ There is no need to randomize treatments; the most important differences can be given the lowest variance.

Comparing super-valid restricted randomization and efficient row-column designs

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r	n					
	5	6	7	8	9	10
3	1.68	1.83	1.86	2.02	2.13	2.27
4	1.26	1.47	1.57	1.71	1.80	1.86
5		1.21	1.37	1.48	1.58	1.64

Use a carefully chosen Latinized design with REML/ANOVA estimates of variance components

Choose a design with the λ_{ij} as equal as possible.
Randomize rows and columns.

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Hence unbiased estimators of ξ_2 and ξ_3 and of

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But this estimator of V does not have a χ^2 distribution,
so **how do we do hypothesis tests?**

Also, there are so few effective df for ξ_2 that
these estimates have **very poor precision**.