

Factorials, Blocking & Repeated Measures

Dewayne E Perry
ENS 623
Perry@ece.utexas.edu

Factorial Design

- ❖ **Factorial design**
 - ★ 2 or more factors
 - ★ Each with discrete values or levels
 - ★ all possible combinations of the levels across all factors
- ❖ **Enables the study of**
 - ★ The effect of each factor on the dependent variable
 - ★ The effects of interactions between the factors on the dependent variable
- ❖ **Advantages**
 - ★ Reduces the possibility of experimental error
 - ★ Reduces the possibility of confounding variables
- ❖ **Disadvantages**
 - ★ Difficulties when more than two factors, or many levels

Comparing Multiple Conditions

	Psychotherapy		
Drug	<i>present</i>	<i>absent</i>	<i>mean</i>
<i>present</i>	8[3]	4[3]	6[6]
<i>absent</i>	4[3]	2[3]	3[6]
<i>mean</i>	6[6]	3[6]	4.5[12]

❖ Comparisons

- ★ Column means: effect of psychotherapy
- ★ Row means: effect of drug therapy
- ★ Number of observations for mean has doubled
- ★ Greater economy:
 - Each condition or group contributes data to several comparisons

Analysis of Variance

❖ Can decomposed 4 basis means into

★ Grand mean

➤ $(8 + 4 + 4 + 2)/4 = 4.5$

★ Residual/interaction effects of group membership

➤	group mean	-	grand mean	=	residual effect
➤	PD 8	-	4.5	=	3.5
➤	D 4	-	4.5	=	-0.5
➤	P 4	-	4.5	=	-0.5
➤	O 2	-	4.5	=	-2.5
➤	18		18		0.0

★ Sum of residual effects is always zero

What Do We Learn

- ❖ **Group Mean** tells us general level of measurements
 - ★ Usually not of great interest
- ❖ **Row Effects**
 - ★ Better to receive drug therapy than not
- ❖ **Column Effects**
 - ★ Better to receive psychotherapy than not
- ❖ **Interaction effects**
 - ★ Better to receive both than either
 - ★ Indication that it is better to receive neither than either is more than offset by row/column effects

Individual Differences

- ❖ Analysis so far does not tell quite the whole story
 - ★ Does not take into account in various scores
 - ★ Variability from mean - deviations
 - ★ Call these deviations errors
 - Error = score - group mean
 - Large error: falls far from mean
 - Small error: falls close to the mean
 - ★ Score = grand mean + row effect + column effect + interaction effect + error
- ❖ Variance
 - ★ Drug therapy and psychotherapy
 - Large eta (.76), and significant ($p = .012$)
 - ★ Interaction effect
 - Not trivial eta (.36), not close to statistically significant ($p = .30$)
 - Important in two way and higher order analyses of variance
 - Often misinterpreted

Interaction Effects

- ❖ $eta = \sqrt{SS_{between} / (SS_{between} + SS_{within})}$
- ❖ *eta*, like *r*, represents square root of proportion of variance accounted for
 - ★ But, *eta* is a very non-specific index of effect size when it is based on a source of variance with $df > 1$
 - ★ Eg, *eta* = .86 based on $df = 3$ is very large, but cannot say why it is large
 - ★ When $df = 1$, *eta* is identical with *r*
 - Drug/psychotherapy: *eta* = *r* = .76
 - Get all the ways of interpreting *r*
- ❖ While not significant ($p = .30$), *eta* is of promising magnitude (*eta* = *r* = .36)
- ❖ We regard each effect size estimate as though it were the only one in the study
- ❖ Remember that when r^2 or *eta*² exceeds 1.00
 - ★ $.574 + .574 + .130 = 1.278$

Testing Grand Mean

- ❖ Lack of interest in the magnitude of grand mean
 - ★ In part due to arbitrary units of measurement often employed
- ❖ Sometimes, the constant of measurement may be of interest
 - ★ When we failed to replicate a relationship obtained in an earlier experiment; compare our sample of subjects with an earlier sample
 - ★ When dependent variable might estimate some skill that might or might not be better than chance
 - ★ When our dependent variable might already be a difference score - eg, the difference between pre and post test
 - GM is then a equivalent to a matched pair t test

Unequal Sizes

- ❖ For one-way or omnibus analysis of variance it does not matter if we have the same number of units per condition or not
- ❖ For two-way or higher order analysis must take special care when number varies from condition to condition
 - ★ One possible approach:
 - discard units til all conditions are equal
 - Almost never justified
- ❖ Multiple regression procedures available
 - ★ Yield identical results when sample sizes equal
 - ★ Vary substantially when samples sizes become increasingly unequal
 - ★ procedure here represents yields closer to the “fully simultaneous multiple regression method” (FSMR) recommended by Overall et al 75
 - ★ For factorial designs of any size, always having 2 levels per factor, yields results identical to FSMR

Higher Order Factorial Designs

- ❖ So far dealt only with 2 way
- ❖ Suppose current example had been done twice, once for females, once for males
 - ★ Benefits: more subjects, more comparisons
 - ★ 2x2x2 factorial design = 2^3 factorial
 - ★ 3 factors: drug, psychotherapy, gender
 - ★ $N = 2 \times 2 \times 2 \times 3 = 24$ if the same twice
 - ★ $MS_{\text{error}} = 2.5$, adjustment factor $1/3 = .833$
- ❖ General strategy
 - ★ Compute main effects first
 - ★ Then two way interactions
 - Ie, residuals when two contributing main effects are subtracted from the variation in the two tables
 - ★ Then the three way interactions
 - Ie, the residuals when the three main effects and the three two way interactions are subtracted from the total variation from the total variation among the 8 conditions

Higher Order Factorial Designs

❖ Summary

- ★ Effect sizes of .76, .76, and .36 are identical in earlier two way analysis
- ★ F scores have all increases
- ★ p values are much smaller
 - As we would expect: study size increased
- ★ Tendency of gender to make some difference
 - gender and drug interactions significant

❖ Generalized strategy

- ★ Eg four way factorial
 - Construct all possible 2 and 3 way tables
 - Compute 4 main effects, 6 two way interactions, 4 three way interactions, and one four way interaction

Nature of Blocking

- ❖ Remember one of the ways to increase power:
 - ★ Increase the size of the effect
- ❖ One way to increase effect size
 - ★ Decrease the size of the within-group or error variation
- ❖ Blocking does this - increases precision
 - ★ Stratifying or subdividing of subjects/samples
 - ★ In such a way that those within a common block are more similar to each other on the dependent variable than they are to subjects/samples in another block/group

Nature of Blocking

❖ Example

- ★ Block according to anxiety level in study of new type of treatment
- ★ Measure anxiety level prior to treatment
- ★ Take top scores, randomly assign to T and C, iteratively

★ Anxiety	hst	h	m	l	lst	sum
T	8	6	3	1	1	19
C	9	7	5	3	1	25
sum	17	13	8	4	2	44

- ★ Mean: T=3.8, C=5.0, GM=4.4

★ Summary of sources

➤	SS	df	MS	F	p	eta
➤ T	3.60	1	3.60	10.29	.04	.85
➤ AB	77.40	4	19.35	55.29	.002	.99
➤ R	1.40	4	0.35			

★ Comparison

- Treatment effect is large and significant at $p < .05$
- blocking variable effect even more so

Nature of Blocking

★ Omitting blocking

➤	SS	df	MS	F	p	eta
➤ T	3.60	1	3.60	0.37	.56	.21
➤ R	78.80	4	9.85			

★ Summary

- Effect of treatment not very significant
- But same mean squares
- Residual variance decomposed into a large between blocks component compared with the small one in the blocked
- Removes the large sources of variation known to be associated with the systematic pre-experimental differences among subjects

Benefits of Blocking

- ❖ Consider the size of the sample needed to achieve the same F ratio for blocked and unblocked analyses

$$\text{reps} = \frac{MS_{\text{error}_{\text{unblocked}}} (\text{no. of blocks})}{MS_{\text{error}_{\text{blocked}}}}$$

$$\text{reps} = (9.85 \times 5) / 0.35 = 140.7$$

- ★ Would need 140 pairs in the unblocked experiment to reach the same F we have in our blocked with 5 pairs
- ★ But a difference in df and p: 4 and .04 vs 279.4 and .002
- ❖ To achieve the same significance level, would need 60.3 subjects/samples - a ratio of about 12 to 1 against blocked
- ❖ Example designed to show dramatic effects of blocking
- ❖ the larger the correlation between the blocking variable and the dependent variable
 - ★ the greater the benefits
 - ★ the greater the precision

Blocking and Covariance

- ❖ Analysis of covariance - a special case of ANOVA
 - ★ observed scores adjusted for individual differences within conditions of
 - some predictor variable, or
 - some covariate known to correlate with the dependent variable
 - ★ typical covariate is the pretest administration of the same (or similar) test that is to be employed as the dependent variable
- ❖ Detecting interactions
 - ★ another benefit besides increasing precision
 - detection of interactions between experimental and blocking variable
 - usually in designs where each block has a number of replications for each treatment condition
 - ★ Example (Treatment, Control, Mean, Residual)
 - above 60: TM=6, CM=7 TR=.33, CR= -.33
 - 40-59: TM=3, CM=6 TR= -.67, CR=.67
 - below 40: TM=6, CM=7 TR=.33, CR= -.33
 - Middle age tend to benefit more than younger or older

Blocking and Covariance

❖ Increasing precision

- ★ Might have used ANOCVA instead of blocking
 - Sometimes better at increasing precision
 - Especially when pre and post test scores are highly correlated
 - Special case of ANOVA when perfectly correlated
- ★ Useful rules of thumb (Cox 57)
 - Blocking better when correlation is .6 or less
 - ANOCVA better when correlation .8 or more
 - neither clearly better when correlation is between .6 and .8
- ★ Blocking equally efficient for both linear and curvilinear; ANCOVA only when linear
- ★ Blocking also useful when the blocks differ in qualitative rather than quantitative ways
- ★ Blocking always imposes some cost in terms of loss of df for error.
 - Cost usually small in relation to decreased MS error
 - if little reduction in MS error, can always unblock and recapture the lost df

Blocking and Repeated Measures

- ❖ Remember matched pairs t test?
 - ★ Example of blocking: each pair of observations is a block
 - ★ simplest form of repeated measures design
- ❖ All repeated measures designs are examples of blocked designs
 - ★ the more positively correlated the successive observations on the same sampling units, the more we benefit from increased precision
 - versus between subjects design
 - typically get greatest precision when block on sampling unit itself

Blocking within Blocks

- ❖ Example: determining accuracy of decoding non-verbal clues from face, body, voice
 - ★ Repeated measures design
 - ★ 30 students, 60 clips, 20 of each
 - might have face items over-represented in last half, eg
 - ★ Alternative: divide into 20 blocks of 3 each
 - randomly present one of each
 - doesn't necessarily increase precision
 - does eliminate possible confounding effects of the order of presentation
 - does allow us to learn from blocks x channels interaction
 - ✓ extent to which differences change over time

Use of Repeated Measures

- ❖ **Between subjects designs**
 - ★ Sampling units only observed once
 - ★ Variation based on individual differences between subjects
 - ★ subjects nested with in their treatment conditions
- ❖ **Within Subjects Designs**
 - ★ very efficient to administer two or more treatments to same sampling units
 - ★ sampling units serve as their own control
 - ★ subjects crossed by treatment conditions
 - ★ the more correlated, the more advantageous this approach
- ❖ **Intrinsic nature of experiment might call for repeated measures type of design**
 - ★ effect of practice on learning a task
 - ★ effects in a longitudinal study of development
 - ★ series of tests or subtests for a variety of reasons
- ❖ **Simplest type: subjects measured twice**
 - ★ compare scores under each condition
 - ★ use non-independent t test to compare correlation

Fixed and Random Effects

- ❖ Distinction to help us employ the appropriate error term
 - ★ Fixed
 - Selected particular levels of the factor
 - Cannot generalize to other levels
 - Includes most factors involving experimental manipulations, various organismic variables and repeated measures factors
 - ★ Random
 - Randomly sampled from population of levels
 - ✓ Most common is that of sampling units, especially people
- ❖ In previous example
 - ★ If we regard between subjects as random
 - We can test its significance only very conservatively
 - ★ If we regard it as fixed
 - Restrict inferences to this four subjects
 - Can test subject factors against sessions x subjects interactions
 - ★ Will consider all combinations
 - Fixed and random
 - For between and within subjects factors

Fixed and Random Effects

❖ Examples

- ★ 4 countries as our between sampling units factor
 - If only interested in these 4, fixed
 - If view as a sample from which we want to generalize, random
- ★ Longitudinal design with a summary score for each country for each of 3 decades
 - Scores are repeated measures, or within sampling units factors
 - Regard as fixed if we have chosen them specifically
 - Regard as random if we view as samples from which to generalize
- ❖ General principle that helps in determining the appropriateness of the error term
 - ★ The effect (fixed or random) we to test are properly tested by dividing MS for that effect by the MS for a random source of variation

Latin Squares

- ❖ Consider three drugs and 4 patients
 - ★ Suppose each subject given three drugs in the same sequence
 - Confound drug and order
 - Suppose A is best
 - Rival hypothesis is the first is best
 - ★ Use counterbalancing to avoid confounding
 - Sequence is systematically varied
 - Essential in organization and sequencing presentation
 - ✓ Primacy: opinions influenced by arguments presented 1st
 - ✓ Recency: opinions influenced by what is presented last

Latin Squares

- ❖ Latin squares has counterbalancing built in
 - ★ Nr of rows equals the nr of columns
 - ★ The letter presenting treatments appears in each column and row only once
 - ★ Effects of treatment, order and sequence are isolated - systematic counterbalancing

Order	1	2	3
seq 1	A	B	C
seq 2	B	C	A
seq 3	C	A	B

Latin Squares

❖ Analysis

- ★ Sequence effects tell how sequencings differ
- ★ Order effects tell how orders differ
- ★ Where is the treatment effect in latin squares?
 - In a 2×2 latin square, treatment effect is the sequence \times order interaction effect
 - Compares the two diagonals
 - Sources of variation:
 - ✓ Sequences: 1 df
 - ✓ Orders: 1 df
 - ✓ Treatments ($s \times o$): 1 df
- ★ Significance testing is a problem
 - No df available for error terms for the 3 sources of variation
 - Can use the mean of the MS 's of the other two in computing very conservative F 's
- ★ As the size of the latin square increases, the df for an error term increases
 - Get less conservative F 's
 - More accurate in the sense of less Type II errors
 - On average F 's will be too small

Other Counterbalancing Designs

- ❖ What if we have an unequal number of subjects and treatments?
- ❖ Useful strategies
 - ★ Multiple squares
 - ★ Rectangular arrays
- ❖ Rectangular arrays
 - ★ Eg, 3 treatments and 6 subjects
 - Could do two 3x3 squares
 - Could assign each subject a unique sequence of treatments since $3! = 6$
 - ★ If 4 treatments, then would need $4! = 24$ subjects
 - ★ Call such designs $t \times t!$ designs ($t = 2$ is latin squares)
 - ★ Fewer than $t!$ sample units?
 - Multiple latin squares
 - Random assignment from $t!$ Sequences
 - ✓ Constraint: ensure maximum degree of balancing in the resulting samples
 - ★ More than $t!$ Sample units?
 - Multiple rectangular arrays
 - Subjects-within sequences designs

Other Counterbalancing Designs

❖ Subjects-within sequences Designs

★ Suppose $2 \times t!$ Subjects

➤ Could randomly assign half the subjects to each of two rectangular arrays

✓ Treat each array as a different and replicated experiment

➤ Could randomly assign from $t!$ Sequences

✓ Constraint: ensure maximum counterbalancing

★ Suppose 18 subjects, 3 treatments

➤ $3! = 6$, assign 3 subjects at random to each sequence

➤ Subjects are not confounded with sequence as in latin squares

➤ Subjects are nested within sequences so sequences can be tested

Other Counterbalancing Designs

★ Noteworthy features of this design

➤ More than a single error term in the design

- ✓ Previously only one error term
- ✓ Usually associate with differences among subjects
- ✓ Have that: subjects within sequences
 - Used to test whether sequences differ from each other
 - Note: error is within conditions but between subjects
- ✓ Other error term: orders x subjects within-sequences interaction
 - Test all within-subjects sources of variation
 - Is itself a within-subjects source of variation
 - Formed by crossing repeated measures factors by the random factor of sampling units

Other Counterbalancing Designs

- To test for treatments, we must reach into the order \times sequences interactions and pull out the variation of the treatment means around the grand mean
- Analysis - pretty much same as already discussed
 - ✓ Between-subjects SS is broken down into a sequences SS and a subjects-within-sequences SS
 - ✓ Later is the difference between the between-subjects SS and the sequences SS
 - ✓ etc

3 or more Factors

- ❖ So far considered only two factors
 - ★ Between-subjects factor
 - ★ Within-subjects or repeated measures factor
 - ★ Often have two or more of each
- ❖ 2 or more between-subjects factors
 - ★ Does not increase complexity of the design as much increasing the number of within-subjects factors
 - ★ Eg, 4 subtests of personality test of three age levels and two genders

Fixed vs Random Factors

- ❖ Examples so far have assumed that all factors other than subjects within conditions have been fixed
 - ★ The most common situation
- ❖ Example:
 - ★ 5 f and 5 m teachers, 4 schools, teach lesson to 3 pupils, one designated as bright
 - ★ If fixed, 2 error terms
 - ★ If school is random factor, 5 error terms

Do repeated measures help?

- ❖ Basic utility: subjects are their “own control”
- ❖ High correlation yields advantages
- ❖ If low correlations, little advantage
- ❖ Note on Assumptions
 - ★ For F and t tests
 - Independence of errors
 - Homogeneity of variance
 - Normality
 - ★ Additional needed for repeated measures
 - Homogeneity of correlation coefficients among the various levels of the repeated measures factors
 - Patterns of inter-correlations is consistent among various levels is consistent from level to level