#### **True Experiments**

## **Cause and Effect Relationships**

- Can be clear relationship but not a causal one - e.g., between getting dressed up and having a headache the next morning
- Four basic elements to establish cause and effect:
  - 1. Manipulation
  - 2. Measurement
  - 3. Comparison
  - 4. Control



# Defining Characteristics of the Experiment

- Manipulation of the independent variable
  - Experimenter creates the conditions to be studied
  - If IV is measured rather than manipulated, it is *not* a true experiment
- Holding all other variables constant (Control)
- Participants in all conditions have as similar an experience as possible, except for IV



# Defining Characteristics of the Experiment

- > Ensuring that participants in all conditions
  - have equivalent personal characteristics
     are equivalent with respect to the DV before
  - are equivalent with respect to the DV before taking part in the study
  - attained by
  - holding the characteristics constant
  - random assignment to condition
     matching experimental and control particity
  - matching experimental and control participants on selected characteristics
  - $\cdot$  having the same participants take part in all conditions



#### Forms of Experiment

- In its simplest form, an experiment has two conditions
- To test how people behave in presence of treatment, researcher compares
- experimental condition, which represents IV
   control condition, which provides a baseline against which effect of treatment is assessed
- To compare the effects of two treatments, two comparison conditions are used



#### Mill's Methods

Experimental Group

Control Group

If X, then Y (Method of Agreement)

If not-X, then not-Y (Method of Difference)

Method of agreement

- If X (drink), then Y (headache)
  - $\boldsymbol{X}$  is a sufficient condition of  $\boldsymbol{Y}$
- > Method of difference
  - If not-X, then not-Y
  - X is a necessary condition of Y

## When Are More Than Two Experimental Conditions Needed?

- > When hypothesis being tested has more than one component
- When the baseline control condition is not sufficient to rule out all alternative explanations
- E.g., Does watching violence on TV cause more aggression?
- Would violent TV versus no TV suffice?



#### Characteristics of a Good Manipulation

Has good construct validity

- The manipulation accurately represents the construct
- > E.g., Does having confederate flirt with partner create threat of infidelity?



#### Two Forms of Manipulation Checks

#### Post-experimental interview

- Participants are asked questions that indicate whether manipulation had intended effect
- Checking it against a measure of the construct E.g., those in infidelity threat condition worried more about infidelity in self report measure
- DVs that assess the construct being manipulated Ensuring that participants in different experimental conditions have different experiences (related to IV)
  - Ave different experiences (related to IV) E.g., Participants primed for aggression should be more likely to complete word strings with synonyms of aggression than participants primed for altruism E.g., In study where a perpetrator's motives are manipulated, participants report what they believed the motive to be If participants report a different motive than the manipulation, construct validity is poor

#### Characteristics of a Good Manipulation

Reliability: Manipulation is applied in the same way every time

- Every participant in experiment experiences it in the same way
- Manipulations with low reliability have low validity



#### Characteristics of a Good Manipulation

Salience: Manipulation is noticeable

- Stands out from background
- If manipulation is complex, should be presented in different ways to ensure it is understood

#### Characteristics of a Good Manipulation

Strength: With a strong manipulation, the conditions of IV differentially affect behavior

- Best if extreme levels of IV are also
- realistic
- similar to situations in everyday life
- ethical





#### Extraneous Variance

- Variance in DV not caused by IV is error variance
  - variations in how the experimenter treats the research participants
  - factors systematically related to DV, but not of interest to researcher
  - extraneous variables that are part of the research situation but can be controlled
  - Hold absolutely constant or
  - · Limit to restricted range

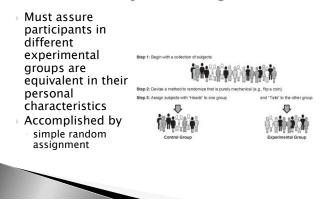


#### **Between-Subjects Designs**

- Participants take part in only one experimental condition, so contribute only one data point (between groups)
- Data points should be independent (independent samples)



#### **Between-Subjects Designs**



#### Methods of Controlling Extraneous Variables

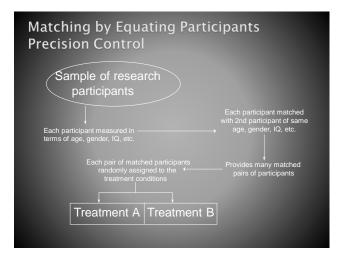
- Randomization
  - Disrupts any systematic relation between EVs and IV
     prevents EVs from becoming CVs
  - Unpredictable, unbiased procedure to distribute different values of each EV across treatment conditions
  - All possible outcomes equally likely
  - But chance CAN produce biased outcomes e.g., all heads with 10 coin tosses



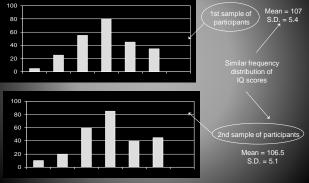
#### Matched Random Assignment

- Ensures that the members of the experimental and control groups are equivalent on one (or more) characteristic(s)
  - Researcher measures the characteristic to be controlled via a pretest
  - Participants are rank-ordered on their scores
  - Members of adjacent pairs of participants are randomly assigned to experimental or control condition





## Matching by Equating Participants Frequency Distribution Control



## Participants' Level of Introversion by Condition (with Random Assignment)

Participant	Introversion Level (0–100)	Condition	Average Introversion Level by Condition
А	36	Experimental	
В	45	Experimental	50
С	69	Experimental	
D	29	Control	
E	51	Control	50
F	70	Control	

## Matching by Yoked Control

- Controls for possible influence of temporal relationship between event and response
- Match on events that happen within experiment
   Ulcers due to physical or
- psychological stress of shock in monkeys (Brady, 1958)





#### Sample of participants with IQs of 90 - 119 Subsamples with specified IQ values identified Participants with 90-99 IQ Participants with 100-109 IQ Participants with 100-109 IQ Participants with 100-109 IQ Participants With 100-109 IQ

#### **Multiple Groups Design**

- Experiments that consist of more than the traditional two groups, experimental and control
- > Can involve quantitative IVs
  - · Conditions referred to as levels
  - Represent greater or lesser amount of the IV
  - Amount of alcohol consumed
- Can also involve qualitative IVs
- Conditions of IV represent different types or aspects of IV
- Nature of video; graphic, comedic, cartoon

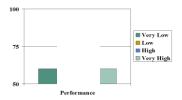
#### **Multiple Groups Design**

When additional levels of IV are examined, results can distinguish between two general categories of relationships between IV and DV

- Linear relationships: Scores on DV change constantly with level of IV
  - Positive relationship: Scores on DV increase as level of IV increase
  - Negative relationship: Scores on DV decrease as levels of IV increase

#### **Multiple Groups Design**

- Curvilinear relationships: Relationship between IV and DV takes a form other than a straight line
  - If only two levels of IV are examined, these relationships can be overlooked



#### Interpreting Results of Multiple Group Experiments

- Results of ANOVA do not tell you which of all possible two-group comparisons are statistically significant
  - Only know that at least one is
- To determine significant difference, you conduct a follow-up analysis
- When you do not have a specific a priori hypothesis, use a post-hoc test
- If you have an a priori hypothesis, use a priori contrasts

#### Advantages of Between Designs

- > Each score is independent
- > Not susceptible to:
  - Practice or experience gained in other treatments
  - Fatigue or boredom
  - · Contrast or carry-over effects



#### Disadvantages

- Large # of participants
  - Esp. problematic with special populations
- Environmental Confounds
   Characteristics of environment that might vary between groups
- Individual Differences
   Can become confounding variables
  - Assignment bias
  - Can produce high variability in scores



## Threats to Internal Validity

- Biased assignment of participants to conditions
- > Differential attrition
- > Diffusion of treatment
- Compensatory behavior
- Rivalry (John Henry effect)
- Equalization
- Resentful Demoralization



#### Within-Subjects Designs

- Each research participant experiences all of the conditions/levels of the IV (Repeated Measures)
- Results in perfect equivalence of participants across conditions (Dependent samples)
   Reduces error variance



## Advantages of Within Subjects Designs cont.

- Reduces number of participants needed
- When individual differences are consistent across treatments, can measure them and separate effects from the rest of the variance
  - Treatment effects easier to see when individual differences removed
- So within design more powerful than between design



#### Between vs. Within

Student	Speeded Test	Untimed Test
John	78	
Mary		74
Peter		68
Paul	80	
Average	79	71



#### Between vs. Within

Student	Speeded Test	Untimed Test
John	78	88
Mary	62	74
Peter	60	68
Paul	80	93
Average	70 (79)	81 (71)



#### Disadvantages of Within Subjects Designs

- > Time demand
- Participant Attrition
  - Volunteer Bias
- Time Related Factors/Threats to IV
  - History
  - Maturation
  - Instrumentation
  - Regression
- Pretest sensitization
- Testing
- Order Effects Carry-Over

#### Solutions to Time-Related Threats

- Reducing time between treatments
- But can increase risk of carry-over etc.
- Switch to between design
- Counterbalancing
- $\,\circ\,$  Matching treatments with respect to time

#### Counterbalancing

- > Treatments given in different orders
- Balances but hides order effects
- NOTE: does NOT make it a between design
   Groups balanced on order but NOT on IV itself

Group 1	Treatment A	Treatment B
Group 2	Treatment B	Treatment A





#### Intrasubject (within subject) Counterbalancing

- The ABBA Technique
- Administer treatment conditions to each participant in more than one order
- › Coke pepsi pepsi coke
- Based on assumption that order effects are linear
- If not linear use each treatment condition in every possible position in sequence
  - Also use BAAB pepsi coke coke pepsi
  - Half participants assigned to each sequence

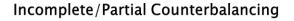
#### Intragroup Counterbalancing

- > Less time-consuming
- Groups of participants rather than individuals counterbalanced
- Different groups take each of the sequences



#### **Complete Counterbalancing**

- $\,\circ\,$  All possible treatment sequences are presented.
- You can calculate the number of sequences by using the formula *n!* (*n* factorial).
- With n = 6, *n*! = 720!
- 6 X 5 X 4 X 3 X 2 X 1
- Might require too many participants

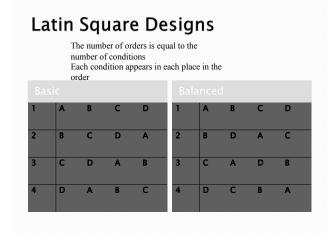


- Only a portion of all possible sequences are presented
  - Three basic requirements:
    - Each treatment must be presented to each participant an equal number of times.
  - Each treatment must occur an equal number of times at each testing or practice session.
  - Each treatment must precede and follow each of the other treatments an equal number of times.





Cour	nterbalanci	ing		
Α	В	Α	В	С
В	A	Α	С	В
		В	Α	С
		В	С	Α
		С	Α	В
		С	В	Α



#### Counterbalancing

- Sequence or Order Effects Sequence or order effects are produced by the participants being
  - exposed to the sequential presentation of the treatments.
  - The sequence or order effect depends on where in the sequential presentation of treatments the participant's performance is evaluated, not which treatment is experienced.

### Counterbalancing

**TABLE** 5-7
 EXAMPLE OF SEQUENCE OR ORDER EFFECTS IN A COUNTERBALANCED EXPERIMENT

The performance increase shown in parentheses below each sequence indicates the effect of testing reaction time to red (R), green (G), and flashing white (FW) lights on an instrument panel at that particular point in the sequence. Thus, second and third testings result in increases of 4 and 3, respectively, regardless of the experimental task.

	ORDER OF	TASK PRESEN	TATION
	R	G	FW
Performance Increase $\rightarrow$	(0	4	3)
	R	FW	G
	(0	4	3)
	G	R	FW
	(0	4	3)
	G	FW	R
	(0	4	3)
	FW	R	G
	(0	4	3)
	FW	G	R
	(0	4	3)

## Counterbalancing

 Carryover Effects
 The effects of one treatment persist or carry over and influence responses to the next treatment. 
 State
 State
 Description

 Convert effects our whom a specific preceding treatment influences the performance in a laborator treatment, in this avarple, expense risk presenter (2 prior to freatment 1 prior to the streament 1. The streament 2 prior to the streament 3 prior to the streament 4 prior to the s

next treatment.	Effect on Performance -+	(0	-2	+3)
next treatment.		G	FW	R
		(0)	0	-3)
		R	G	FW
		(0	+2	0)
		R	FW	G
		(0	+3	0)
		FW	G	R
		(0	0	-2)
		FW	R	G
		(0	-3	+2)

#### **Categories of Order Effects**

- Sensitization effects: A form of reactivity in that experiencing one condition affects performance in another condition
  - Example: Participants evaluate the résumés of both a physically attractive and physically unattractive job applicant
  - Results in demand characteristics if participants: form hypothesis about purpose of experiment (Does
  - physical attractiveness affect evaluations?)
  - respond in socially desirable way
  - Harder to control than practice or carryover effects

#### **No Order Effects**

- No difference if treatment is presented first or second
- Where for Group A, Treatment 1 occurred 1<sup>st</sup> and for Group B Treatment 2 occurred 1<sup>st</sup>
- Difference of 5 points between Treatments regardless of when presented

1 (A, B)	20	15
2 (B, A)	20	15
1 (A, B)	20 Treatment 1	15 Treatment 2
2 (B, A)	15 Treatment 2	20 Treatment 1

## Symmetrical Order Effects

- > Order matters and is the same regardless of what the treatment is
- E.g., second treatment score always raised by 10 points regardless of which treatment it is

Group		
Group 1 (A, B)	20	30
Group 2 (B, A)	34	24
Group		
Group 1 (A, B)	20 Treatment 1	30 Treatment 2
Group 2 (B, A)	24 Treatment 2	34 Treatment 1

## Nonsymmetrical Order Effects

Specific treatments determine the type of order effects, e.g., fatigue vs. practice (Differential order effects)

Group 1 does better on Treatment B when receiving it second, but Group 2 does the same on both treatments when receiving treatment B first

Group		Treatment 2
Group 1 (A, B)	20	30
Group 2 (B, A)	24	24
Group	Order 1	Order 2
Group 1 (A, B)	20 Treatment 1	30 Treatment 2

# Example of Differential Order Effects

Condition					
Experimental Control Difference					
True performance level	40	30	10		
Order Effect for E to C	0	10			
Order Effect for C to E	20	0			
Observed Score	60	40	20		