<ul> <li>Introduction to Meta-Analysis</li> <li>a bit of history</li> <li>definitions, strengths &amp; weaknesses</li> <li>what studies to include ???</li> <li>"choosing" vs. "coding &amp; comparing" studies</li> <li>what information to code along with each effect size ???</li> </ul>	<ul> <li>The two events that seem to have defined &amp; stimulated meta-analysis in Psychology</li> <li>In 1952, Hans J. Eysenck reviewed the available literature and concluded that there were no favorable effects of psychotherapy – guess how that went over</li> <li>20 additional years of empirical research "failed to resolve the debate"</li> <li>In 1978, Gene V. Glass statistically aggregated the findings of 375 psychotherapy outcome studies</li> <li>Glass (and colleague Smith) concluded that psychotherapy did indeed work</li> <li>Glass called his method "meta-analysis"</li> </ul>
<ul> <li>The Emergence of Meta-Analysis</li> <li>The statistical ideas behind meta-analysis predate Glass's work</li> <li>R. A. Fisher (1944)</li> <li>"When a number of quite independent tests of significance have been made, it sometimes happens that although few or none can be claimed individually as significant, yet the aggregate gives an impression that the probabilities are on the whole lower than would often have been obtained by chance"</li> <li>Source of the idea of aggregating probability values</li> <li>W. G. Cochran (1953)</li> <li>Discusses a method of averaging means across independent studies</li> <li>Laid-out much of the statistical foundation that modern meta-analysis is built upon (e.g., inverse variance weighting and homogeneity testing)</li> </ul>	

What got all this started?

# The Logic of Meta-Analysis

- Traditional methods of review focus on statistical significance testing to decide "whether or not" there is an effect (though we really don't "believe" in the H0:")
- Significance testing is not well suited to this task
  - highly dependent on sample size
  - Most errors are Type II errors (e.g., Butcher's 59%)
  - question of comparability of studies of "same study"
- Meta-analysis changes the focus to the direction and magnitude of the effects across studies
  - Isn't this what we are interested in anyway?
  - Direction and magnitude represented by the effect size

## When is meta-analysis applicable?

- Meta-analysis is applicable to collections of research that...
  - are empirical, rather than theoretical
  - produce quantitative results, rather than qualitative findings (need means and variances)
  - have findings that can be configured in a comparable statistical form (e.g., as effect sizes, correlation coefficients, odds-ratios, etc.)
  - examine constructs and relationships that are "comparable" given the question at hand
  - Can compute, approximate, or estimate an effect size (ES)

Kinds of Research Amenable to Meta-Analysis

- Central Tendency Research
  - prevalence rates & averages
- Between Group Contrasts
  - Experimental designs
  - Non-experimental & Natural Groups designs
- Within-Groups Contrasts
  - Experimental designs
  - Non-experimental & Pre-Post designs
- Studies of Statistical Association Between Variables
  - measurement research (e.g., reliability & validitty)

O

- individual differences research

The "Parts" of a meta-analysis

- Each study / analysis is a "case" in the meta analysis
  - simple studies will have single analysis giving a single ES
  - more complex studies may yield several ESs
- Effect Size (ES) is the "dependent variable" in the meta analysis
  - is comparable across studies
  - represents the magnitude & direction of the effect of interest
  - is independent of sample size
- Other "important" attributes of the study / analysis producing the effect size are the "independent variables" in the meta analysis

   these have to be coded into the database

What are the strengths of meta-analysis ?

- A disciplined and quantitative approach to combining and comparing empirical research findings
- Is a non-hierarchical approach doesn't favor earlier or later studies as a "starting place" to which we compare other studies
- · Protects against over-interpreting differences across studies
- Can handle a large numbers of studies (this would overwhelm traditional approaches to review)
- Allows us to evaluate what attributes of a study are related to smaller vs. larger effect sizes
- Allows us to better balance concerns about "maximum effect size" and "maximum representativeness" when designing studies
- Allows us to plan smarter, more sensitive, and more useful studies!

What are the weaknesses\* of meta-analysis ?

- Requires a huge amount of effort
- "Apples and oranges"; comparability of studies is often in the "eye of the beholder" (Wilson)
- Most meta-analyses include "blemished" studies
- Various forms of subjectivity...
  - What studies to include in the meta analyses
  - What study attributes to code
  - Coding of those attributes
- Often can't obtain study results or can't summarize as effect sizes
- Analysis of between study differences is fundamentally correlational
- \* None of these should impress you!

### Which Studies to Include?

A bit of an aside...

- The main meta analytic question <u>used</u> to be...
   "What is the size of the effect under study?"
- Leading to the question → "What studies should we include?"
- The answer <u>used</u> to be "all <u>comparable</u> studies"
- You might imagine that answer led to much argument...
- Are studies using... ... comparable?
  - ...different operationalizations / measures of the DV...
  - ...experimental and non-experimental designs...
  - ...different populations (or subpopulations)...
  - ...different → tasks ... stimuli ... equipment... settings ...

# The Replication Continuum

Pure Replications Conceptual Replications

You have to be able to argue that the collection of studies chosen for meta-analysis examine the same relationship. This may be at a broad level of abstraction, such as the relationship between criminal justice interventions and recidivism or between school-based prevention programs and problem behavior. Alternatively it may be at a narrow level of abstraction and represent pure replications.

The closer to pure replications your collection of studies, the easier it is to argue comparability.

(Lipsey & Wilson, 1993)

### Which Studies to Include?

- The main meta analytic question is now more commonly... "What things influence the size of the effect under study?"
- Leading to the answer → Every study of "the effect"
- Leading to the question → "What attributes should we include?"
- The answer is → "all important attributes"
- Lots of coding, from careful methodological evaluation of each study!!! This is often the hardest part of the meta analysis!!!!

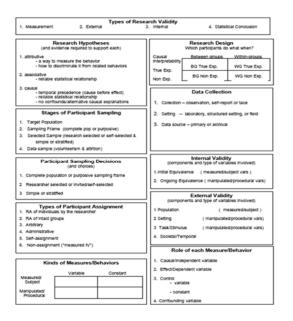
## Said differently...

Meta analyses were primarily used in the past to "combine effect sizes from comparable studies," usually to ask if the effect was "non-zero" (e.g., Glass & Smith).

Meta analyses are primarily used currently to "examine relationships between how a study is conducted and the effect sizes obtained from that study."

So (finally) ... Which Studies to Include? Which Information to Include about each study? Coding your database, so that you know "all the important stuff" You must have explicit criteria for what studies you include about each study has 4 purposes. The coding will help you ,,, • Those criteria must conform to the "standards of your people" identify groups of studies that are "replications" "Published studies" won't hack it ! · compare studies to understand what design elements are - Because studies retaining Null are less likely to be related to the size of the effect found published, including only published studies biases effect size estimates away from 0. Adjust/correct individual effect sizes to give more useful values Potential sources for identification of documents. • It will give you a better understanding of the research literature than you can possibly imagine!!! - computerized bibliographic databases - Folks who survive the meta-analysis process often say that - authors working in the research domain this was the most valuable result of their study - conference programs - You will see details, similarities, differences, genus & dissertations mistakes in a literature that you thought you knew!!!! - review articles - hand searching relevant journals What attributes of the study to code ??? - government reports, bibliographies, clearinghouses Everything that may be influencing the results & effect size !!! Q On the next several pages are summaries of materials we've used in previous classes to characterize and evaluate research designs - any of the attributes listed could add value to your meta-analysis. Put differently... Everything that can influence study results & statistical conclusion validity can also influence the effect size found!!! • All-the-Words page - organizing the design/validity jargon Validity Net - organizing the jargon around article critiquing • Researcher Choices - organizing jargon around designing studies Relationships among the types of research validity Variance sources in research designs & procedures

## Suggested Data to Code Along with the Effect Size



# Suggested Data to Code Along with the Effect Size

Measurement error → bad data →     worthless results     Is the IV properly manipulated?     Is the IV properly measured?     Are the values we have for every     measure/behavior correct?     No	causes DV mporal Precedence liable statistical ationship alternative potheses/confounds	Statistical sgnificance tests
Observational, Self-report or Trace?     Primary or Archival data?	Ci	every measure/benavior Plays a "Role" in a Study ausal Variable (IV) 4 – Ongoing Eq fect Variable (DV) 2 – Ongoing Eq
Every Measure/Behavior is either Constant Variable Measured 1 2 Manipulated 3 4	Toles' in a design	ontrol Constant 1 Inilial Eq 3 Ongoing Eq ontrol Variable 2 - Initial Eq 4 Ongoing Eq onfounding Variable 2 - Initial Eq 4 Ongoing Eq
		Internal Validity Design
External Validity	Choices we make influence Internal and External Validity !!	BG WG True Experiment © © Non-experiment © ©
Population Participant Sampling Transpropulation Sampling Finane Selected Sample Data Sample Data Sample	Participants representation vs. control	Initial Equivence – Participant Assignment • RA el individual participants by the researcher before manipulation of the IV – best hor not a guarante • Without proper RA all subject variables are potential contourd • Subject constante can't be anonymolify extended • Subject variables that are equivalent across IV conditions are control variables
Setting Laboratory, Structured or Field ?	Setting representation vs. control	<ul> <li>Subject variables that are nonequivalent across IV conditions are confounding variables – even if RA was used (remember R doesn't always work)</li> </ul>
Task/Stimulus Familiar/Representative or Unfamiliar/Control ?	Task-Stimulus representation vs. control	Ongoing Equivalence – Procedural Standardization Only the IV is different across IV conditions Procedural variables that are equivalent across IV condition are control variables
Societal/Temporal • Relationships among variables change over time in a society		<ul> <li>are control visitables that are nonequivalent across IV conditions are confounding variables</li> <li>Ongoing equivalence is harder to maintain in field settings</li> <li>Ongoing equivalence is harder to maintain during longer procedures</li> </ul>

### Suggested Data to Code Along with the Effect Size

#### Research Processes, Choices & Validity Consequences

Who participates in what ondition(s), when?" Assign ran ran ran sele	lation (Representative or easy to control) pg (Laboratory, Structured or Field) ment Procedure dom assignment of individuals by the researcher ○ dom assignment or groups ◎ dom assignment - provide the structure of the structure dom assignment - conditions set by "administrator © assignment (e.g., natural or pre-existing groups) ◎	Internal → Initial Equivalence Validity (MeasuredSubject variables) Choices of 'who' and 'where' can influence ability to perform proper random assignment Research Design BG WG True Experiment © © Non-experiment © ©
ask Completion & Settir Jata Collection Stack/ / manipulation happens first. equence and timing of the thers can vary greatly. How I	lation (Representative or easy to control) g (Laboratory, Structured or Field) Stimulus (Representative or easy to control) th of manipulation (shorter or longer) IV is manipulated DV is measured	Internal → Ongoing Equivalence Validity ManipulateRPocedural variables Choices of 'ano', "where', "what doing', and for how long' can influence the ability to maritaining control & get good measures External → Population, Setting, Task/Sitmututa Choices of 'ano', "where', "what doing', and 'for how long' can influence the ability to generalize research findings Measurement Validity Most all if ymanipulations and DV measures are 'equally good' =should know 'conventions' and 'traditions' Choices of 'who', "where', "what doing', and 'traditions' Choices of who', "where', "what doing', and 'traditions' Choices of who', "where', "what doing' and 'traditions'

