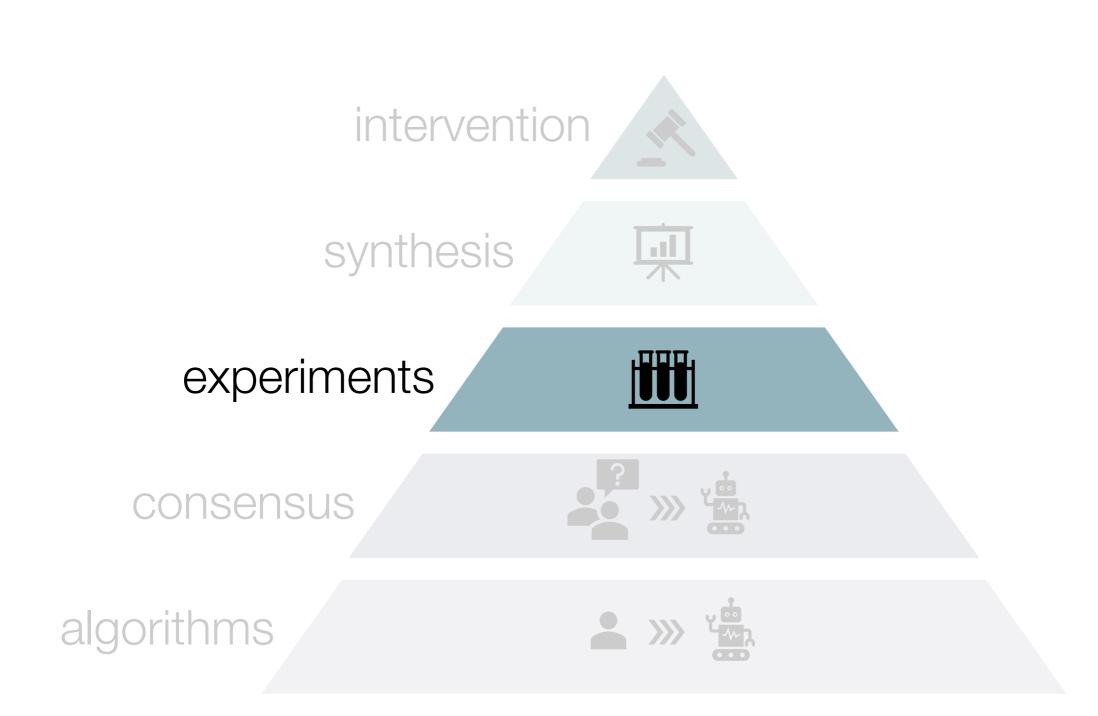
Evidence-based Decision Making Counterfactuals: Experiments

Loreen Tisdall, FS 2024

Version: February 23, 2024



Goals for today

- Understand the nature of causal inference as the comparison of treatment to some counterfactual
- Understand that experiments, and in particular RCTs, have desirable properties for causal inference – but also have limitations...
- Consider alternatives to RCTs to establish the counterfactual

Causality

: a causal quality or <u>agency</u>

: the <u>relation between a cause and its effect or</u> between regulatory correlated events or phenomena

: someone or something <u>responsible for a result</u>

Causal relations as counterfactual relations

"D shoots at V, but only grazes him, leaving V with a slightly bleeding flesh wound. X then comes along and shoots V through the heart, killing him instantly. D's act is clearly not a "cause in fact" of V's death, since V would have died, and in just the manner he did, even if D had not shot him."

- Singular judgment of causation: a single cause (e.g., A) is necessary and sufficient for effect (Y) to occur
- In reality: conjunctive plurality of causes (A&B&C → Y), disjunctive plurality of causes (A|B|C → Y)
- Complex regularities (e.g., A&B&C → Y) are rarely (if ever) fully known, thus we formulate propositions which entail the probability of a variable being causally connected with an effect

Evidence-based decision making





Francis Bacon (1561-1626)

1620

Bacon suggests that one can draw up a list of all things in which the phenomenon to explain occurs, as well as a list of things in which it does not occur. Then one can rank the lists according to the degree in which the phenomenon occurs in each one. Then one should be able to deduce what factors match the occurrence of the phenomenon in one list and do not occur in the other list, and also what factors change in accordance with the way the data had been ranked.

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"The critical step in any causal analysis is estimating the counterfactual—a prediction of what would have happened in the absence of the treatment."

Varian, H. R. (2016). Causal inference in economics and marketing. *Proceedings of the National Academy of Sciences of the United States of America, 113*(27), 7310–7315. <u>http://doi.org/10.1073/pnas.1510479113</u>

Shorter, E. (2011). A brief history of placebos and clinical trials in psychiatry. *Canadian Journal of Psychiatry*, 56(4), 193–197.

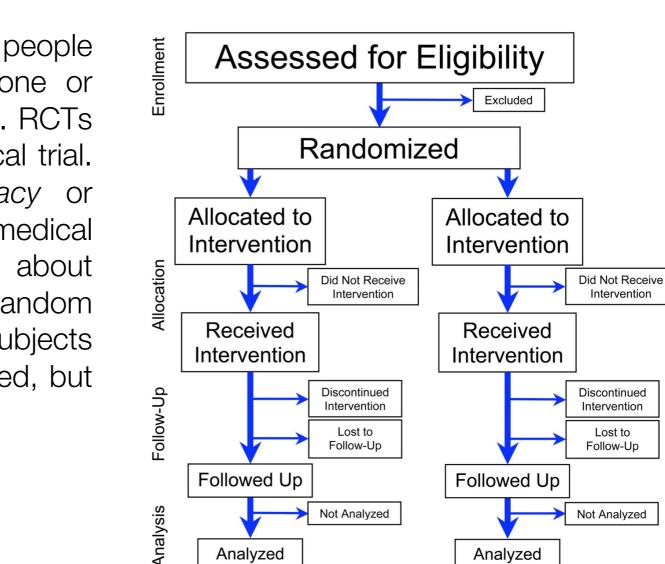
The gold standard...

Experiments/Randomised control trials (RCT)

A type of scientific experiment, where the people being studied are randomly allocated to one or other of the different treatments under study. RCTs are considered the gold standard for a clinical trial. RCTs are often used to test the efficacy or effectiveness of various types of medical intervention and may provide information about adverse effects, such as drug reactions. Random assignment of intervention is done after subjects have been assessed for eligibility and recruited, but before the intervention to be studied begins.

Efficacy:

Effectiveness:



Analyzed

"To find out what happens when you change something, it is necessary to change it." (Box et al., 2005)

Analyzed

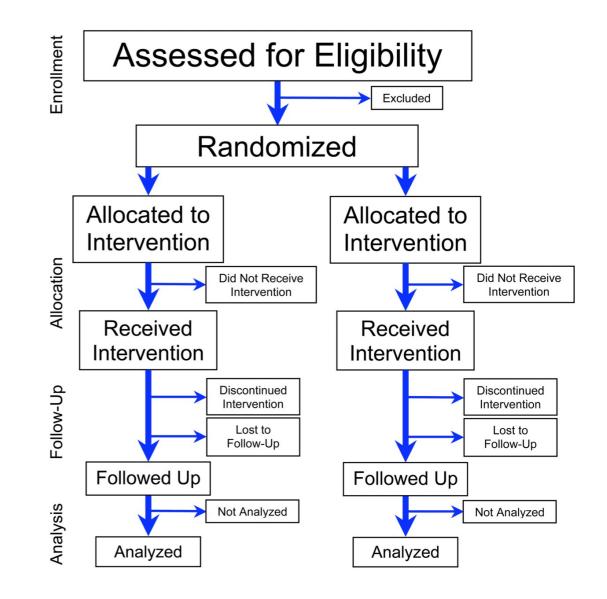
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Efficacy: how well a treatment/intervention works under ideal, controlled (laboratory) settings

Effectiveness: how well a treatment/intervention works in real-world (clinical) settings

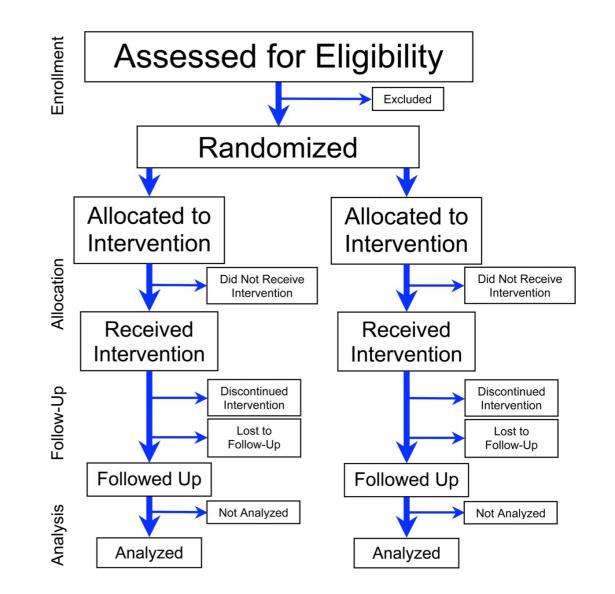


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 $Y = B_0 + B_1 group$



The gold standard...

Consolidated Standards of Reporting Trials



Schulz, K. F., Altman, D. G., & Moher, D. (2010). CONSORT 2010 statement: updated guidelines for reporting parallel group randomised trials. *Journal of Pharmacology and pharmacotherapeutics, 1*(2), 100-107..

The Salk Polio Vaccine Trial & the Cutter Incident

• The 1954 Salk Polio vaccine trial was the largest RCT (a doubleblind, randomized, and placebo-controlled study) ever conducted, involving over 1.8 million children, to test the safety and efficacy of a polio vaccine developed by Jonas Salk.

• The results showed that the vaccine was safe and effective in preventing polio.

• In 1955, shortly after the Salk polio vaccine was licensed, a manufacturing error at one of 5 licensed laboratories, Cutter Laboratories, resulted in the contamination of some batches of the vaccine with live polio virus, which led to an outbreak that affected a few hundred children, including some deaths and cases of permanent paralysis, known as the Cutter incident.

• The Cutter incident led to significant changes in vaccine regulation including the creation of oversight agencies and legislation.

 \rightarrow The Cutter incident is an example of the problems that may arise from generalizing RCTs – and the continued need for evaluation (also their legal repercussions)...



A manufacturing error at Cutter Laboratories resulted in the contamination of some batches of the vaccine with live polio virus

Offit, P.A. (2005). The Cutter incident, 50 years later. N Engl J Med. 352, 1411-1412.

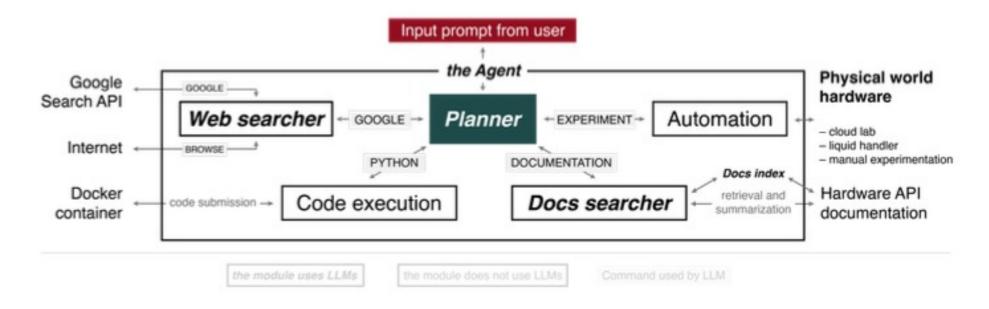
Dawson, L. (2004). The Salk polio vaccine trial of 1954: Risks, randomization and public involvement in research. *Clinical Trials, 1*, 122–130.

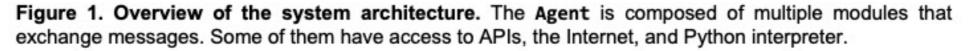
The gold standard is not always gold...

Experiments/Randomised control trials (RCT)

- Efficacy vs. effectiveness: Trials may not be widely applicable in realworld conditions....
- Generalizability: Results may not always generalize to other samples (e.g. inclusion /exclusion criteria)
- Ethical limitations: randomisation requires experimental equipoise: one cannot ethically randomise participants to some treatments (no-schooling condition)

On the horizon: Autonomous Scientific Agents

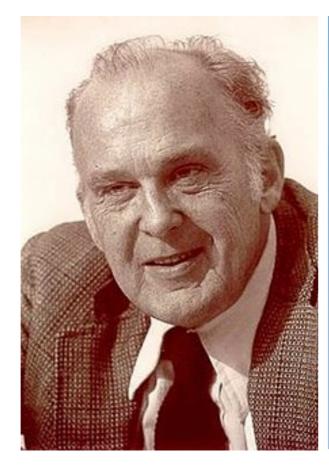




In this paper, we presented an Intelligent Agent system capable of autonomously designing, planning, and executing complex scientific experiments. Our system demonstrates exceptional reasoning and experimental design capabilities, effectively addressing complex problems and generating high-quality code. However, the development of new machine learning systems and automated methods for conducting scientific experiments raises substantial concerns about the safety and potential dual use consequences, particularly in relation to the proliferation of illicit activities and security threats. By ensuring the ethical and responsible use of these powerful tools, we can continue to explore the vast potential of large language models in advancing scientific research while mitigating the risks associated with their misuse.

Boiko, D.A., MacKnight, R. & Gomes, G. L. (2023). Emergent autonomous scientific research capabilities of large language models. https://doi.org/10.48550/arXiv.2304.05332

There are alternatives...



Donald Campbell 1916-1996

2 THE CAMPBELL COLLABORATION

Systematic reviews of the effects of interventions in education, crime and justice, and social welfare, to promote evidence-based decision-making.

What helps?

What harms?

Based on what evidence?



Does education work?



YOUR TURN!

How could you try to find out if education has an effect on intelligence?

Image created with AI (Bing), January 31, 2024

Quasi-Experimental Designs: Educational effects on intelligence

control prior intelligence =

longitudinal studies in which cognitive testing data were collected before and after variation in the duration of education (e.g., before and after university vs. no university)

policy change =

study of the effects of a change in educational duration (e.g., increase of compulsory education by I year) on mental testing **school-age cutoff** = studies use regression-discontinuity analysis to leverage the fact that school districts implement a date-of-birth cutoff for school entry (example: compare 3.9year olds that did not attend ''Kindsgi'' vs. 4.0 year-olds that did)

Table 1. Descriptive Statistics for Each Study Design

Design	Control prior intelligence	Policy change	School age cutoff
k studies	7	11	10
k data sets	10	12	20
k effect sizes	26	30	86
N participants	51,645	456,963	107,204
Mean age at early test in years (SD)	12.35 (2.90)	_	_
Mean time lag between tests in years (SD)	53.17 (15.47)		_
Mean age at policy change in years (SD)	_	14.80 (2.59)	_
Mean age at outcome test in years (SD)	63.48 (18.80)	47.92 (19.39)	10.36 (1.60)
n outcome test category (composite:fluid:crystallized)	5:20:1	2:23:5	3:67:16
n achievement tests (achievement:other)	1:25	7:23	38:48
Male-only estimates (male only:mixed sex)	2:24	8:22	0:86
Publication status (published:unpublished)	22:4	21:9	64:22

Note: To estimate N from studies with multiple effect sizes with different ns, we averaged sample sizes across effect sizes within each data set and rounded to the nearest integer. "Unpublished" refers to any study not published in a peer-reviewed journal.

Ritchie, S. J., & Tucker-Drob, E. M. (2018). How much does education improve intelligence? A meta-analysis. *Psychological Science*, 29(8), 1358–1369. <u>http://doi.org/10.1177/0956797618774253</u>

Quasi-Experimental Designs: Educational effects on intelligence

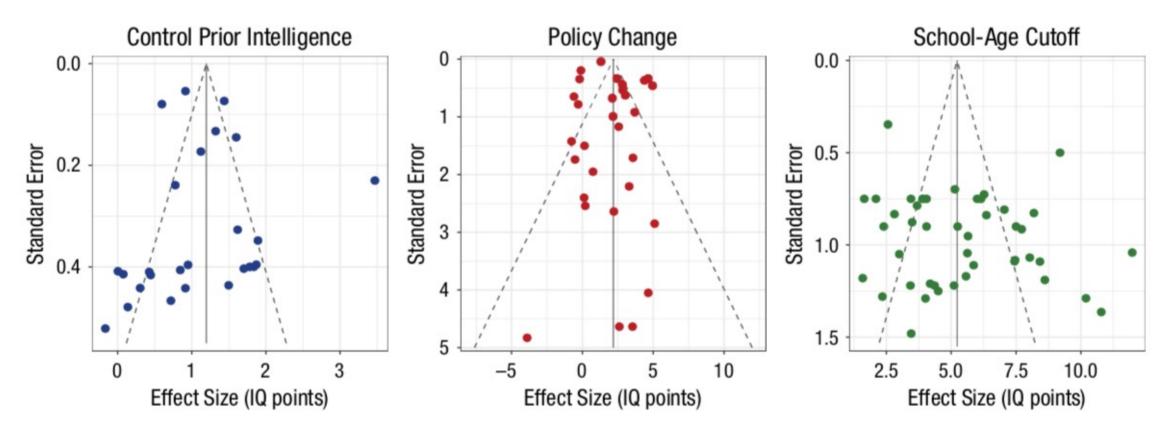


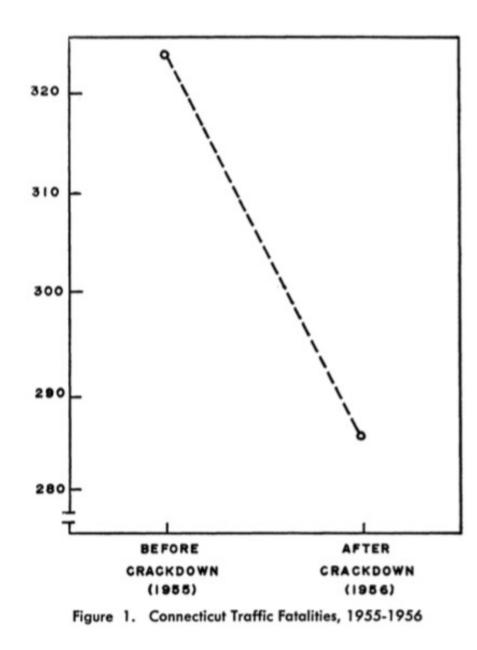
Fig. 2. Funnel plots showing standard error as a function of effect size, separately for each of the three study designs. The dotted lines form a triangular region (with a central vertical line showing the mean effect size) where 95% of estimates should lie in the case of zero within-group heterogeneity in population effect sizes. Note that 42 of the total 86 standard errors reported as approximate or as averages in the original studies were not included for the school-age-cutoff design.

"[...] we found highly consistent evidence that longer educational duration is associated with increased intelligence test scores. [...] Thus, the results support the hypothesis that education has a causal effect on intelligence test scores. The effect of 1 additional year of education—contingent on study design, inclusion of moderators, and publication-bias correction—was estimated at approximately 1 to 5 standardized IQ points."

Ritchie, S. J., & Tucker-Drob, E. M. (2018). How much does education improve intelligence? A meta-analysis. *Psychological Science, 29*(8), 1358–1369. <u>http://doi.org/10.1177/0956797618774253</u>

Quasi-experimental designs

Before-and-after measures

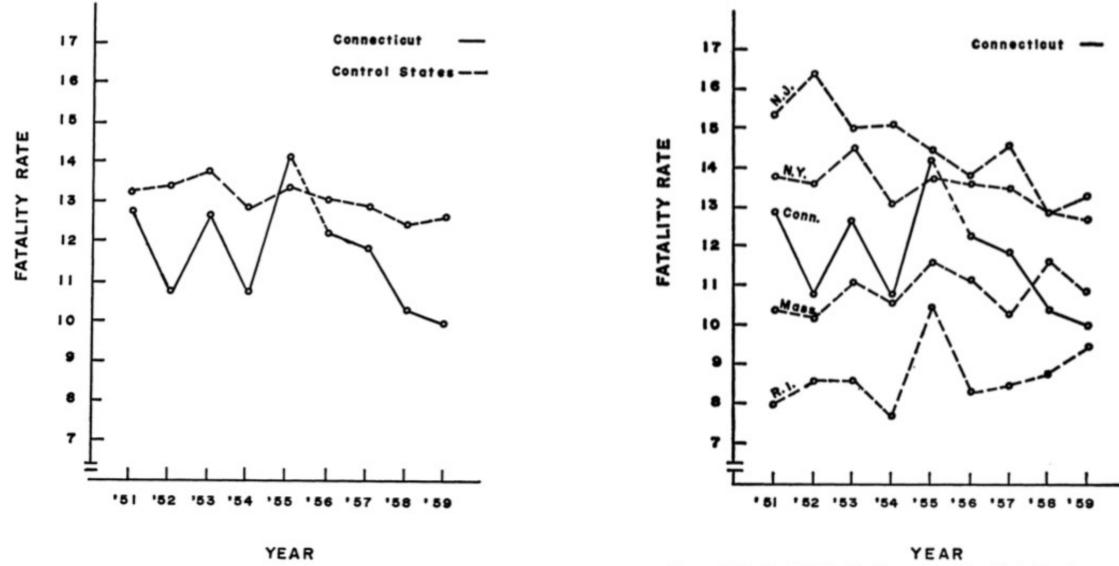


- was 1956 a dry year? (history)
- overall trends in road safety? (maturation)
- did publicising of death rates have an effect? (testing)
- were fatalities counted differently? (instrumentation)
- was this a big decrease? (instability)
- was 1955 an extreme year? (regression)

Campbell, D. T., Ross, H. L. (1968). The Connecticut crackdown on speeding: Time-series data in quasi-experimental analysis. *Law and Society Review, 3*(1), 33. <u>http://doi.org/10.2307/3052794</u>

Quasi-experimental designs

Multiple time series



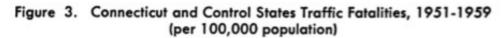
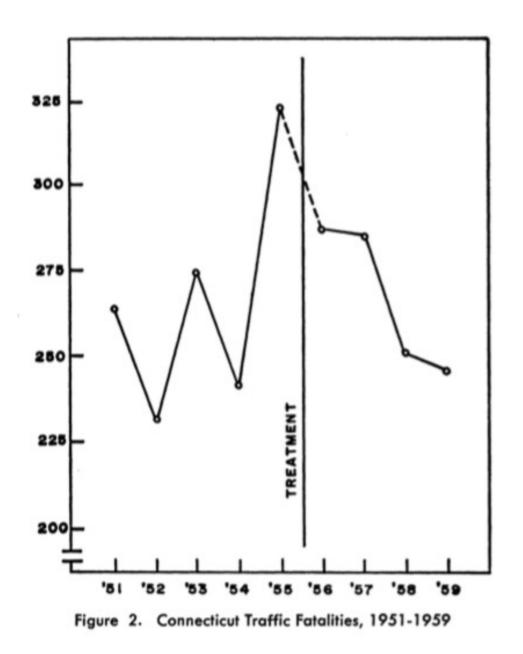


Figure 4. Traffic Fatalities for Connecticut, New York, New Jersey, Rhode Island, and Massachusetts (per 100,000 persons)

Campbell, D. T., Ross, H. L. (1968). The Connecticut crackdown on speeding: Time-series data in quasi-experimental analysis. *Law and Society Review, 3*(1), 33. <u>http://doi.org/10.2307/3052794</u>

Quasi-experimental designs

Interrupted time series



- was publicising of death rates similar across years? (testing)
- were fatalities counted differently before and after the intervention? (instrumentation)

Campbell, D. T., Ross, H. L. (1968). The Connecticut crackdown on speeding: Time-series data in quasi-experimental analysis. *Law and Society Review, 3*(1), 33. <u>http://doi.org/10.2307/3052794</u>

Experimental and Quasi-Experimental Designs for Research

Donald T. Campbell Julian C. Stanley

 TABLE 1

 Sources of Invalidity for Designs 1 through 6

			Sources of Invalidity										
• X = treatment / event		_				Inter	nal				Exter	nal	
 • O = observation of outcome / effect 		History	Maturation	Testing	Instrumentation	Regression	Selection	Mortality	Interaction of Selection and Maturation, etc.	Interaction of Testing and X	Interaction of Selection and X	Reactive Arrangements	Multiple-X Interference
	Pre-Experimental Designs: 1. One-Shot Case Study X O	_	-		1		-	-	9		-		
	2. One-Group Pretest- Posttest Design 0 X 0	-	-	-	-	?	+	+	-	-	-	\$	
	3. Static-Group Comparison <u>X</u> 0 0	+	?	+	+	+	-	-	T.				а. С
	True Experimental Designs: 4. Pretest-Posttest Con- trol Group Design R O X O R O O	+	+	+	+	+	+	+	+	-	?	?	8
	5. Solomon Four-Group Design R O X O R O O R X O R X O R O	+	+	+	+	+	+	+	+	+	?	?	
	6. Posttest-Only Control Group Design R X O R O	+	+	+	+	+	+	+	+	+	?	?	

Note: In the tables, a minus indicates a definite weakness, a plus indicates that the factor is controlled, a question mark indicates a possible source of concern, and a blank indicates that the factor is not relevant.

It is with extreme reluctance that these summary tables are presented because they are apt to be "too helpful," and to be depended upon in place of the more complex and qualified presentation in the text. No + or - indicator should be respected unless the reader comprehends why it is placed there. In particular, it is against the spirit of this presentation to create uncomprehended fears of, or confidence in, specific designs.

TABLE 2

Г									Sources of	Invalidit	7		
• X = treatment / event						Inter	nal				Exter	mal	
• O = observation of					ion				d d.	4×	X	52	
outcome / effect			uo		intat	U	-		on on,	ono	on o	nent	X S
	2	Å	urati	Bu	ume	essi	tion	alit	action	ng	action	ngen	iple
		History	Maturation	Testing	Instrumentation	Regression	Selection	Mortality	Interaction of Selection and Maturation, et	Interaction of Testing and	Interaction of Selection and A	Reactive Arrangements	Multiple-X Interference
	Quasi-Experimental Designs:	_	-		_	-	•,	-	101		10		
	7. Time Series		+	+	?	+	+	+	+	-	?	2	
	0 0 0 0X0 0 0 0 8. Equivalent Time	+	+	+	+	+	+	+	+	_	2	_	_
	Samples Design					1	1	1			•	-	_
	X10 X00 X10 X0, etc. 9. Equivalent Materials	+	+	+	+	+	+	+	+	-	>	>	_
	Samples Design					'					•	•	
	$M_{a}X_{1}O M_{b}X_{0}O M_{c}X_{1}O M_{d}$ 10. Nonequivalent Con-		, et	с. +	+	?	+	+	-	-	2	2	
	trol Group Design											•	
	$\frac{0 \times 0}{0 0}$									2			
	11. Counterbalanced	+	+	+	+	+	+	+	?	?	?	?	~
	$\begin{array}{c} \text{Designs} \\ X_1 O X_2 O X_4 O X_4 O \end{array}$,					
	X10 X10 X10 X10	•											
	X_{10} X_{10} X_{40} X_{30}												
	X40 X40 X10												
	12. Separate-Sample Pretest-Posttest	-	-	+	2	+	+	-	-	+	+	+	
	Design												
	RO(X) R XO												
	$\begin{array}{cccc} 12a.R & O & (X) \\ R & X & O \end{array}$	+	-	+	?	+	+	-	+	+	+	+	
	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$								3				
	$\begin{array}{ccc} R & O(X) \\ R & X & O \end{array}$												
	12b. $R O_1$ (X)	-	+	+	?	+	+	-	?	+	+	+	
	$\begin{array}{ccc} R & O_{1}\left(X\right) \\ R & X & O_{2} \end{array}$	3								1			
	12c. R O1 X O2	-	-	+	?	+	+	+		+	+	+	
	R X Os												

Sources of Invalidity for Quasi-Experimental Designs 7 through 12

TABLE 3
SOURCES OF INVALIDITY FOR QUASI-EXPERIMENTAL DESIGNS 13 THROUGH 16

Internal

Sources of Invalidity

External

• X = treatment / event
 O = observation of
outcome / effect

	Internal											
	History	Maturation	Testing	Instrumentation	Regression	Selection	Mortality	Interaction of Selection and Maturation, etc.	Interaction of Testing and X	Interaction of Selection and X	Reactive Arrangements	Multiple-X Interference
Quasi-Experimental Designs Continued: 13. Separate-Sample Pretest-Posttest Control Group Design R O (X) R X O R O R O R O	+	+	+	+	+	+	+	-	+	+	+	
$ \begin{cases} R' \\ R' \\$	+	+ ***	+	+	+	+	+	+	+	+	+	
14. Multiple Time-Series 0 0 0X0 0 0 0 0 0 0 0 0	+	+	+	+	+	+	+	+	-	-	?	
15. Institutional Cycle Design Class $A \times O_1$ Class $B_1 RO_2 X O_3$ Class $B_2 R X O_4$ Class $C O_5 X$ Gen. Pop. Con. Cl. $B \ O_6$ Gen. Pop. Con. Cl. $C \ O_7$ $O_2 < O_1$ $O_5 < O_4$ $O_2 < O_3$	- - - - - -	-	+ -	+ ?	?	-+	? +		+	2	++++	
$O_2 < O_4$ $O_6 = O_7$ $O_{2y} = O_{2o}$ 16. Regression	-	+	+	?	?	+	+ ?	-	+	?	2	
Discontinuity	+	+	+	1	+	+	?	+	+	-	+	+

• General Population Controls for Class B, etc.

Campbell & Stanley (1963)

Experimental and Quasi-experimental Designs

Experimental and Quasi-Experimental Designs for Research

Donald T. Campbell Julian C. Stanley "In conclusion, in this chapter we have discussed alternatives in the arrangement or design of experiments, with particular regard to the problems of control of extraneous variables and threats to validity. (...) Throughout, attention has been called to the possibility of creatively utilizing the idiosyncratic features of any specific research situation in designing unique tests of causal hypotheses." (p. 71)

A colorful bouquet of creating counterfactuals

"The stronger the demonstrated consistency of an association under conditions that rule out alternative hypotheses and the stronger the evidence regarding a mechanism that can explain the observed association, the more likely we are to accept the causal hypothesis. Usually the evidence required to confirm a causal hypothesis is cumulated across multiple studies, many of which are, of necessity, observational. Although a wide variety of research designs and analytic techniques are available to assist in gathering evidence to support a causal inference, they are helpful only to the extent that their use is guided and constrained by appropriate subject-matter considerations. No method or set of methods defines causality."



- Importance of counterfactuals: "The critical step in any causal analysis is estimating the counterfactual—a prediction of what would have happened in the absence of the treatment."
- Limitations for RCTs: RCTs are great but do not guarantee effectiveness, generalizability, or ethical treatment of participants.
- Alternatives to RCTs: Automation is on the rise, but ethical and safety issues will be crucial! Quasi-experimental designs come in many different forms with different threats to internal and external validity.