

Tutorial: March 18, 2018

**IEEE VR 2018
REUTLINGEN**

The Replication Crisis in Empirical Science: Implications for Human Subject Research in Virtual Environments

J. Edward Swan II

Mississippi State University

Outline

- **The Replication Crisis**
- **Reproducibility and Inferential Statistics**
 - Hypothesis Testing
 - Power, Effect Size, p -value
- **Reproducibility Project: Psychology**
- **What Does it Mean?**
- **What Should We Do?**

The Replication Crisis

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The Replication Crisis (Reproducibility Crisis)

Dr. John Ioannidis Exposes the Bad Science of Colleagues - The Atlantic

The

CORRESPONDENCE

LINK TO ORIGINAL ARTICLE

The screenshot shows a web browser displaying an article on Salon.com. The page features a dark header with the Salon logo and navigation links for News, Politics, Entertainment, Life, Business, Innovation, and Video. Below the header, the article title "Big Science is broken" is prominently displayed in a large, bold font. The author's name, "Pascal-Emmanuel Gobry", is shown below the title. The article's date, "April 16, 2016", is visible on the left side. The main text of the article begins with a large "S" and discusses the replication crisis in science, mentioning William A. Wilson and the pharmaceutical company Bayer. Social media sharing icons for Facebook, Twitter, and YouTube are located on the left side of the article. A "SHARE!" label is positioned above these icons. The article's content is partially obscured by a stack of overlapping window frames, suggesting it is part of a sequence of images.

[Hen Thom 2017]

The Problem

- Failure to replicate many published findings, even textbook findings
- Research biases
 - **Publication bias**: only significant ($p \leq 0.05$) results published
 - **Selection bias**: only significant results selected for analysis
 - **Reporting bias**: only significant results reported in paper
- Replication studies rarely funded, rarely published
 - Little incentive to do them
 - Therefore, most conducted studies are exploratory in nature

Evidence

- **Cancer Biology**
 - 2011 Analysis: 95% of cancer drugs fail in clinical trials
 - Led to replication studies on drug effectiveness (2011–2012)
- **In other fields, additional replication studies followed**

	Sponsor	%Replicated	Number Replicated
	Bayer	21%	14/67
	Amgen	11%	6/53
National Institute for Neurological Disorders and Stroke		8%	1/12
ALS Therapy Development Institute		0%	0/47
	Reproducibility Project: Psychology	36%	35/97

[Hen Thom 2017]

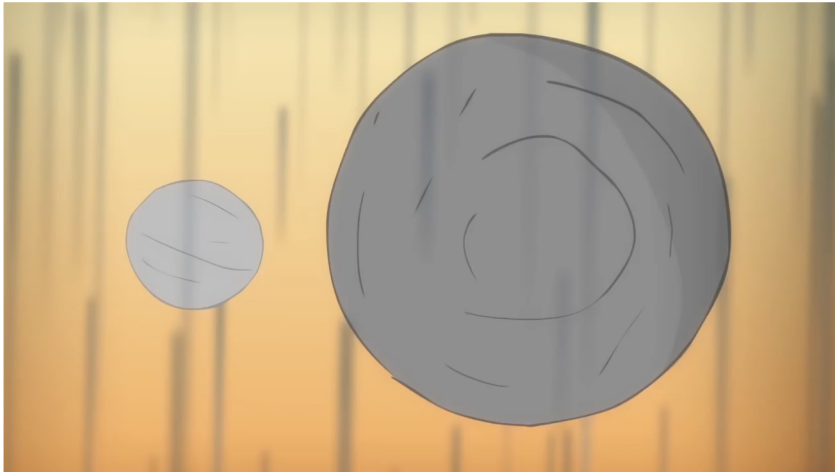
Evidence

- Replication studies conducted in **biomedicine, psychology**
- Survey data, based on question:
 - “Have you failed to reproduce somebody else’s experiment?”

Field	% Yes
Chemistry	87%
Biology	77%
Physics / Engineering	69%
Medicine	67%
Earth / Environment	64%
Other	62%

[Hen Thom 2017]

The Importance of Replication



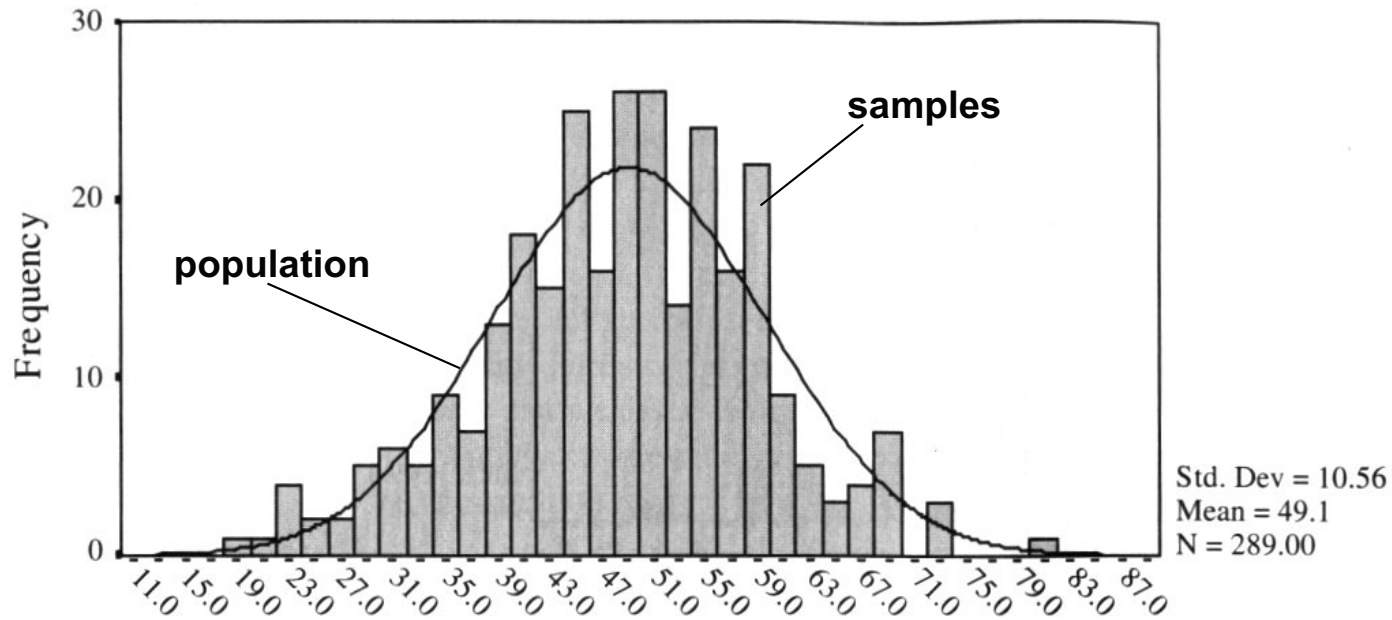
[Hen Thom 2017]

Reproducibility and Inferential Statistics

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Hypothesis Testing

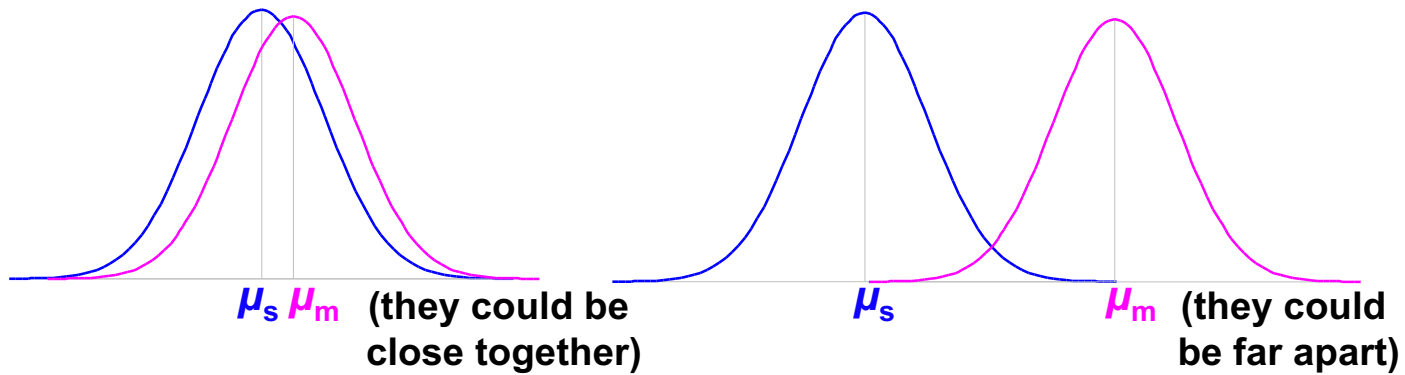
- Goal is to infer population characteristics from sample characteristics



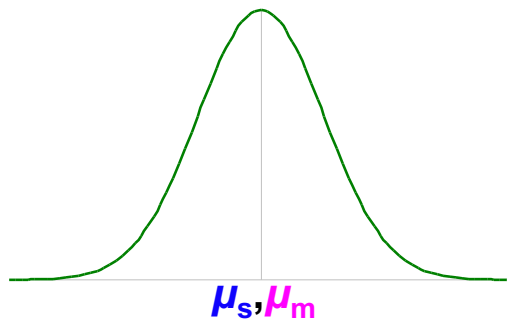
[Howell 2002, p 78]

What Are the Possible Alternatives?

- Let time to navigate be μ_s : stereo time; μ_m : mono time
 - Perhaps there are two populations: $\mu_s - \mu_m = d$



- Perhaps there is one population: $\mu_s - \mu_m = 0$



Hypothesis Testing Procedure

1. Develop testable hypothesis $H_1: \mu_s - \mu_m = d$
 - (E.g., subjects faster under stereo viewing)
2. Develop null hypothesis $H_0: \mu_s - \mu_m = 0$
 - Logical opposite of testable hypothesis
3. Construct sampling distribution assuming H_0 is true.
4. Run an experiment and collect samples; yielding sampling statistic X .
 - (E.g., measure subjects under stereo and mono conditions)
5. Referring to sampling distribution, calculate conditional probability of seeing X given $H_0: p(X | H_0)$.
 - If probability is low ($p \leq 0.05$), we are unlikely to see X when H_0 is true. We reject H_0 , and embrace H_1 .
 - If probability is not low ($p > 0.05$), we are likely to see X when H_0 is true. We do not reject H_0 .

Example 1: VE Navigation with Stereo Viewing

1. Hypothesis $H_1: \mu_s - \mu_m = d$
 - Subjects faster under stereo viewing.
2. Null hypothesis $H_0: \mu_s - \mu_m = 0$
 - Subjects same speed whether stereo or mono viewing.
3. Constructed sampling distribution assuming H_0 is true.
4. Ran an experiment and collected samples:
 - 32 participants, collected 128 samples
 - $X_s = 36.431$ sec; $X_m = 34.449$ sec; $X_s - X_m = 1.983$ sec
5. Calculated conditional probability of seeing 1.983 sec given $H_0: p(1.983 \text{ sec} | H_0) = 0.445$.
 - $p = 0.445$ not low, we are likely to see 1.983 sec when H_0 is true. We do not reject H_0 .
 - This experiment did not tell us that subjects were faster under stereo viewing.

[Swan et al. 2003]

Example 2: Effect of Intensity on AR Occluded Layer Perception

1. Hypothesis $H_1: \mu_c - \mu_d = d$
 - Tested constant and decreasing intensity. Subjects faster under decreasing intensity.
2. Null hypothesis $H_0: \mu_c - \mu_d = 0$
 - Subjects same speed whether constant or decreasing intensity.
3. Constructed sampling distribution assuming H_0 is true.
4. Ran an experiment and collected samples:
 - 8 participants, collected 1728 samples
 - $X_c = 2592.4$ msec; $X_d = 2339.9$ msec; $X_c - X_d = 252.5$ msec
5. Calculated conditional probability of seeing 252.5 msec given $H_0: p(252.5 \text{ msec} | H_0) = 0.008$.
 - $p = 0.008$ is low ($p \leq 0.01$); we are unlikely to see 252.5 msec when H_0 is true. We reject H_0 , and embrace H_1 .
 - This experiment suggests that subjects are faster under decreasing intensity.

[Living Swan et al. 2003]

Some Considerations...

- The conditional probability $p(X | H_0)$
 - Much of statistics involves how to calculate this probability; source of most of statistic's complexity
 - Logic of hypothesis testing the same regardless of how $p(X | H_0)$ is calculated
 - If you can calculate $p(X | H_0)$, you can test a hypothesis
- The null hypothesis H_0
 - H_0 usually in form $f(\mu_1, \mu_2, \dots) = 0$
 - Gives hypothesis testing a double-negative logic: assume H_0 as the opposite of H_1 , then reject H_0
 - Philosophy is that can never prove $f = 0$, because 0 is point value in domain of real numbers
 - H_1 usually in form $f(\mu_1, \mu_2, \dots) \neq 0$; we don't know what value it will take, but main interest is that it is not 0

When We Reject H_0

- Calculate $\alpha = p(X | H_0)$, when do we reject H_0 ?
 - In science generally, $\alpha = 0.05$
 - But, just a social convention
- What can we say when we reject H_0 at $\alpha = 0.008$?
 - “If H_0 is true, there is only an 0.008 probability of getting our results, and this is unlikely.”
 - **Correct!**
 - “There is only a 0.008 probability that our result is in error.”
 - **Wrong**, this statement refers to $p(H_0)$, but that’s not what we calculated.
 - “There is only a 0.008 probability that H_0 could have been true in this experiment.”
 - **Wrong**, this statement refers to $p(H_0 | X)$, but that’s not what we calculated.

When We Don't Reject H_0

- What can we say when we don't reject H_0 at $\alpha = 0.445$?
 - “We have proved that H_0 is true.”
 - “Our experiment indicates that H_0 is true.”
 - **Wrong**, hypothesis testing cannot prove $H_0: f(\mu_1, \mu_2, \dots) = 0$.
- Statisticians do not agree on what failing to reject H_0 means.
 - Conservative viewpoint (Fisher):
 - We must suspend judgment, and cannot say anything about the truth of H_0 .
 - Alternative viewpoint (Neyman & Pearson):
 - We can accept H_0 if we have sufficient experimental power, and therefore a low probability of **type II error**.

Probabilistic Reasoning

- If hypothesis testing was **absolute**:
 - If H_0 is true, then X **cannot occur**...however, X has occurred...therefore H_0 is **false**.
 - e.g.: If a person is a Martian, then they are not a member of Congress (**true**)...this person is a member of Congress...therefore they are not a Martian. (**correct result**)
 - e.g.: If a person is an American, then they are not a member of Congress (**false**)...this person is a member of Congress...therefore they are not an American. (**incorrect result, but correct logical reasoning**)

p	q	$p \rightarrow q$	$\neg q \rightarrow \neg p$		
T	T	T	T	$\left. \begin{array}{l} p \rightarrow q \\ \neg q \\ \hline \rightarrow \neg p \end{array} \right\} \text{modus tollens}$	
T	F	F	F		
F	T	T	T		
F	F	T	T		

[Cohen 1994]

Probabilistic Reasoning

- However, hypothesis testing is **probabilistic**:
 - If H_0 is true, then X is **highly unlikely**...however, X has occurred...therefore H_0 is **highly unlikely**.
 - e.g.: If a person is an American, then they are probably not a member of Congress (**true, right?**)...this person is a member of Congress...therefore they are probably not an American.
(**incorrect result, but correct hypothesis testing reasoning**)

p	q	$p \rightarrow q$	$\neg q \rightarrow \neg p$
T	T	T	T
T	F	F	F
F	T	T	T
F	F	T	T

$$p \rightarrow q$$

$$\neg q$$

$$\rightarrow \neg p$$

}

modus tollens

[Cohen 1994]

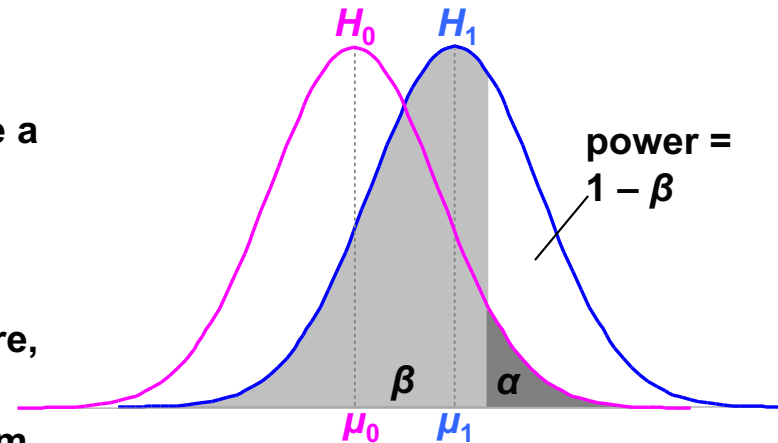
Reproducibility and Inferential Statistics

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Interpreting α , β , and Power

		Decision	
		Reject H_0	Don't reject H_0
True state of the world	H_0 false	a result! $p = 1 - \beta = \text{power}$	type II error $p = \beta$
	H_0 true	type I error $p = \alpha$	argue H_0 ? $p = 1 - \alpha$

- If H_0 is true:
 - α is probability we make a **type I error**: we think we have a result, but we are wrong
- If H_1 is true:
 - β is probability we make a **type II error**: a result was there, but we missed it
 - **Power** is a more common term than β



Increasing Power by Increasing α

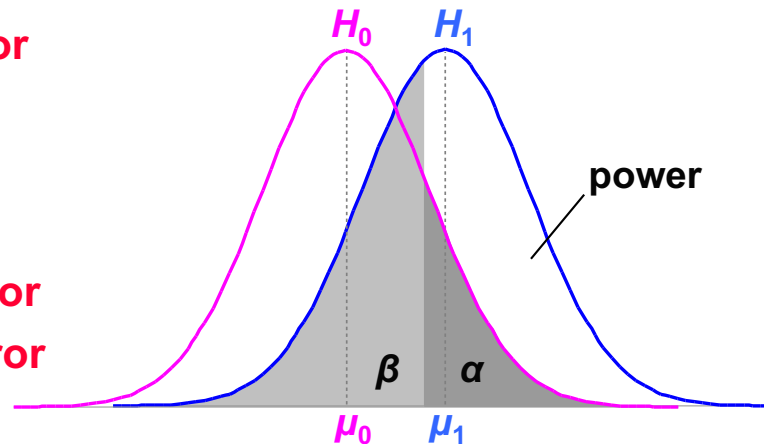
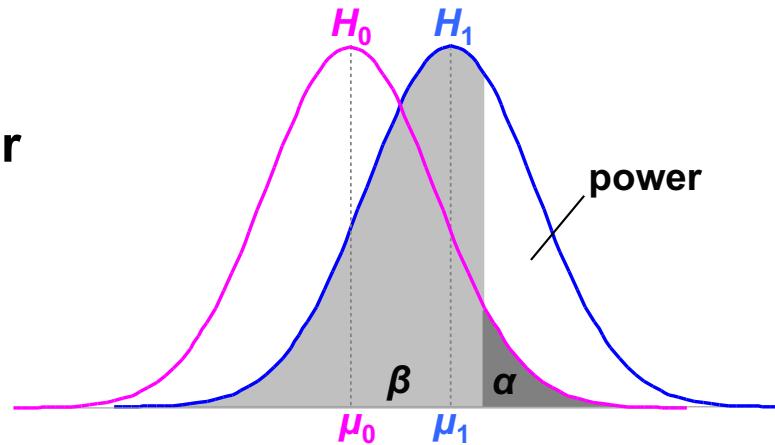
- Illustrates α / power tradeoff

- Increasing α :

- Increases power
- Decreases **type II error**
- Increases **type I error**

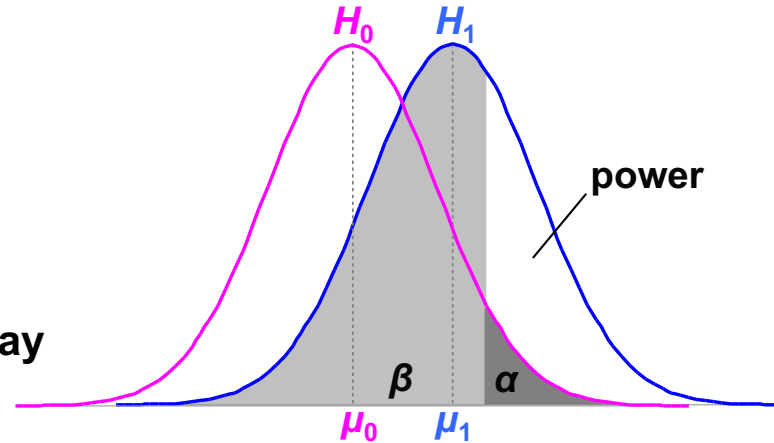
- Decreasing α :

- Decreases power
- Increases **type II error**
- Decreases **type I error**

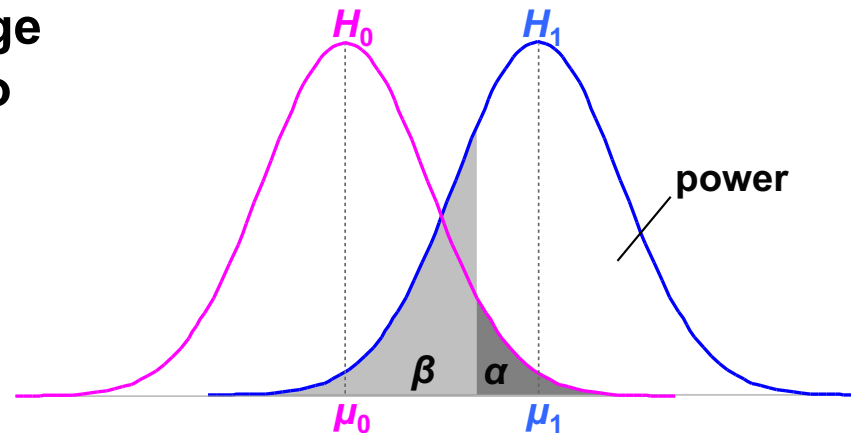


Increasing Power by Measuring a Bigger Effect

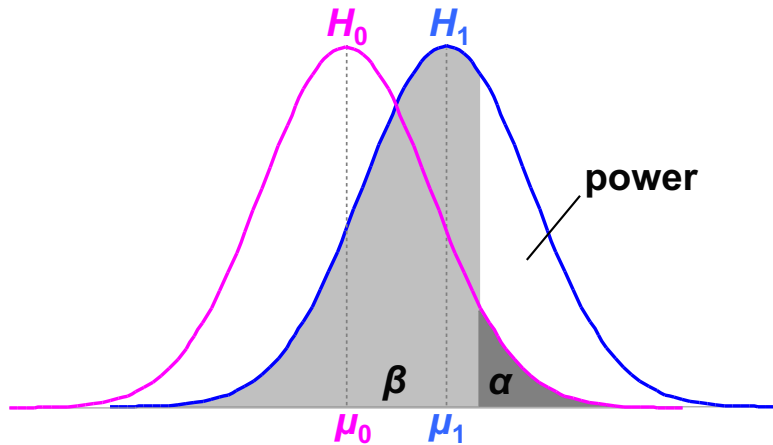
- If the effect size is large:
 - Power increases
 - **Type II error** decreases
 - α and **type I error** stay the same



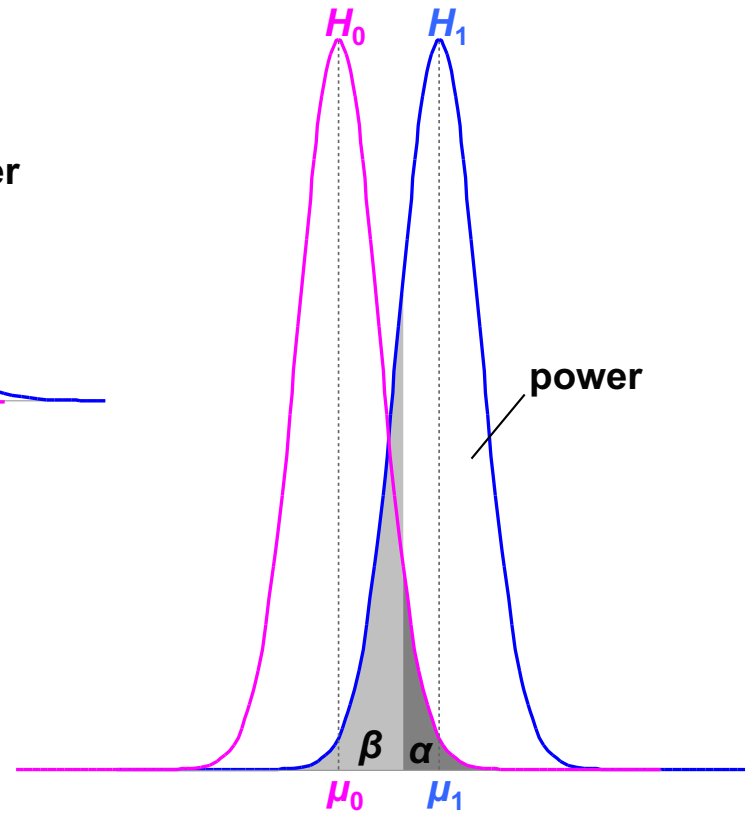
- Unsurprisingly, large effects are easier to detect than small effects



Increasing Power by Collecting More Data

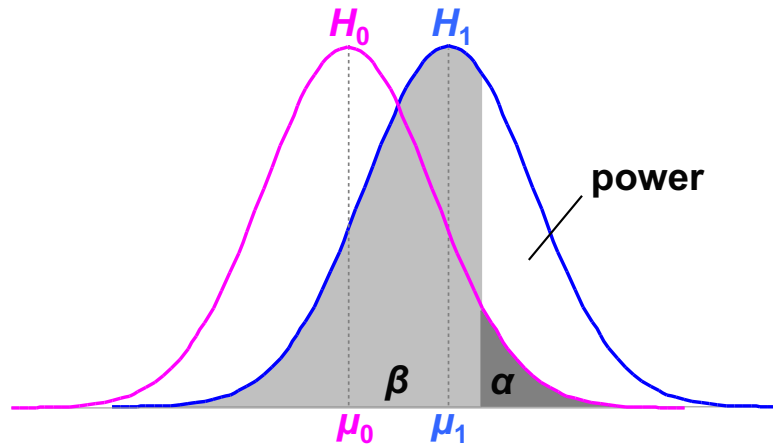


- Increasing sample size (N):
 - Decreases variance
 - Increases power
 - Decreases **type II error**
 - α and **type I error** stay the same
- There are techniques that give the value of N required for a certain power level.

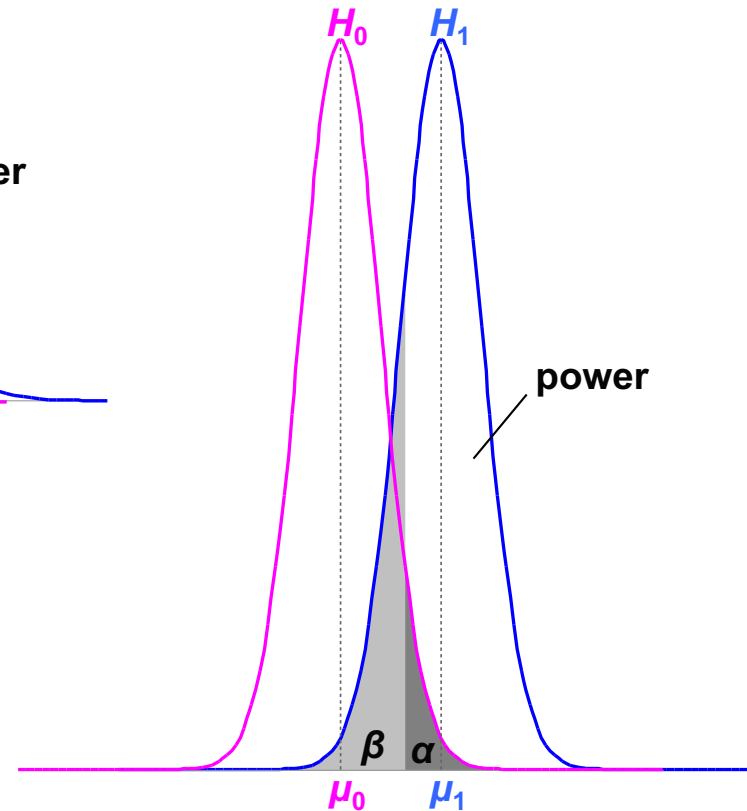


- Here, effect size remains the same, but variance drops by half.

Increasing Power by Decreasing Noise



- Decreasing experimental noise:
 - Decreases variance
 - Increases power
 - Decreases **type II error**
 - α and **type I error** stay the same
- More careful experimental results give lower noise.



- Here, effect size remains the same, but variance drops by half.

Using Power

- Need α , effect size, and sample size for power:

$$\text{power} = f(\alpha, |\mu_0 - \mu_1|, N)$$

- Problem for VR / AR:

- Effect size $|\mu_0 - \mu_1|$ hard to know in our field
 - Population parameters estimated from prior studies
 - But our field is so new, not many prior studies
- Can find effect sizes in more mature fields

- Post-hoc power analysis:

$$\text{effect size} = |X_0 - X_1|$$

- Then, calculate power for experiment
- But this makes statisticians grumble (e.g. [Howell 2002] [Cohen 1988])
- Same information as p value

Other Uses for Power

1. Number samples needed for certain power level:

$$N = f(\text{power}, \alpha, |\mu_0 - \mu_1| \text{ or } |X_0 - X_1|)$$

- Number extra samples needed for more powerful result
- Gives “rational basis” for deciding N
- Cohen [1988] recommends $\alpha = 0.05$, power = 0.80

2. Effect size that will be detectable:

$$|\mu_0 - \mu_1| = f(N, \text{power}, \alpha)$$

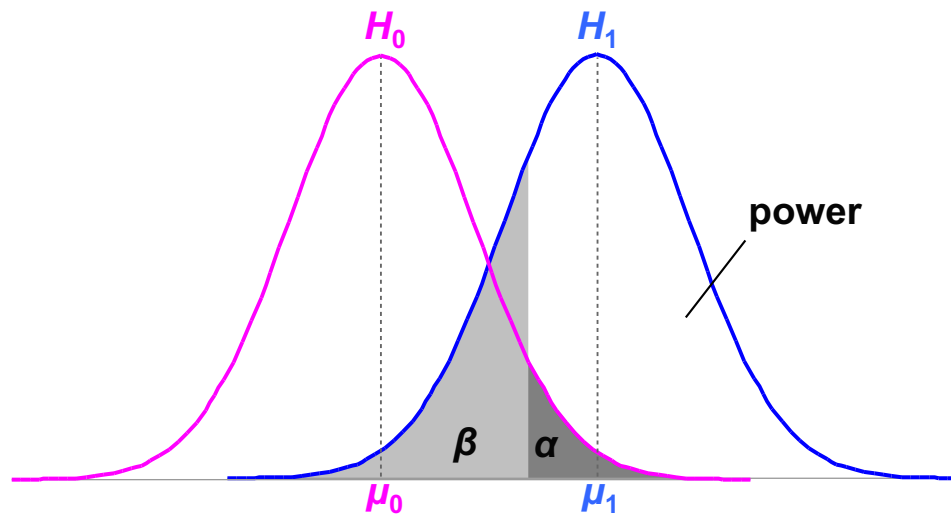
3. Significance level needed:

$$\alpha = f(|\mu_0 - \mu_1| \text{ or } |X_0 - X_1|, N, \text{power})$$

(1) is the most common power usage

Arguing the Null Hypothesis

- Cannot directly argue $H_0: \mu_s - \mu_m = 0$.
But we can argue that $|\mu_0 - \mu_1| < d$.
 - Thus, we have bound our effect size by d .
 - If d is *small*, effectively argued null hypothesis.
 - Cohen recommends $\alpha = 0.05$, power = 0.20



[Cohen 1988, p 16]

Reproducibility Project: Psychology

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Reproducibility Project: Psychology

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	Reproducibility Project: Psychology	36%	35/97

Reproducibility Project: Psychology

- Begun by Brian Nosek, University of Virginia, 2011
- Replicated 100 published studies
- Recruited very large team
 - Final paper has 270 coauthors
- Which studies to replicate?
 - Goal: minimize selection bias
 - Goal: maximize generalizability
- Published **sampling frame** and **selection criteria**



[OSC 2015, 2012]

Sampling frame and selection criteria

- Covered 3 leading journals
 - Psychological Science
 - Journal of Personality and Social Psychology
 - Journal of Experimental Psychology: Learning, Memory, and Cognition
- First 20 articles in each journal, then 10 more; begin with first 2008 issue
- Replicate last study in article (unless infeasible); 84% were last study
- Result must be a single inference test, usually *t*-test, *F*-test, *r* correlation
- If available, use original materials
- Seek design feedback from original authors
- Enough participants for high statistical power ($1 - \beta$ (power) ≥ 0.80)

Article selection results

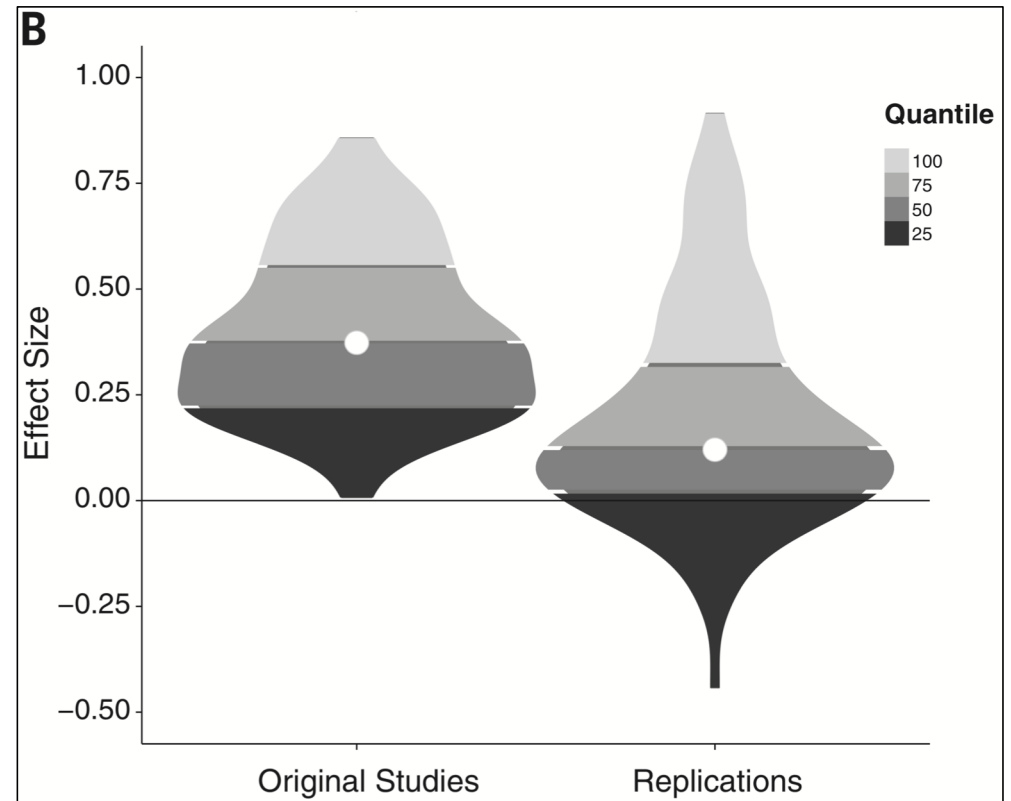
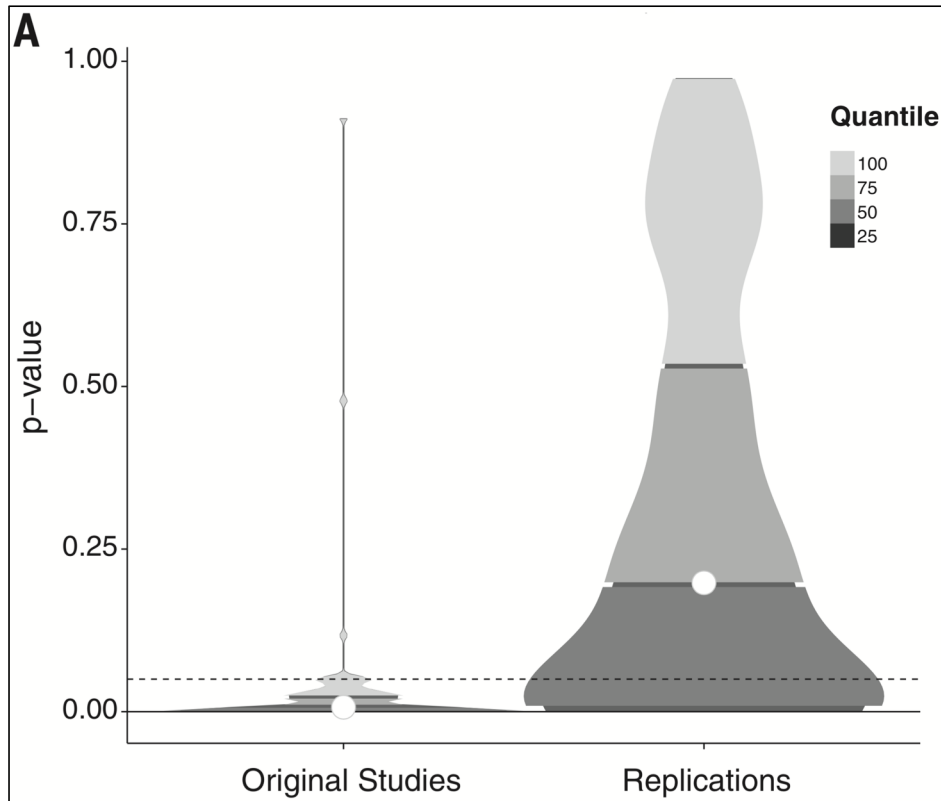
- 488 articles in 2008 issues of the 3 journals
- 158 available for replication
- 113 replications selected
- 100 completed by deadline

Data collection and processing

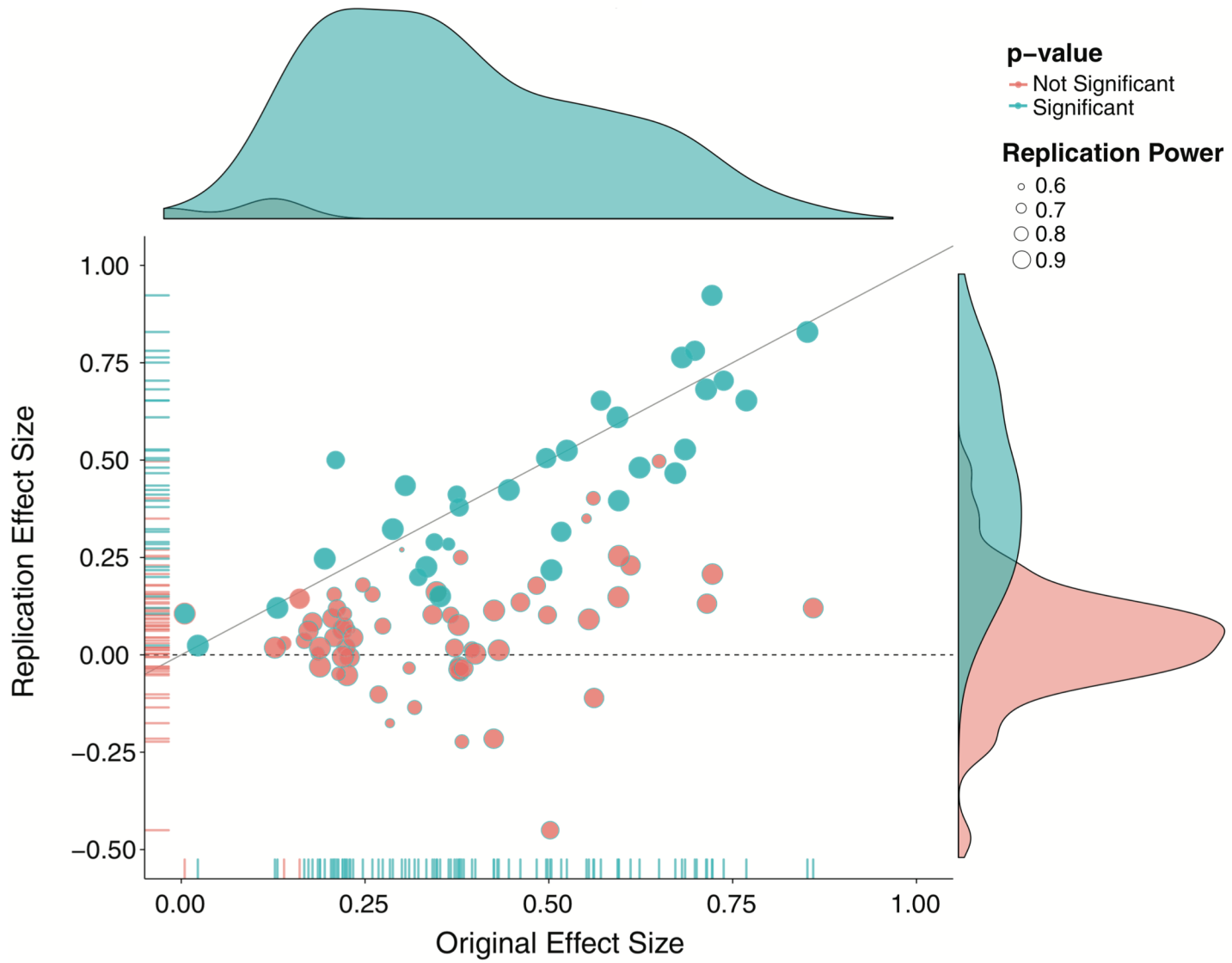
- How to measure a replication?
- How to quantify a series of replications?
- Each experiment analyzed with standard R packages
- Each analysis performed independently by 2nd team

Original Study Result Characteristics	Replication Study Result Characteristics
<i>p</i> value	<i>p</i> value
effect size	effect size
<i>df</i> or sample size	<i>df</i> or sample size
result importance rating	power
result surprisingness rating	replication challenge rating
experience, expertise rating of original team	experience, expertise rating of replicating team
	replication quality rating

Results



Results



Results by %Replicated ($p \leq 0.05$)

- Initial strength of evidence predicts replication success

Original Strength of Evidence	%Replicated ($p \leq 0.05$)	Number Replicated
$p \leq 0.001$	63%	20/32
$p \leq 0.02$	41%	26/63
$0.02 \leq p \leq 0.04$	26%	6/23
$0.04 \leq p$	18%	2/11

- Cognitive psychology more successful than social psychology

Sub-Discipline	%Replicated ($p \leq 0.05$)	Number Replicated
Cognitive Psychology	50%	21/42
Social Psychology	25%	14/55

- Weaker original effects in social psychology
- More within-subject, repeated measures designs in cognitive psychology

Results by %Replicated ($p \leq 0.05$)

- Main effects more successful than interactions

Effect Type	%Replicated ($p \leq 0.05$)	Number Replicated
Main Effect	47%	23/49
Interaction Effect	22%	8/37

Results by Correlation with replications ($p \leq 0.05$, original direction)

- **Surprising effects** were less reproducible ($r = -0.244$)
- **Challenging experiments** less reproducible ($r = -0.219$)
- **Original result importance** had little effect ($r = -0.105$)
- **Team experience and expertise** had almost no effect
 - Original ($r = -0.072$); Replication ($r = -0.096$)
- **Replication quality** had almost no effect ($r = -0.069$)

- **Larger original effect sizes** were more reproducible ($r = 0.304$)
- **Larger replication effect sizes** were more reproducible ($r = 0.731$)
- **More powerful replications** were more reproducible ($r = 0.731$)

Summary

- Even though the replications:
 - Used materials from original authors
 - Were reviewed in advance for methodological fidelity
 - Had high statistical power to measure original effect size
 - **replications produced weaker evidence for original findings**
- The strength of initial evidence (p value, effect size)
 - predicted replication success
- The characteristics of the teams, and the original finding
 - no impact on replication success

Why so few replications?

- **Publication, selection, reporting biases**
 - effect sizes of original studies inflated
- **Replications**
 - All results reported
 - no **publication bias**
 - All confirmatory tests based on pre-analysis plans
 - no **selection, reporting bias**
- Lack of biases likely big part of the reason

What Does it Mean?

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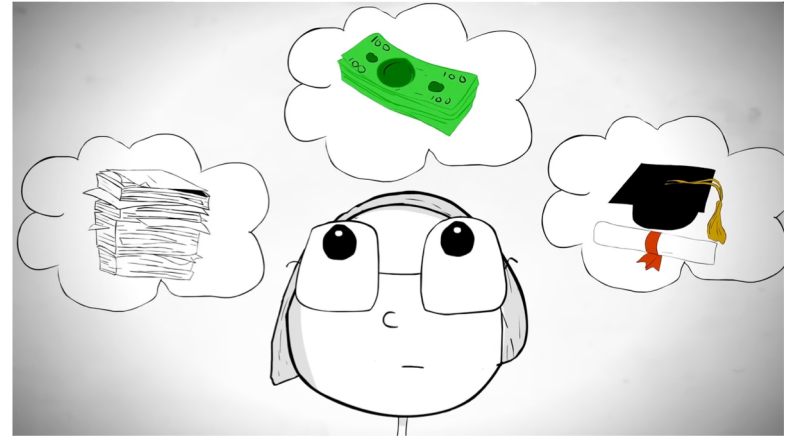
Reasons for Irreproducibility

- A study **finds A**, but the replication study **does not find A**. Why?
 1. The original study study is wrong → **A** is not true
 2. The replication study is wrong → **A** is true
 3. Both original and replication study are correct → **A** could be true or false
- How could #3 be the case?



Reasons for Irreproducibility

- First impressions are often false
- Can be hard to detect difference between real result and noise
- If enough hypothesis tests are conducted, can usually find something
 - Can be controlled by adjusting familywise α level [Howell 2002, ch 12]
- Incentive structure of science does not maximize yield of true results
 - Incentives result in many exploratory studies
 - True for every field of science
- If a finding is spurious, won't find evidence until replication is attempted



Considering Reproducibility

- A study **finds A**, and the replication study **finds A**.
What does this mean?
 - **A** is a reliable finding
- What about theoretical explanation for **A**?
 - Explanation might still be wrong
- Understanding the reasons for **A** requires multiple investigations
 - Provide converging support for the true theory
 - Rule out alternative, false theories



How Many Studies Should Be Reproducible?

- Is 36% reproducibility too small?
- What would 100% reproducibility mean?
- Progress requires both
 - **Exploratory studies**: innovative, new ideas
 - **Confirmatory studies**: replications
- Innovation points out ideas that are possible
- Replication points out ideas that are likely
 - **Progress requires both**
- Scientific incentives—**funding, publication, awards, advancement**—should be tuned to encourage an optimal balance, in a collective effort of discovery

What Should We Do?

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Value (Accept) Replication Studies

- Value confirmation (replication) studies
- Value exploratory studies
 - Value studies that are well done, regardless of type or results
- Requires changing our incentive system
- Less emphasis on surprise
 - “...but rather a reduction in the available cues, which makes the reduced performance **not terribly surprising.**”
 - “...this experiment tells us something important about depth perception in AR, **most of which isn't especially surprising,** it is not clear that this will help very much...”
 - “**It is not entirely surprising** that participants became more accurate in ‘feedback’ condition...”

Recommendations

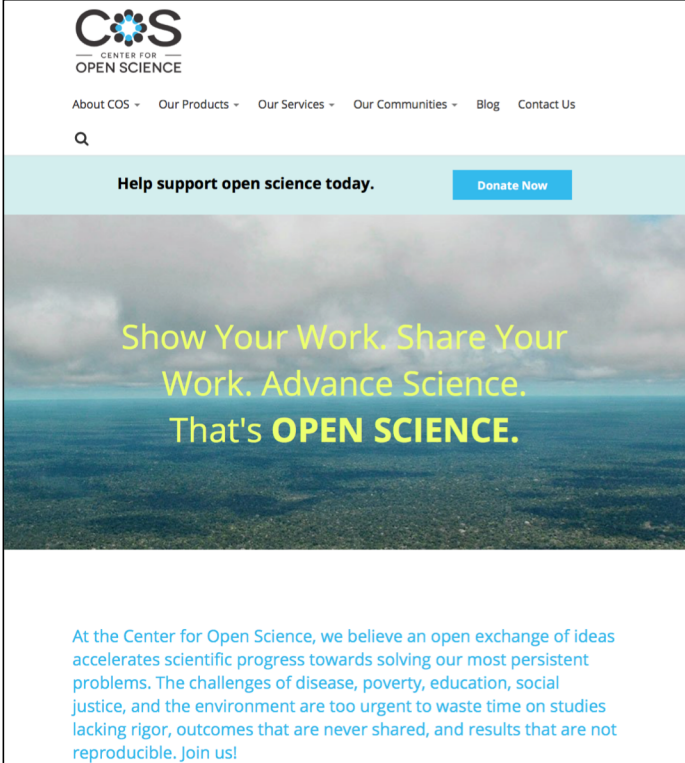
- **Value (accept) replication studies**
 - If accepted, they will come
- **Pre-register research plans**
 - Before collecting data, create detailed, written plan:
 - hypothesis, methods, analysis
 - Removes possibility of **p-hacking**
 - Even better: publically pre-register the plan
 - e.g., Center for Open Science (<https://cos.io>) → Preregistration Challenge (<https://cos.io/prereg/>)
- **Run larger studies**
 - more participants == more experimental power
 - **BUT:** more expensive

Recommendations

- **Describe methods in more detail → easier replication**
 - Problem in our field: limited pages
 - Solutions:
 - Additional details in supplementary material, or in associated thesis / dissertation
 - We could adopt longer page limits
 - Main paper in bigger font, methods in smaller font (e.g., *Nature*)
- **Upload materials to open repositories → easier replication**
 - Data, materials, code
 - Center for Open Science (<https://cos.io>)
 - Virtual Reality Knowledgebase (Doug Bowman, <https://research.cs.vt.edu/3di/node/201>)
 - arXiv, many other preprint servers
 - Other repositories...

Conclusion: Reasons for Optimism

- Current zeitgeist among **journals, funders, scientists:** paying more attention to **replication, statistical power, p-hacking, etc.**
- **In Psychology:**
 - Journals have begun publishing pre-registered studies
 - Scientists from many labs have collaboratively replicated earlier studies
- **Center for Open Science:**
 - Established 2013
 - Developing standards for transparency and openness
 - Channeling 1M USD to pre-registration challenge



The image shows a screenshot of the Center for Open Science (COS) website. At the top left is the COS logo, which consists of the letters 'COS' in a bold, sans-serif font, with a stylized blue and white flower-like icon to the right. Below the logo, the text 'CENTER FOR OPEN SCIENCE' is written in a smaller, all-caps font. To the right of the logo is a navigation menu with links: 'About COS', 'Our Products', 'Our Services', 'Our Communities', 'Blog', and 'Contact Us'. Below the navigation menu is a search icon (a magnifying glass). A light blue banner across the top of the main content area contains the text 'Help support open science today.' on the left and a blue button with the text 'Donate Now' on the right. The main content area features a large background image of a vast, green, hilly landscape under a cloudy sky. Overlaid on this image is the text 'Show Your Work. Share Your Work. Advance Science. That's **OPEN SCIENCE.**' in a yellow, sans-serif font. At the bottom of the page, there is a paragraph of text in a small, blue font: 'At the Center for Open Science, we believe an open exchange of ideas accelerates scientific progress towards solving our most persistent problems. The challenges of disease, poverty, education, social justice, and the environment are too urgent to waste time on studies lacking rigor, outcomes that are never shared, and results that are not reproducible. Join us!'

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Contact Information

J. Edward Swan II

Professor and Interim Department Head

Computer Science and Engineering

Mississippi State University

swan@acm.org

+1-662-325-7507

Slide Location:

<http://web.cse.msstate.edu/~swan/teaching/tutorials/Swan-VR2018-Replication-Crisis.pdf>