Statistical Methods for Replicability Assessment

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Introduction: Replicability Crisis

Many Psychology Findings Not as Strong as Claimed, Study Says

Bv Benedict Carev

Aug. 27, 2015











The past several years have been bruising ones for the credibility of the social sciences. A star social psychologist was caught fabricating data, leading to more than 50 retracted papers. A top journal published a study supporting the existence of ESP that was widely criticized. The journal Science pulled a political science paper on the effect of gay canvassers on voters' behavior because of concerns about faked data.

Now, a painstaking yearslong effort to reproduce 100 studies published in three leading psychology journals has found that more than half of the findings did not hold up when retested. The analysis was done by

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Introduction: Replicability crisis

Social psychology facing urgent crisis in replicability of results

Commonly attributed to varied factors

- selection for significance
- p-hacking, questionable research practices (QRPs)
- fraud
- infidelity of replication experimental designs
- flaws in original experimental designs
- "Hidden moderators": subtle, uncontrollable differences in experimental conditions

Reproducibility Project: Psychology

- Preregistered replications of 100 studies published in 2008 in three top psych. journals
- Massive collaborative effort by hundreds of researchers

Results from RP:P

RP:P reported descriptive statistics:

- 36% of replications significant in same direction as original study
- 47% of original point estimates in replication studies' 95% Cls
- ullet 83% of the effect size estimates declined $(\hat{ heta}_{i,R}/\hat{ heta}_{i,O}<1)$

Widely reported as damning result:

- Washington Post: "... affirms that the skepticism [of published results] was warranted" (Achenbach, 2015)
- Economist: "... managed to replicate satisfactorily the results of only 39% of the studies investigated" (Ano, 2016)
- New York Times: "more than half of the findings did not hold up when retested" (Carey, 2015)

Debate

What should we make of these numbers?

• e.g. 47% of original point estimates in replication studies' 95% CIs

Gilbert et al. (2016a) critiqued 47% number:

- Confidence interval \neq predictive interval
- No replication is exact $(\theta_{i,O} \neq \theta_{i,R})$
- Low fidelity of some replications (e.g. race questionnaire in Italy)
- "OSC seriously underestimated the reproducibility of psychological science"

Further debate between defenders (Anderson et al., 2016; Srivastava, 2016; Nosek and Gilbert, 2016), critics (Gilbert et al., 2016c,b)

What estimand?

Did OSC "underestimate replicability?"

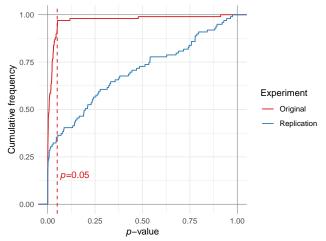
First need to answer: what is the estimand?

RP:P offers

- descriptive statistics
- tiny p-values for tests of very strong nulls
 - e.g. McNemar's test of whether orig. studies more likely to be significant at level 0.05
- no attempt to define target of inference
- no attempt to disentangle sources of error

Selection for significance

RP:P data: unmistakable sign of selection at $\alpha = 0.05$



Can this alone explain all results?

Can selection bias alone explain RP:P's descriptive statistics?

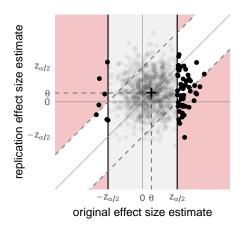
Simulation experiment

- ullet all original / replication studies: same effect size heta
- Gaussian estimators $\hat{\theta}_{i,O}, \hat{\theta}_{i,R}$, s.e. = 1
- ullet observe pair only when $|\hat{ heta}_{i,O}|>z_{lpha/2}$

Plot descriptive statistics as function of θ

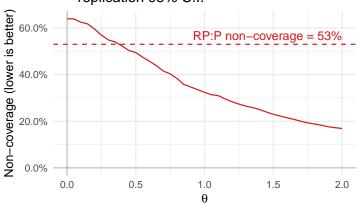
Question: with $\theta = 1/2$, what fraction of repl. Cls cover orig. estimates?

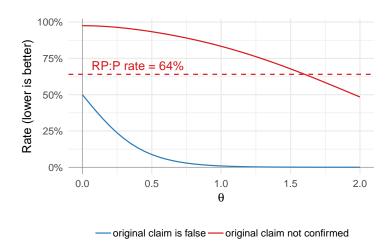
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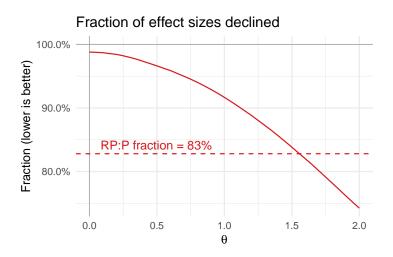


Answer: $\approx 50\%$

Non-coverage of original point estimate by replication 95% C.I.







Selection bias

Selection bias is basic feature of data

- Qualitatively, can explain RP:P metrics
- Can't learn anything else from data unless we adjust for it

Different explanations suggest different priorities for reform, e.g.:

- Selection bias: publish negative results, post-hoc stat. adjustment
- Infidelities: more detailed methods sections?
- Hidden moderators: abandon experimental psychology?

Key tools:

- conditional post-selection inference (Lee et al., 2016; Fithian et al., 2014, many others)
- ideas from multiple testing (Benjamini and Hochberg, 1995; Storey, 2002; Heller et al., 2007, many others)

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A model for replications

Model for original (O) / replication (R) study pair i = 1, ..., m:

$$\hat{\theta}_{i,O} \sim N\left(\theta_{i,O}, \ \sigma_{i,O}^2\right) \mathbf{1}_{\left\{|\hat{\theta}_{i,O}| > c\right\}} \quad \text{ and } \quad \hat{\theta}_{i,R} \sim N\left(\theta_{i,R}, \ \sigma_{i,R}^2\right) \tag{1}$$

Can formalize three definitions in terms of parameters of model (1)

• Some hypotheses defined in terms of $S_i = \operatorname{sign}(\hat{ heta}_{i,O})$

Finite population model:

- ullet no assumptions on dist. of $(heta_{i,O}, heta_{i,R})$
- agnostic to why $\theta_{i,O} \neq \theta_{i,R}$

Defining replicability

What do we estimate when we "estimate replicability?"

RP:P statistic 1: 36% of replications significant in same direction as original study

Definition 1: False directional claims

- What fraction of original directional claims were wrong?
- Psychology as enterprise in large-scale multiple testing
- Type S error: true effect 0 or opposite sign as claimed (Gelman and Tuerlinckx, 2000)
- Would it be different if we used a lower publication threshold?

Answers question: "Would an exact replication with huge n affirm the directional claim?"

Formalizing replicability

Definition 1: False directional claims

What fraction of original directional claims were wrong?

Null hypothesis for Type S error:

$$H_i^{S,O}: S_i \cdot \theta_{i,O} \leq 0$$

Directional FDP for all experiments with $p_{i,O} < \alpha$:

$$FDP_{\alpha} = \frac{\#\{i : H_i^{S,O} \text{ true}, p_{i,O} < \alpha\}}{\#\{i : p_{i,O} < \alpha\}}$$

Does it improve if $\alpha=0.005$ were used instead? (Benjamin et al., 2018)

Defining replicability

What do we estimate when we "estimate replicability?"

RP:P statistic 2: 47% of orig. point estimates in repl. studies' 95% Cls

Definition 2: Effect shift of replication

- How much do effect sizes shift from original to replication?
- Stability across direct replications
- Bare minimum form of external validity
- Identify studies where effect definitely shifted, produce CIs for shifts

Answers question: "Can psychologists successfully replicate experimental conditions?"

Formalizing replicability

Definition 2: Effect shift of replication

How much do effect sizes shift from original to replication?

Construct CIs for $\theta_{i,O} - \theta_{i,R}$

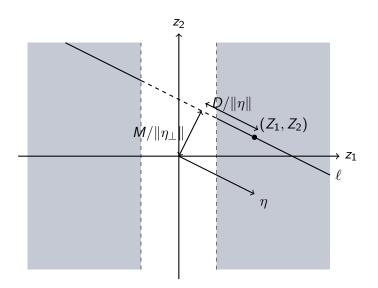
Linear in natural parameters for truncated bivariate Gaussian family

Invert selective z-test (Lee et al., 2016) of

$$H_i^{E,\delta}: \theta_{i,O} - \theta_{i,R} = \delta$$

Which / how many CIs exclude 0 (after multiplicity adjustment)?

Selective *z*-test



Defining replicability

What do we estimate when we "estimate replicability?"

RP:P statistic 3: 83% of the effect size estimates declined from original to replication

Definition 3: Overall effect decline

- What fraction of effect sizes declined by at least 20%?
- Refers to true effect sizes, not point estimates

Answers question: "Do effects systematically attenuate in replications?"

Defining replicability

Definition 3: Overall effect decline

What fraction of effect sizes declined by at least 20%?

Did replication *i* show decline by at least $\rho \in [0, 1]$?

$$H_i^{D,\rho}: S_i \cdot \theta_{i,R} \geq S_i \cdot (1-\rho)\theta_{i,O}$$

After conditioning on S_i , this is a linear hypothesis in $(\theta_{i,R}, \theta_{i,O})$

- Use Lee et al. (2016) test for individual $H_i^{D,\rho}$
- \bullet Aggregate to estimate / bound fraction that declined by ρ

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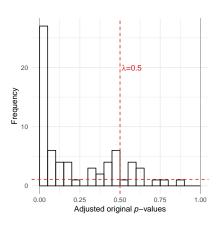
Reproducibility Project: Psychology

Pormalizing replicability

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False directional claims

Work with adjusted *p*-values: conditionally, $p_{i,O}/0.05 \ge_{\text{st}} U[0,1]$, under $H_i^{S,O}$



NB: no spike near 1 ($p_{i,O} \approx 0.05$)

Inference on FDP_{0.05}

Adjusted <i>p</i> -value	$H_i^{S,O}$ true	$H_i^{S,O}$ false	Total
$20p_{i,O} \leq \lambda$	*	*	*
$20p_{i,O} > \lambda$	U	*	В
Total	$V_{0.05}$	*	$R_{0.05}=m$

Inferences based on:

$$B \ge U \ge_{\mathsf{st}} \mathsf{Binom}(V, 1 - \lambda)$$
 (2)

 $\mathbb{E}B \ge (1-\lambda)V_{0.05}$ leads to estimator (Storey, 2002):

$$\widehat{\text{FDP}}_{0.05} = \frac{B}{R_{0.05}(1-\lambda)} = 2B/m \text{ if } \lambda = 1/2$$

(2) also gives UCB $V_{0.05}^*$, leads to UCB

$$\mathsf{FDP}^*_{0.05} = V^*_{0.05}/R_{0.05}$$

Related questions

So far, asking about Type S error:

• Does S_i is correctly describe $sign(\theta_{i,O})$? (true orig. effect)

Two related questions:

- Does S_i correctly predict sign $(\theta_{i,R})$? (true repl. effect)
- Would Type S error be better if we'd used a different threshold, e.g. 0.005?

Requires slightly more subtle methods, similar in spirit

Results: False directional claims

Note:

- Estimate FDP \approx 32% for 0.05 threshold
- For $\alpha = 0.005$, estimate is 7%, UCB 18%
- But not clear that FDP for replications is improved
- Numbers overestimate Type S error ($\theta = 0.001$ similar to 0).

Estimates /	/ Cls	for	false	directional	claims
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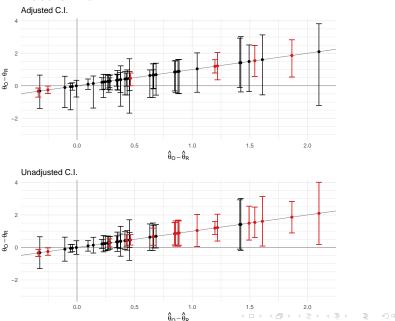
α	Orig. Est.	Orig. U.C.B.	Repl. Est.	Repl. U.C.B.
0.001	2%	9%	27%	55%
0.005	7%	18%	36%	61%
0.01	11%	22%	39%	61%
0.05	32%	47%	47%	63%

Results: Effect shift

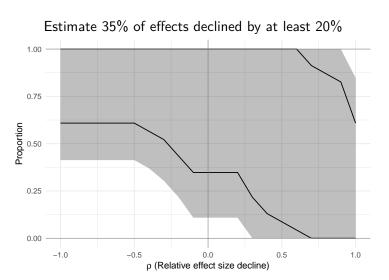
We construct 95% CIs for $\theta_{i,O} - \theta_{i,R}$:

- 15% of studies exclude 0 (exact replication), adjusting for selection
- 39% of studies rejected, otherwise
- 11% exclude 0 after BH_{0.1} correction, 1 after Bonferroni

Results: Effect shift



Results: Effect decline



Takeaways

Selection bias is a powerful force

- Leads to many predictable pathologies
- Can't learn anything else without accounting for it
- Truncated Gaussian model opens many avenues for inference

Replicability has many possible meanings

Precisely specifying estimand is essential for meaningful discussions

Rate of Type S errors in experimental psych is high: (\approx 32% of publ.?)

- Reducing threshold to 0.005 seems to improve FDP w.r.t. orig. effects
- Doesn't mean results will be replicable in new experiments

Evidence in a few studies that true effect sizes differ substantially

Some evidence of systematic effect decline (need more data)

Future work

- Preregistration
 - Better publication bias model
 - Less conservative estimate: information of nonsignificant studies are useful still!
- Higher powered design, e.g. Camerer et al. (2018); Klein et al. (2018)
- More formal criteria
- Clearer picture of the replicability crisis

Thanks!

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