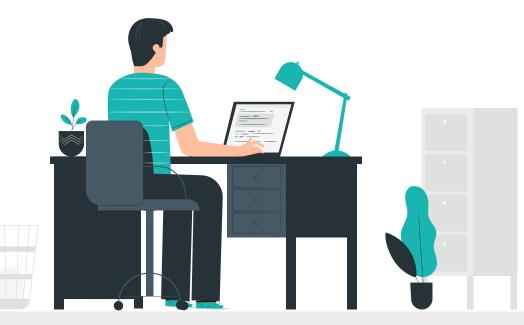
# **Registered Reports:** Why, What, & How

Melissa Thye PPLS Open Research 3<sup>rd</sup> March 2023



# Why registered reports

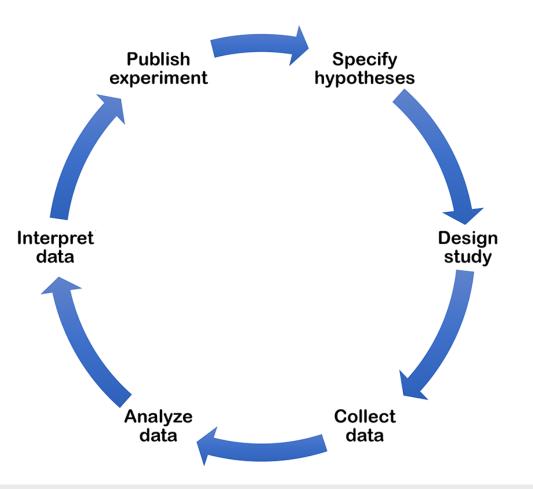


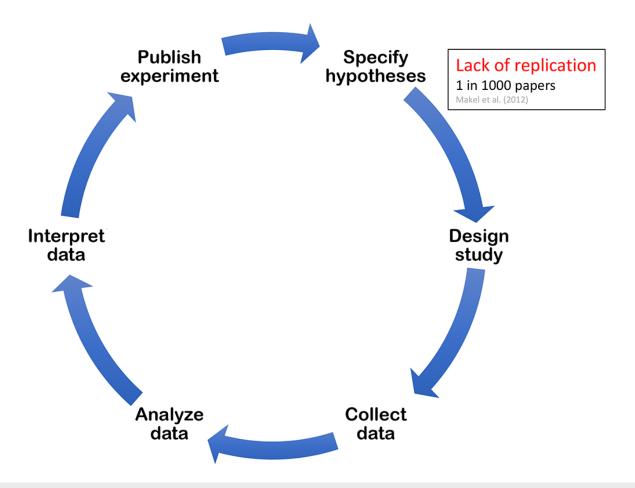
### File Drawer Problem (Publication Bias)

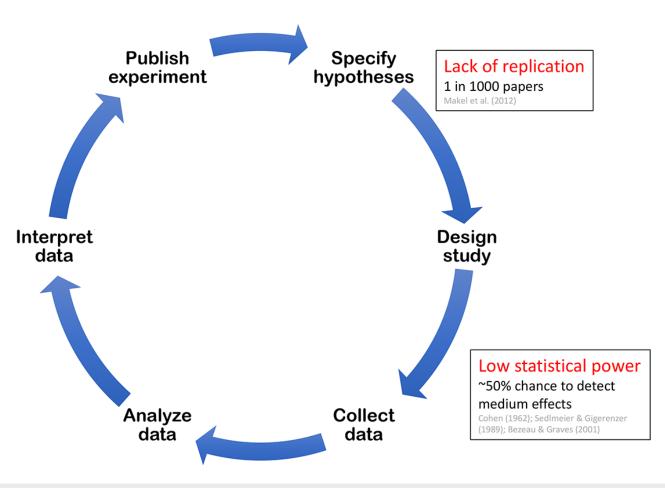
- Selective reporting of findings based on experiment outcome
- Tendency to publish positive (statistically significant) results
- Negative (non-significant) results are put into the file drawer and are unknown to other researchers
- Limits utility of meta-analytic approaches

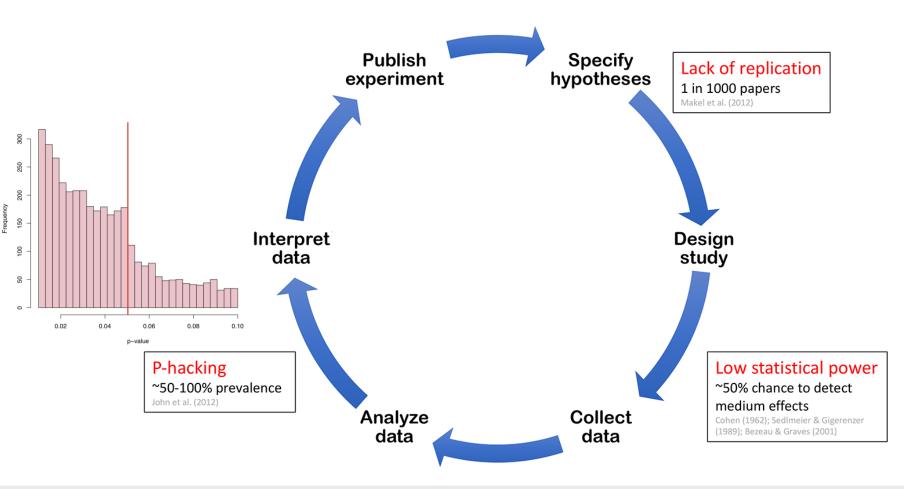
### Why?

- Rejection by journals, editors, or reviewers
- Emphasis on novelty
- Competing interests
- Viewed as a "failed" experiment

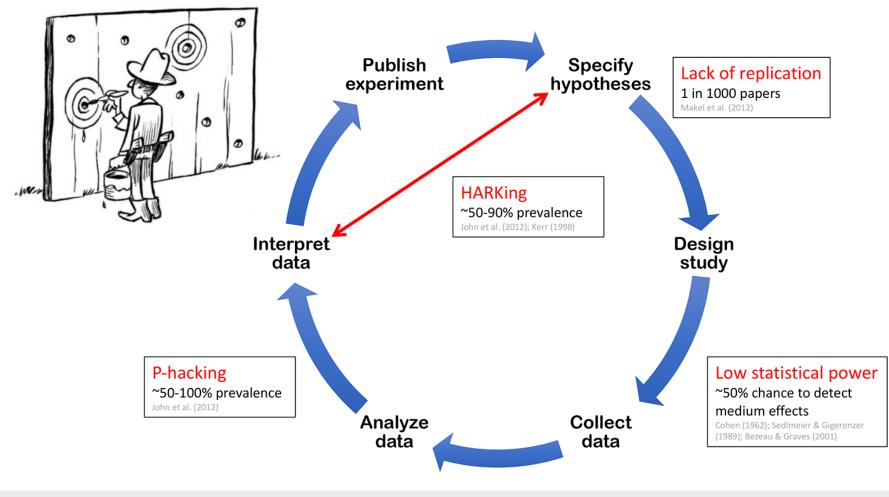


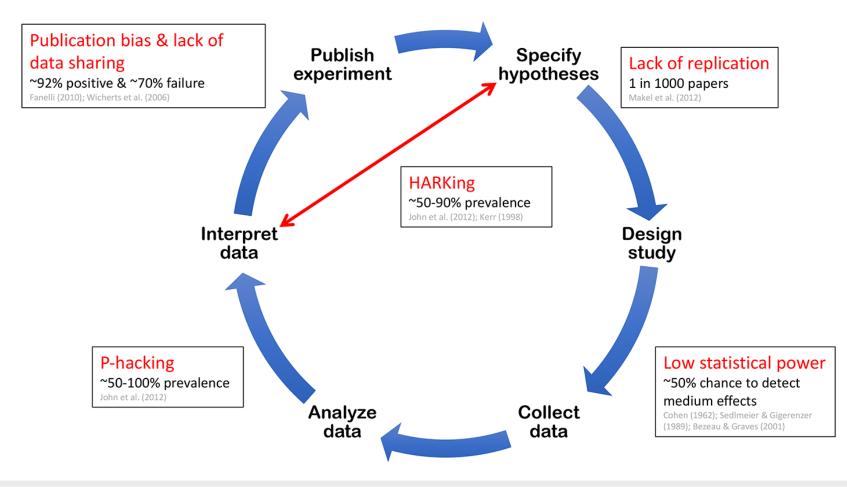






Center for Open Science Masicampo & Lalande (2012), Quarterly Journal of Experimental Psychology





# What are registered reports

### **Pre-registration**

Submitting a study plan (often including analysis plan) to a registry before conducting the study

- Increases transparency of planned versus exploratory analyses
- Reduces HARKing & P-hacking
- Time-stamped (addresses scooping concerns)

But what happens when the results are hard to publish?



# What are registered reports

### **Registered Reports**

- A peer-reviewed preregistration
- Provide sufficient detail for an independent researcher to replicate study and analysis plan
- Over 300 journals currently accept Registered Reports

- 1. Peer review occurs before outcomes are known
- 2. In-principle acceptance (IPA) will not be revoked based on outcomes







# How to do registered reports

- Primary data collection
- Secondary data analyses
  - Additional constraints on prior data access
- Systematic reviews + meta-analyses
- Allows for exploratory analyses <u>but not at Stage 1</u>
  - Any analysis that cannot be planned precisely should be withheld until **<u>Stage 2</u>**
  - $\circ~$  Be clear which analyses were pre-registered and which were exploratory
  - $\circ~$  Allows others to weigh the evidence appropriately



### Can you answer these 10 questions....

- 1. What is the main question being addressed in your study?
- 2. Can you describe the key independent and dependent variable(s), specifying how they will be measured?
- 3. What are your hypotheses?
- 4. How many and which conditions will participants/samples be assigned to?
- 5. How many observations will be collected and what rule will you use to terminate data collection?
- 6. What are your study inclusion criteria?
- 7. What are your data exclusion criteria?
- 8. What positive controls or quality checks will confirm that the obtained results are able to provide a fair test of the stated hypothesis?
- 9. Can you specify exactly which analyses you will conduct to examine the main question/hypotheses?
- 10. Are you proposing to collect new data or analyse existing data?

### ...then you're ready to do a Registered Report!

https://osf.io/93znh

### What to Include in a Registered Report

### Introduction

- Review relevant literature
- Describe research questions & hypotheses

### Methods

- Ethics information
- Pilot data (if applicable)
- Study design (all experimental procedures)
- Sampling plan
- Power analysis
- Inclusion/exclusion criteria
- How outliers will be handled
- Analysis plan
- All planned analysis (no exploratory analysis)

### What to Include in a Registered Report

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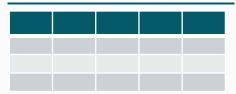
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### Data/Code Availability

### **Design Table**



# ?

### Hypotheses,

### predictions, and interpretations

GOAL: limit space of potential post-hoc interpretation

- Hypotheses should be testable and concrete
- Describe how they will be tested
- Describe how possible outcomes will be interpreted

### Power analyses and sample size

GOAL: minimize false positives and negatives

- Estimate expected effect size
- Determine required sample size with power analysis
- Identify critical tests that will be used to test predictions

# Reproducible methods and exclusion criteria plan

GOAL: control experimenter degrees of freedom

- Explicitly define variables
- Exhaustively describe inclusion and exclusion criteria
- Could someone repeat the study based on the description?

#### Is the proposed protocol doable?

- Could a simpler design address the same question?
- Is every criterion well-motivated?
- Is sample size practical?

# **Design Table**

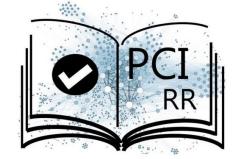
Question	Hypothesis	Sampling Plan	Analysis Plan	Interpretation given to different outcomes
RQ 1				
RQ 2				

- Gives readers an overview of analysis plans and predictions
- Number of rows will depend on number of research questions (RQ)
- Ensure exact correspondence between each hypothesis and statistical test
  - **NOT** acceptable: "Condition A will affect performance differently than Condition B"
  - Instead define: (1) the performance measure (e.g., RT), (2) the predicted direction of the difference, and (3) the anticipated effect sizes or smallest effect size of interest (if possible)

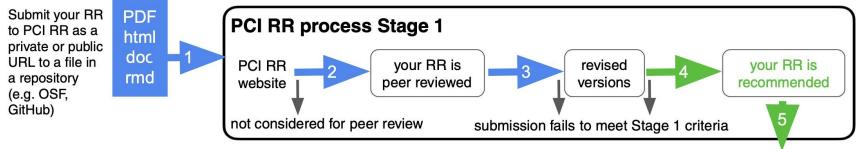
# Peer Community In Registered Reports

A community, not a journal → manages peer review of Registered Report preprints

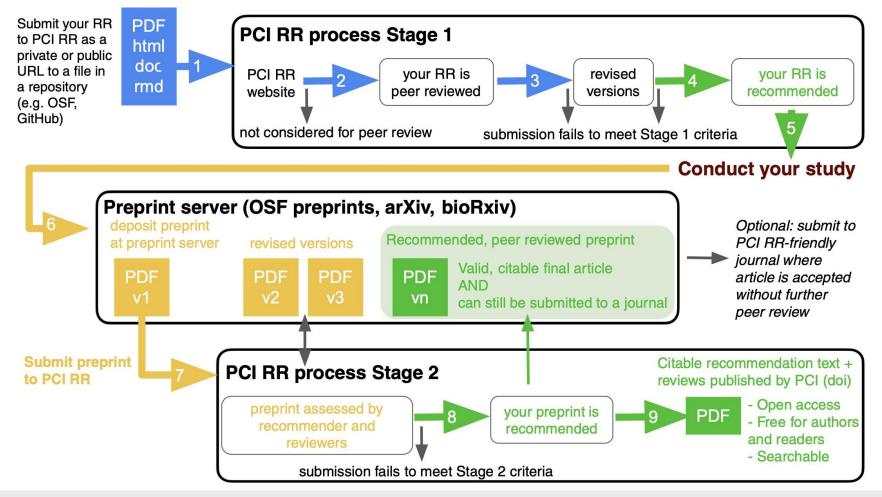
- The review process is managed by accredited recommenders (i.e., editors)
- Peer review occurs independent of journal
  - $\circ~$  Decide which journal to publish in after Stage 2 recommendation
  - Currently 28 PCI RR-friendly journals
- Peer reviews published for increased transparency
- Offer scheduled review
  - Reduces Stage 1 review from weeks to days



Free and transparent pre- and post-study recommendations across research fields



### **Conduct your study**



https://rr.peercommunityin.org/

### Pros

# Cons

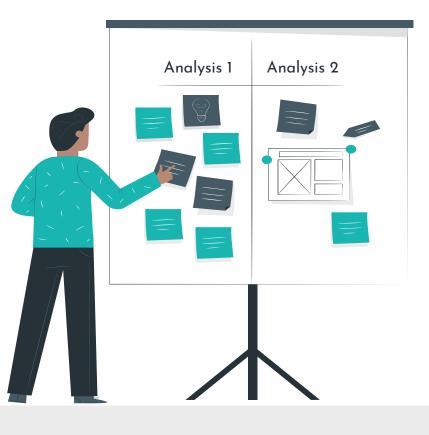
Reduces publication bias		Stage 1 IPA can take awhile		
	Pre-study review when feedback is most useful	Not all journals accept secondary data analyses		
Acceptance after IPA regardless of results		Journal policies, timelines, editing requirments vary		
	Beneficial for PhD students & ECR	Not guaranteed to get useful reviews or informed reviewers		
Increases credibility of study results		Challenging when proposing new (to you) analyses		
	Easier write-up after IPA (work frontloaded)	Significant workload shift to start of project		

# Approaches to registered reports

### **Familiar Territory**

- Confident proposing study design or methods before implementation
- Anticipate possibility of null or complex results
- Guaranteed to be added to scientific record
- Reviewer feedback is more about interpretation of outcomes and threshold for significance
- Everyone agrees with what will count as evidence for an effect

# Approaches to registered reports



### **Unfamiliar Territory**

- Less confidence in study design, methods or approach
- Anticipate that the results might be hard to publish
- Reviewer feedback might help ensure that the methods actually test the research question
- Useful for students & ECR who might be less familiar with methods
  - Thesis proposal model → hear from several experts before starting study
- Great chance to get feedback which many students want earlier in the process

## **Resources** to get started

- Pre-Registration Templates: <u>https://osf.io/zab38/</u>
- Registered Report criteria across journals: <u>https://docs.google.com/spreadsheets/d/1D4\_k-8C\_UENTRtbPzXfhjEyu3BfLxdOsn9j-otrO870/edit#gid=0</u>
- Registered Report Checklist: <u>https://osf.io/93znh</u>
- Zotero library of published Registered Reports: <u>https://www.zotero.org/groups/479248/osf/collections/KEJP68G9</u>
- Webinars (including pre-registering qualitative research): <u>https://www.cos.io/events</u>