



BERKELEY INITIATIVE FOR TRANSPARENCY  
IN THE SOCIAL SCIENCES

Implementation  
Registration  
and  
Pre-Analysis  
Plans

Garret  
Christensen

Outline

Publication  
Bias

Registrations

P-Hacking

Pre-Analysis  
Plan

Conclusion

# Registration and Pre-Analysis Plans

## Making research more transparent and reproducible

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MCA Zambia, March 2016

## Overview

- Publication bias is a problem.
- Registration can help with publication bias.
- P-hacking/specification searching is a problem.
- Pre-analysis plans can help with specification searching.
- What should we include in our PAP, and where should we post it?

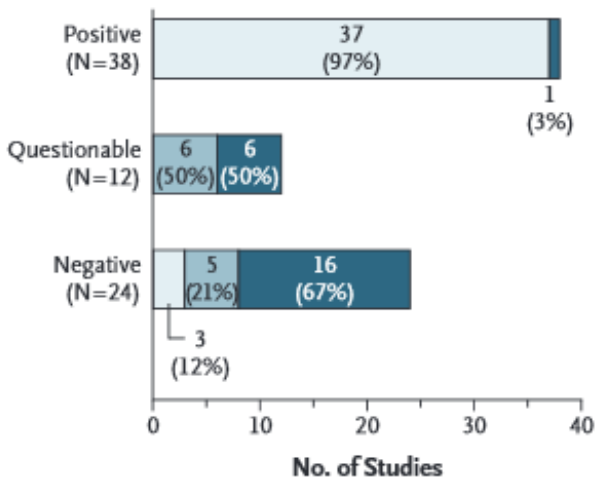
## Existence of the problem:

- Effect sizes diminish with sample size (Gerber, Green, Nickerson 2001)
- There is a higher fraction of rejected hypothesis tests in social compared to hard sciences (Fanelli 2010).
- Published null results are disappearing over time, in all disciplines (Fanelli 2011).
- Data on the complete set of experiments run shows strong results are 40pp more likely to be published, and 60pp more likely to be written up. The file drawer problem is large. (Franco, Malhotra, Simonovits 2014)

- Published, agrees with FDA decision
- Published, conflicts with FDA decision
- Not published

# A Studies (N=74)

## FDA Decision





# Publication Bias

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If we only write up/publish significant results, and we have no record of all the insignificant results, we have no way to tell if our 'significant' results are real, or if they're the 5% we should expect due to noise.



# Registrations-Medicine

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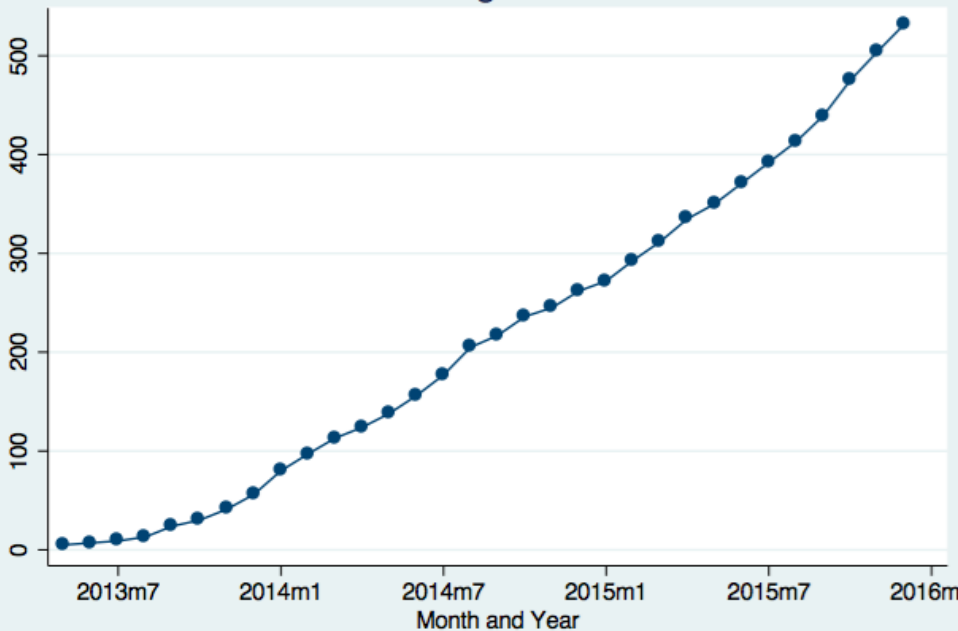
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- Publicly stating all research you will do, and what hypotheses you will test, prospectively.
- Store this statement in a public registry.
- Near universal adoption in medical RCTs. Top journals (ICMJE) won't publish if it's not registered.
- Largest: <http://clinicaltrials.gov>
- Even better if registry requires registering outcomes after study. Currently limited, and poor compliance (Anderson et al. 2015) but NIH is moving on this. [Link](#)

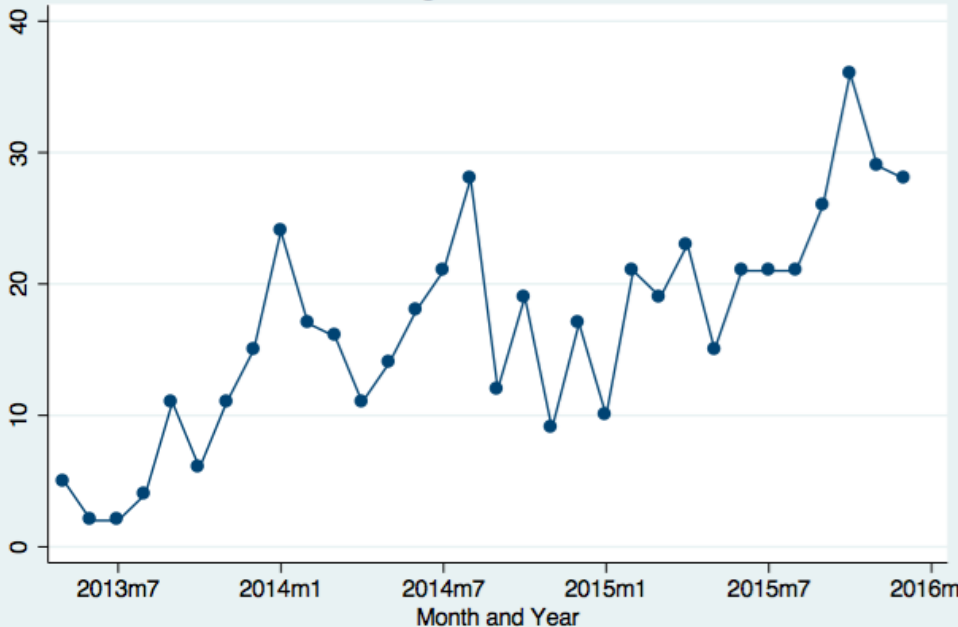
- Newer to social sciences, but good locations for several fields.
- AEA registry, currently only for RCTs.  
<http://socialscienceregistry.org>
- “J-PAL supports the American Economic Association’s (AEA) registry for randomized controlled trials in economics (<http://socialscienceregistry.org>). It is a free, easy-to-use database that makes access to trial results more transparent, aims to address the growing number of requests for registration by funders and referees, and helps solve the problem of publication bias by providing a single place where all trials are registered in advance of their start. As of May 31, 2015, the AEA Registry had a total of 379 registered controlled trials in 70 different countries. See how the registry continues to grow below.”

# Total AEA Trial Registrations over Time





## New AEA Registrations Each Month





# Registrations-Social Sciences

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- EGAP registry

<http://egap.org/design-registration>

- 3ie registry, for developing country evaluations.

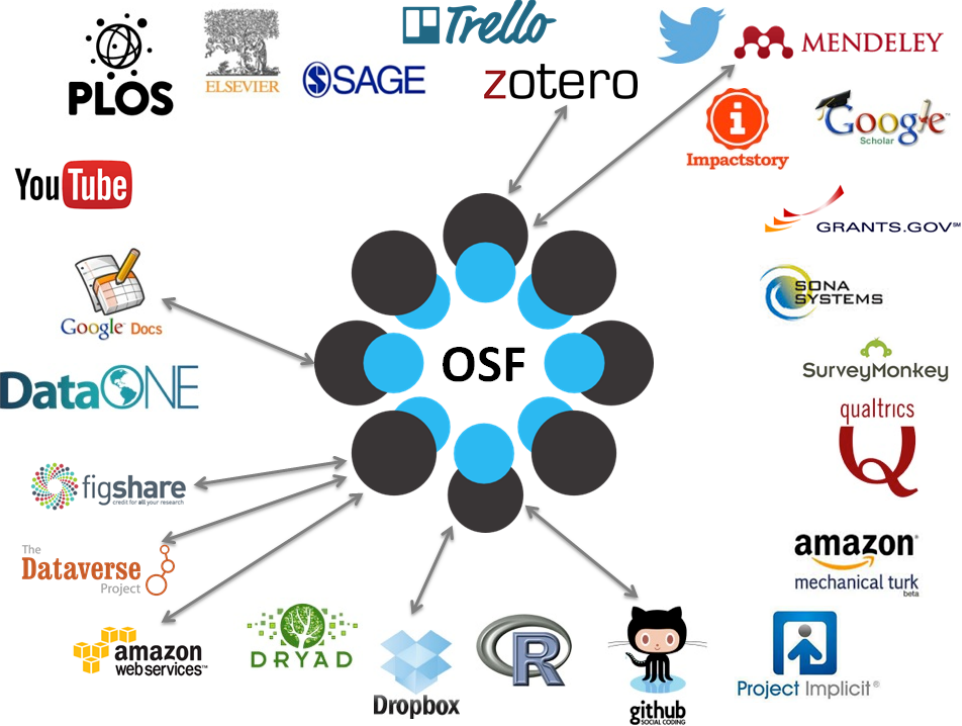
<http://ridie.3ieimpact.org>

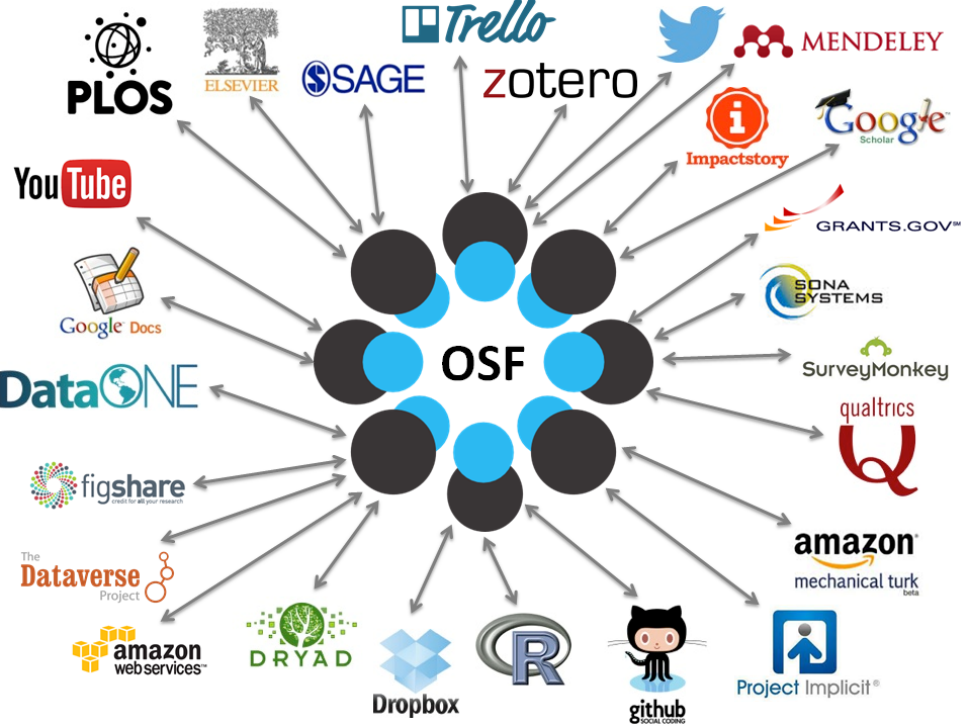
- Open Science Framework <http://osf.io>

- Open format

- Will soon sync with above

- Version Control!









# P-Hacking

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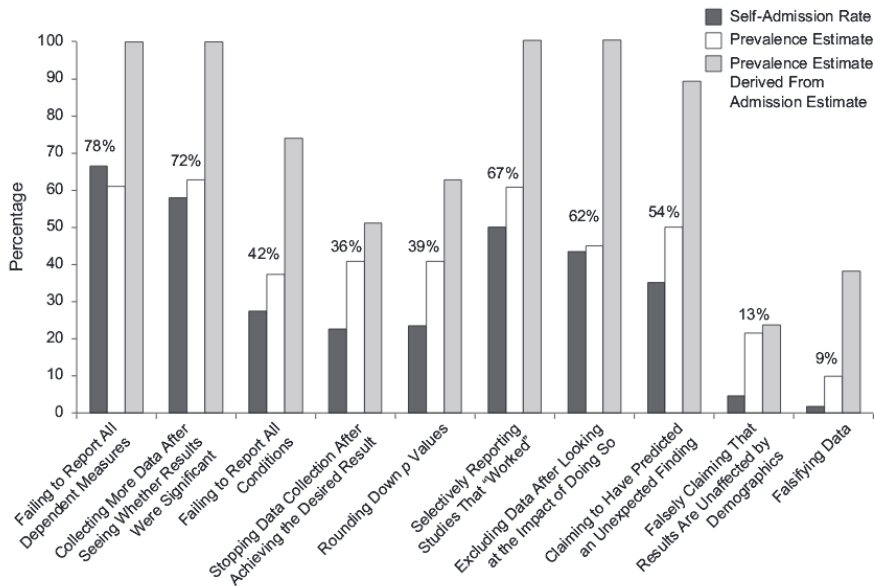
Do people actually do this? (John, Loewenstein, Prelec  
2011)

1. In a paper, failing to report all of a study's dependent measures
2. Deciding whether to collect more data after looking to see whether the results were significant
3. In a paper, failing to report all of a study's conditions
4. Stopping collecting data earlier than planned because one found the result that one had been looking for
5. In a paper, "rounding off" a  $p$  value (e.g., reporting that a  $p$  value of .054 is less than .05)
6. In a paper, selectively reporting studies that "worked"
7. Deciding whether to exclude data after looking at the impact of doing so on the results
8. In a paper, reporting an unexpected finding as having been predicted from the start
9. In a paper, claiming that results are unaffected by demographic variables (e.g., gender) when one is actually unsure (or knows that they do)
10. Falsifying data

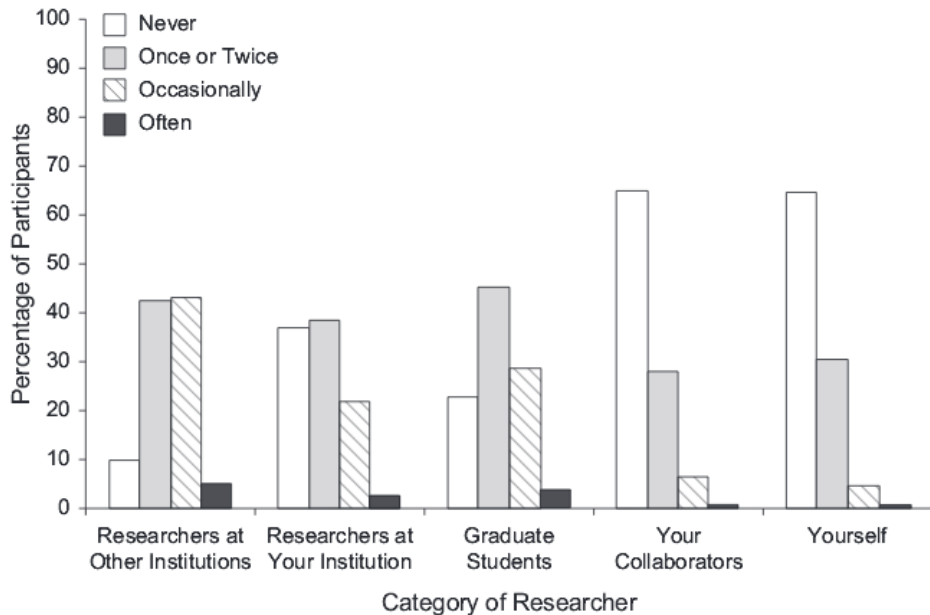
1. In a paper, failing to report all of a study's dependent measures	63.4
2. Deciding whether to collect more data after looking to see whether the results were significant	55.9
3. In a paper, failing to report all of a study's conditions	27.7
4. Stopping collecting data earlier than planned because one found the result that one had been looking for	15.6
5. In a paper, "rounding off" a <i>p</i> value (e.g., reporting that a <i>p</i> value of .054 is less than .05)	22.0
6. In a paper, selectively reporting studies that "worked"	45.8
7. Deciding whether to exclude data after looking at the impact of doing so on the results	38.2
8. In a paper, reporting an unexpected finding as having been predicted from the start	27.0
9. In a paper, claiming that results are unaffected by demographic variables (e.g., gender) when one is actually unsure (or knows that they do)	3.0
10. Falsifying data	0.6



	Admission rate	Defensibility rate
1. In a paper, failing to report all of a study's dependent measures	63.4	1.84 (0.39)
2. Deciding whether to collect more data after looking to see whether the results were significant	55.9	1.79 (0.44)
3. In a paper, failing to report all of a study's conditions	27.7	1.77 (0.49)
4. Stopping collecting data earlier than planned because one found the result that one had been looking for	15.6	1.76 (0.48)
5. In a paper, "rounding off" a p value (e.g., reporting that a p value of .054 is less than .05)	22.0	1.68 (0.57)
6. In a paper, selectively reporting studies that "worked"	45.8	1.66 (0.53)
7. Deciding whether to exclude data after looking at the impact of doing so on the results	38.2	1.61 (0.59)
8. In a paper, reporting an unexpected finding as having been predicted from the start	27.0	1.50 (0.60)
9. In a paper, claiming that results are unaffected by demographic variables (e.g., gender) when one is actually unsure (or knows that they do)	3.0	1.32 (0.60)
10. Falsifying data	0.6	0.16 (0.38)



**Fig. 1.** Results of the Bayesian-truth-serum condition in the main study. For each of the 10 items, the graph shows the self-admission rate, prevalence estimate, prevalence estimate derived from the admission estimate (i.e., self-admission rate/admission estimate), and geometric mean of these three percentages (numbers above the bars). See Table 1 for the complete text of the items.



**Fig. 2.** Results of the main study: distribution of responses to a question asking about doubts concerning the integrity of the research conducted by various categories of researchers.





BITSS

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# Pre-Analysis Plan

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- Often part of a registration
- From 3ie: “A pre-analysis plan is a detailed description of the analysis to be conducted that is written in advance of seeing the data on impacts of the program being evaluated. It may specify hypotheses to be tested, variable construction, equations to be estimated, controls to be used, and other aspects of the analysis. A key function of the pre-analysis plan is to increase transparency in the research. By setting out the details in advance of what will be done and before knowing the results, the plan guards against data mining and specification searching. Researchers are encouraged to develop and upload such a plan with their study registration, but it is not required for registration.”

- Often part of a registration
- From 3ie: “A pre-analysis plan is a detailed description of the analysis to be conducted that is written in advance of seeing the data on impacts of the program being evaluated. It may specify hypotheses to be tested, variable construction, equations to be estimated, controls to be used, and other aspects of the analysis. A key function of the pre-analysis plan is to increase transparency in the research. By setting out the details in advance of what will be done and before knowing the results, the plan guards against data mining and specification searching. Researchers are encouraged to develop and upload such a plan with their study registration, but it is not required for registration.”

## “E9 Statistical Principles for Clinical Trials” (1998) [▶ Link](#) §V Data Analysis Considerations

- 1 Prespecification of the Analysis
- 2 Analysis Sets
- 3 Missing Values and Outliers
- 4 Data Transformation
- 5 Estimation, Confidence Intervals, and Hypothesis Testing
- 6 Adjustment of Significance and Confidence Levels
- 7 Subgroups, Interactions, and Covariates
- 8 Integrity of Data and Computer Software Validity



# Glennerster, Takavarasha Suggestions

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## *Running Randomized Evaluations*

- 1 the main outcome measures,
- 2 which outcome measures are primary and which are secondary,
- 3 the precise composition of any families that will be used for mean effects analysis,
- 4 the subgroups that will be analyzed,
- 5 the direction of expected impact if we want to use a one-sided test, and
- 6 the primary specification to be used for the analysis.



## World Bank Development Impact Blog

- 1 Description of the sample to be used in the study
- 2 Key data sources
- 3 Hypotheses to be tested throughout the causal chain
- 4 Specify how variables will be constructed
- 5 Specify the treatment effect equation to be estimated
- 6 What is the plan for how to deal with multiple outcomes and multiple hypothesis testing?
- 7 Procedures to be used for addressing survey attrition
- 8 How will the study deal with outcomes with limited variation?
- 9 If you are going to be testing a model, include the model
- 10 Remember to archive it

Wide range of when to write and how detailed to make the plan. At the extreme level of detail you would have your entire code already written before you got any data.

- J-PAL Hypothesis Registry (11), see <http://www.povertyactionlab.org/Hypothesis-Registry>
- AEA Registry has relatively few, plentiful in EGAP.
- Casey, Glennerster, Miguel, “Reshaping Institutions: Evidence on Aid Impacts Using a Pre-Analysis Plan” *QJE* 2012. (Paper, Plan)
  - Government-sponsored program.
  - Broad program (Community Driven Development)
  - Broad outcomes (trust, public goods, public services, community groups, information, participation, crime, welfare, attitudes)

Outcome variable	(1) Mean for controls	(2) Treatment effect
Panel A: GoBifo “weakened” institutions		
Attended meeting to decide what to do with the tarp	0.81	−0.04 <sup>+</sup>
Everybody had equal say in deciding how to use the tarp	0.51	−0.11 <sup>+</sup>
Community used the tarp (verified by physical assessment)	0.90	−0.08 <sup>+</sup>
Community can show research team the tarp	0.84	−0.12 <sup>*</sup>
Respondent would like to be a member of the VDC	0.36	−0.04 <sup>*</sup>
Respondent voted in the local government election (2008)	0.85	−0.04 <sup>*</sup>
Panel B: GoBifo “strengthened” institutions		
Community teachers have been trained	0.47	0.12 <sup>+</sup>
Respondent is a member of a women’s group	0.24	0.06 <sup>**</sup>
Someone took minutes at the most recent community meeting	0.30	0.14 <sup>*</sup>
Building materials stored in a public place when not in use	0.13	0.25 <sup>*</sup>
Chiefdom official did not have the most influence over tarp use	0.54	0.06 <sup>*</sup>
Respondent agrees with “Responsible young people can be good leaders” and not “Only older people are mature enough to be leaders”	0.76	0.04 <sup>*</sup>
Correctly able to name the year of the next general elections	0.19	0.04 <sup>*</sup>

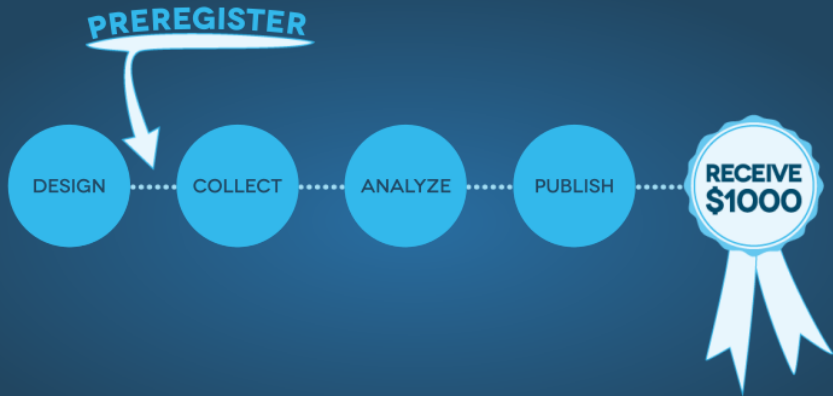


## Conclusion

- It's good science; you can distinguish between confirmatory and exploratory analysis.
- You get a badge.
  - Project Page. [Link](#)
  - Fertile women aren't more racist (Hawkins, Fitzgerald, Nosek 2015).
- You get \$1000. [Link](#)



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- Register your work to reduce publication bias.
- Include a pre-analysis plan to reduce researcher degrees of freedom.
- Register in most appropriate site for your work.
  - AEA
  - EGAP
  - RIDIE
  - OSF will hold anything, and link to your entire workflow.