### Towards a reliable and cumulative psychological science

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A little bit about the problem, and then on to some solutions.

One of the bigger problems of *p*-values is their use as a threshold to publish.



n valua

### And this is where we put the nonsignificant results.

### Most published findings confirm the hypothesis (Fanelli, 2010)



# Non-significant studies should be expected:

### 0.8×0.8×0.8×0.8=0.41

As long as a research area doesn't share all results, it's not a quantitative science.

#### Study Registry



Enabling research excellence

Promoting excellence in parapsychological research and education Koestler Parapsychology Unit

### Cosych FileDrauer Archive of Replication Attempts in Experimental Psychology

#### ClinicalTrials.gov PRS Protocol Registration and Results System

#### + All Trials

There can be 200 published studies with p < 0.05, but no true effect.

## Publication bias can not be corrected, but it can be detected.

#### Hagger et al, 2010



Effect size

# P-curve Analysis





# What do *p*-values look like from 100 studies with an effect size of 0?



# What do *p*-values look like from 100 studies with a true effect?



Small *p*-values are more likely

# P-curve analysis: Test whether a set of *p*-values has evidential value.

## Key to the filedrawer: Test is only performed on *p* < 0.05!

### *P*-curve.com



Does the *p*-value distribution look like one with or one without an effect?



## Looking at Elderly Priming and Professor Priming.

#### **P-Curve Elderly Priming**



#### **P-Curve Professor Priming** 60 50 **Percentage of p-values** 00 10 20 100 0.01 0.02 0.03 0.04 0.05

•••null of zero effect –•null of 33% power –observed

You can use this technique for small sets of *p*-values (with care!).

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#### Reading Literary Fiction Improves Theory of Min

#### David Comer Kidd<sup>\*</sup>, Emanuele Castano<sup>\*</sup>

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#### Table 1. RMET and DANVA2-AF analyses.

Experiment	Independent variable	Test	Р	ω <sub>p</sub> ²
Exp. 1 RMET	Condition	$F_{1,82} = 6.40$	0.01	0.05
	Author Recognition Test	β <b>= 0.36</b>	0.0003	0.13
	Author Recognition Test x Condition	$F_{1.82} = 1.06$	0.30	0.00
Exp. 2 DANVA2-AF	Condition	$F_{2,108} = 2.57$	0.08	0.02
	Author Recognition Test	$\beta = -0.16$	0.08	0.01
	Author Recognition Test x Condition	$F_{2.108} = 1.17$	0.31	0.00
Exp. 3 RMET	Condition	$F_{1,65} = 4.07$	0.04	0.04
	Author Recognition Test	$\beta = -0.01$	0.90	-0.01
	Author Recognition Test x Condition	$F_{1.65} = 0.01$	0.90	-0.01
Exp. 4 RMET	Condition	$F_{1,68} = 4.39$	0.04	0.04
	Author Recognition Test	β <b>= 0.39</b>	<0.001	0.15
	Author Recognition Test x Condition	$F_{1,68} = 1.50$	0.22	0.00
Exp. 5 RMET	Condition	$F_{2,352} = 3.10$	0.04	0.01
	Author Recognition Test	β <b>= 0.28</b>	<0.001	0.07
	Author Recognition Test x Condition	$F_{2,352} = 1.37$	0.25	0.00



# A theory might be true, the data just don't provide evidence for it.

# P-curve tells you if significant *p*-values look more like a true or null effect.

# Sample Size Justification

## How do you determine the sample size for a new study?

Small samples have large variation, more Type 2 errors, and inaccurate estimates.



# Power failure: why small sample size undermines the reliability of neuroscience

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Do Studies of Statistical Power Have an Effect on the Power of Studies?

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Studies in psychology often have low power. Estimates average around 50%.

Cohen, 1962; Fraley & Vazire, 2014

One reason for low power is that people use heuristics to plan their sample size.

You need to justify the sample size of a study. What goal do you want to achieve?

#### Correlation = 0.7



#### Correlation = 0.61



### **Planning for accuracy** Select a sample size based on the width of the confidence interval

Maxwell, Kelley, & Rausch, 2008

#### P-value Distribution with 50 % Power



#### P-value Distribution with 95 % Power



Planning for power Select a sample size based on probability of finding p < 0.05.

**Planning for power** Yesterday's "Sample size in psychology research" workshop



Take care when using effect sizes from the literature. Publication bias inflates effects.

If effect sizes are uncertain sequential analyses let you look at data as it comes in.

**Optional stopping:** Collecting data until p < 0.05 inflates the Type 1 error.

### A user of NHST could **always** obtain a significant result through optional stopping.

Wagenmakers, 2007

#### **Optional stopping with 5 looks**



Sequential analysis controls Type 1 error rates (e.g., Pocock correction).

Because of the substantial savings in the expected number of observations effected by the sequential probability ratio test, and because of the simplicity of this test procedure in practical applications, the National Defense Research Committee considered these developments sufficiently useful for the war effort to make it desirable to keep the results out of the reach of the enemy, at least for a certain period of time. The author was, therefore, requested to submit his findings in a restricted report [7] which was dated September, 1943.<sup>3</sup> In this

### Wald, 1945



<b>Pocock Boundary</b>			
Number of	<i>p</i> -value		
analyses	threshold		
2	0.0294		
3	0.0221		
4	0.0182		
5	0.0158		

### **Increase power by:** 1) Decreasing measurement error

**Increase power by:** 2) Using within designs (when within-measure correlation > 0.5)

**Increase power by:** 3) Increasing variability (e.g., use 7 or 9 instead of 3 or 5 answer scales)

### **Increase power by:** 4) Use one-sided tests (if you have a directional prediction)









### Share non-significant results (for novel and replication studies).

### Share data, code and materials to make science more efficient 불**PRO** INITIATIVE for open science





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### Thanks!

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