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MetaLab: A platform for cumulative meta-meta-analyses

- <sup>2</sup> Christina Bergmann<sup>1</sup>, Sho Tsuji<sup>2</sup>, Page Piccinini<sup>3</sup>, Molly Lewis<sup>4</sup>, Mika Braginsky<sup>5</sup>, Michael C. Frank<sup>4</sup>, & Alejandrina Cristia<sup>1</sup>
- <sup>4</sup> Ecole Normale Supérieure, PSL Research University, Département d'Etudes Cognitives,
- Laboratoire de Sciences Cognitives et Psycholinguistique (ENS, EHESS, CNRS)
- <sup>2</sup> University of Pennsylvania, Department of Psychology
- <sup>3</sup> Ecole Normale Supérieure, PSL Research University, Département d'Etudes Cognitives,
- 8 Neuropsychologie Interventionnelle (ENS, EHESS, CNRS)
- <sup>4</sup> Stanford University, Department of Psychology, Language and Cognition Lab
- $^{5}$  Massachusetts Institute of Technology, Department of Brain and Cognitive Sciences

Author Note

Correspondence concerning this article should be addressed to Christina Bergmann,
Ecole Normale Supérieure, Laboratoire de Sciences Cognitives et Psycholinguistique, 29, rue
d'Ulm, 75005 Paris, France.. E-mail: chbergma@gmail.com

## MetaLab: A platform for cumulative meta-meta-analyses

#### 6 Todo list

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- shorten introduction
- remove redundancies
  - Collapse over non-independent rows (?)
- Suggestions I am not sure how to implement:
- Show how infants and children are inherently a more "noisy" population
- 22 Would exclusion rates do the trick?
- Re title comments: I went with the title we submitted in the proposal
- Psychology has recently seen a "crisis of confidence", as recent findings challenge both
- the validity of key findings (Klein et al., 2014) as well as the general replicability rate of
- <sup>26</sup> published findings CITE-MANYLABS2. In this process, some practices have been discussed
- 27 as introducing bias and error, starting at data collection, over analysis to publication,
- <sup>28</sup> (Ioannidis, 2005), SCIUTOPIA1, 2. In other words, psychology is today facing the same issues
- that caused substantial changes in research practices within the medical sciences a decade
- ago (De Angelis et al., 2004). The present paper discusses to what extent these issues are
- 31 present in developmental psychology, with a focus on early studies of language
- comprehension as one example of a largely consistent subfield of child development research.

## Relevance of the confidence crisis for studies on child development

The problems underlying recent confidence crises are thought to be true of empirical sciences at large, given the current reward structure CITE-UTOPIA. Specifically, at present researchers are valued on the basis of the quantity and (some measure of) impact of their publications, and publication in a high-impact venue is dependent, among other factors, on (a) the topic being "hot"; (b) the result being surprising; and (c) the result being statistically significant. One of the obvious consequences of this reward structure is that replications and

even conceptual extensions are less likely to be undertaken (since they are not rewarded as much), and if they are, they are unlikely to be published, particularly if they reveal a non-significant result. Furthermore, modeling suggests that these issues may be exacerbated depending on the characteristics of a given subfield (Ioannidis, 2005), specifically in fields where both the underlying effect sizes and typical sample sizes are small.

All of these descriptions are highly relevant to developmental studies, particularly those focusing early childhood. Since the population under investigation is costly to recruit and difficult test, there is a pressure towards small sample sizes. Moreover, the sample is typically less consistent, as children differ a great deal in their individual developmental trajectory [CITATION?], leading to larger variance and consequently smaller underlying effect sizes. At the confluence of these two factors, one expects to find habitually underpowered studies, which is problematic both when assessing whether an effect is truly present or absent, and when estimating its magnitude for theory building and study planning.

There is a conceptually separate set of issues, which is nonetheless relevant given the aforementioned warning signs for developmental psychology. One of the habits that has become under attack recently pertains to flexibility in data collection and analysis, as arguably this flexibility exacerbates the incidence of false positives for various reasons (Ioannidis, 2005). [More details here? Avoid redundancy with later section.]

In the next sections, we discuss key issues that might undermine the reliability of child development studies for the reasons mentioned above, followed by a principled assessment of possibly questionable practices in language development research as an example case. The use of a large-scale meta-analytic dataset with unprecedented detail and standardization allows for rich analyses that can diagnose the state of a whole subfield and make principled recommendations for future research. The crucial difference between the present work and single studies resides in the quantitative assessment of study outcomes, instead of a binary result, namely whether the null hypothesis can be rejected with sufficient confidence or not. To preview our results, we find [xxx fill in when results are final xxx]. We end by describing

how our the investigations conducted here can be extended to other topics of developmental research.

# 69 Key concepts for evaluating the state of a field

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# suggestions for a different header welcome

**Replication and replicability.** Replicability is a core concept in the recent crisis, 71 as exactly this property of scientific studies (and potentially whole sub-fields) has turned out 72 to be surprisingly problematic. We define the concept here, as authors vary in their 73 understanding of what constitutes a replication. Replicating a study means (in the context of this paper) conducting a conceptually similar experiment with new stimuli and in a 75 slightly different population but following the same procedure and analyses (based on the published report), tapping into the same phenomenon, and with the same outcome as 77 previously reported (allowing for a margin of error). Being able to (repeatedly) successfully replicate a study can be taken as an indicator that the phenomenon under investigation is true and therefore that theories can be built on it. In addition, varying populations and using different, yet comparable stimuli, assesses generalizability across presumably irrelevant 81 dimensions. If an effect does not generalize across stimuli or populations, it is possible that a 82 previously unknown limitation has been uncovered, which needs to be specified for future 83 replications and within all theories building on the general effect. It is alternatively possible that the initially reported effect does not exist, in that case most attempts to replicate, especially with appropriate power (as discussed in the next section) given the expected size of an effect, should fail. Replicability can be assessed when aggregating all studies that aim to tap into a given phenomenon and assessing whether – taking all evidence together – the effect is statistically different from 0. A next step consists of comparing the direction and magnitude of initial reports and replications.

Effect sizes. The classical experimental result is measured in terms of a binary distinction: Was the *p*-value significant or not? Beyond answering that simple question,

p-values cannot offer a quantification of a phenomenon nor can they directly be compared.

However, both properties are desirable, especially in a developmental context. We would like
to address questions such as how much children improve as they mature or when an ability
emerges. Comparing significant and non-significant results to determine the onset of an
ability is not a suitable use of p-values. The null hypothesis, for example that two groups do
not differ, is in fact not being tested, so it cannot be considered confirmed when observing a
high p-value. Instead, p-values can only support rejections of the null hypothesis with a
certainty that the data at hand are incompatible with it below a pre-set threshold.

Effect sizes quantify a phenomenon by standardizing differences between groups (either 101 of observations in within-participant designs or of participants) by the variance of each 102 group. Thereby, statements such as improvements with age become statistically founded. 103 Two counter-intuitive fact about effect sizes should be noted. First, a large effect can coincide with a non-significant p-value, for example in the presence of large variance around 105 the mean and conversely a significant p-value might indicate a small effect, provided the estimate is sufficiently precise. Second, and somewhat related, numeric differences between 107 group means do not determine effect sizes. Again, the variance around those means plays a 108 crucial role, scaling the effect size accordingly. 109

Sample size and statistical power. Power refers to the probability of detecting
an effect and correctly rejecting the null hypothesis given that it is present in a population.
The larger the sample size, the higher power. Due to its dependence on the effect size under
investigation, and given that infant and child studies are expected to be more noisy
compared to adult experiments, we expect lower effect sizes. In tandem with the constraints
placed upon researchers in testing this sensitive population, it can be assumed that
underpowered studies of child development are the norm.

In developmental psychology, reasons for sample size decisions are rarely reported, it is

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<sup>&</sup>lt;sup>1</sup>The effect size defined here is one of three types, we focus on standardized mean differences for clarity, but note that other paradigms might require a different effect size family.

thus unclear how (or if) decisions about sample size are made before commencing data collection. Formal prospective power calculations are as yet rare, especially those based on multiple studies. Alternatively, it is possible that resource limitations determine sample size, as recruitment can be difficult and is very costly.

Underpowered studies specifically pose a problem for branches of developmental studies 122 that interpret both significant and non-significant findings; for example when tracking the 123 emergence of an ability as children mature or when examining the boundary conditions of an 124 ability. Even in the most rigorous study design and execution, null results will occur ever so 125 often; for example in a study with 80% power (a number typically deemed sufficient), every 126 fifth result will not reflect that there is a true effect present in the population. Disentangling 127 whether a non-significant finding indicates the absence of a skill, random measurement noise, or the lack of experimental power to detect this skill reliably and with statistical support is impossible based on p-values. 130

A second problem emerges when underpowered studies yield significant outcomes, as 131 the effects reported in such cases will be over-estimating the true effect. This makes 132 appropriate planning for future research which aims to build on this report more difficult, as 133 sample sizes will be too small, leading to null-results which do not speak to the phenomenon 134 under investigation. This poses a serious hindrance for work building on seminal studies, 135 including replications across languages and extensions. However, aggregating over such 136 null-results using a graded estimate, i.e. a standardized effect size, can reveal whether a 137 phenomenon is present in the population and correct for the initial over-estimation. In short, 138 even a true positive result is insufficient in the quest for the truth when it is underpowered. 139

To investigate the status quo, we first compute typical power across a range of
phenomena in early language acquisition, and explore which effect sizes are detectable with
sample sizes in the included studies. Further, we investigate how researchers might
determine sample sizes (for example by following the first paper in a literature), and whether
they take into account sensitivity of methods used.

**Procedural variability.** Improving our procedures can be considered both an 145 economical and ethical necessity, because our population is difficult to recruit and test. For 146 this reason, developmentalists often "tweak" paradigms and develop new ones to increase 147 reliability and robustness, all with the aim of obtaining a clear signal. Especially given the 148 time constraints, we aim for a maximum of data in the short time span infants and children 149 are willing to participate in a study. Emerging technologies, such as eye-tracking and tablets, 150 have been eagerly adopted (M. C. Frank, Sugarman, Horowitz, Lewis, & Yurovsky, 2016). As 151 a result, multiple ways to tap into the same phenomenon have been developed, consider for 152 example the fact that both headturn-based paradigms and the measurement of eye 153 movements have been employed to measure infant-directed speech preference (Dunst, 154 Gorman, & Hamby, 2012), Manybabies 1. It remains an open question to what extent these 155 different methods lead to comparable results. It is possible that some are more robust, but it remains difficult to extract such information based on studies that use different materials and test various age groups. Aggregating over experiment results, however, allows us to 158 extract general patterns of higher or lower noise via comparison of effect sizes, which are 159 directly affected by the variance of the measurement. 160

We will assess in how far the different methods used to test the same construct vary in their sensitivity. Further, taking possible resource limitations into account, we consider drop-out rates as a potential measure of interest and discuss whether higher exclusion rates coincide with more precise measures, yielding higher effect sizes.

P-hacking. Undisclosed flexibility during data collection and analysis is a problem independent of the availability of various methods to conduct infant studies. During data collection, a number of practices can inflate the number of significant p-values, effectively rendering p-values and the notion of statistical significance meaningless (Ioannidis, 2005). First, flexible stopping rules, including adding observations when a test statistic is "promising" or stopping data collection when a result is "significant" increases the likelihood to obtain a "significant" outcome. Another form of p-hacking is measuring several dependent

variables and conducting multiple significance tests with each variable and with a 172 combination of the variables. In developmental research, this problematic practice 173 encompasses computing several dependent variables (such as mean scores, difference scores, 174 percentages, and so on) based on the same measured data as well as selectively excluding 175 trials and re-testing the new data. Next, multiple conditions that selectively can be dropped 176 from the final report increase the number of significance tests. Finally, it is problematic to 177 post hoc introduce covariates, most prominently gender, and test for an interaction with the 178 main effect. Finally combining two or more of these strategies again inflated the number of 179 significant results. All these practices might seem innocuous and geared towards "helping" 180 an effect to emerge that the researcher believes to be real. 181

A related issue is that there is little standardization in data analyses practices. The 182 same type of data, such as eye movements, can be assessed with various statistical 183 procedures and little consensus has emerged over time methods have been applied. While 184 the availability of multiple analyses can be tempting to adopt practices that undermine the 185 reliability of results, it it worthwhile to adopt new analyses methods to reduce noise. 186 However, researchers might try out multiple statistical analyses and only report those with a 187 p-value below the significance threshold. From a single report it is not possible to assess 188 whether such analytic flexibility led to an inflation of significant results, but we use tools 189 that are based on cumulative science in the present work to distinguish between optimizing 190 informativeness and unsavory practices. 191

A "symptom" of such practices is a distribution of *p*-values with increased frequency just below the significance threshold. P-curves test for this problem, but they come with some limitations and only consider statistically significant reports [p-curve citation, limitation citation].

Publication biases. As mentioned previously, current incentives including
publication of data in a prestigious journal, are geared towards surprising and statistically
significant studies. However, even when an effect is robust and tested with sufficiently high

participant numbers, null results are expected to occur. This becomes even more pressing in a field with small effect sizes and low numbers of participants. For an accurate estimate of the true effect it is crucial to have access to all results, to avoid overestimations. However, we expect that studies remain in the file-drawer and never see the light of day. One reason is that a failure to obtain an expected significant result is often ascribed to the researcher's skill or an unknown flaw in the experiment [citation?].

We will investigate publication biases in our data with standard meta-analytic tool and further discuss how meta-analyses in general, and repositories such as MetaLab in particular, can be a "home" for null results. A cumulative view can help isolate factors that systematically lead to effect sizes closer to zero, or what is often called boundary conditions.

Prerequisite for such analyses is a systematic and consistent reporting of such factors; if they are relevant this should however be the case according to standard scientific practice.

211 Methods

#### 212 Source data: MetaLab

In this paper, we extract measures of interest from meta-analyses of child language 213 development. Meta-analyses are built on a collection of standardized effect sizes on a single, 214 well-defined phenomenon. By accumulating effect sizes and weighting them by their 215 reliability (effectively the sample size), it is possible to compute an estimate of the 216 population effect, as well as its structured variance. By harnessing data from hundreds of 217 studies, we can quantify patterns important for experimental practices. Furthermore, 218 combining multiple meta-analysis – each centered on a different research question – allows us 219 to assess whether current practices differ across different topics. 220

Given that all 11 meta-analyses we discuss in this paper focus on language acquisition in early childhood, our suggestions will be most relevant to this sub-field. We present our methods and results to researchers on developmental psychology in general to encourage others to build similar meta-meta-analyses, thus allowing them to explore the state of their

own sub-fields and to improve their practices if necessary. The analyses in this paper are
based on MetaLab, an online collection of meta-analyses on early language development.

Currently, MetaLab contains 11 meta-analyses, but it is open to submissions and updates.

The present analyses thus are a snapshot; through dynamic reports on the website, and by
downloading the freely available data, it is continuously possible to obtain the most recent
results.

In MetaLab, parts of each meta-analysis are standardized to allow for the computation 231 of common effect size estimates and for analyses that span across different phenomena. 232 These standardized variables include study descriptors (such as citation and peer review 233 status), participant characteristics (including mean age, native language), methodological 234 information (for example what dependent variable was measured), and information necessary 235 to compute effect sizes (number of participants, if available means and standard deviations of 236 the dependent measure, otherwise test statistics, such as t-values or F scores). This way, the 237 analyses presented in this paper become possible. 238

MetaLab contains datasets that address phenomena ranging from infant-directed 239 speech preference to mutual exclusivity, sampled opportunistically based on data collected 240 with involvement of (some) authors of this paper (n=9 datasets) or they were extracted from 241 previously conducted meta-analyses related to language development (n=2, i.e. Colonnesi, 242 Stams, Koster, and Noom (2010); Dunst et al. (2012)). In the former case, we attempted to 243 document as much detail as possible for each entered experiment (note that a paper can 244 contain many experiments). Detailed descriptions of all phenomena covered by MetaLab, 245 including which papers and other sources have been considered, can be found on the 246 companion website at http://metalab.stanford.edu and in the supporting information. 247

## 48 Statistical approach

As dependent measure, we report Cohen's d, a standardized effect size based on comparing sample means and their variance. This effect size was calculated when possible

from means and standard deviations across designs with the appropriate formula. When these data were not available, we used test statistics, more precisely t-values or F scores of 252 the test assessing the main hypothesis. We also computed effect size variance, which allows 253 to weigh each effect size when aggregating across studies. The variance is mainly determined 254 by the number of participants; intuitively effect sizes based on larger samples will be 255 weighted higher. Note that for research designs testing participants in two conditions that 256 need to be compared (for example exposing the same infants to infant- and adult-directed 257 speech), correlations between those two measures are needed to estimate the effect size 258 variance. This measure is usually not reported, despite being necessary for effect size 259 calculation. Some correlations could be obtained through direct contact with the original 260 authors (see e.g., (C. Bergmann & Cristia, 2016) for details), for others we estimated this 261 factor based on the information in our database. We report all details of effect size calculation in the supplementary materials. 263

Meta-analytic model. To aggregate effect sizes within a phenomenon, we used a 264 multilevel approach, which takes into account not only the effect sizes and variance of single 265 studies, but also that effect sizes from the same paper will be based on more similar studies 266 than effect sizes from different papers (Konstantopoulos, 2011).. We relied on the 267 implementation in the metafor package (Viechtbauer, 2010) of R (R Core Team, 2016). 268 Excluded as outliers were effect sizes more than three standard deviations away from the 260 median effect size within each dataset, thus accounting for the difference in median effect 270 size across phenomena. 271

Power. Based on the meta-analytical effect size and the median number of participants, we calculated typical power (using the pwr package (Champely, 2015)). We remind the reader that recommendations are for this value to be above 80%, which refers to a likelihood that 4 out of 5 studies show a significant outcome for an effect truly present in the population.

For analyses involving p-values, we re-computed p-values from our 277 effect-size estimates. This is due to two main reasons: First, we did not have the same 278 information available for all data points, even within the same meta-analysis. For example, 279 in XXX, XXX% of the effect sizes were calculated based on t-values, XXX% based on 280 group-level means and standard deviations, and XXX% based on F-scores. In addition, some 281 datasets only contain effect sizes, because they are based on extant meta-analyses. Second, 282 p-values are not always computed and reported correctly or consistently (Nuijten, Hartgerink, 283 Assen, Epskamp, & Wicherts, 2016). To ensure a consistent relationship between p-values 284 and effect sizes, we thus opted for recalculation. The recalculation pipeline is as follows: We 285 transform Cohen's d into Pearson's r, from which it is possible to calculate a t-value.

Assessing publication bias. There are numerous ways to estimate whether the published literature is biased. The most common and straightforward is an assessment of funnel plot asymmetry. A funnel plot displays effect sizes against their variance (with 0 being plotted up). The expectation in the absence of biases is that effect sizes are equally distributed around the meta-analytic mean, and that the are spread out more the larger their variance, creating a triangle-like shape. Biases can lead to distortions in this distribution.

The large the asymmetry, the more likely a bias is. We quantify funnel plot asymmetry with a rank correlation test implemented in the metafor package (Viechtbauer, 2010).

A second analysis pertains to the relationship between observed effect sizes in single studies and the associated sample size. The smaller the effect size, the larger the sample needed for a significant p-value. If sample size decisions are made before data collection and all results are published, we expect no relation between observed effect size and sample size. A significant non-parametric correlation indicates that only those studies with significant outcomes were published (Begg & Mazumdar, 1994).

Results 301

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# Measures of a typical study on early language acquisition

Table 1 provides a summary of typical sample sizes and effect sizes by phenomenon, 303 but before discussing those descriptors in detail, we begin by observing general trends. 304 Sample sizes are small across the board, with the overall median in the MetaLab database being 18. With such a sample size, and assuming a paired t-test based on within-participant comparisons (the most frequent experiment design and test statistic in MetaLab) it is 307 possible to detect an effect in 80% of all studies when Cohen's d = 0.70, in other words when 308 investigating a medium to large effect. When comparing two independent groups, this 309 number increases to Cohen's d = 0.96, a large effect. The observation concerning sample 310 sizes and which effect size could be detected is in stark contrast with the effect sizes we 311 actually observe, which tend to fall into ranges of small to medium effects. It turns out that 312 by and large, studies are underpowered. 313

Phenomena in MetaLab differ in the age groups typically tested and the age range covered, with the mean age ranging between 4.5 months (infant directed speech preference) and 2.5 years (mutual exclusivity). One might expect a relationship between effect sizes and infant age both for theoretical and practical reasons. On one hand, younger infants might show a smaller effect in general because they are not yet as proficient in their native language, having had less experience, and because they are a more immature in terms of their information processing abilities [CITE]. On the practical side, methods – a topic we will investigate in depth in an upcoming section – might be more noisy for younger infants and they could be a more difficult population to recruit.

While there is no strict linear relationship between infant age and sample size, effect 323 size, and the derived power, we observe a difference between studies typically testing infants younger than one year and those testing older infants. First, sample sizes are much lower for younger infants, which do usually not test more than 20 infants (although all datasets contain 326 studies with larger samples). This is not the case for older children. The only exception is

the dataset addressing mutual exclusivity, which habitually tests around 16 children, which is off-set by a comparatively large effect size. Additionally, the number of participants tested within each dataset ranges a great deal. This might indicate that researchers are mostly limited by their resources and participant availability in planning their studies.

Turning to effect size, we see a similar split by age group in our data. Younger infants 332 show both a greater range and include lower effect sizes which fall into the classical range of 333 small effects (Cohen's d below .5), which is not the case for older children. Power is directly related to sample size and effect size, so it is not surprising that typical power is greater for 335 older children. Interestingly, however, there seems to be little to no relationship between 336 effect sizes and number of participants typically tested. For phenomena with large effects, 337 this means that studies are very high-powered (see gaze following, online word recognition, 338 as two examples). For younger children, because sample sizes and effect sizes are both small, 339 power is habitually very low, and the only dataset which typically achieves appropriate 340 power near 80% is non-native vowel discrimination. For older children, power is solely caused 341 by lower effect sizes. The lack of a relationship between overall meta-analytic power and 342 sample size might indicate that researchers' experiment planning is not impacted by the 343 phenomenon under investigation. Studies might instead be designed and conducted with 344 pragmatic considerations in mind, such as participant availability. 345

Besides this very general point, we refrain here from strong conclusions based on the above-discussed observations, since the present dataset is not exhaustive and topics typically investigated in younger children are over-represented. However, we sampled in an opportunistic and thus to some degree random fashion and the phenomena covered span very different aspects of language acquisition and linguistic processing.

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How does effect size relate to *p*-values?. The key measure in this paper is, as
described in the Introduction, an effect size. In contrast, single experiments are often
evaluated by their associated *p*-value, despite the frequent criticisms and well-documented
shortcomings of that measure [citations]. Effect sizes are a continuous measure, while

Table 1

Descriptions of meta-analyses currently in MetaLab.

Topic	Age	Sample Size (Range)	N Effect Sizes	N Papers	Cohen's
Phonotactic learning	10.69	18 (8, 40)	47	15	0.1
Label advantage in concept learning	12.36	13 (9, 32)	48	15	0.4
Gaze following	14.24	23 (12, 63)	32	11	1.0
Online word recognition	18.00	25 (16, 95)	14	6	1.2
Mutual exclusivity	23.99	16 (8, 72)	58	19	0.8
Infant directed speech preference	4.34	20 (10, 60)	48	16	0.7
Vowel discrimination (native)	6.54	12 (6, 50)	112	29	0.6
Vowel discrimination (non-native)	7.69	16 (8, 30)	46	14	0.7
Sound symbolism	7.89	20 (11, 40)	44	11	0.2
Statistical sound category learning	8.16	14.75 (5, 35)	16	9	-0.2
Word segmentation	8.29	20 (4, 64)	284	68	0.1

p-values are largely used in a binary way, namely to either reject the null hypothesis or fail
 to do so. Figure 1 illustrates this binary distinction with a vertical line.

A common intuition might be that very high effect sizes are related to significant p-values and very low effect sizes near zero lead to non-significant p-values. For extreme values, namely effect sizes near zero and above 2, this intuition is borne out, as Figure 1 shows. However, the majority of effect sizes we observe falls in a range that with sufficient power, in other words a sufficiently large sample, can lead to a significant outcome.

Underpowered studies, in contrast, might tap into a similar sized effect but fail to reach significance. In Figure 1, the grey horizontal band illustrates such a region where both significant and non-significant results are observed. Further, 5 of the 12 meta-analytical

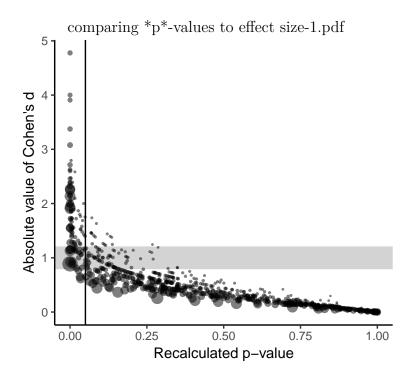


Figure 1. (#fig:Plot comparing p-values to effect size)Comparison of a study's effect size and the according p-values. Point size reflects sample size. The typical significance threshold of .05 is indicated by a vertical line.

effect sizes fall in a range where single studies are not significant, that is below d = 0.61.

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#### Comparing meta-analytic effect size and oldest paper to estimate power.

As Table 1 and Figure 1 both show, experimenters are frequently not including a sufficient 367 number of participants to observe a given effect, assuming the meta-analytic estimate is 368 accurate. It might, however, be possible, that power has been determined based on a seminal 369 paper to be replicated and/or built on. Initial reports tend to overestimate effect sizes 370 (Jennions & Møller, 2002), possibly explaining the lack of power in some datasets and studies. We extracted for each dataset the oldest paper and therein the largest reported 372 effect size and re-calculated power accordingly, using the median sample size of a given 373 dataset. The results are shown in the table below. It turns out that in some cases, such as 374 native and non-native vowel discrimination, sample size choices match well with the oldest 375 report. The difference in power, noted in the last column, can be substantial, with native 376

Table 2

For each meta-analysis, largest d from first paper and power, along with the difference between power base

Meta-analysis (MA)	Oldest d	Meta-analytic d	Sample Size	Power based on first re
Statistical sound category learning	0.56	-0.26	15	
Word segmentation	0.56	0.16	20	
Mutual exclusivity	0.70	0.81	16	
Label advantage in concept learning	0.86	0.45	13	
Vowel discrimination (non-native)	1.02	0.79	16	
Phonotactic learning	0.98	0.12	18	
Sound symbolism	0.95	0.22	20	
Online word recognition	0.89	1.24	25	
Vowel discrimination (native)	1.87	0.69	12	
Gaze following	1.29	1.08	23	
Infant directed speech preference	2.39	0.73	20	

vowel discrimination and phonotactic learning being the two most salient examples. Here,
sample sizes match well with the oldest report and studies would be appropriately powered if
this estimate were representative of the true effect.

In general, it turns out that the larger the original effect size, the more likely is an overestimation of the meta-analytic effect size. Researchers might thus be wary of reports implying a strong, robust effect with infants and toddlers in the absence of corroborating data, as is the case for online word recognition, one example where power is appropriate.

There are also datasets where sample sizes seem unrelated to either seminal reports or meta-analytic effect sizes. In the next sections we consider resources as a factor in experiment design, which might also be at play here as well.

### 7 Procedure comparison

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In this section we address how methods might be chosen, adopting two angles. We first take a pragmatic, resource-oriented approach and compare methods with respect to their dropout rate. Then we compare how effect size across phenomena is relating to method choice.

Drop-out rates across methods. Choosing a robust method can help increase the

power of studies, such that more precise measurements lead to larger effects and thus require 393 fewer participants to be tested. However, the number of participants relates to the final 394 sample and not how many infants had to be invited into the lab. We thus first quantify 395 whether methods differ in their typical drop-out rate, as the available participant pool might 396 inform method choice. To this end we consider all methods across datasets in MetaLab 397 which have more than 10 associated effect sizes and for which information on the number of 398 dropouts was reported; this information is not always available in the published report, and 399 in the case of the two meta-analyses we added based on published reports, the information 400 was not added. Therefore, the following analyses only cover 6 methods and 224 data points. The results of the linear mixed effect model predicting dropout rate by method and 402 mean participant age (while accounting for the different effects being tested) are summarized 403 in the table below. The results show that, taking the most frequently used central fixation as 404 baseline, conditioned headturn and stimulus alternation have significantly more drop-outs. 405 Figure XXX underlines this observation, and illustrates the relationship of drop-out rate 406 with age. Overall, stimulus alternation leads to the highest drop-out rates, which lies at 407 around 50% (see Figure XXX). While age is not significantly impacting drop-out rates, it 408 interacts with the different methods. We observe an increase in drop-out rates, which is most 400 prominent in conditioned headturn (a significant interaction) and headturn preference 410 procedure (where the interaction approaches significance). 411

Interestingly, the methods with lower drop-out rates, namely central fixation and headturn preference procedure, are among the most frequent ones in MetaLab and certainly

more frequent than those with higher drop-out rates, indicating that drop-out rate might inform researchers' choices. Being able to retain more participants as a factor in method choice points to the mentioned limitations regarding the participant pool, as more participants will have to be tested to arrive at the same sample size.

Table 3
(#tab:Method vs excluded)Caption.

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	Estimate	Std. Error	t value
(Intercept)	31.21	4.46	7.00
methodconditioned head-turn	30.62	5.61	5.45
methodforced-choice	-26.42	9.40	-2.81
methodhead-turn preference procedure	-2.33	4.75	-0.49
methodlooking while listening	-6.42	5.37	-1.19
methodstimulus alternation	21.34	4.10	5.21
ageC	0.42	0.44	0.95
methodconditioned head-turn:ageC	2.88	1.16	2.47
methodforced-choice:ageC	-0.22	0.65	-0.33
methodhead-turn preference procedure:ageC	0.96	0.72	1.34
methodlooking while listening:ageC	-0.57	0.80	-0.71
methodstimulus alternation:ageC	-0.26	0.91	-0.29

The effect of method choice on effect sizes (and thus power). Methods
which retain a higher percentage of participants might either be more suitable to test infants,
decreasing noise as most participants are on task, or less selective, thus increasing noise as
participants who for example are fussy are more likely to enter the data pool. We analyze
the same methods investigated above with respect to their drop-out rates here, this time
considering more data (as we do not need to rely on reported drop-out rates).

We built a meta-analytic model with the effect size measure Cohen's d as the

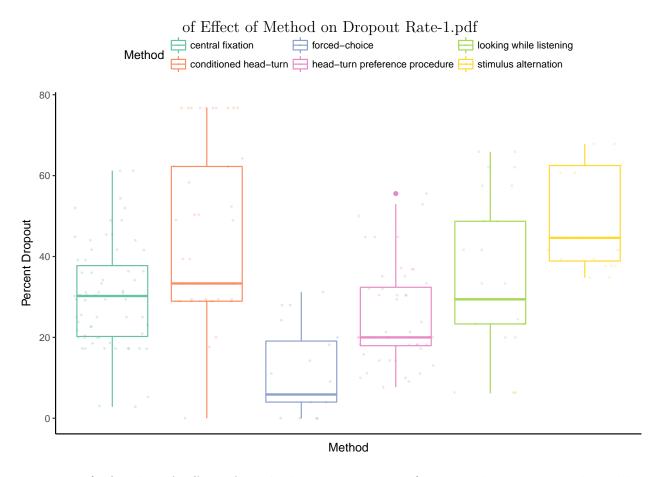


Figure 2. (#fig:Plot of Effect of Method on Dropout Rate)Percent dropout as explained by different methods.

dependent variable, method and mean age centered as independent variables. The model also includes the variance of d for sampling variance, and paper within meta-analysis as a random effect (because we assume that within a paper experiments and thus effect sizes will be more similar to each other than across papers). We again selected central fixation as baseline method.

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The model results in Table XXX show that compared to central fixation only
conditioned headturn and looking while listening yield reliably higher effect sizes, all other
methods do not statistically differ from this baseline. When factoring in age, forced choice
and looking while listening show significant interactions, indicating an improvement as
infants mature. Age is marginally above the significance threshold, indicating that as infants

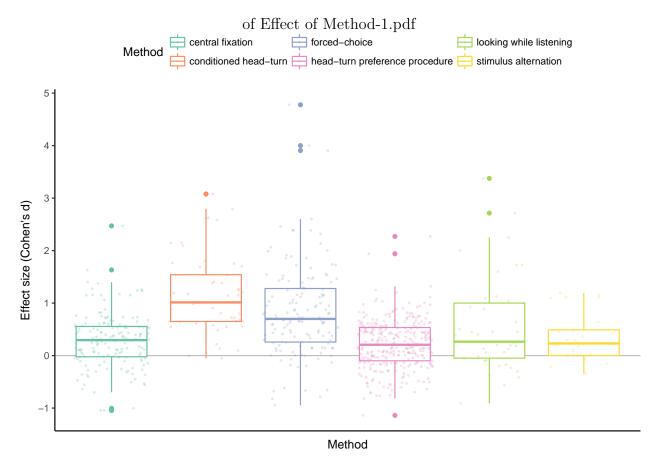


Figure 3. (#fig:Plot of Effect of Method)Effect size as explained by different methods.

mature effect sizes increase across methods – an observation consistent with the view that infants and toddlers become more proficient language users and are increasingly able to react appropriately in the lab, yielding higher effect sizes.

Comparing our analyses (Table XXX) and Figure YY in this section with those in the previous section, it seems that high drop-out rates might be offset by high effect sizes in the case of conditioned headturn. While drop-out rates are around 40-50%, effect sizes are above 1. Stimulus alternation does not fall into this pattern of high drop-out rates being correlated with high effect sizes, as the observed outcomes are in the range typical for meta-analyses in our dataset. There is an important caveat to this interpretation that some methods, specifically conditioned headturn, which have higher dropout rates, are better at generating high effect sizes due to decreased noise (e.g., by excluding infants that are not on task).

Studies with fewer participants (thanks to higher drop-out rates) might simply be underpowered, and thus any significant finding is likely to over-estimate the effect. Due to publication biases, we might not have access to all null results using the same method, and the resulting overestimation is directly reflected in our effect size estimate. As a next step, we thus quantify the possible publication biases in our data.

# 51 P-hacking

The figure below shows the distribution of p-values around the significance threshold of 452 .05 that were recalculated based on effect sizes for consistency (see Nuijten et al. (2016), cite bishop, etc?). For reliability purposes, we only discuss datasets with more than 10 p-values between 0 and .2. In the absence of questionable research practices and the presence of an effect, we expect a distribution tilted towards small values. In the absence of both p-hacking 456 and an effect, the distribution should be flat. Unexpected "bumps" towards higher p-values 457 in contrast can indicate severe p-hacking, including adding and removing samples and/or 458 predictors, and conducting multiple statistical analyses (Ioannidis, 2005). Out of the 459 datasets that we could include in this analysis, the majority exhibits the expected right-skew. 460 There are two exceptions, however, where we observe an unexpected distribution in the 461 absence of questionable research practices. 462 #to be expanded 463 http://journals.plos.org/plosbiology/article?id=10.1371/journal.pbio.1002106#sec008 464 https://peerj.com/articles/1935/https://peerj.com/articles/1715/ 465

## 66 Publication biases

Funnel plots, displaying effect sizes of single studies against their variance, show
whether observed results are evenly spread around their median. Across datasets, the
difference in distributions and range covered in effect sizes is striking, as is the variance in
observed precision (points plotted upwards close to zero). The indicated relationship
between effect sizes and their variance was assessed with a nonparametric test and turnd out

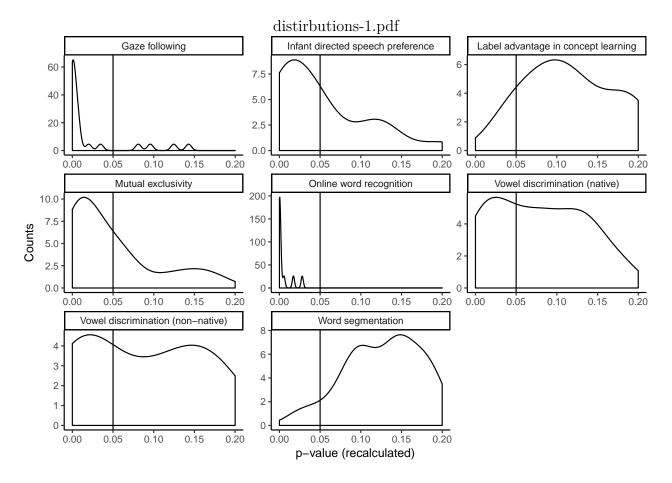


Figure 4. (#fig:P-value distirbutions)Caption.

to be significant (see supplementary materials for details) – indicating publication bias in favor of significant results – for four datasets.

Only two datasets turn out to have a significant negative relationship between sample size and effect size, indicating bias, both assessing infants' ability to discriminate vowels. As discussed in the methods section, however, a number of alternative explanations are possible. As soon as researchers are aware that they are measuring a more subtle effect (in this case for example when selecting a contrast that is acoustically more difficult to distinguish) and adjust sample sizes, we expect to observe this negative correlation. However, in both datasets, funnel plot asymmetry was also significant. Thus, we might conclude that in two out of the 11 datasets in the present paper, we find significant publication bias.

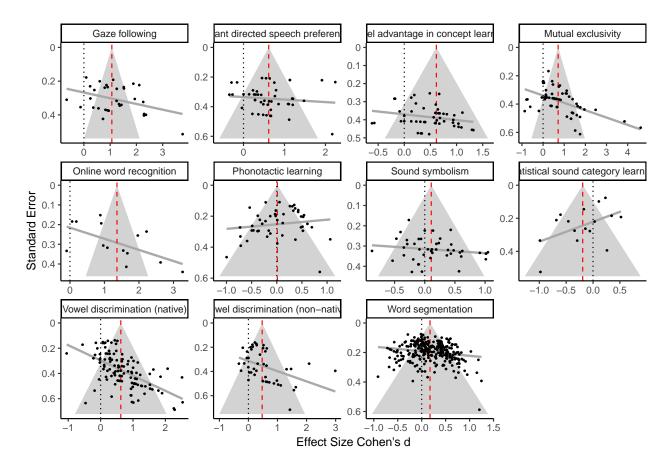


Figure 5. Funnel plots of all datasets with linear regression line indicated in grey. The dashed red line indicates the median effect size, the dotted black line zero. The grey triangle denotes the expected range area of effect sizes in the absence of bias or heterogeneity.

482 Discussion

483 Concrete recommendations for developmental psychologists

- 1. Calculate power prospectively.
- 2. Study planning.

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3. Increase availability and use of meta-analyses. To support the improvement current practices, we propose to make meta-analyses available in the form of ready-to-use online tools, dynamic reports, and as raw data. These different levels allow researchers with varying interest and expertise interests to make the best use of the extant record on infant language development, including study planning by choosing robust

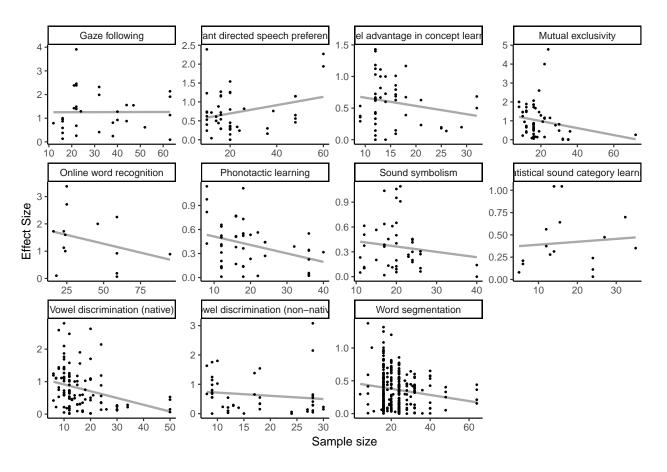


Figure 6. Caption.

methods and appropriate sample sizes. There are additional advantages for interpreting 491 single results and for theory building that emerge from our dataset. On one hand, 492 researchers can easily check whether their study result falls within the expected range of 493 outcomes for their research question – indicating whether or not a potential moderator 494 influenced the result. On the other hand, aggregating over many data points allows for the 495 tracing of emerging abilities over time, quantifying their growth, and identifying possible 496 trajectories and dependencies across phenomena (for a demonstration see M. Lewis et al. (2016)). Finally, by making our data and source code open, we also invite contributions and can update our data, be it by adding new results, file-drawer studies, or new datasets. Our 499 implementation of this proposal is freely online available at metalab.stanford.edu.

We have shown that power varies greatly across phenomena and that method choice is important. It turns out, however, that researchers do not choose the most robust methods.

This might to be due to a lack of consideration of meta-analytic effect size estimates. One of 503 the reasons for this is a lack of information on and experience in how to interpret effect size 504 estimates and use them for study planning (Mills-Smith, Spangler, Panneton, & Fritz, 2015). 505 Meta-analyses on infant language development are also rare, as showcased by the fact that 506 the present dataset relied on the authors' involvement, and only two out of XXX 507 meta-analyses used could be extracted from the extant work, and an extensive search in the 508 present literature did not yield additional candidates (excluding clinical contexts). 500 Conducting a meta-analysis is a laborious process, particularly according to common 510 practice where only a few people do the work, with little support tools and educational 511 materials available. Incentives for creating meta-analyses are low, as public recognition is 512 tied to a single publication. The benefits of meta-analyses for the field, for instance the 513 possibility to conduct power analyses, are often neither evident nor accessible to individual researchers, as the data are not shared and traditional meta-analyses remain static after publication, aging quickly as new results emerge. 516

A different application of meta-analytic tools is within a paper reporting on several studies. [expand]

#### 3. Report everything and ideally add anonymized supplementary

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materials. A possible reason for prospective power calculations and meta-analyses being
rare lies in the availability of data in published reports. To be able to draw the conclusions
we did in this paper, be it in the form of reported effect sizes within paper or as ready-to-use
dataset. As noted elsewhere, researchers would ideally always report effect sizes. Despite
long-standing recommendations to move beyond the persistent focus on p-values (such as
American Psychological Association (2001)), a shift towards effect sizes or even the reporting
of them is not (yet) widely adopted (Mills-Smith et al., 2015). A final impediment to
meta-analyses in developmental science are current reporting standards, which make it
difficult and at times even impossible to compute effect sizes from the published literature.

- Best practices when creating new meta-analyses
- Data standardization and sharing of materials.
- Visualization.
- 532 Community-augmented meta-analyses.
- Metalab as a model for other domains in child development research.
- 534 Conclusion

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