Quantifying Sources of Variability in Infancy Research Using the Infant-Directed-Speech Preference

The ManyBabies Consortium*
*All consortium members are listed in the Transparency section at the end of the article.

Abstract

Psychological scientists have become increasingly concerned with issues related to methodology and replicability, and infancy researchers in particular face specific challenges related to replicability: For example, high-powered studies are difficult to conduct, testing conditions vary across labs, and different labs have access to different infant populations. Addressing these concerns, we report on a large-scale, multisite study aimed at (a) assessing the overall replicability of a single theoretically important phenomenon and (b) examining methodological, cultural, and developmental moderators. We focus on infants' preference for infant-directed speech (IDS) over adult-directed speech (ADS). Stimuli of mothers speaking to their infants and to an adult in North American English were created using seminaturalistic laboratory-based audio recordings. Infants' relative preference for IDS and ADS was assessed across 67 laboratories in North America, Europe, Australia, and Asia using the three common methods for measuring infants' discrimination (head-turn preference, central fixation, and eye tracking). The overall meta-analytic effect size (Cohen's d) was 0.35, 95% confidence interval = [0.29, 0.42], which was reliably above zero but smaller than the meta-analytic mean computed from previous literature (0.67). The IDS preference was significantly stronger in older children, in those children for whom the stimuli matched their native language and dialect, and in data from labs using the head-turn preference procedure. Together, these findings replicate the IDS preference but suggest that its magnitude is modulated by development, native-language experience, and testing procedure.

The recent focus on power, replication, and replicability has had important consequences for many branches of psychology. Confidence in influential theories and classic psychological experiments has been shaken by demonstrations that many of the studies reported in the experimental literature have been underpowered (Button et al., 2013), that surprisingly few empirical claims have been subject to direct replication (Makel, Plucker, & Hegarty, 2012), and that the direct replication attempts that do occur often fail to substantiate original findings (Open Science Collaboration, 2015). As disturbing as these demonstrations may be, they have already led to important positive consequences in psychology, encouraging scientific organizations, journals, and researchers to work to improve the transparency and replicability of psychological science.

To date, however, infancy researchers have remained relatively silent on issues of replicability. This silence is not because infancy research is immune from the issues raised. Indeed, the statistical power associated with experiments on infant psychology is often unknown (and presumably too low; Oakes, 2017), and the replicability of many classic findings is uncertain. Instead, one reason for the infancy field's silence is likely related to the set of challenges that come with collecting and interpreting data on infants—and development more generally. For example, it can be quite costly to test

Corresponding Author:
The ManyBabies Consortium, Department of Psychology, 450 Serra Mall, Stanford, CA 94305
E-mail: mcfrank@stanford.edu

large samples of infants or to replicate past experiments. Another challenge for infancy researchers is that it is often difficult to interpret contradictory findings in developmental populations, given how children's behavior and developmental timing vary across individuals, ages, contexts, cultures, languages, and socioeconomic groups. Although these challenges may make replicability in infancy research particularly difficult, they do not make it any less important.

Indeed, it is of primary importance to evaluate replicability in infancy research (see Frank et al., 2017). But how can this evaluation be done? Here we report the results of a large-scale, multilab, preregistered infant study. This study was inspired by the ManyLabs studies (e.g., Klein et al., 2014), in which multiple laboratories have attempted to replicate various social- and cognitive-psychology studies and have assessed moderators of replicability systematically across labs. For the reasons just discussed, it would be prohibitively difficult to examine the replicability of a large number of infant studies simultaneously. Instead, we chose to focus on what developmental psychology can learn from testing a single phenomenon, assessing its overall replicability, and investigating the factors moderating it. As a positive side effect, this approach leads to the standardization of decisions concerning data collection and analysis across a large number of labs studying similar phenomena or using similar methods. For this first "ManyBabies" project, we selected a finding that the field has good reason to believe is robust—namely, infants' preference for infant-directed speech (IDS) over adult-directed speech (ADS)—and tested it in 67 labs around the world. This phenomenon has the further advantage that it uses a dependent measure, looking time, that is ubiquitous in infancy research. In the remainder of this introduction, we briefly review the literature on the relevance of IDS in development and then discuss our motivations and goals in studying a single developmental phenomenon at a larger scale than is typical in developmental research.

Infant-Directed-Speech Preference

IDS is a descriptive term for the characteristic speech that caregivers in many cultures direct toward infants. Compared with ADS, IDS is typically higher pitched, has greater pitch excursions, and is characterized by shorter utterances, among other differences (Fernald et al., 1989). Although caregivers across many different cultures and communities use IDS, the magnitude of the difference between IDS and ADS varies (Englund & Behne, 2006; Farran, Lee, Yoo, & Oller, 2016; Fernald et al., 1989; Newman, 2003). Nevertheless, the general acoustic pattern of IDS is readily identifiable to adult listeners (Fernald, 1989; Grieser & Kuhl, 1988; Katz, Cohn, & Moore, 1996; Kitamura & Burnham, 2003).

A substantial literature reporting studies using a range of stimuli and procedures has demonstrated that infants prefer IDS over ADS. For example, Cooper and Aslin (1990), using a contingent visual-fixation auditory preference paradigm, showed that infants fixate on an unrelated visual stimulus longer when hearing IDS than when hearing ADS, even as newborns. Across a variety of ages and methods, other studies have also found increased attention to IDS compared with ADS (Cooper, Abraham, Berman, & Staska, 1997; Cooper & Aslin, 1994; Fernald, 1985; Hayashi, Tamekawa, & Kiritani, 2001; Kitamura & Lam, 2009; Newman & Hussain, 2006; Pegg, Werker, & McLeod, 1992; Santesso, Schmidt, & Trainor, 2007; Singh, Morgan, & Best, 2002; Werker & McLeod, 1989). In a meta-analysis by Dunst, Gorman, and Hamby (2012), which included 34 experiments, the IDS preference had an effect size (Cohen's *d*) of 0.67, 95% confidence interval (CI) = [0.57, 0.76]—quite a large effect size for an experiment with infants (Bergmann et al., 2018).

The evidence suggests that IDS augments infants' attention to speakers (and presumably what speakers are saying) because of its highly salient acoustic qualities, such as frequency modulation (Cusack & Carlyon, 2003). In addition, it is hypothesized that the IDS preference plays a pervasive supporting role in early language learning. For example, young infants are more likely to discriminate speech sounds when they are pronounced with typical IDS prosody rather than ADS prosody (Karzon, 1985; Trainor & Desjardins, 2002). There are also reports that infants show preferences for natural phrase structure in narratives spoken in IDS but not in ADS (cf. Fernald & McRoberts, 1996; Hirsh-Pasek et al., 1987). In addition, word segmentation (Thiessen, Hill, & Saffran, 2005) and word learning (Graf Estes & Hurley, 2013; Ma, Golinkoff, Houston, & Hirsh-Pasek, 2011) are reported to be facilitated in IDS compared with ADS. Naturalistic observations confirm that the amount of speech directed to U.S. 18-month-olds (which likely bears IDS features), rather than the amount of speech they overhear (which is likely predominantly ADS), relates to the efficiency of their word processing and to their expressive vocabulary at age 24 months (Weisleder & Fernald, 2013). Finally, infants show increased neural activity in response to familiar words in IDS compared with the same words in ADS, and also compared with unfamiliar words in either register (Zangl & Mills, 2007). From a theoretical perspective, the IDS register has been claimed to trigger specialized learning mechanisms (Csibra & Gergely, 2009), as well as to boost social preferences and perhaps attention in general (Schachner & Hannon, 2011), as it

has been reported to improve even nonlinguistic associative learning (e.g., Kaplan, Jung, Ryther, & Zarlengo-Strouse, 1996).

The Current Study: Motivations and Goals

Despite the large body of research on infants' preference for IDS and the positive effects of IDS on the processing of linguistic and nonlinguistic stimuli, a number of open questions remain. This study was designed to answer some of these IDS-specific questions, as well as questions about methods for assessing infants' cognition, including questions about the interaction between statistical power and developmental methodologies. In this section, we describe the key questions for our study (as well as our predictions, where applicable), in rough order of decreasing specificity, highlighting methodological decisions that followed from particular goals.

What is the magnitude of the IDS preference?

First and foremost, our study was intended to provide a large-scale, precise measurement of IDS preference across a large number of labs. Given the evidence summarized in a previous meta-analysis (Dunst et al., 2012), we expected that the preference would be nonzero and positive. We suspected, however, that this phenomenon, like many others, suffers from a file-drawer effect, in which studies with low effect sizes (or large *p* values) often do not get published. Also, there was reason to believe that effect sizes in infancy research are often incorrectly reported; for example, partial eta-squared is often misreported as eta-squared. This confusion is likely to inflate the practical significance of findings, leading to an overestimation of the statistical magnitude and importance of effects (Mills-Smith, Spangler, Panneton, & Fritz, 2015). Therefore, we believed that the mean effect size of 0.67 reported by Dunst et al. (2012) was likely an overestimate of the real effect size.

How does IDS preference vary across ages?

We could plausibly predict that, all else being equal, older infants can more effectively process ADS than younger infants, and so the attraction of IDS over ADS might attenuate with age (Newman & Hussain, 2006). On the other hand, older infants might show a stronger preference for IDS over ADS, given that older infants have had more opportunity to experience the positive social interactions that likely co-occur with IDS,

including, but not limited to, eye contact, positive facial expressions, and interactive play.

How does IDS preference vary with linguistic experience and language community?

Preference for IDS might be affected by infants' language experience. Across many areas of language perception, infants show a pattern of perceptual narrowing. They begin life as "universal listeners" ready to acquire any language, but with experience gain sensitivity to distinctions in their native language and lose sensitivity to nonnative distinctions (Maurer & Werker, 2014). If preference for IDS follows a similar pattern, then older infants would be expected to show a stronger preference for IDS over ADS in their native language than in a nonnative language.

Faced with several competing concerns, we made the decision that all infants in our study, regardless of their native language, would be exposed to ADS and IDS stimuli in North American English (NAE). This design choice had several practical advantages. Most important, it allowed every infant to be tested with the same stimulus set. Creating different stimulus sets in different languages would have added methodological variability across labs that would be statistically indistinguishable from lab identity and language environment. Further, creating a single high-quality stimulus set shared across labs reduced the time and cost of conducting the study.

This decision had both advantages and drawbacks. A limitation of our design is that NAE stimuli are unfamiliar to infants from other language or dialect communities; thus, these infants might show less interest for NAE speech overall or might have a harder time recognizing IDS features as such when those features differ from those used in their native language or dialect. In fact, previous work even suggests that infants' IDS preference depends on the characteristics of the type of IDS addressed to children their age (McRoberts, McDonough, & Lakusta, 2009). Although this was a relevant concern, previous research had documented some IDS preference in the face of language and age mismatches (McRoberts et al., 2009; Werker, Pegg, & McLeod, 1994), and corpus studies suggested that, if anything, the distinction between IDS and ADS is more salient in NAE than in other linguistic variants (e.g., Fernald et al., 1989; Shute, 1987). Further, although this design did not allow us to disentangle the effects of stimulus language (native vs. nonnative) from the effects of infants' cultural background, we were able to explore how aspects of these factors influence infants' preference for IDS.

After weighing these considerations, we adopted NAE stimuli in order to have the maximal chance of recovering a positive effect, ensure that stimuli were not a source of variance across labs, allow comparability with previous work, and also minimize participating labs' barriers to entry (i.e., the need to create lab-specific stimuli). So as to be able to assess children's language background at the group level, we also chose to focus our primary analyses on monolingual infants (a separate effort focused on IDS preferences in bilingual children; Byers-Heinlein et al., 2020).

What is the impact of methodology on measurement of infants' preference?

We focused on three primary methods for assessing infants' interest: single-screen central fixation, eye tracking, and the head-turn preference procedure (HPP). All three methods are widely used in the field of infant language acquisition and yield measurements of preference for a given type of auditory stimulus, indexed by looking to an unrelated visual stimulus. In labs using the single-screen central-fixation method, infants were shown an uninformative image (a checkerboard) on a single, centrally located monitor, while they listened to either IDS or ADS, and the time they spent looking toward the monitor was manually coded via a closed-circuit video camera. In labs using the eye-tracking method, infants saw a similar display, but looking times were measured automatically via a remote corneal-reflection eye tracker. In labs using the HPP method, infants saw an attractor visual stimulus (often a flashing light bulb) appear to either their left or their right, and the duration of their head turn toward the attractor while IDS or ADS played was manually coded via a closed-circuit video camera (Nelson et al., 1995). Labs using the eye-tracking and central-fixation methods (and some of those using the HPP) employed an attention getter between trials to orient the infant's gaze toward the display.

Each lab tested the same phenomenon, using the same stimuli and the same general experimental parameters (including, e.g., trial order, maximum trial length); only the method of measuring preference varied. We therefore could analyze whether this theoretically irrelevant methodological choice influenced the observed effect size, which could help guide future decision making.

What are the effects of testing infants in multiple experiments during a single lab visit?

Labs vary in whether each infant visiting the lab completes a single experiment only, or whether some infants participate in a second study as well. These "second session" experiments are thought by some researchers to yield greater dropout rates and less reliable measurements, but the existence and magnitude of a second-session effect has not been tested, to our knowledge. In our study, a number of participating labs ran the IDS-preference study with some infants who had already participated in an additional study; measurements from these infants could inform future lab administration practices.

What should expectations be regarding replicability and statistical power in studies of infancy?

Although we replicated only a single phenomenon, the importance and assumed robustness of the IDS preference meant that our study would provide data relevant to developing a more nuanced understanding of replicability and power in infancy research. Because of the large number of participating labs, it was expected that data from some labs would not support an IDS preference (i.e., the data would yield a small, or even negative, effect size when analyzed individually). Some variability was to be expected given the mathematics of estimating an effect at so many independent sites. Nonetheless, we inspected whether there was systematic variability explained by lab effects.

In addition, by providing an unbiased estimate of effect size for an important developmental phenomenon (plus estimates of how that effect varies across ages, language backgrounds, and methods), this project would set a rough baseline for other scientists to use when planning studies. Existing attempts to estimate the statistical power of experiments with infants have been contaminated by publication bias, which has led to an overestimation of typical effect sizes in infancy research. Such overestimates can lead subsequent studies to be underpowered (researchers expect to see larger effects than are truly present). Though we planned to estimate the effect size for a particular developmental preference, we also planned to compare our unbiased estimate, calculated both across all three methods and for each method, with the meta-analytic effects extracted from previously published studies.

How should researchers think about the relationships among experimental design, statistical significance, and developmental change?

Previous work has often employed a contrast between two ages to suggest a developmental change, for example, by showing that 7-month-old infants exhibit a statistically reliable preference in a task, but 5-month-old

infants do not. Such a finding (the pairing of a significant difference and a nonsignificant difference) is not sufficient to show a difference between two time points (Nieuwenhuis, Forstmann, & Wagenmakers, 2011). Moreover, even when a significant difference between two age groups is found, such a result is not sufficient to elucidate the developmental pattern underlying this discrete test. By measuring how effect sizes changed with age using a much denser sampling approach, we aimed to illustrate what stands to be gained with a more gradient approach to testing behavior over development.

Summary

This broad replication of the IDS preference was aimed at helping to answer basic questions about the replicability of developmental-psychology findings and also at providing useful benchmarks for how to design infant cognition studies going forward. Projects such as ManyLabs have led to important improvements in research practices in cognitive and social psychology, and we hoped that ManyBabies would play a similar role for developmental cognitive science.

Disclosures

Preregistration

Prior to data collection, our manuscript was reviewed, and we registered our instructions and materials on the Open Science Framework (OSF; see https://osf.io/gf7vh/).

Data, materials, and online resources

All materials, data, and analytic code are available on OSF (https://osf.io/re95x/). The specific code and data required to render our submitted manuscript are also available there (https://osf.io/zaewn/).

Reporting

We report how we determined our sample size, all data exclusions, all manipulations, and all measures in the study.

Ethical approval

All the labs collected data under their own independent ethical approval via the appropriate governing body for their institution. Central data analyses used exclusively de-identified data. Identifiable video recordings of individual infant participants were coded and archived locally at each lab. Where institutional review boards'

protocols permitted, video recordings were also uploaded to Databrary, a central controlled-access database accessible to other researchers (http://databrary.org).

Method

Participation details

Time frame. We issued an open call for labs to participate in our study, which we designated "ManyBabies 1," on February 2, 2017. Data collection began on May 1, 2017. Data collection was scheduled to end on April 30, 2018 (1 year later). In order to allow labs to complete their sample, however, a 45-day extension was granted, and data collection officially ended on June 15, 2018. Data collection from one laboratory extended beyond this time frame (see Deviations From the Registered Protocol, later in the Method section).

Age distribution. Each participating lab was asked to recruit participants in one or more of four age bins: 3 months 0 days through 6 months 0 days, 6 months 1 day through 9 months 0 days, 9 months 1 day through 12 months 0 days, and 12 months 1 day through 15 months 0 days. Each lab was tasked with ensuring that, for each age bin contributed, the mean age fell close to the middle of the range and the sample was distributed across the bin. We selected 3-month bins as a compromise, on the assumption that tighter bins would make recruitment more difficult whereas broader bins would lead to more variability and would blur developmental trends (i.e., by introducing possible interactions between age and lab-specific effects, for instance, if a particular method turned out to be most appropriate for a subset of the ages tested). The ability to contribute to one or more age bins at the laboratory's discretion was necessary because labs differ in their ability to recruit infants of different ages.

Criterion for lab participation. During study planning, we used data from MetaLab (Bergmann et al., 2018) to compute the meta-analytic mean effect size for IDS preference; the resulting Cohen's *d* was 0.72. A power analysis indicated that 95% power to detect this effect in a paired-samples *t* test would require 27 participants, and 80% power would require 17. On the basis of these calculations, we asked participating labs to commit to a minimum sample of 32 in each age group they targeted. However, given that for many of our analyses, power across labs was more critical than power within a lab (Judd, Westfall, & Kenny, 2017), we allowed labs to contribute a "half sample" of 16, with the assumption that this would increase the number of laboratories capable of participating and allow more laboratories to contribute

samples from multiple age bins. We specified that in their recruitment efforts, labs should specifically target infants with the desired demographic characteristics outlined in the study's protocol (e.g., full-term infants; the full list of inclusion criteria is discussed later in the Method section). Given this recruitment strategy, however, we asked that sample Ns be calculated on the basis of the total number of infants tested, not the number of infants retained after exclusions (which were performed centrally as part of the broader data analysis, not at the lab level).

We included data from a lab in our analysis if the lab achieved the minimum N (16) required for a half-sample in an age bin by the end date of testing and if, after exclusions, the lab contributed data for 10 or more infants. If a lab collected more than their required sample, we included the extra data as well. Laboratories were cautioned not to consider the data (e.g., whether a statistically significant effect was evident) in their internal decision making regarding how many infants to recruit and when to stop recruitment.

Participants

Our final sample comprised 2,329 monolingual infants from 67 labs (mean sample size per lab = 34.76, SD = 20.33, range: 10–93). Demographic exclusions were implemented primarily during recruitment; despite this, additional infants were tested and excluded on the basis of preset criteria (see Inclusion Criteria for percentages). In addition, 2 labs registered to participate but failed to collect data from at least 10 included infants, and so their data were not included in analyses. Information about all the included labs is given in Table 1.

The mean age of infants included in the study was 291.99 days (range: 92–456). There were 310 infants in the 3- to 6-month-old bin (23 labs), 772 infants in the 6- to 9-month-old bin (49 labs), 554 infants in the 9- to 12-month-old bin (35 labs), and 693 infants in the 12- to 15-month-old bin (42 labs). Forty-five labs collected data in more than one bin. Of the total sample, 1,066 infants (from 30 labs) were acquiring NAE, and 1,263 infants (from 37 labs) were acquiring a language other than NAE. As noted earlier, a separate sample of bilingual children was tested in a parallel investigation, but these data are not reported here.

Materials

Visual stimuli. In labs using the central-fixation or eye-tracking method, a brightly colored static checkerboard was the fixation stimulus, and a small engaging video (an animation of colorful rings decreasing in size) was the attention getter. Labs using the HPP were asked to use their typical visual stimulus, which varied considerably across labs. Some used flashing lights as the visual fixation stimulus (as in the original protocol developed in the 1980s), whereas others used a variety of other visual displays on video screens (e.g., a looming circle).

Speech stimuli. The goal of our stimulus-creation effort was to construct a set of recordings of naturalistic IDS and ADS gathered from a variety of mothers. To do so, we made recordings of mothers speaking to their infants and to experimenters, selected a subset of individual utterances from these recordings, and then constructed stimulus items from this subset. All characteristics of the recordings other than register (IDS vs. ADS) were as balanced as possible across clips. On the basis of our intuitions and the data from the norming ratings described later, we consider these stimuli to be representative of naturally produced IDS and ADS across middle- and high-socioeconomic-status mothers in North America. Although future studies could vary particular aspects of the IDS systematically (e.g., age of the mother, age of the infant being spoken to, dialect), we did not do so in the current study. Our stimulus-elicitation method was designed to meet the competing considerations of laboratory control and naturalism.

Source recordings were collected in two laboratories, one in central Canada and one in the northeastern United States. The recorded mothers had infants whose ages ranged from 122 to 250 days. The same recording procedures were followed in the two laboratories. In an infant-friendly greeting area or testing room, recordings were collected using a simple lapel clip-on microphone connected to a smartphone (iPhone 5s or 6s), with the Voice Record or Voice Record Pro app (Dayana Networks Ltd.) in the Canadian lab and the Voice Memos app (Apple Inc.) in the U.S. lab. The targets for conversation were objects in an opaque bag: five familiar objects (a ball, a shoe, a cup, a block, a train) and five unfamiliar objects (a sieve, a globe, a whisk, a flag, and a bag of yeast). To ensure that the mothers used consistent labels, we affixed to each object a small sticker showing its name. Each object was taken out of the bag separately, and the mother was asked to talk about the object, either to her baby (for the IDS samples) or to an experimenter (for the ADS samples), until she ran out of things to say; at this point, the next object was taken out of the bag. Recording stopped when all the objects had been removed from the bag and had been talked about. The order of IDS and ADS recording was counterbalanced across mothers. A total of 11 mothers were recorded in Canada and 4 in the United States.

There were a total of 179 unedited minutes of recording from Canada and 44 from the United States. A

Table 1. Summary of the Included Labs and Their Samples

Lab	Infants		Language	Method	Country
	Mean age (days)	N^a			
babylabbrookes	255	53	English	Central fixation	United Kingdom
babylabvuw	224	15	English	Central fixation	New Zealand
babylabyork	268	32	English	Central fixation	United Kingdom
baldwinlabuoregon	320	16	English	Central fixation	United States
bchdosu	269	67	English	Central fixation	United States
bcrlunlv	411	29	English	Central fixation	United States
bounbcl	411	31	Turkish	Central fixation	Turkey
icclbc	222	15	English	Central fixation	United States
infantcoglablouisville	325	35	English	Central fixation	United States
ldlottawa	276	59	English	Central fixation	Canada
madlabucsd	234	10	English	Central fixation	United States
minddevlabbicocca	158	15	Italian	Central fixation	Italy
udssaarland	332	43	German	Central fixation	Germany
unlvmusiclab	138	20	English	Central fixation	United States
weescienceedinburgh	213	32	English	Central fixation	United Kingdom
wsigoettingen	274	88	German	Central fixation	Germany
infantcogubc	165	39	English	Central fixation, eye tracking	Canada
lancaster	326	42	English	Central fixation, eye tracking	United Kingdom
babylablangessex	289	27	English	Eye tracking	United Kingdom
babylablmu	368	62	German	Eye tracking	Germany
babylabshimane	195	28	Japanese	Eye tracking	Japan
babylabuclajohnson	408	22	English	Eye tracking	United States
babylabumassb	308	30	English	Eye tracking	United States
babylingoslo	227	31	Norwegian	Eye tracking	Norway
callab	369	30	English	Eye tracking	United States
cdcceu	272	27	Hungarian	Eye tracking	Hungary
cfnuofn	298	15	English	Eye tracking	Australia
childlabmanchester	269	26	English	Eye tracking	United Kingdom
cogdevlabbyu	161	29	English	Eye tracking	United States
dcnlabtennessee	345	19	English	Eye tracking	United States
earlysocogfm	310	35	English	Eye tracking	United States
escompicbsleipzig	159	14	German	Eye tracking	Germany
ethosrennes	187	90	French	Eye tracking	France
irlconcordia	310	37	English	Eye tracking	Canada
jmucdl	340	17	English	Eye tracking	United States
kokuhamburg	305	25	German	Eye tracking	Germany
kyotobabylab	281	30	Japanese	Eye tracking	Japan
labunam	302	36	Spanish	Eye tracking	Mexico
lcdfsu	354	23	English	Eye tracking	United States
lcduleeds	413	14	English	Eye tracking	United Kingdom
lllliv	302	36	English	Eye tracking	United Kingdom
lscppsl	404	14	French	Eye tracking	France
pocdnorthwestern	409	30	English	Eye tracking	United States
socialcogumiami	131	19	English	Eye tracking	United States
weltentdeckerzurich	414	30	German	Eye tracking	Switzerland
nusinfantlanguagecentre	337	21	Mandarin	Eye tracking, central fixation	Singapore
babylabkingswood	312	32	English	HPP	Australia
babylabkonstanz	235	15	German	HPP	Germany

Table 1. (Continued)

Lab	Infants		Language	Method	Country
	Mean age (days)	N^{a}			
babylableiden	319	15	Dutch	HPP	Netherlands
babylabnijmegen	279	49	Dutch	HPP	Netherlands
babylabparisdescartes1	403	16	French	HPP	France
babylabplymouth	332	34	English	HPP	United Kingdom
babylabprinceton	307	24	English	HPP	United States
babylabutrecht	276	61	Dutch	HPP	Netherlands
bllumanitoba	281	79	English	HPP	Canada
chosunbaby	313	77	Korean	HPP	Korea
infantlanglabutk	323	65	English	HPP	United States
infantllmadison	316	93	English	HPP	United States
infantstudiesubc	228	20	English	HPP	Canada
islnotredame	411	28	English	HPP	United States
isplabmcgill	411	11	French	HPP	Canada
langlabucla	250	63	English	HPP	United States
lppparisdescartes2	241	30	French	HPP	France
musdevutm	229	31	English	HPP	Canada
purdueinfantspeech	355	58	English	HPP	United States
trainorlab	241	24	English	HPP	Canada
babylabpotsdam	306	46	German	HPP, central fixation	Germany

Note: Full identifying information for the labs is available in the metadata folder of the analysis code on the Open Science Framework. HPP = head-turn preference procedure.
[a]The numbers in this column refer to the number of infants in the final sample.

first-pass selection of low-noise IDS and ADS samples yielded 1,281 utterances, with a total duration of 4,479 s. From this first pass, we selected 238 utterances that were considered to be the best examples of IDS and ADS and met other basic stimulus selection criteria (e.g., they did not contain laughter or the baby's name).

This library of 238 utterances was then normed on five variables: accent, affect, naturalness, noisiness, and IDS-ness. The goal of this norming was to gather intuitive judgments so that we would have a principled basis for identifying utterances that were clearly anomalous in some respect (e.g., odd affect or background noise) and excluding them. Naive, NAE-speaking adults recruited from Amazon Mechanical Turk listened to all 238 utterances and rated them on 7-point Likert scales. Raters were assigned randomly to one of the five variables; the number of raters assigned to a particular task ranged from 8 to 18 because of variability in random assignment. Affect and IDS ratings were made using low-pass-filtered recordings (a 120-Hz filter with standard roll-off was applied twice using the *sox* software package, available at http://sox.sourceforge.net). In general, with the exception of IDS-ness, ratings were not highly variable across clips (the largest *SD* was 0.85, for noise ratings).

These ratings were used to produce a set of utterances in which accent was rated similar to "standard English" (ratings < 3; 1 = completely standard), naturalness was rated high (ratings > 4; 7 = completely natural), noisiness was rated low (ratings < 4; 1 = noiseless), and IDS and ADS clips were consistently distinguished (ratings > 4 for IDS clips and ratings < 4 for ADS clips; 7 = clearly directed at a baby or child). This procedure resulted in a total of 163 utterances that met our inclusion criteria.

Our next goal was to create eight IDS and eight ADS stimuli that were exactly 18 s in length, each containing utterances from the set we created. First, the amplitudes of all the clips were root mean square normalized to 70 dB sound-pressure level (SPL). Stimuli were then assembled from the normalized clips, and finally, the amplitudes of the stimuli were renormalized to 70 dB SPL. We assembled the final stimuli considering the following issues:

- *Identity:* Each audio stimulus was constructed using clips from more than one mother. The number of different mothers included in a given stimulus was matched across the IDS and ADS stimuli. In addition, multiple clips from the same mother

Table 2. Characteristics of the Infant-Directed Speech (IDS) and Adult-Directed Speech (ADS) Stimuli

Characteristic	IDS		ADS	
	Mean	*SD*	Mean	*SD*
Number of mothers speaking per stimulus	4.00	0.00	3.75	0.46
Number of clips per stimulus	6.88	1.13	4.50	0.76
Number of objects mentioned per stimulus	2.75	0.71	2.75	0.71
F0 per stimulus (Hz)				
Mean	206.90	19.50	174.90	13.20
10th percentile	131.40	26.10	139.00	17.70
90th percentile	340.00	21.50	232.00	13.80
Mean number of utterances per stimulus	7.75	1.04	6.63	0.92
Mean duration (s) of utterances	1.58	0.74	2.12	1.41
Mean interutterance interval (s)	0.75	0.30	0.59	0.33

Note: F0 = fundamental frequency.

were grouped together within a given stimulus in order to match the number of "mother transitions" across registers.

- *Lexical items:* We matched the frequency of object labels in the clips across the IDS and ADS contexts. We also ensured an even distribution of the order in which each particular word was presented across stimuli and registers (ADS vs. IDS).

- *Questions:* IDS tends to include a much higher proportion of questions compared with ADS (Snow, 1977; Soderstrom, Blossom, Foygel, & Morgan, 2008). However, because the nature of the recording task may have served to inflate this difference, we preferentially selected declarative sentences over questions in the IDS sample. In the final stimulus set, 47% of the utterances in the IDS samples and 3% of the utterances in the ADS samples were questions. We felt that retaining this naturally occurring difference between IDS and ADS within our stimuli was more appropriate than precisely and artificially controlling for utterance type across registers.

- *Duration of individual clips:* As expected, the utterances in IDS were much shorter than those in ADS, so it was not possible to match the IDS and ADS stimuli on duration or number of clips. Because there were more clips per stimulus in the IDS samples, there were also more utterance boundaries. This property is consistent with what has been reported in the literature on the natural characteristics of IDS (Martin, Igarashi, Jincho, & Mazuka, 2016).

- *Total duration:* We fixed all stimuli to have a total duration of 18 s by concatenating individual utterance files into single audio files that were more than 18 s in length, trimming these down to 18 s, and fading the audio in and out with 0.5-s half-cosine windows.

Table 2 and Figure 1 provide additional details regarding the final stimulus set. Measurements were made using STRAIGHT (Kawahara & Morise, 2011), using default values for extraction of the fundamental frequency (F0).

Table 3 provides a comparison of our stimuli with a sample of others that have been used in previous IDS-preference studies. Across studies in the broader literature, the only measure that we found to be reported consistently was F0 for IDS and ADS, and even this statistic was reported for only about half the studies we examined (those studies listed in Table 3). Various measures of variability in F0 were reported in some cases (e.g., range within each sample, range across samples, standard deviation), but because of variation in the length and number of samples used, and a lack of systematicity in reporting, it was difficult to compare studies directly. Numerically, the average pitch difference between IDS and ADS in our materials was less extreme than that in previous studies.

To confirm that our composite IDS and ADS stimuli were perceived to be natural and that the pitch difference between registers was sufficient to lead to the two sets of stimuli being categorized differently, we conducted another norming study. Using the same basic paradigm as in the first norming study, we collected a new sample of judgments from Mechanical Turk participants. They were randomly assigned to listen to all 16 stimuli and judge either whether they were directed at infants or children or at adults (*N* = 22) or else whether they sounded natural (*N* = 27). All IDS clips were judged extremely likely to be directed at infants or children (*M* = 6.74, *SD* = 0.09, on a rating scale from
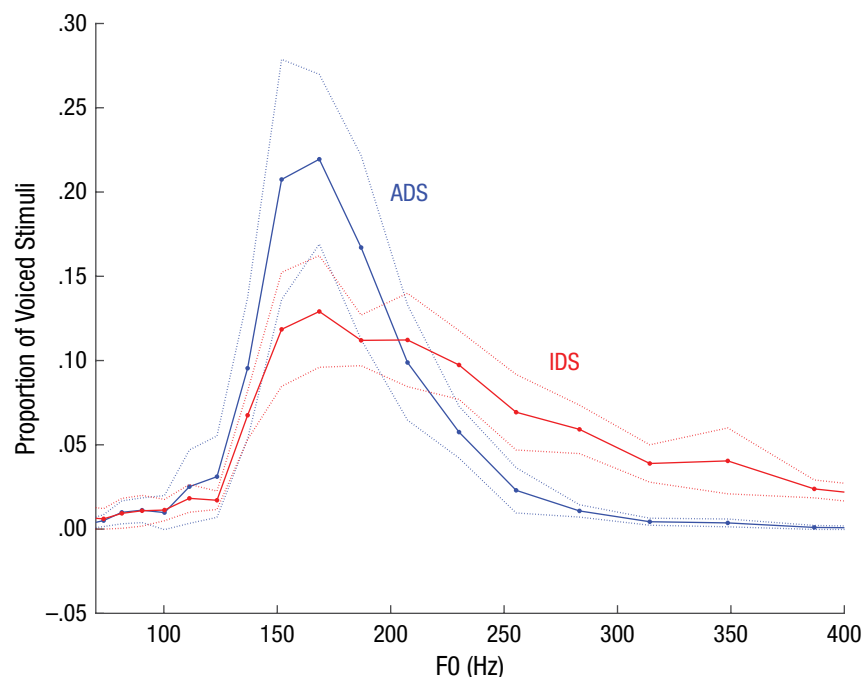
Fig. 1. Distribution of fundamental frequency (F0) values in the stimulus set. The graph shows the proportion of voiced segments of infant-directed speech (IDS) and adult-directed speech (ADS) that fell in each bin of a logarithmically spaced series of F0 bins within a search range of 32 through 650 Hz. The solid and dotted lines show means ±1 *SE* across stimuli.

Table 3. Comparison of the Present Study and Previous Studies on Infant-Directed-Speech Preferences

Study	Age group or groups targeted (months)	Nature of the speech recorded	Number and description of stimuli	Mean F0 (Hz) IDS	Mean F0 (Hz) ADS	Difference in mean F0 (IDS − ADS; Hz)	Ratio of IDS F0 to ADS F0
Present study	3–15	Semistructured speech directed toward a 4- to 8-month-old child or an adult experimenter	Eight full trials for each type of speech	207	175	32	1.2
Cooper and Aslin (1990)	0, 1	Scripted speech read with no infant present	Four sentences in each type of speech	316	260	56	1.2
Newman and Hussain (2006)	4.5, 9, 13	Scripted speech read with no infant present	Four passages in each type of speech	226	190	36	1.2
Thiessen, Hill, and Saffran (2005)	7	Nonsense strings of syllables read with no infant present	Twelve strings in each type of speech	292	230	62	1.3
Cooper, Abraham, Berman, and Staska (1997)	1, 4	Mothers' naturalistic speech directed toward their own infants	Twenty seconds of speech in each type of speech	219	184	35	1.2
Schachner and Hannon (2011)	5	Speech elicited by asking adults to describe a picture they were looking at	Two 1-min videos in each type of speech	273.0	224.7	48.3	1.2

Note: F0 = fundamental frequency; IDS = infant-directed speech; ADS = adult-directed speech.

1 through 7), whereas all ADS clips were judged highly likely to be directed at adults ($M = 2.12$, $SD = 0.38$). Both sets of clips were judged to be relatively natural; if anything, the ADS clips were rated as slightly more natural ($M = 5.18$, $SD = 0.19$) than the IDS clips ($M = 4.47$, $SD = 0.31$). In sum, because our IDS stimuli were created from naturalistic productions from a wide range of mothers, they were less extreme in their intonation than the stimuli used in previous studies, but they were judged as natural and were easily identified as infant directed.

Procedure

Basic procedure. Each lab used the testing paradigm (or paradigms) with which they were most familiar, among variants of three widely used paradigms: Twenty laboratories used the HPP, 16 used the single-screen central-fixation preference procedure, and 27 used single-screen central-fixation with fixations recorded by a corneal-reflection eye tracker; 4 labs contributed data using two different methods. All procedural instructions to participant labs can be found on OSF (https://osf.io/s3jca/).

To minimize researcher degrees of freedom, we asked participating labs to adhere to our instructions closely. Deviations from the basic protocol for each paradigm were necessary in some cases because of variation in the software and procedures used in each laboratory and were documented for future analysis.

First versus second test session. In some laboratories, infants were sometimes tested in an unrelated experiment during their visit, either prior to or following the IDS-preference experiment. Each lab noted whether infants completed the IDS-preference experiment as their first (and possibly only) or second test session.

Onset of each trial. At the beginning of each trial, a centrally positioned visual stimulus (typically the study's standard attention getter, or a light in some HPP labs) was used to attract the infant's attention. When the infant fixated on this attention getter, a visual stimulus appeared (a checkerboard on a screen in the central-fixation and eye-tracking paradigms, a flashing light or fixation stimulus on a screen—e.g., a flashing circle—in the HPP paradigm). The stimulus appeared to the left or right of the infant in HPP setups and in the center in the other two setups.

Trials. At the beginning of each session, there were two warm-up trials that familiarized the infant with the general procedure. The auditory stimulus for warm-up trials was an 18-s clip of piano music, and the visual stimulus was identical to that shown on the test trials. In addition to familiarizing the infant with the general experimental

setup, the warmup trials highlighted the contingency between looking at the visual display and the onset of the auditory stimulus. We did not analyze data from these trials. The warmup trials were followed by 16 test trials presenting the IDS and ADS auditory stimuli.

Minimum looking time. There was no minimum required looking time during data collection (i.e., trials were never repeated). A minimum looking time of 2 s was the criterion for inclusion of a trial in analysis. This 2-s minimum was chosen after the laboratories discussed their typical standards of practice regarding minimum trial length, which varied considerably. The 2-s criterion was selected to ensure that the infants had sufficient time to hear enough of the stimulus to discriminate IDS from ADS.

Maximum looking time. On each test trial, infants could hear speech for a maximum of 18 s (i.e., the duration of one sound file). At labs using software that could implement infant-controlled trial lengths, a trial ended if the infant looked away from the visual stimulus for two consecutive seconds. Otherwise, the trial continued until the stimulus ended.

Randomization. Four pseudorandom trial orders were created. Each order contained four blocks, and each block contained two IDS and two ADS trials, in alternation. Two blocks in each order began with IDS, and the other two began with ADS. The same IDS and ADS stimuli were always paired with one another, to facilitate analyses of preference scores by item.

Volume. Each lab was asked to use a stimulus volume level that was consistent with their general lab practices; this decision was not standardized across labs. Labs were instead instructed to measure and report their average volume (dB SPL) with and without a white-noise reference audio clip playing, though not all contributing labs reported these measurements (measurements were reported by 47 labs). From these values, we calculated a signal-to-noise ratio for each lab, $M = 1.95$, $SD = 0.43$, range: 1.25–3.30.

Minimizing caregiver bias. A custom masking stimulus containing a blend of instrumental music and a pastiche of speech stimulus materials triggered at random times and with random amplitude was available as part of the study materials. This masking stimulus was played to the caregiver over noise-attenuating headphones, to mask the stimuli that the infant was hearing via external loudspeakers. Experimenters were instructed to play the masking music at a high (but comfortable and safe) volume.

Coding. Coding of looking times was conducted via the standard procedure in each lab. There were three methods

of coding infants' eye gaze: on-line coding by an experimenter via button press during the experimental session, off-line coding of a video after the experimental session, and automatic coding by an eye tracker. When both on-line and off-line coding were reported, we used the off-line coding.

Minimizing experimenter bias. Experimenters making on-line coding decisions (in the central-fixation and HPP methods) were blind to the particular stimulus presented during each test trial, as they either were located in a room separate from the infant or were in the same room but were wearing noise-attenuating headphones playing the same masking stimulus presented to the infant's caregiver. Off-line coding was conducted without direct access to the auditory stimuli.

Demographics. All labs were instructed to collect basic demographic information about participants: sex, date of birth, estimated proportion of language exposure for each language heard in their daily life, race-ethnicity (using categories appropriate for the cultural and geographic context), preterm-versus-full-term status, history of ear infections, known hearing or visual impairments, and known developmental concerns (e.g., developmental disorders). Parents were also asked to report information about themselves (sex, level of education, and native language or languages) and the infant's siblings (sex and date of birth). A standard recommended questionnaire was distributed to participating labs as part of the instructions, although labs were permitted to use their own forms as long as they gathered the necessary information. A subset of laboratories provided extensive additional information about participants and testing circumstances (not analyzed here), for use in planned follow-up projects.

General lab practices

Training of research assistants. Each lab was responsible for maintaining good practices for training experimenters and was expected to use the same rigor with the ManyBabies study as with any other study in their lab. Laboratories reported on which research assistant ran each infant, using pseudonyms or numerical codes. Each laboratory completed a questionnaire regarding its training practices, the experience and academic status of each experimenter, and its basic practices for greeting participants.

Reporting of technology mishaps and infants' and parents' behavior. Laboratories were asked to record relevant concerns, anomalies, and comments according to their standard lab practices, and these were converted to a standardized form during the main analysis. Relevant concerns included infants crying during testing, parents intervening in a way that would affect infants' looking behavior (e.g., talking or pointing), and technical problems that prevented the normal presentation of experimental stimuli.

Videos

All laboratories provided a walk-through video that detailed their basic processes, including procedures for greeting participants and caregivers, obtaining consent, and collecting data, and that showed the physical characteristics of their laboratory. (Our preregistration stated that further procedural documentation would be collected and made available, but standardized reporting for procedural decision making proved difficult to develop and implement.) In addition, labs were strongly encouraged to collect and share video recordings of their data collection, within the limits of what was permissible given their ethics approval and participants' consent. If labs could not provide videos of participants, they were asked to provide a video showing a run-through of their procedure, pictures and information regarding the study setup, or both. A number of laboratories contributed these video recordings to Databrary, where they can be found by searching for "ManyBabies."

Inclusion criteria for participants

All data collected for the study (i.e., data for every infant for whom a data file was generated, regardless of how many trials were completed) were given to the analysis team for confirmatory analyses. Participants were included in analysis only if they met all the criteria described in this section. All exclusion rules were applied sequentially, and the reported percentages of excluded infants reflect this sequential application to an initial sample (prior to exclusions) of 2,754. The following list describes the inclusion criteria, in the order of their application. Note that the first three criteria preemptively prevented participation (although some infants were run erroneously).

- *Monolingual:* Only monolingual infants, of any language background, were included in the final sample. Monolingualism was defined as a minimum of 90% exposure to the native language, as reported by the infant's parent. This cutoff score struck a balance between including most infants who would typically be considered monolingual in infant language studies and excluding those who

might be considered bilingual (Byers-Heinlein, 2015). Of the initial sample, 162 (5.88%) infants did not meet this criterion.

- *Full-term:* We defined *full term* as gestation time greater than or equal to 37 weeks. Of the remaining sample, 62 (2.39%) infants did not meet this criterion.

- *No diagnosed developmental disorders:* We excluded infants with parent-reported developmental disorders (e.g., chromosomal abnormalities) or diagnosed hearing impairments. Of the remaining sample, 2 (0.08%) infants were tested but did not meet this criterion. Because of concerns about the accuracy of parent reports, infants whose parents reported that they had experienced ear infections were not excluded unless the parents also reported medically confirmed hearing loss.

- *Contributed usable data:* To be included in the study, an infant was required to have contributed nonzero looking times for at least one pair of test trials (i.e., IDS and ADS trials from the same stimulus pair), after trial-level exclusions were applied. Of the remaining sample, 78 (3.09%) infants did not meet this criterion. We adopted this relatively liberal inclusion criterion even though it is at variance with the more stringent standards that are typically used in infancy research. We were interested in maximizing the amount of data from each lab to be included in the initial analysis, and our paradigm was, by design, less customized for any particular age group than previously used paradigms were (and hence likely to produce greater data loss, especially for older infants, who tend to habituate more quickly). In the exploratory analyses we report, we considered how exclusion decisions affected our effect-size estimates.

After these rules were applied, participants could also be excluded from analysis because of session-level errors, including equipment error (e.g., no sound or visuals on the first pair of trials), experimenter error (e.g., an unblinded experimenter, if looking time was measured by live button press), or reported evidence of consistent outside interference (e.g., talking or pointing by parents, construction noise, sibling pounding on the door). Session-level error resulted in 78 (3.18%) infants for whom we had other reported data being dropped from analysis. This number is likely an underestimate, however, because many participating labs did not provide data for all children with session-level errors. In addition, session-level errors were not classified consistently across labs, so an accurate classification of the proportion of different types of errors was not possible.

Trial exclusions

We excluded individual trials for which issues were reported (e.g., fussy infant, incorrect stimulus, single instance of parent or sibling interference). A total of 4,471 (10.61%) trials were affected by such errors, which resulted in the complete removal of 29 infants (1.22%). As with session-level errors, classification of these concerns was inconsistent across participating labs, but the most common source of trial-level errors was infant fussiness.

Our trial-length minimum of 2 s of looking time resulted in the exclusion of an additional 6,027 (16.13%) trials. These trials were analyzed as missing in our planned analysis. This trial-level exclusion led to the exclusion of 3 additional infants who had no usable trial pairs (0.13%).

Inclusion of a lab's data

We included a lab's data if they were able to achieve the minimum *N* required for a half-sample and if, after exclusions, they contributed data from 10 or more infants. This criterion led to 11 (0.47%) infants from two labs being excluded from the final sample.

Deviations from the registered protocol

Given that this was the first experimental cross-laboratory infant study of such a large scale, there were a number of unanticipated issues that arose during data collection within individual labs and at the study level, and these issues resulted in deviations from our registered protocol. Necessary decisions were made without consideration of their impact on the results, and all such cases were documented. Fuller documentation can be found accompanying our shared data in the OSF repository; here we summarize the nature and extent of these deviations. Note that some were the result of typical within-laboratory protocol deviations (experimenter error, etc.), whereas others stemmed from the additional challenges inherent in harmonizing methodology and data format across such a large number of laboratories with different internal protocols and standards.

The protocol deviations include the following:

- Before labs had commenced data collection, we altered our attention-getter stimulus to be a precessing annulus accompanied by chimes (to address the concern that a laughing baby, our

original attention getter, might be a factor in observing a preference for IDS rather than ADS, e.g., via presenting an infant as part of the content of the experiment); however, some labs used the original stimulus.

- For a variety of reasons, labs' protocols deviated from the registered protocol in ways that resulted in trials longer than the assumed maximum of 18 s. In all cases, looking times on these trials were truncated to 18 s.

- A number of labs provided data from infants who were within the 3- to 15-month age range, but outside the individual lab's preregistered age bin. These infants were included in the analyses.

- Many labs deviated from their preregistered sample size because of constraints on testing resources. We included these labs provided they met the minimum inclusion criteria for the study as a whole. All such labs certified that they did not make decisions regarding sample size on a data-dependent basis.

- A number of laboratories marked participants for exclusion due to session-level errors for reasons other than equipment error, experimenter error, or outside interference.

Deviations regarding exclusions bear further discussion. Some labs marked participants for exclusion because of trial-level errors (e.g., fussiness, parental interference), even though sufficient trial-level data were available for analysis. Similarly, individual trials were sometimes marked as error trials for reasons related to session-level issues. All trial-level and session-level errors were reviewed centrally by at least two coders using all available information in the spreadsheet to determine whether the error was more appropriately categorized as a trial-level or session-level error. Specific information about each error coding that was changed during this process is available in the metadata directory within the data-analysis codebase, available on our OSF project page.

In total, 313 participants (from 50 labs) who had been marked for session-level exclusions were retained for further processing and analysis, for the following reasons: The session-level exclusion was based solely on the existence of trial-level errors (190 infants), the exclusion was based on a different exclusion criterion (e.g., the participant was out of the age range or was preterm; 93 infants), or the central analysis team decided that the issue identified by the lab did not warrant exclusion (e.g., the lab implemented a look-away criterion slightly different from the preregistered one; 30 infants). Note that many of these retained participants were subsequently excluded at other points

in the analysis pipeline because, although they did not meet the criteria for exclusion on the basis of session-level errors, they did meet other conditions for exclusion (e.g., as noted, some of these participants were out of the age range or were preterm).

In addition to recoding session-level errors, we corrected the coding of trial-level errors when appropriate. In total, 778 such errors, involving 62 participants in 16 different labs, were recoded. The majority of these cases involved a lab coding a session-level error (e.g., outside the age range) as a trial-level error (584 trials) or coding a trial-level error as a session-level error (e.g., a lab recorded a session-level error when an infant was fussy on a specific trial, but did not code the affected trial as an error trial; 133 trials). Other trials were corrected when subsequent investigation of lab notes and discussion with lab members revealed that the original code needed to be changed (61 trials).

A variety of additional errors were found (e.g., participants who were identified as having been run in a lab's pilot test but who were not properly excluded) and fixed within the spreadsheets. Video data were not reviewed centrally, although in some cases when a question arose, the laboratory reviewed its own video in order to respond. Our corrections of trial- and session-level errors have been carefully documented, and this information can be accessed upon request, but because the documentation in some cases includes identifiable information about participants, it is not possible to share it publicly.

Other reported protocol deviations included failing to submit a preregistration form (one lab), setting the trial-end (look-away) criterion to 3 s rather than 2 s for some participants (one lab), temporarily changing location during data collection (one lab), making minor technical changes in the protocol after the start of data collection (two labs), alternating left- and right-side presentation of stimuli and testing skin conduction during the procedure (one lab), implementing procedural deviations related to high-chair usage (one lab), using attention getters other than the preregistered stimulus (four labs), and using a pinwheel rather than a checkerboard as the main visual fixation stimulus in the HPP (one lab).

We also detected a large number of other kinds of errors in the submitted data as a result of the comprehensive checking process conducted during analysis. These typographical and other errors were resolved when necessary by contacting the submitting lab. In general, we included data obtained with minor protocol deviations, and erred on the side of excluding data, when necessary, at the trial rather than session level. A

few demographic variables required greater central scrutiny than originally anticipated. Most notably, there was considerable variability in the interpretation of the *preterm* and *bilingual* designations (despite centrally dictated standards). When necessary, we recoded lab data so as to conform to the definitions in the original protocol.

The instructions in our registered protocol were ambiguous as to whether the inclusion criterion for labs was contributing data for 10 or more infants or for more than 10 infants. The more liberal of these two criteria was used.

Finally, two labs submitted data after the deadline. In one case, this was because of a communication error; in the other case, the lab continued data collection after the deadline, and 8 additional infants were tested. Both data sets were included in the final analysis reported here.

Results

Confirmatory analyses

Data processing and analytic framework. All planned analyses were preregistered in our initial Registered Report submission (available at https://osf.io/vd789/). Our primary dependent variable of interest was looking time during test trials. Looking time was defined as time spent fixating the screen (central-fixation and eye-tracking methods, some HPP setups) or light (other HPP setups); looking time did not include any time spent looking away from the screen, even when that time was below the threshold for terminating a trial. Because looking times are nonnormally distributed, we followed Csibra, Hernik, Mascaro, Tatone, and Lengyel's (2016) recommendation and log-transformed all looking times prior to statistical analysis.

We adopted two complementary analytic frameworks: meta-analysis and mixed-effects regression. In the meta-analytic framework, we conducted standard analyses within each lab and then estimated variability in the results of these analyses across labs. The meta-analytic approach has a number of advantages over the mixed-effects approach, including the use of simple within-lab analyses, the ability to estimate cross-lab variability directly, and the possibility of making direct comparisons with the standardized effect sizes estimated in previous meta-analyses. However, the standard random-effects meta-analytic model is designed for cases in which the raw data are unavailable and procedures and data types are not standardized. In contrast, in our situation, procedures and data were standardized across labs, and relevant moderators were recorded. The availability of trial-by-trial

data for all labs allowed us to use mixed-effects models, which account for the nesting and crossing of random effects (e.g., participants nested within labs, items crossed across labs) and can provide more accurate estimates of the main effect and moderators. We report both analyses to allow for the most comprehensive understanding of the variance in the data.

Our meta-analyses were conducted as follows. Each lab's data set was considered a separate study. For each such study, we computed individual infants' IDS preference by (a) subtracting the looking time on each IDS trial from the looking time on its paired ADS trial (excluding trial pairs with missing data) and (b) computing a mean difference score across trial pairs. Then we computed a group IDS preference for each lab and age group within that lab using d_z, a version of Cohen's standard d statistic, computed as the average of infants' IDS preference scores divided by the standard deviation of those scores. We then used standard random-effects meta-analysis fit using restricted maximum likelihood (REML) with the *metafor* package (Viechtbauer, 2010).

Although we did not anticipate this in our initial analysis plan, a large number of labs collected data outside of their planned samples. For example, many labs contributed data from a sample of children within a specific age bin as well as several children outside of that age bin, or from a sample of children using one method and from a handful of children using another. Although we included these children in the mixed-effects analyses described next, we worried that the inclusion of many unplanned samples of just one or a few infants in the meta-analytic models would excessively increase lab-level variance. Thus, for the meta-analyses, we included only samples (e.g., age, language, or method groups) with 10 or more infants.

Our mixed-effects models, fit to the entire data set collected from the 67 labs, were specified as follows:

$$DV \sim IV_1 + IV_2 + \ldots + (\ldots | \text{participant}) + (\ldots | \text{item}) + (\ldots | \text{lab})$$

The goal of this approach was to examine effects of the independent variables (IV) on the dependent variable (DV), while controlling for variation in both the DV (random intercepts) and the relationship of the IV to the DV (random slopes) due to relevant grouping units (participants, items, and labs). The use of mixed-effects models also allowed us to move away from using a difference score as the dependent variable of interest. Although difference scores simplified the process of calculating effect sizes for the metaregression, their use required that trials be paired, so some collected data

(i.e., from unpaired trials) could not be analyzed. In the mixed-effects framework, in contrast, looking time on individual trials was the dependent measure, so all trials could be included.

In our mixed-effects models, we planned a maximal random-effects structure (Barr, Levy, Scheepers, & Tily, 2013), which entailed specifying all random effects that were appropriate for the experimental design (e.g., IDS/ADS trial type could be nested within participants—because each infant heard stimuli in both conditions—but could not be nested within items because each item was unique to its trial type). In cases of mixed-effects models that failed to converge, we pursued an iterative pruning strategy. We began by removing random slopes nested within items (as that grouping was of least theoretical interest) and next removed random slopes nested within participants and then random slopes nested within labs. We then removed random intercepts from groupings in the same order, retaining effects of trial type until last because these were of greatest theoretical interest. We fit all models using the *lme4* package (Version 1.1-21; Bates, Mächler, Bolker, & Walker, 2015) and computed *p* values using the *lmerTest* package (Kuznetsova, Brockhoff, & Christensen, 2017).

IDS preference. What was the overall magnitude of the IDS preference we observed? This question was answered in the cross-lab meta-analysis by fitting the main-effect model specified by $d_z \sim 1$ to the 108 separate group means and variances (after aggregating the data by lab and age group). The mean effect-size estimate was 0.35 (95% CI = [0.29, 0.42], $z = 10.67$, $p < .001$). A forest plot for this meta-analysis is shown in Figure 2. Further, 1,373 of the 2,329 infants (58.95%) showed a numerical preference for IDS.

Independent relationship of IDS preference to moderating variables. We next fitted a set of moderated meta-analytic models. We began by examining the relationship of IDS preferences to age, using the average age in months for each lab's sample as the moderator value. Labs that contributed samples from two age bins had a separate value added for each age (because of the small number of such cases, we did not model this dependency between labs). For ease of interpretation, we centered age in this analysis. The age-moderated model, $d_z \sim 1 + \text{age}$, yielded an estimated main effect of 0.35 (95% CI = [0.29, 0.41], $z = 11.47$, $p < .001$) and an age effect of 0.05 (95% CI = [0.03, 0.07], $z = 4.89$, $p < .001$). This positive age coefficient indicated that the measured IDS preference was on average larger for older children. The age trends for the NAE and non-NAE samples are plotted in Figure 3.

We next investigated effects of experimental method, with method dummy coded using single-screen central fixation as the reference level. The method-moderated model ($d_z \sim 1 + \text{method}$) yielded a reference-level intercept of 0.29 (95% CI = [0.18, 0.41], $z = 4.98$, $p < .001$), which is the mean effect size for single-screen presentation. The HPP yielded an additional effect of 0.21 (95% CI = [0.06, 0.37], $z = 2.74$, $p = .006$), a substantial gain in measured IDS preference for those labs using the HPP as compared with single-screen central fixation. In contrast, eye tracking yielded an effect of –0.06 (95% CI = [–0.21, 0.10], $z = –0.71$, $p = . 479$); thus, there was a slight, nonsignificant decrease in measured effect size when the method used was eye tracking, rather than single-screen central fixation.

The language-moderated model ($d_z \sim 1 + \text{language}$) was fitted with language group coded as a categorical variable indicating whether infants were tested in a lab in which NAE was the standard language (i.e., in the United States or Canada). The reference-level effect (i.e., not NAE) was 0.29 (95% CI = [0.20, 0.37], $z = 6.56$, $p < .001$). For infants in North American labs, the effect was increased by 0.15 (95% CI = [0.02, 0.27], $z = 2.26$, $p = .024$). Thus, measured IDS preferences were higher in those infants for whom the stimuli were native-language congruent.

Moderating variables' joint relationships with IDS preference. Because age group, language, and method were confounded across labs (labs with particular methods also chose specific sample age ranges, and these choices were not independent), we next turn to the mixed-effects modeling framework to estimate participant-level age effects and lab-level method effects. Figure 4a shows the spread of participant-level IDS preference, and Figure 4b shows estimated trends in IDS preference. The mean looking time across all trials was 8.21 s for the IDS condition and 7.38 s for the ADS condition.

Our main model was as follows:

log looking time ~ trial type * method + trial type
  * trial number + age * trial number + trial type
  * age * language + (trial type * trial number|participant)
  + (trial type * age|lab)
  + (method + age * language|item)

Trial type, language, and method were dummy coded with ADS trials, non-NAE community, and single-screen method as the reference levels; thus, coefficients are interpretable such that positive effects of trial type indicate longer looking on IDS trials, positive effects of language indicate longer looking in NAE communities,
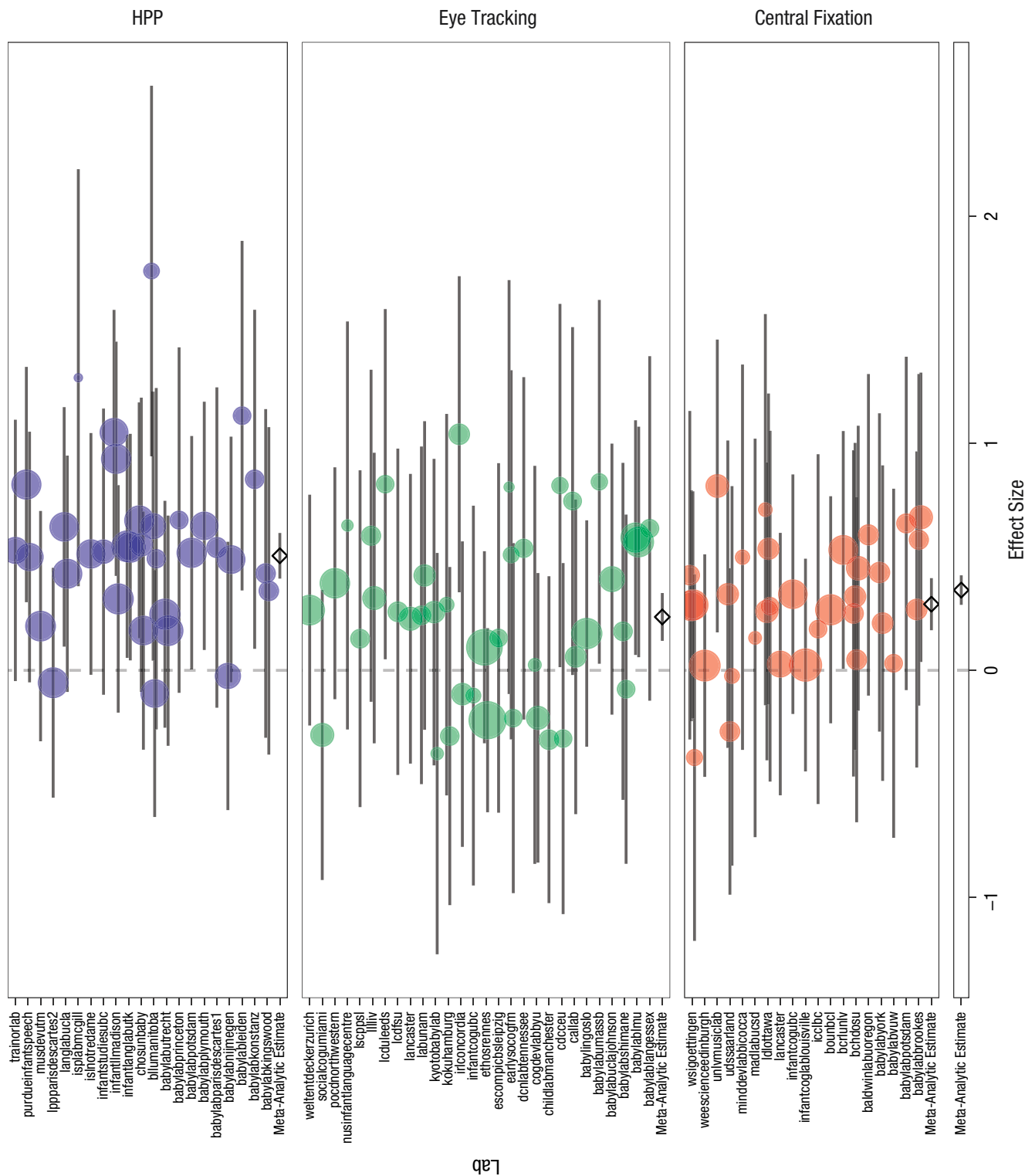
Fig. 2. Forest plot of the meta-analytic results for infant-directed-speech preference. The standardized effect size is shown for each lab; error bars indicate 95% confidence intervals. Labs are grouped into separate panels by method: head-turn preference procedure (HPP), eye tracking, or central fixation. Larger plotted points correspond to greater inverse variance. In each panel, the diamond and associated error bars represent the meta-analytic estimate from the method-moderated model and the estimate's 95% confidence interval. The bottom panel shows the global meta-analytic estimate and 95% confidence interval from the unmoderated model.
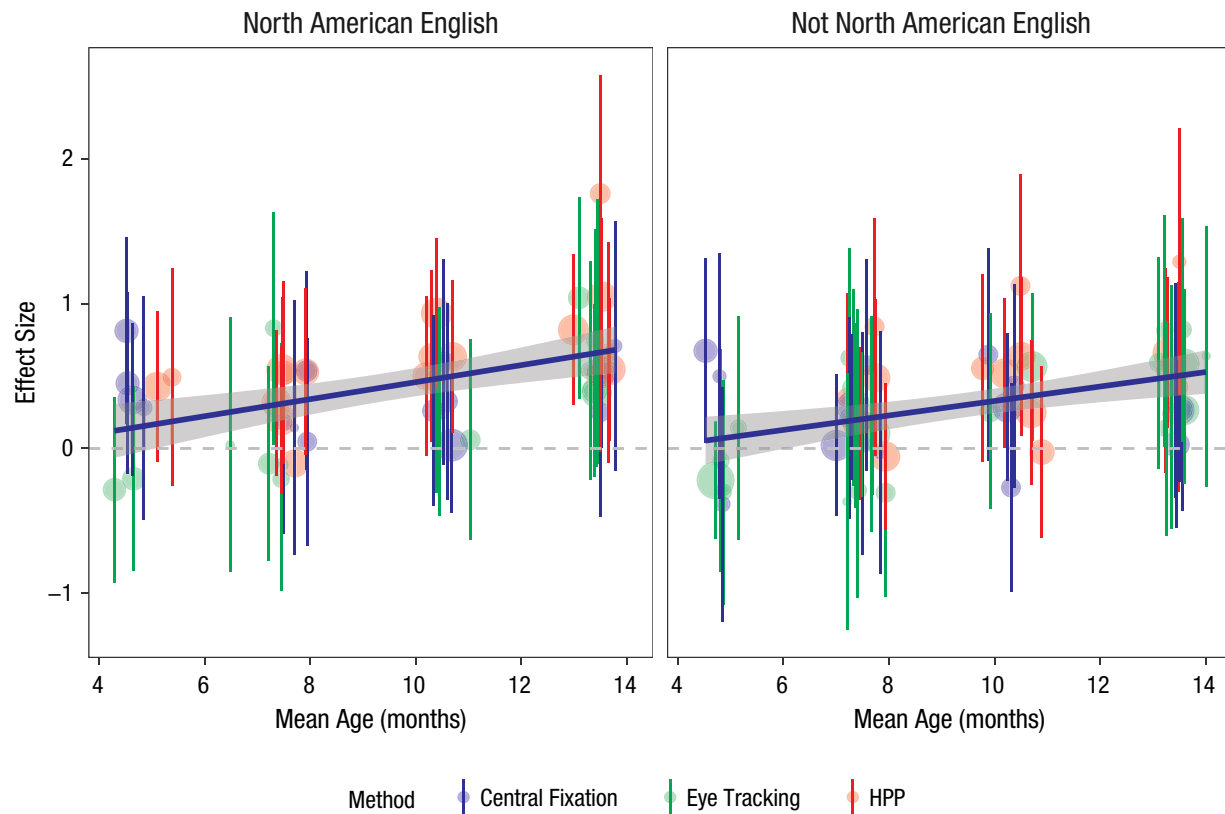
Fig. 3. Individual labs' standardized effect-size estimates as a function of age group. Error bars indicate 95% confidence intervals, and larger plotted points correspond to larger samples. The color coding indicates the method used: central fixation, eye tracking, or head-turn preference procedure (HPP). Also shown are regression smoothing lines, with gray bands indicating 95% confidence intervals. Results are shown separately for infants learning North American English (left panel) and infants learning other languages and dialects (right panel).

and positive effects of method indicate longer looking with eye tracking and the HPP. To increase the interpretability of coefficients, we centered age (in months) and coded trial number with Trial 1 as the reference level.

We specified this model to minimize higher-order interactions but preserve theoretically important interactions. We included main effects of trial type, method, language, age, and trial number, capturing the basic effects of each on looking time (e.g., longer looking times for IDS, shorter looking times on later trials). In addition, we included two-way interactions of trial type with method (modeling the possibility that some methods would show larger IDS preferences than others) and trial type with trial number (modeling the possibility of faster habituation to ADS), as well as an interaction of age and trial number (modeling faster habituation for older children). We also included two- and three-way interactions of age, trial type, and language (modeling possible developmental changes in IDS preference and

developmental differences in IDS preference across language groups). Both developmental effects and trial effects were treated linearly in this model; although they likely had nonlinear effects, adding quadratic or other effects would have substantially increased the model's complexity. After pruning random effects for nonconvergence, our final model specification was

log looking time ~ trial type * method + trial type *
    trial number + age * trial number + trial type * age *
    language + (1|participant) + (1|lab) + (1|item)

Table 4 shows the coefficient estimates from this model.

Overall, the fitted coefficients of the mixed-effects model were consistent with the results of the individual meta-analyses. Given the structure of the mixed-effects model, positive coefficients for the IDS predictor indicated greater IDS preference (i.e., greater looking times on IDS trials). The fitted model showed a significant
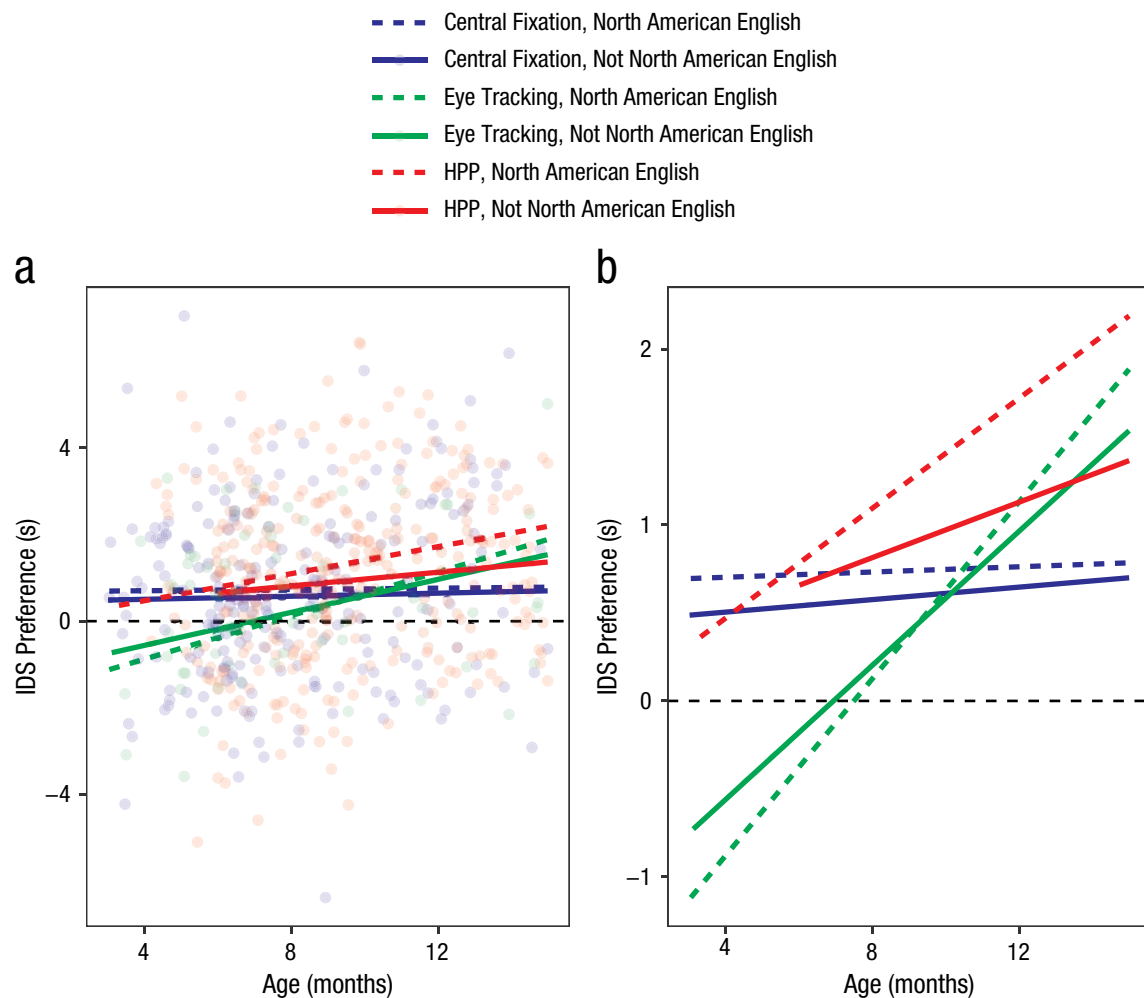
Fig. 4. Simple linear trends for infant-directed-speech (IDS) preference by age, language group, and method: central fixation, eye tracking, or head-turn preference procedure (HPP). Individual participants' preferences are plotted in (a) but omitted in (b) to show the trends more effectively.

positive effect of IDS stimuli, consistent with a global IDS preference. Results were also consistent with the age- and language-moderated meta-analyses, as there were significant and positive two-way interactions of IDS with age and with NAE; preferences for IDS were greater among older children and children in NAE contexts. Further, there was a positive interaction of IDS with the HPP method, consistent with the method-moderated model. There was not a significant three-way interaction of IDS, age, and NAE, however; in other words, there was not a reliable differential change in IDS preference for older children in NAE contexts over and above that expected given each of these factors alone.

In addition, a number of other factors were significant predictors of looking time. Looking time decreased

across trials and was shorter among older children. This result generally confirmed that all infants habituated to our experimental stimuli, and older infants did so more quickly. Further, eye tracking led to shorter looking times overall across stimulus classes.

Effect of second-session testing on IDS preference. We preregistered an analysis of whether the pattern of IDS preference was different for second-session infants than for first-session infants (i.e., those who completed the IDS-preference experiment as their first, and possibly only, session). Only six labs contributed data for second-session infants, however, and only 41 infants were represented. Thus, we did not fit the full, preregistered mixed-effects model for this variable, as we did not have enough variability on the important covariates. We note that 46.34%

Table 4. Coefficient Estimates From the Linear Mixed-Effects Model Predicting Log Looking Time

Predictor	Estimate	t	p
Intercept	2.180 (0.051)	43.100	.000
IDS	0.099 (0.036)	2.740	.010
Eye tracking	−0.265 (0.046)	−5.790	.000
HPP	−0.052 (0.051)	−1.020	.308
Trial number	−0.038 (0.002)	−25.000	.000
Age	−0.035 (0.004)	−7.950	.000
NAE	−0.016 (0.049)	−0.335	.738
IDS * Eye Tracking	−0.009 (0.017)	−0.548	.584
IDS * HPP	0.034 (0.015)	2.270	.023
IDS * Trial Number	−0.003 (0.002)	−1.370	.172
Trial Number * Age	0.001 (0.000)	3.140	.002
IDS * Age	0.012 (0.003)	4.300	.000
IDS * NAE	0.039 (0.013)	3.060	.002
Age * NAE	0.001 (0.006)	0.198	.843
IDS * Age * NAE	0.004 (0.004)	1.050	.292

Note: Numbers in parentheses are standard errors. IDS = infant-directed speech; HPP = head-turn preference procedure; NAE = North American English.

(19/41) of second-session infants (95% CI = [31.65%, 61.30%]) showed a numerical preference for IDS. This percentage was numerically different, but not distinguishable statistically, from the 58.95% of first-session infants who showed an IDS preference, likely because of the small sample of second-session infants.

Sex and IDS preference. In order to investigate effects of biological sex on IDS preference, we fitted the mixed-effects model specified earlier with the addition of a main effect of sex and a trial-type-by-sex interaction. Female was coded as the reference level. The main effect of sex was not significant, $\beta = 0.01$ ($SE = 0.02$, $p = .63$), and neither was the interaction of sex with trial type, $\beta = −0.01$ ($SE = 0.01$, $p = .51$). These small, nonsignificant coefficients suggest that sex was not a strong determinant of measured IDS preferences in our data.

Moderator effects on missing data. One further question regarding our data was whether particular moderator variables affected not only the amount of looking time recorded, but also whether children looked at all during a trial. To test for effects of moderators on the presence of missing data, we constructed a categorical variable (missing), which was true if a trial had no included looking time (i.e., no looking recorded, a look under 2 s, or no looking because the experiment was already terminated) and false otherwise. We fitted a logistic mixed-effects model with all two-way interactions between method, age, and trial number, using the following specification:

$$\text{missing data point} \sim \text{method} * \text{age} + \text{method} *$$
$$\text{trial number} + \text{age} * \text{trial number} + (1|\text{participant}) +$$
$$(\text{trial number} * \text{age}|\text{lab}) + (\text{method} + \text{age}|\text{item})$$

After pruning for nonconvergence, our final model specification was

$$\text{missing data point} \sim \text{method} * \text{age} +$$
$$\text{method} * \text{trial number} + \text{age} * \text{trial number} + (1|\text{lab})$$

Table 5 shows the coefficient estimates from this model. To aid convergence, we centered and scaled age and trial number, and set single-screen presentation as the reference level. Positive coefficients indicate a higher probability of missing data. More data were missing for older children and later trials, a pattern consistent with the idea that all children habituated to the stimuli, but that older children habituated faster. There was also a significant negative interaction of age and eye tracking, suggesting that data loss for eye tracking was substantially greater in younger children and lower in older children. The other coefficients were relatively small and nonsignificant.

Exploratory analyses

Meta-analytic heterogeneity. One question of interest was whether we observed any meta-analytic heterogeneity in the data. A finding of meta-analytic heterogeneity indicates the presence of unexplained variance in effect size over and above that due to sampling variation. We calculated τ^2 as an estimate of the total heterogeneity in

Table 5. Coefficient Estimates From the Linear Mixed-Effects Model Predicting Whether an Observation Was Missing

Predictor	Estimate	z	p
Intercept	−1.090 (0.152)	−7.140	.000
Eye tracking	0.167 (0.130)	1.290	.198
HPP	−0.178 (0.195)	−0.913	.361
Age	0.356 (0.038)	9.380	.000
Trial number	0.663 (0.030)	22.100	.000
Eye Tracking * Age	−0.238 (0.047)	−5.090	.000
HPP * Age	−0.059 (0.051)	−1.150	.251
Eye Tracking * Trial Number	0.068 (0.036)	1.850	.064
HPP * Trial Number	0.046 (0.040)	1.130	.257
Trial Number * Age	−0.003 (0.014)	−0.208	.835

Note: Numbers in parentheses are standard errors. HPP = head-turn preference procedure.

Table 6. Meta-Analytic Effect Size (d_z) and Percentage of Included Participants for Three Different Inclusion Criteria

	Minimum number of trials					
	2 trials		4 trials		8 trials	
Method	Estimate	Included participants (%)	Estimate	Included participants (%)	Estimate	Included participants (%)
Central fixation	0.29 (0.06)	98	0.34 (0.06)	88	0.40 (0.06)	73
Eye tracking	0.24 (0.06)	85	0.33 (0.06)	59	0.41 (0.10)	36
HPP	0.51 (0.06)	98	0.56 (0.06)	92	0.63 (0.07)	78

Note: Numbers in parentheses are standard errors. HPP = head-turn preference procedure.

our models. In addition, we assessed heterogeneity using the I^2 statistic (Higgins, Thompson, Deeks, & Altman, 2003), which quantifies the proportion of total variation in estimates that is due to heterogeneity. We also report the results of a standard hypothesis test for heterogeneity, the Cochran Q test; a statistically significant Q test indicates that the null hypothesis of homogeneity of variance can be rejected (Huedo-Medina, Sanchez-Meca, Marin-Martinez, & Botella, 2006).

In our primary, intercept-only meta-analytic model, we found nonsignificant heterogeneity, $\tau^2 = 0.01$, $I^2 = 12.39\%$, and $Q(107) = 122$, $p = .15$. In the language-moderated model, heterogeneity was also nonsignificant, $\tau^2 = 0.01$, $I^2 = 7.76\%$, and $Q(106) = 116.18$, $p = .23$. In the age-moderated model, heterogeneity was even lower, $\tau^2 = 0.00$, $I^2 = 0.00\%$, and $Q(106) = 98.06$, $p = .70$. Finally, in the method-moderated model, heterogeneity was also low, $\tau^2 = 0.00$, $I^2 = 3.20\%$, and $Q(105) = 106.78$, $p = .43$. For none of these models could we reject the null hypothesis of no heterogeneity beyond sampling variation, and in no case was the magnitude of observed heterogeneity large. Although there were reliable moderators (see the meta-analytic results reported earlier), the effects of these moderators were quite small in magnitude relative to the sampling variation in individual labs' effect-size estimates (because of the small median sample size within each lab).

Usable-data inclusion criterion. Because our criterion for including infants in the analysis was so liberal (i.e., infants needed to contribute data from only two trials to be included), we next explored the effects of different inclusion rules on the results. In particular, we calculated the meta-analytic effect size with four trials and eight trials as the minimum inclusion criterion. With a minimum of four trials, the effect size was 0.42 (95% CI = [0.35, 0.48], $z = 12.05$, $p < .001$), and with a minimum of eight trials, the effect size was 0.48 (95% CI = [0.40, 0.57], $z = 11.23$, $p < .001$). In comparison, our original analysis

yielded a meta-analytic effect size of 0.35 (95% CI = [0.29, 0.42], $z = 10.67$, $p < .001$). Furthermore, we computed the effect size for each method for each of these alternative inclusion criteria (see Table 6). Overall, more stringent inclusion criteria yielded substantially larger effects, although they also led to substantial data loss (especially for labs using the eye-tracking method).

General Discussion

We designed a large-scale, multilab study of infants' preference for IDS and invited infancy researchers to participate. Our call for participation resulted in contributions from 69 labs, representing a total of 2,845 infants from 16 countries. The final sample used for analysis included 2,329 infants (see Table 1). We believe that this is the largest laboratory study of infancy to date. We begin our discussion by summarizing the principal results of the study with respect to four analytic questions and then discuss limitations of the study as well as future directions.

Summary of findings

Our first goal was to address the issue of the replicability of infants' preference for IDS over ADS by conducting a preregistered study to obtain an unbiased measure of the magnitude of this preference. We expected to replicate prior demonstrations of the existence of an IDS preference in infant listeners, and our study indeed confirmed this expectation. Our overall meta-analytic mean was smaller in size than the effect found in a preceding meta-analysis of the literature, however (Bergmann et al., 2018; Dunst et al., 2012).

Although one possible interpretation of this difference in magnitude is that previously obtained effect sizes were inflated by publication bias, there are other possible explanations as well. In an individual laboratory, the methodology is tailored to the specific research

question and the age range and other characteristics of the infants tested (or, conversely, research questions are tailored to the existing methodological expertise of the laboratory). The approach used here, namely, applying multiple methodologies to the same research question across diverse age ranges and samples of infants, including non-English-learning infants, may have led to an underestimate of the true effect size (i.e., because the ideal presentation details that would maximize effect sizes might differ across methods and ages). Further, several of our methodological decisions might have decreased the effect size. For example, our stimuli had less extreme acoustic characteristics than the stimuli used in previous work and were generated by multiple speakers. In addition, our criterion for including participants in the final sample was less stringent than the criteria used previously.

Our second goal was to examine possible age effects in the preference for IDS. As did the prior published meta-analysis (Dunst et al., 2012), we found an increase in IDS preference across development. This trend is consistent with the idea that preference for IDS grows in response to experience with positive social interactions, but contrasts with some other reports in the literature (e.g., Hayashi et al., 2001; Newman & Hussain, 2006; Segal & Newman, 2015). Further, the magnitude of the positive developmental change we observed was considerable, at 0.05 standard deviations per month. This finding suggests that the preference for IDS is at a minimum modulated by experience, maturation, or both.

Any developmental trend, however, may be driven by changes in factors other than the underlying construct. As we discuss later, the stimuli we used may have been best suited for the older age ranges in our study. In addition, stronger effects among older infants may result from a more robust or more measurable behavioral response, independently of an underlying preference. Some evidence in favor of this possibility can be found in the data in MetaLab, an online data bank for meta-analysis in infancy research: Most extant meta-analyses show an increase in absolute effect size as infants mature, regardless of the research question (see, e.g., Bergmann et al., 2018).

Our third goal was to examine how the preference for IDS varies with the differing linguistic experiences of infants growing up in different linguistic communities. We found a preference for NAE IDS over NAE ADS even among participants whose native language or dialect was not NAE. This finding replicates previous work (Werker et al., 1994). However, in our study, NAE-exposed infants showed the strongest preference. Note that our findings do not support the idea of a simple attentional effect (i.e., greater attention to speech

overall when presented in the native language): The effect of language background on overall (as opposed to preferential) looking times was not large in our regression models.

There are several possible interpretations of the native-language effect we observed. One possibility is that as infants become experts in their native language's phonology and begin to acquire word meanings, they listen to speech in their own language differently, starting to process what is being said not just as "speech" or "register" per se, but as meaningful language (Gervain & Mehler, 2010; Johnson, 2016). For infants hearing a foreign language or even a foreign dialect of their native language, the ability to listen in this deeper, or more predictive, way is not available. Another possibility is that processing speech in an unfamiliar language requires more attentional resources, leaving fewer attentional resources to process some of the characteristics that may differentiate IDS and ADS. Regardless of which of these possibilities is true, preference for IDS may depend in part on the similarity of the IDS to one's native-language experiences with IDS. This idea is somewhat supported by the age effect we observed; however, we did not observe a three-way interaction among age, stimulus type, and language background, which this interpretation would predict. Companion data are currently being collected in several non-NAE language communities using native-language stimuli created using the ManyBabies 1 protocol, and these data may shed further light on this issue.

A fourth goal was to examine whether the measured experimental effect varied with the methodological approach. We found a stronger effect with the HPP than with the central-fixation or eye-tracking approaches. One potential interpretation of this finding is that the greater effort on the part of the infant in the HPP (i.e., turning the head, as opposed to making small eye movements) leads to stronger engagement in the task and therefore to stronger effects.

It is important to keep in mind, however, that methodology was not randomly assigned to laboratories, and the characteristics of laboratories probably varied systematically with their methodological choices. It may well be, for example, that laboratories with more expertise in investigating infants' language acquisition were more likely to use the HPP. Furthermore, these findings should not be interpreted as suggesting that the HPP would be best suited for all research questions. Instead, a more modest interpretation is simply that a theoretically irrelevant variable related to laboratories and their methodological decisions appears to have a substantial and systematic effect on measured effect size (see Bergmann et al., 2018, for a similar conclusion based on meta-analytic data). We hope to undertake secondary

analyses of our data set to better understand factors that may have covaried with methodological choices. Moreover, further large-scale projects that include methodological contrasts of this type—perhaps with random assignment—may allow more specific conclusions about the sources of methodological variability and their interactions with the investigated phenomenon and participants' age.

Another methodological contribution of this project was our investigation of how different infant-level inclusion criteria affect the magnitude of the obtained effect size. For our main analysis, we included all infants who completed at least one IDS and one ADS trial. This is something of a departure from the norm, as most participating labs reported using a stricter inclusion criterion in their own independent work. Our meta-analytic effect size was 0.35 when we included all infants with a minimum of two trials, grew to 0.42 when the minimum was raised to four trials, and increased further to 0.48 when the minimum was eight trials. Moreover, the effect of age on the amount of missing data was substantially larger in the eye-tracking paradigm compared with the other methods. The amount of missing data increased across the length of the experiment; this increase was numerically most prevalent for eye tracking. Setting stricter inclusion criteria necessarily decreases final sample size if the total sample tested remains constant, but at the same time, stricter criteria appear to lead to more robust effects in this paradigm.

Challenges and limitations

As with any study, the current experiment required specific methodological choices, several of which influence the generalizability of our results. Two aspects of the decision making regarding the stimuli in particular are worth further discussion. The first is the choice to use NAE (as opposed to, say, the native language or dialect for each infant group tested). This choice was based on the need to use consistent stimuli across laboratories, in order to limit cross-lab variation and ensure feasibility of the overall project, and to use stimuli from a language in which there was robust evidence of a strong IDS-preference effect, both in native and non-native settings. However, our design necessarily complicates the interpretability of our findings from laboratories outside of North America. They confound effects of native language or dialect (infants prefer listening to their native language) and true cultural variation in IDS preference. Further, substantial diversity in the non-NAE samples was obscured in our preregistered analyses. Together with the previously mentioned native-language follow-up study using the ManyBabies

1 protocol, further analyses of our data set focusing on specific subsamples with sufficient sample size (e.g., French, German, Dutch, British English) will shed additional light on how the differences between the North American and other infants in the current study should be interpreted.

The second challenging decision hinged on the elicitation of the IDS stimuli. Stimuli used in previous IDS-preference studies range from scripted speech with no infant present (e.g., Cooper & Aslin, 1990; Newman & Hussain, 2006), which maximizes control over the experimental stimuli, to more naturalistic samples collected in free-play, unscripted contexts (e.g., Hayashi et al., 2001; Werker et al., 1994), which maximize generalizability to real-world contexts. We opted for a relatively naturalistic approach, with an elicitation protocol using real mothers and their infants and centering around concrete objects. It is likely that this approach led to the observed reduction in the distinctiveness of the acoustic characteristics of our IDS samples, and it limited our ability to fully control the characteristics of the samples. Other aspects of our elicitation approach are important to keep in mind in interpreting our finding, including our developmental effects. In particular, our speech stimuli were elicited from mothers speaking to 4- to 8-month-old infants in the context of an objects-focused task (which was likely best suited to infants at the older range of our age bins). The extent to which these age-related characteristics of the IDS stimuli affected the magnitude of infants' IDS preference across development merits further inquiry. Further, the use of multiple speakers in the stimuli may have increased the processing load for infants.

As the first collaboration of its kind, ManyBabies 1 revealed a number of important challenges in conducting multilab infancy research. As any researcher who has tested infant participants knows, data collection with infants is slow and labor intensive. In the current project, over a period of approximately 13 months, 69 labs were able to collect data from 2,845 infants. In contrast, in ManyLabs 1, a similar initiative with adult participants (Klein et al., 2014), data were collected from more than 6,000 participants tested in 36 labs over just a handful of months. Moreover, whereas adults can often be tested in multiple studies in a single session, this option is very limited for infants.

We expected challenges in implementing a standardized data-collection procedure across labs, but the depth of these challenges, and the diversity of methodological implementation across laboratories, was surprising. Laboratories that conduct infancy research are highly diverse in both the software and the hardware they have available to implement testing methods. We planned flexibility in the specific setup (eye

tracking, HPP, central fixation) given this known variability, but despite this flexibility, several labs were forced to deviate from aspects of the protocol, for example, because of limitations in how stimuli could be presented (e.g., the ability to implement infant-controlled trial lengths, software settings for repeating trials). One important conclusion from our work, as evidenced in the walk-through videos laboratories provided to illustrate their protocols (see the next paragraph), is that typical Method sections fail to provide the detail necessary to capture the methodological diversity in the ways that different labs implement a paradigm with the same name.

Additional benefits of large-scale collaboration

Although our primary goals were empirical, the ManyBabies 1 project offers numerous additional benefits to both individual researchers and the field at large. All of the questionnaires, how-tos, and stimuli (e.g., attention getters) used in the project are freely available for reuse in future studies. Each participating lab created a walk-through video that showed the lab and study setup. These videos provide an unprecedented peek "behind the curtain" of other infancy labs—something that previously was possible only through visiting labs in person. Such information could be a particularly helpful resource for investigators setting up an infancy lab for the first time. It also provides a unique data set whereby infancy researchers can begin to understand the variety of lab setups and study implementations.

This large-scale collaborative effort also has provided broader benefits for the field. It created a strong collaborative network of infancy researchers. Informal "ManyBabies" gatherings are now organized at developmental conferences, enabling researchers who have previously collaborated only virtually to meet in person. This project also was many participating researchers' introduction to open-science practices and tools, such as preregistration and OSF.

Finally, ManyBabies 1 has launched several "knock-on" projects. For example, ManyBabies Bilingual (Byers-Heinlein et al., 2020) is comparing bilingual infants' preference for IDS with our results for monolingual infants. Other projects will examine the test-retest reliability of infants' IDS preference, examine whether IDS preference predicts vocabulary size at 18 and 24 months (Soderstrom et al., 2020), and test whether lab-specific variables affect infants' performance and attrition. We believe that these additional benefits are not unique to infancy research, and that other scientific communities embarking on large-scale collaborative projects will garner similar benefits.

Conclusion

Replication research can go far beyond simply asking whether an effect is present: It can allow for an assessment of how an effect varies and how it changes over development. We observed a robust and statistically significant preference for IDS over ADS, confirming previous observations in the literature. Yet the value of our experiment lies not purely in this binary result—or even in the quantitative estimate of the overall magnitude of the IDS preference—but also in the further theoretical and methodological opportunities that the data afford. By measuring the relationship of IDS preference to age and language community, this experiment provides a starting point for developing a more nuanced theory of how IDS preference relates to children's language experiences. Further, by revealing the substantial contributions of methodological decision making to observed effect sizes, our study points the way toward developing best-practices templates in further infancy work of this kind. In sum, we hope our work reported here illustrates the power of large-scale collaboration for the study of developmental variation and change.

Transparency

Action Editor: Daniel J. Simons
Editor: Daniel J. Simons
Consortium Members
 Michael C. Frank, Stanford University
 Katherine Jane Alcock, Lancaster University
 Natalia Arias-Trejo, Universidad Nacional Autónoma de México (UNAM)
 Gisa Aschersleben, Saarland University
 Dare Baldwin, University of Oregon
 Stéphanie Barbu, Université de Rennes 1 and Centre national de la recherche scientifique
 Elika Bergelson, Duke University
 Christina Bergmann, Max Planck Institute for Psycholinguistics, Nijmegen, The Netherlands
 Alexis K. Black, Haskins Laboratories, New Haven, Connecticut
 Ryan Blything, University of Bristol
 Maximilian P. Böhland, Technische Universität Dresden
 Petra Bolitho, Victoria University of Wellington
 Arielle Borovsky, Purdue University
 Shannon M. Brady, UCLA
 Bettina Braun, University of Konstanz
 Anna Brown, University of Liverpool
 Krista Byers-Heinlein, Concordia University
 Linda E. Campbell, University of Newcastle, Australia
 Cara Cashon, University of Louisville

Mihye Choi, University of Massachusetts Boston
Joan Christodoulou, UCLA
Laura K. Cirelli, University of Toronto Mississauga
Stefania Conte, University of Milano-Bicocca
Sara Cordes, Boston College
Christopher Cox, University of York
Alejandrina Cristia, PSL University and Centre national de la recherche scientifique
Rhodri Cusack, Trinity College Dublin
Catherine Davies, University of Leeds
Maartje de Klerk, Utrecht University
Claire Delle Luche, University of Essex
Laura de Ruiter, University of Manchester
Dhanya Dinakar, Western Sydney University
Kate C. Dixon, University of Louisville
Virginie Durier, Université de Rennes 1 and Centre national de la recherche scientifique
Samantha Durrant, University of Liverpool
Christopher Fennell, University of Ottawa
Brock Ferguson, Strong Analytics, Chicago, Illinois
Alissa Ferry, University of Manchester
Paula Fikkert, Radboud University
Teresa Flanagan, Franklin & Marshall College
Caroline Floccia, University of Plymouth
Megan Foley, Florida State University
Tom Fritzsche, University of Potsdam
Rebecca L. A. Frost, Max Planck Institute for Psycholinguistics, Nijmegen, The Netherlands
Anja Gampe, University of Zurich
Judit Gervain, Université Paris Descartes
Nayeli Gonzalez-Gomez, Oxford Brookes University
Anna Gupta, Leiden University
Laura E. Hahn, Radboud University
J. Kiley Hamlin, University of British Columbia
Erin E. Hannon, University of Nevada, Las Vegas
Naomi Havron, PSL University and Centre national de la recherche scientifique
Jessica Hay, University of Tennessee, Knoxville
Mikołaj Hernik, Central European University
Barbara Höhle, University of Potsdam
Derek M. Houston, The Ohio State University
Lauren H. Howard, Franklin & Marshall College
Mitsuhiko Ishikawa, Kyoto University
Shoji Itakura, Kyoto University
Iain Jackson, University of Manchester
Krisztina V. Jakobsen, James Madison University
Marianna Jarto, University of Hamburg
Scott P. Johnson, UCLA
Caroline Junge, Utrecht University
Didar Karadag, Boğaziçi University
Natalia Kartushina, University of Oslo
Danielle J. Kellier, Stanford University
Tamar Keren-Portnoy, University of York
Kelsey Klassen, University of Manitoba
Melissa Kline, Massachusetts Institute of Technology
Eon-Suk Ko, Chosun University
Jonathan F. Kominsky, Harvard University
Jessica E. Kosie, University of Oregon

Haley E. Kragness, McMaster University
Andrea A. R. Krieger, Saarland University
Florian Krieger, University of Luxembourg
Jill Lany, University of Notre Dame
Roberto J. Lazo, University of Miami
Michelle Lee, University of California, San Diego
Chloé Leservoisier, Université de Rennes 1 and Centre national de la recherche scientifique
Claartje Levelt, Leiden University
Casey Lew-Williams, Princeton University
Matthias Lippold, University of Goettingen
Ulf Liszkowski, University of Hamburg
Liquan Liu, Western Sydney University
Steven G. Luke, Brigham Young University
Rebecca A. Lundwall, Brigham Young University
Viola Macchi Cassia, University of Milano-Bicocca
Nivedita Mani, University of Goettingen
Caterina Marino, Université Paris Descartes
Alia Martin, Victoria University of Wellington
Meghan Mastroberardino, Concordia University
Victoria Mateu, UCLA
Julien Mayor, University of Oslo
Katharina Menn, Radboud University
Christine Michel, Max Planck Institute for Human Cognitive and Brain Sciences, Leipzig, Germany
Yusuke Moriguchi, Kyoto University
Benjamin Morris, University of Chicago
Karli M. Nave, University of Nevada, Las Vegas
Thierry Nazzi, Université Paris Descartes
Claire Noble, University of Liverpool
Miriam A. Novack, Northwestern University
Nonah M. Olesen, University of Louisville
Adriel John Orena, McGill University
Mitsuhiko Ota, University of Edinburgh
Robin Panneton, Virginia Polytechnic Institute and State University
Sara Parvanezadeh Esfahani, University of Tennessee, Knoxville
Markus Paulus, Ludwig Maximilian University of Munich
Carolina Pletti, Ludwig Maximilian University of Munich
Linda Polka, McGill University
Christine Potter, Princeton University
Hugh Rabagliati, University of Edinburgh
Shruthilaya Ramachandran, National University of Singapore
Jennifer L. Rennels, University of Nevada, Las Vegas
Greg D. Reynolds, University of Tennessee, Knoxville
Kelly C. Roth, University of Tennessee, Knoxville
Charlotte Rothwell, Lancaster University
Doroteja Rubez, The Ohio State University
Yana Ryjova, University of Nevada, Las Vegas
Jenny Saffran, University of Wisconsin–Madison
Ayumi Sato, Shimane University
Sophie Savelkouls, Boston College
Adena Schachner, University of California, San Diego
Graham Schafer, University of Reading
Melanie S. Schreiner, University of Goettingen
Amanda Seidl, Purdue University

Mohinish Shukla, University of Massachusetts Boston
Elizabeth A. Simpson, University of Miami
Leher Singh, National University of Singapore
Barbora Skarabela, University of Edinburgh
Gaye Soley, Boğaziçi University
Megha Sundara, UCLA
Anna Theakston, University of Manchester
Abbie Thompson, University of Notre Dame
Laurel J. Trainor, McMaster University
Sandra E. Trehub, University of Toronto Mississauga
Anna S. Trøan, University of Oslo
Angeline Sin-Mei Tsui, University of Ottawa
Katherine Twomey, University of Manchester
Katie Von Holzen, Université Paris Descartes
Yuanyuan Wang, The Ohio State University
Sandra Waxman, Northwestern University
Janet F. Werker, University of British Columbia
Stephanie Wermelinger, University of Zurich
Alix Woolard, University of Newcastle, Australia
Daniel Yurovsky, University of Chicago
Katharina Zahner, University of Konstanz
Martin Zettersten, University of Wisconsin–Madison
Melanie Soderstrom, University of Manitoba

Author Contributions

The order in which authors are listed here reflects the authorship order, rather than the order of their contribution to each of the individual elements. M. C. Frank, E. Bergelson, C. Bergmann, K. Byers-Heinlein, B. Ferguson, J. Gervain, J. K. Hamlin, M. Kline, C. Levelt, C. Lew-Williams, C. Marino, T. Nazzi, R. Panneton, H. Rabagliati, A. Seidl, and M. Soderstrom contributed to the study concept. M. C. Frank, C. Bergmann, K. Byers-Heinlein, C. Floccia, J. Gervain, N. Gonzalez-Gomez, J. K. Hamlin, E. E. Hannon, M. Kline, C. Lew-Williams, T. Nazzi, R. Panneton, H. Rabagliati, J. L. Rennels, S. Waxman, D. Yurovsky, and M. Soderstrom contributed to the study's design. M. C. Frank, R. Cusack, C. Floccia, D. J. Kellier, K. Klassen, C. Lew-Williams, R. Panneton, M. Shukla, and M. Soderstrom contributed to the creation of the stimuli. N. Gonzalez-Gomez, J. K. Hamlin, and D. J. Kellier contributed to pilot testing. M. C. Frank, C. Bergmann, R. Blything, K. Byers-Heinlein, C. Delle Luche, L. de Ruiter, B. Ferguson, I. Jackson, M. Kline, J. F. Kominsky, M. Mastroberardino, K. Twomey, and D. Yurovsky contributed to the final protocol. M. C. Frank, C. Bergmann, K. Byers-Heinlein, J. Gervain, M. Kline, C. Lew-Williams, M. Mastroberardino, and M. Soderstrom contributed to the documentation of the study. M. C. Frank, C. Bergmann, K. Byers-Heinlein, R. L. A. Frost, J. K. Hamlin, M. Kline, C. Lew-Williams, K. Twomey, and M. Soderstrom contributed to study management. K. J. Alcock, N. Arias-Trejo, G. Aschersleben, D. Baldwin, S. Barbu, A. K. Black, M. P. Böhland, P. Bolitho, A. Borovsky, S. M. Brady, B. Braun, A. Brown, K. Byers-Heinlein, L. E. Campbell, C. Cashon, M. Choi, J. Christodoulou, L. K. Cirelli, S. Conte, S. Cordes, C. Cox, A. Cristia, C. Davies, M. de Klerk, C. Delle Luche, L. de Ruiter, D. Dinakar, K. C. Dixon, V. Durier, S. Durrant, C. Fennell, A. Ferry, P. Fikkert, T. Flanagan, C. Floccia, M. Foley, T. Fritzsche, R. L. A. Frost, A. Gampe, J. Gervain, N. Gonzalez-Gomez, A. Gupta, L. E. Hahn, J. K. Hamlin, E. E. Hannon, N. Havron, J. Hay, M. Hernik, B. Höhle, D. M. Houston, L. H. Howard, M. Ishikawa, S. Itakura, I. Jackson, K. V. Jakobsen, M. Jarto, S. P. Johnson, C. Junge, D. Karadag, N. Kartushina, T. Keren-Portnoy, K. Klassen, E.-S. Ko, J. E. Kosie, H. E. Kragness, A. A. R. Krieger, F. Krieger, J. Lany, R. J. Lazo, M. Lee, C. Leservoisier, C. Levelt, U. Liszkowski, L. Liu, S. G. Luke, R. A. Lundwall, V. Macchi Cassia, N. Mani, C. Marino, A. Martin, M. Mastroberardino, V. Mateu, J. Mayor, K. Menn, C. Michel, Y. Moriguchi, B. Morris, K. M. Nave, C. Noble, M. A. Novack, N. M. Olesen, A. J. Orena, M. Ota, R. Panneton, S. Parvanezadeh Esfahani, M. Paulus, C. Pletti, L. Polka, C. Potter, H. Rabagliati, S. Ramachandran, J. L. Rennels, G. D. Reynolds, K. C. Roth, C. Rothwell, D. Rubez, Y. Ryjova, J. Saffran, A. Sato, S. Savelkouls, A. Schachner, G. Schafer, M. S. Schreiner, A. Seidl, E. A. Simpson, L. Singh, B. Skarabela, G. Soley, M. Sundara, A. Theakston, A. Thompson, L. J. Trainor, S. E. Trehub, A. S. Trøan, A. S.-M. Tsui, K. Twomey, K. Von Holzen, Y. Wang, S. Waxman, J. F. Werker, S. Wermelinger, A. Woolard, D. Yurovsky, K. Zahner, M. Zettersten, and M. Soderstrom contributed to data collection. M. C. Frank, C. Bergmann, A. Cristia, M. Kline, J. E. Kosie, M. Lippold, H. Rabagliati, A. S.-M. Tsui, A. Woolard, M. Zettersten, and M. Soderstrom contributed to data analysis. M. C. Frank, E. Bergelson, C. Bergmann, K. Byers-Heinlein, A. Cristia, R. Cusack, C. Floccia, J. Gervain, N. Gonzalez-Gomez, J. K. Hamlin, E. E. Hannon, M. Kline, C. Lew-Williams, R. A. Lundwall, T. Nazzi, H. Rabagliati, J. L. Rennels, and M. Soderstrom contributed to the Stage 1 manuscript. M. C. Frank, C. Bergmann, K. Byers-Heinlein, A. Cristia, J. Gervain, J. K. Hamlin, M. Kline, M. Lippold, C. Marino, J. L. Rennels, and M. Soderstrom contributed to the Stage 2 manuscript.

Declaration of Conflicting Interests

The author(s) declared that there were no conflicts of interest with respect to the authorship or the publication of this article.

Open Practices

    Open Data: https://osf.io/re95x/
    Open Materials: https://osf.io/re95x/
    Preregistration: https://osf.io/gf7vh
    All data and materials have been made publicly available via the Open Science Framework and can be accessed at https://osf.io/re95x/. The design and analysis plans were preregistered at the Open Science Framework and can be accessed at https://osf.io/gf7vh/. The complete Open Practices Disclosure for this article can be found at http://journals.sagepub.com/doi/suppl/10.1177/2515245 919900809. This article has received badges for Open Data, Open Materials, and Preregistration. More information about the Open Practices badges can be found at http://www.psychologicalscience.org/publications/badges.

ORCID iD

Michael C. Frank 🄳 https://orcid.org/0000-0002-7551-4378

Prior Versions

A previous version of this manuscript was posted at https://psyarxiv.com/s98ab/.

References

Barr, D. J., Levy, R., Scheepers, C., & Tily, H. J. (2013). Random effects structure for confirmatory hypothesis testing: Keep it maximal. *Journal of Memory and Language*, *68*, 255–278. doi:10.1016/j.jml.2012.11.001

Bates, D., Mächler, M., Bolker, B., & Walker, S. (2015). Fitting linear mixed-effects models using lme4. *Journal of Statistical Software*, *67*(1). doi:10.18637/jss.v067.i01

Bergmann, C., Tsuji, S., Piccinini, P. E., Lewis, M. L., Braginsky, M., Frank, M. C., & Cristia, A. (2018). Promoting replicability in developmental research through meta-analyses: Insights from language acquisition research. *Child Development*, *89*, 1996–2009. doi:10.1111/cdev.13079

Button, K. S., Ioannidis, J. P. A., Mokrysz, C., Nosek, B. A., Flint, J., Robinson, E. S. J., & Munafò, M. R. (2013). Power failure: Why small sample size undermines the reliability of neuroscience. *Nature Reviews Neuroscience*, *14*, 365–376. doi:10.1038/nrn3475

Byers-Heinlein, K. (2015). Methods for studying infant bilingualism. In J. Schwieter (Ed.), *The Cambridge handbook of bilingual processing* (pp. 133–154). Cambridge, England: Cambridge University Press.

Byers-Heinlein, K., Bergmann, C., Black, A., Carbajal, J. M., Fennell, C. T., Frank, M. C., . . . Tsui, A. S. M. (2020). *A multi-lab study of bilingual infants: Exploring the preference for infant-directed speech*. Manuscript in preparation, Concordia University, Department of Psychology.

Cooper, R. P., Abraham, J., Berman, S., & Staska, M. (1997). The development of infants' preference for motherese. *Infant Behavior & Development*, *20*, 477–488. doi:10.1016/S0163-6383(97)90037-0

Cooper, R. P., & Aslin, R. N. (1990). Preference for infant-directed speech in the first month after birth. *Child Development*, *61*, 1584–1595. doi:10.1111/j.1467-8624.1990.tb02885.x

Cooper, R. P., & Aslin, R. N. (1994). Developmental differences in infant attention to the spectral properties of infant-directed speech. *Child Development*, *65*, 1663–1677. doi:10.1111/j.1467-8624.1994.tb00841.x

Csibra, G., & Gergely, G. (2009). Natural pedagogy. *Trends in Cognitive Sciences*, *13*, 148–153. doi:10.1016/j.tics.2009.01.005

Csibra, G., Hernik, M., Mascaro, O., Tatone, D., & Lengyel, M. (2016). Statistical treatment of looking-time data. *Developmental Psychology*, *52*, 521–536. doi:10.1037/dev0000083

Cusack, R., & Carlyon, R. P. (2003). Perceptual asymetries in audition. *Journal of Experimental Psychology: Human Perception and Performance*, *29*, 713–725. doi:10.1037/0096-1523.29.3.713

Dunst, C., Gorman, E., & Hamby, D. (2012). *Preference for infant-directed speech in preverbal young children*. Retrieved from Center for Early Literacy Learning website: http://www.earlyliteracylearning.org/cellreviews/cellreviews_v5_n1.pdf

Englund, K., & Behne, D. (2006). Changes in infant directed speech in the first six months. *Infant and Child Development*, *15*, 139–160. doi:10.1002/icd.445

Farran, L. K., Lee, C.-C., Yoo, H., & Oller, D. K. (2016). Cross-cultural register differences in infant-directed speech: An initial study. *PLOS ONE*, *11*(3), Article e0151518. doi:10.1371/journal.pone.0151518

Fernald, A. (1985). Four-month-old infants prefer to listen to motherese. *Infant Behavior & Development*, *8*, 181–195. doi:10.1016/S0163-6383(85)80005-9

Fernald, A. (1989). Intonation and communicative intent in mothers' speech to infants: Is the melody the message? *Child Development*, *60*, 1497–1510. doi:10.2307/1130938

Fernald, A., & McRoberts, G. W. (1996). Prosodic bootstrapping: A critical analysis of the argument and the evidence. In J. L. Morgan & K. Demuth (Eds.), *Signal to syntax: Bootstrapping from speech to grammar in early acquisition* (pp. 365–388). Mahwah, NJ: Erlbaum.

Fernald, A., Taeschner, T., Dunn, J., Papousek, M., de Boysson-Bardies, B., & Fukui, I. (1989). A cross-language study of prosodic modifications in mothers' and fathers' speech to preverbal infants. *Journal of Child Language*, *16*, 477–501.

Frank, M. C., Bergelson, E., Bergmann, C., Cristia, A., Floccia, C., Gervain, J., . . . Yurovsky, D. (2017). A collaborative approach to infant research: Promoting reproducibility, best practices, and theory-building. *Infancy*, *22*, 421–435. doi:10.1111/infa.12182

Gervain, J., & Mehler, J. (2010). Speech perception and language acquisition in the first year of life. *Annual Review of Psychology*, *61*, 191–218.

Graf Estes, K., & Hurley, K. (2013). Infant-directed prosody helps infants map sounds to meanings. *Infancy*, *18*, 797–824. doi:10.1111/infa.12006

Grieser, D. L., & Kuhl, P. K. (1988). Maternal speech to infants in a tonal language: Support for universal prosodic features in motherese. *Developmental Psychology, 24*, 14–20. doi:10.1037/0012-1649.24.1.14

Hayashi, A., Tamekawa, Y., & Kiritani, S. (2001). Developmental change in auditory preferences for speech stimuli in Japanese infants. *Journal of Speech, Language, and Hearing Research, 44*, 1189–1200. doi:10.1044/1092-4388 (2001/092)

Higgins, J. P., Thompson, S. G., Deeks, J. J., & Altman, D. G. (2003). Measuring inconsistency in meta-analyses. *British Medical Journal, 327*, 557–560.

Hirsh-Pasek, K., Nelson, D. G. K., Jusczyk, P. W., Cassidy, K. W., Druss, B., & Kennedy, L. (1987). Clauses are perceptual units for young infants. *Cognition, 26*, 269–286. doi:10.1016/S0010-0277(87)80002-1

Huedo-Medina, T., Sanchez-Meca, J., Marin-Martinez, F., & Botella, J. (2006). Assessing heterogeneity in meta-analysis: Q statistic or I^2 index? *Psychological Methods, 11*, 193–206.

Johnson, E. K. (2016). Constructing a proto-lexicon: An integrative view of infant language development. *Annual Review of Linguistics, 2*, 391–412.

Judd, C. M., Westfall, J., & Kenny, D. A. (2017). Experiments with more than one random factor: Designs, analytic models, and statistical power. *Annual Review of Psychology, 68*, 601–625. doi:10.1146/annurev-psych-122414-033702

Kaplan, P. S., Jung, P. C., Ryther, J. S., & Zarlengo-Strouse, P. (1996). Infant-directed versus adult-directed speech as signals for faces. *Developmental Psychology, 32*, 880–891. doi:10.1037/0012-1649.32.5.880

Karzon, R. G. (1985). Discrimination of polysyllabic sequences by one- to four-month-old infants. *Journal of Experimental Child Psychology, 39*, 326–342. doi:10.1016/0022-0965(85)90044-X

Katz, G. S., Cohn, J. F., & Moore, C. A. (1996). A combination of vocal f_0 dynamic and summary features discriminates between three pragmatic categories of infant-directed speech. *Child Development, 67*, 205–217. doi:10.1111/j.1467-8624.1996.tb01729.x

Kawahara, H., & Morise, M. (2011). Technical foundations of tandem-straight, a speech analysis, modification and synthesis framework. *Sadhana, 36*, 713–727. doi:10.1007/s12046-011-0043-3

Kitamura, C., & Burnham, D. (2003). Pitch and communicative intent in mother's speech: Adjustments for age and sex in the first year. *Infancy, 4*, 85–110. doi:10.1207/S15327078IN0401_5

Kitamura, C., & Lam, C. (2009). Age-specific preferences for infant-directed affective intent. *Infancy, 14*, 77–100. doi:10.1080/15250000802569777

Klein, R. A., Ratliff, K. A., Vianello, M., Adams, R. B., Jr., Bahník, Š., Bernstein, M. J., . . . Nosek, B. A. (2014). Investigating variation in replicability. *Social Psychology, 45*, 142–152. doi:10.1027/1864-9335/a000178

Kuznetsova, A., Brockhoff, P. B., & Christensen, R. H. B. (2017). lmerTest package: Tests in linear mixed effects models. *Journal of Statistical Software, 82*(13). doi:10.18637/jss.v082.i13

Ma, W., Golinkoff, R. M., Houston, D. M., & Hirsh-Pasek, K. (2011). Word learning in infant- and adult-directed speech. *Language Learning and Development, 7*, 185–201.

Makel, M. C., Plucker, J. A., & Hegarty, B. (2012). Replications in psychology research: How often do they really occur? *Perspectives on Psychological Science, 7*, 537–542. doi:10.1177/1745691612460688

Martin, A., Igarashi, Y., Jincho, N., & Mazuka, R. (2016). Utterances in infant-directed speech are shorter, not slower. *Cognition, 156*, 52–59.

Maurer, D., & Werker, J. F. (2014). Perceptual narrowing during infancy: A comparison of language and faces. *Developmental Psychobiology, 56*, 154–178. doi:10.1002/dev.21177

McRoberts, G. W., McDonough, C., & Lakusta, L. (2009). The role of verbal repetition in the development of infant speech preferences from 4 to 14 months of age. *Infancy, 14*, 162–194. doi:10.1080/15250000802707062

Mills-Smith, L., Spangler, D. P., Panneton, R., & Fritz, M. S. (2015). A missed opportunity for clarity: Problems in the reporting of effect size estimates in infant developmental science. *Infancy, 20*, 416–432. doi:10.1111/infa.12078

Nelson, D. G. K., Jusczyk, P. W., Mandel, D. R., Myers, J., Turk, A., & Gerken, L. (1995). The head-turn preference procedure for testing auditory perception. *Infant Behavior & Development, 18*, 111–116. doi:10.1016/0163-6383(95)90012-8

Newman, R. S. (2003). Prosodic differences in mothers' speech to toddlers in quiet and noisy environments. *Applied Psycholinguistics, 24*, 539–560. doi:10.1017/S0142716403000274

Newman, R. S., & Hussain, I. (2006). Changes in preference for infant-directed speech in low and moderate noise by 4.5- to 13-month-olds. *Infancy, 10*, 61–76. doi:10.1207/s15327078in1001_4

Nieuwenhuis, S., Forstmann, B. U., & Wagenmakers, E.-J. (2011). Erroneous analyses of interactions in neuroscience: A problem of significance. *Nature Neuroscience, 14*, 1105–1107. doi:10.1038/nn.2886

Oakes, L. M. (2017). Sample size, statistical power, and false conclusions in infant looking-time research. *Infancy, 22*, 436–469.

Open Science Collaboration. (2015). Estimating the reproducibility of psychological science. *Science, 349*, Article aac4716. doi:10.1126/science.aac4716

Pegg, J. E., Werker, J. F., & McLeod, P. J. (1992). Preference for infant-directed over adult-directed speech: Evidence from 7-week-old infants. *Infant Behavior & Development, 15*, 325–345. doi:10.1016/0163-6383(92)80003-D

Santesso, D. L., Schmidt, L. A., & Trainor, L. J. (2007). Frontal brain electrical activity (EEG) and heart rate in response to affective infant-directed (ID) speech in 9-month-old infants. *Brain and Cognition, 65*, 14–21. doi:10.1016/j.bandc.2007.02.008

Schachner, A., & Hannon, E. E. (2011). Infant-directed speech drives social preferences in 5-month-old infants. *Developmental Psychology, 47*, 19–25. doi:10.1037/a0020740

Segal, J., & Newman, R. S. (2015). Infant preferences for structural and prosodic properties of infant-directed speech in the second year of life. *Infancy*, *20*, 339–351.

Shute, H. B. (1987). Vocal pitch in motherese. *Educational Psychology*, *7*, 187–205. doi:10.1080/0144341870070303

Singh, L., Morgan, J. L., & Best, C. T. (2002). Infants' listening preferences: Baby talk or happy talk? *Infancy*, *3*, 365–394. doi:10.1207/S15327078IN0303_5

Snow, C. E. (1977). The development of conversation between mothers and babies. *Journal of Child Language*, *4*, 1–22. doi:10.1017/S0305000900000453

Soderstrom, M., Blossom, M., Foygel, R., & Morgan, J. L. (2008). Acoustical cues and grammatical units in speech to two preverbal infants. *Journal of Child Language*, *35*, 869–902. doi:10.1017/S0305000908008763

Soderstrom, M., Werker, J., Tsui, A., Skarabela, B., Seidl, A., Searle, A., & Anderson, L. (2020). *Testing the relationship between preferences for infant-directed speech and vocabulary development: A multi-lab study*. Manuscript in preparation, University of Manitoba, Department of Psychology.

Thiessen, E. D., Hill, E. A., & Saffran, J. R. (2005). Infant-directed speech facilitates word segmentation. *Infancy*, *7*, 53–71. doi:10.1207/s15327078in0701_5

Trainor, L. J., & Desjardins, R. N. (2002). Pitch characteristics of infant-directed speech affect infants' ability to discriminate vowels. *Psychonomic Bulletin & Review*, *9*, 335–340. doi:10.3758/BF03196290

Viechtbauer, W. (2010). Conducting meta-analyses in R with the metafor package. *Journal of Statistical Software*, *36*(3). doi:10.18637/jss.v036.i03

Weisleder, A., & Fernald, A. (2013). Talking to children matters: Early language experience strengthens processing and builds vocabulary. *Psychological Science*, *24*, 2143–2152. doi:10.1177/0956797613488145

Werker, J. F., & McLeod, P. J. (1989). Infant preference for both male and female infant-directed talk: A developmental study of attentional and affective responsiveness. *Canadian Journal of Psychology/Revue Canadienne de Psychologie*, *43*, 230–246. doi:10.1037/h0084224

Werker, J. F., Pegg, J. E., & McLeod, P. J. (1994). A cross-language investigation of infant preference for infant-directed communication. *Infant Behavior & Development*, *17*, 323–333. doi:10.1016/0163-6383(94)90012-4

Zangl, R., & Mills, D. L. (2007). Increased brain activity to infant-directed speech in 6- and 13-month-old infants. *Infancy*, *11*, 31–62. doi:10.1207/s15327078in1101_2