Limited evidence of test-retest reliability in infant-directed speech preference in a large 1 preregistered infant experiment 2 Melanie S. Schreiner^{1,2}, Martin Zettersten^{3,4}, Christina Bergmann⁵, Michael C. Frank⁶, Tom Fritzsche⁷, Nayeli Gonzalez-Gomez⁸, Kiley Hamlin⁹, Natalia Kartushina¹⁰, Danielle J. Kellier¹¹, Nivedita Mani^{1,2}, Julien Mayor¹⁰, Jenny Saffran³, Mohinish Shukla¹², Priya Silverstein^{13, 14}, Melanie Soderstrom¹⁵, & Matthias Lippold^{1,2} ¹ University of Goettingen ² Leibniz Science Campus PrimateCognition ³ University of Wisconsin-Madison ⁴ Princeton University 10 ⁵ Max Planck Insitute for Psycholinguistics 11 ⁶ Stanford University 12 ⁷ University of Potsdam 13 ⁸ Oxford Brookes University ⁹ University of British Columbia ¹⁰ University of Oslo ¹¹ University of Pennsylvania 17 ¹² Università di Padova 18 ¹³ Institute for Globally Distributed Open Research 19

¹⁴ Ashland University

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¹⁵ University of Manitoba

22 Author Note

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- Correspondence concerning this article should be addressed to Melanie S. Schreiner,
 Gosslerstr. 14, 37073 Göttingen. E-mail: melanie.schreiner@psych.uni-goettingen.de

41 Abstract

Test-retest reliability — establishing that measurements remain consistent across multiple testing sessions — is critical to measuring, understanding, and predicting individual differences in infant language development. However, previous attempts to establish 44 measurement reliability in infant speech perception tasks are limited, and reliability of frequently-used infant measures is largely unknown. The current study investigated the test-retest reliability of infants' preference for infant-directed speech (hereafter, IDS) over adult-directed speech (hereafter, ADS) in a large sample (N=158) in the context of the ManyBabies1 collaborative research project (hereafter, MB1; Frank et al., 2017; ManyBabies Consortium, 2020). Labs were asked to bring in participating infants for a second appointment retesting infants on their IDS preference. This approach allows us to estimate test-retest reliability across three different methods used to investigate preferential listening in infancy: the head-turn preference procedure, central fixation, and eye-tracking. 53 Overall, we find no consistent evidence of test-retest reliability in measures of infants' 54 speech preference (overall r = .09, 95% CI [-.06,.25]). While increasing the number of trials 55 that infants needed to contribute for inclusion in the analysis revealed a numeric growth in 56 test-retest reliability, it also considerably reduced the study's effective sample size. 57 Therefore, future research on infant development should take into account that not all experimental measures may be appropriate for assessing individual differences between 59 infants. 60

Keywords: language acquisition; speech perception; infant-directed speech; adult-directed speech; test-retest reliability

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Limited evidence of test-retest reliability in infant-directed speech preference in a large

preregistered infant experiment

Obtaining a quantitative measure of infants' cognitive abilities is an extraordinarily difficult endeavor. The most frequent way to assess what infants know or prefer is to track overt behavior. However, measuring overt behavior at early ages presents many challenges: participants' attention span is short, they do not follow instructions, their mood can change instantly, and their behavior is often inconsistent. Therefore, most measurements are noisy and the typical sample size of an infant study is small (around 20 infants per group), resulting in low power (Oakes, 2017). In addition, there is individual and environmental variation that may add even more noise to the data (e.g., Johnson & Zamuner, 2010). Despite these demanding conditions, reliable and robust methods for assessing infants' behavior are critical to understanding development.

In order to address these challenges, the ManyBabies collaborative research 76 consortium was formed to conduct large-scale, conceptual, consensus-based replications of 77 seminal findings to identify sources of variability and establish best practices for 78 experimental studies in infancy (Frank et al., 2017). The first ManyBabies collaborative research project (hereafter, MB1, ManyBabies Consortium, 2020) explored the reproducibility of the well-studied phenomenon that infants prefer infant-directed speech 81 (hereafter, IDS) over adult-directed speech (hereafter, ADS, Cooper & Aslin, 1990). Across 82 many different cultures, infants are commonly addressed in IDS, which typically is characterized by higher pitch, greater pitch range, and shorter utterances, compared to the language used between interacting adults (Fernald et al., 1989). A large body of behavioral studies finds that infants show increased looking times when hearing IDS compared to ADS stimuli across ages and methods (Cooper & Aslin, 1990; see Dunst, Gorman, & Hamby, 2012 for a meta-analysis). This attentional enhancement is also documented in neurophysiological studies showing increased neural activation during IDS compared to

ADS exposure (Naoi et al., 2012; Zangl & Mills, 2007). IDS has also been identified as facilitating early word learning. In particular, infants' word segmentation abilities (Floccia et al., 2016; Schreiner & Mani, 2017; Singh, Nestor, Parikh, & Yull, 2009; Thiessen, Hill, & Saffran, 2005) and their learning of word-object associations (Graf Estes & Hurley, 2013; Ma, Golinkoff, Houston, & Hirsh-Pasek, 2011) are enhanced in the context of IDS. In sum, several lines of evidence suggest that IDS is beneficial for early language development.

Within MB1, 67 labs contributed data from 2,329 infants showing that babies 96 generally prefer to listen to IDS over ADS. Nevertheless, the overall effect size of d=0.3597 was smaller than a previously reported meta-analytic effect size of d = 0.67 (Dunst et al., 98 2012). The results revealed several additional factors that influenced the effect size. First, 99 older infants showed a larger preference of IDS over ADS. Second, the stimulus language 100 was linked to IDS preference, with North American English learning infants showing a 101 larger IDS preference than infants learning other languages. Third, comparing the different 102 methods employed, the head-turn preference procedure yielded the highest effect size, while 103 the central fixation paradigm and eye-tracking methods revealed smaller effects. Finally, 104 exploratory analyses assessed the effect of different inclusion criteria. Across methods, 105 using stricter inclusion criteria led to an increase in effect sizes despite the larger proportion of excluded participants (see also Byers-Heinlein, Bergmann, & Savalei, 2021). 107

However, there is a difference between a result being reliable in a large sample of 108 infants and the measurement of an individual infant being reliable. In studies tracking 109 individual differences, the measured behavior during an experimental setting is often used 110 to predict a cognitive function or specific skill later in life. Individual differences research of this kind often has substantial implications for theoretical and applied work. For example, 112 research showing that infants' behavior in speech perception tasks can be linked to later 113 language development (see Cristia, Seidl, Junge, Soderstrom, & Hagoort, 2014 for a 114 meta-analysis) has the potential to identify infants at risk for later language delays or 115 disorders. However, a necessary precondition for this link to be observable is that 116

individual differences between infants can be measured with high reliability at these earlier stages, in order to ensure that measured inter-individual variation mainly reflects differences in children's abilities rather than measurement error. How reliable are the measures used in infancy research?

Previous attempts to address the reliability of measurements have typically been 121 limited to adult populations (Hedge, Powell, & Sumner, 2018; Oliveira, Hayiou-Thomas, & 122 Henderson, 2023), or have been conducted with small sample sizes (e.g., Colombo, 123 Mitchell, & Horowitz, 1988; Houston, Horn, Qi, Ting, & Gao, 2007). For example, Houston 124 et al. (2007) tested 10 9-month-old infants' speech discrimination in a visual habituation 125 procedure in two test sessions 1-3 days apart and found a large correlation (r = .7). These 126 data were subsequently included in a much larger systematic investigation of test-retest 127 reliability in infant speech perception (Cristia, Seidl, Singh, & Houston, 2016). Cristia et 128 al. (2016) analyzed 13 different experiments assessing test-retest reliability in infant speech 129 perception tasks, with the retest session occurring 0-18 days after the first session. The 130 experiments were conducted at three different labs with different implementations of the individual studies. Hence, it was only after completed data collection that the data was 132 pooled together by the different labs revealing potential confounds. Nevertheless, the 133 results showed that reliability was extremely variable across the different experiments and labs and low overall (meta-analytic r = .07). 135

Against this background, the current study investigates test-retest reliability of infants' performance in a speech preference task. Within MB1, a multi-lab collaboration, we examine whether infants' preferential listening behavior to IDS and ADS is reliable across two different test sessions. In addition, we investigate the influence of various moderators on the reliability of IDS preference.

First, the current study also explores whether there are any differences in test-retest reliability between three widely used methods: central fixation (CF), eye-tracking (ET),

and the head-turn preference procedure (HPP). Exploring differences in CF, ET, and HPP, 143 Junge et al. (2020) provide experimental and meta-analytic evidence in favor of using the 144 HPP in speech segmentation tasks. Similarly, the MB1 project reported an increase in the 145 effect size for HPP compared to CF and ET (ManyBabies Consortium, 2020). HPP 146 requires gross motor movements relative to other methods, such as CF and ET paradigms, 147 for which subtle eye movements towards a monitor located in front of the child are 148 sufficient. One possible explanation for the stronger effects with HPP may be a higher 149 sensitivity to the contingency of the presentation of auditory stimuli and infants' head 150 turns away from the typical forward-facing position. While these findings suggest that HPP 151 may be a more sensitive index of infant preference, they do not necessarily imply higher 152 reliability for individual infants' performance using HPP. For example, Marimon and Höhle 153 (2022) found no evidence for test-retest reliability when testing infants' prosodic preferences using the HPP method across three testing sessions, each 7-8 days apart on average. 155

Second, we aim to address whether there are age-related differences in test-retest reliability. Older participants of MB1 yielded larger effect sizes. Given that older infants have had more linguistic experience, we predict that their preference may be more stable than that of younger infants.

Third, we will explore if test-retest reliability of the infants' preference measure is
moderated by their linguistic background. All infants are tested using a North American
English (hereafter, NAE) stimulus set, which is either their native or non-native language.
As NAE infants demonstrated larger effects within the original MB1 study, we assume that
infants for whom NAE is not the native language may behave more variable and
demonstrate less reliable results than English-learning infants.

Last, we will assess whether time between test and retest influences the reliability of
the preference measure. Time between test and retest session may vary across participants.
We predict that longer times between test and retest may yield less reliable results.

In sum, it remains an open question whether the same measures that produce larger effect sizes at the group-level also have higher test-retest reliability for individual infants (Byers-Heinlein, Bergmann, et al., 2021). Therefore, assessing the test-retest reliability of the different preference measures and the influence of various moderators is crucial, so that researchers can make informed decisions about the appropriate methods for their particular research question. Critically, only measures with high test-retest reliability should be used for studies of individual differences.

176 Method

77 Preregistration

We preregistered the current study on the Open Science Framework

(https://osf.io/v5f8t). Section S1 in the Supplementary Materials contains additional notes

on the preregistration decisions and any deviations from the preregistered analytic plan.

181 Data Collection

A call was issued to all labs participating in the original MB1 study on January 24th,
2018 (ManyBabies Consortium, 2020). The collection of retest session data was initially set
to end on May 31st, 2018, one month after the end date of the original MB1 project. Due
to the fact that the original MB1 project extended the time frame for data collection and
the late start of data collection for the MB1 test-retest study, we also allowed participating
labs to continue data collection past the scheduled end date.

188 Participants

Contributing labs were asked to recruit monolingual participants between the ages of 6 to 12 months. If participating labs could not commit to test either of these age groups, they were also allowed to recruit participants from the youngest age group of 3- to

6-month-olds and/or the oldest age group of 12- to 15-month-olds. Labs were asked to contribute half (n=16) or full samples (n=32); however, a lab's data was included in the study regardless of the number of included infants. The study was approved by each lab's respective ethics committee and parental consent was obtained for each infant prior to participation in the study.

Our final sample consisted of 158 monolingual infants from 7 different labs (Table 1).

In order to be included in the study, infants needed a minimum of 90% first language

exposure, to be born full term with no known developmental disorders, and normal hearing

and vision. We excluded 18 additional participants (see Data Exclusion section for details).

The mean age of infants included in the study was 245 days (range: 108 – 373 days;

approximately 8.06 months).

Materials

Visual stimuli. The visual stimuli and instructions were identical to MB1. For the

CF paradigm and ET, labs used a multicolored static checkerboard as the fixation stimulus

as well as a multicolored moving circle with a ringing sound as an attention-getter between

trials. For the HPP method, labs used their standard procedure, as in MB1.

Our study was faced with a critical design choice: what stimuli Auditory stimuli. 208 to use to assess test-retest reliability. One constraint on our study was that, since it was a 209 follow-on to MB1, any stimulus we used would always be presented after the MB1 stimuli. 210 One option would be simply to bring back infants and have them hear exactly the same 211 stimulus materials. A weakness of this design would be the potential for stimulus familiarity effects, however, since infants would have heard the materials before. Further 213 complicating matters, infants might show a preference for or against a familiar stimulus depending on their age (Hunter & Ames, 1988). The ideal solution then would be to create 215 a brand new stimulus set with the same characteristics. Unfortunately, because of the 216 process of how MB1 stimuli were created, we did not have enough normed raw recordings 217

available to make brand new stimulus items that conformed to the same standards as the 218 MB1 stimuli. We therefore chose an intermediate path: we reversed the ordering of MB1 219 stimuli. A second set of naturalistic IDS and ADS recordings of mothers either talking to 220 their infant or to an experimenter was created for the retest session by reversing the order 221 of clips within each sequence of the original study. This resulted in eight reordered 222 sequences of natural IDS and eight reordered sequences of natural ADS with a length of 18 223 seconds each. Average looking times in MB1 were always lower than 9s per trial, even for 224 the youngest children on the earliest trials (the group who looked the longest on average), 225 so most children in MB1 did not hear the second half of most trials. Thus, by reversing the 226 order, we had a perfectly matched stimulus set that was relatively unfamiliar to most 227 infants. The disadvantage of this design was that infants who looked longer might be more 228 likely to hear a familiar clip heard in the previous study. If infants then showed a familiarity preference — an assumption which might not be true — the end result could be to inflate 230 our estimates of test-retest reliability slightly, since longer lookers would on average look longer at retest due to their familiarity preference. We view this risk as relatively low, but 232 do note that it is a limitation of our design. In addition to the 16 reversed-order IDS and 233 ADS speech stimuli, we used the identical training stimuli of piano music from MB1.

Procedure. Infants were retested using the identical procedure as during the first testing day: CF, HPP, or ET. Participating labs were asked to schedule test and retest sessions 7 days apart with a minimum number of 1 day and a maximum number of 31 days. However, infants whose time between test and retest exceeded 31 days were still included in the analyses (n = 3). The mean number of days between test and retest was 10 (range: 1 - 49).

A total of 18 trials, including two training, eight IDS, and eight ADS trials, were presented in one of four pseudo-randomized orders. Trial length was either infant-controlled

or fixed depending on the lab's standard procedure¹: a trial stopped either if the infant looked away for 2 seconds or after the total trial duration of 18 seconds. The online coding experimenter and the parent listened to music masked with the stimuli of the study via noise-cancelling headphones. If the experimenter was in an adjacent room separate from the testing location, listening to masking music was optional for the experimenter.

Data exclusion. In total, 18 participants were excluded from the analysis. 4 248 participants were excluded for being preterm (defined as a gestation time of less than 37 249 weeks). 6 participants were excluded due to session errors involving an experimenter error 250 (e.g., inaccurate coding or presentation of retest stimuli on the first test session). 251 Individual trials were excluded if they were marked as trial errors (5.45\% of remaining 252 trials), i.e., if the infant was reported as fussy, an experimental or equipment error 253 occurred, or there was parental interference during the task (e.g., if the parent spoke with 254 the infant during the trial). Trials were also excluded if the minimum looking time of 2 s 255 was not met (12.60% of the remaining trials). If a participant was unable to contribute at 256 least one IDS and one ADS trial for either test or retest after trial-level exclusions, all data 257 of that participant was excluded from the test-retest analyses (12 additional participants). 258

Results

50 IDS preference

First, we conducted confirmatory analyses examining infants' preference for IDS in both sessions. Two-samples t-tests comparing the difference in average looking time between IDS and ADS to zero revealed that infants showed a preference of IDS over ADS in Session 1, t(157) = 6.47, p < .001, and Session 2, t(157) = 4.19, p < .001, replicating the main finding from MB1 (Table 2). 68.35% of infants in Session 1 and 63.29% of infants in Session 2 showed a preference for IDS. In order to test whether there was a difference in

¹ Labs using CF or HPP had infant-controlled trial length whereas labs using ET had fixed trial length.

Table 1
Statistics of the included labs. n refers to the number of infants included in the final analysis.

| Lab | Method | Language | Mean age (days) | N |
|------------------|------------------|-----------|-----------------|----|
| InfantCog-UBC | central fixation | English | 147 | 7 |
| babylab-potsdam | HPP | German | 227 | 22 |
| babyling-oslo | eye-tracking | Norwegian | 249 | 10 |
| brookes-babylab | central fixation | English | 267 | 18 |
| infantll-madison | HPP | English | 230 | 30 |
| lancslab | eye-tracking | English | 236 | 16 |
| wsi-goettingen | HPP | German | 242 | 16 |
| wsi-goettingen | central fixation | German | 280 | 39 |

the strength of the preference effect across sessions, we fit a linear mixed-effects model predicting infants' average difference in looking time between IDS and ADS from test session (1 vs. 2), including by-lab and by-participant random intercepts. There was no significant difference in the magnitude of infants' preference between the two sessions, β =-0.30, SE=0.24, p=.208.

272 Reliability

We assessed test-retest reliability in two planned, confirmatory analyses. First, we fit
a linear mixed-effects model predicting IDS preference in Session 2 from IDS preference in
Session 1, including a by-lab random intercept. The results revealed no significant
relationship between IDS preference in Session 1 and 2 (Table 3). Second, we calculated
the Pearson correlation coefficient. While a simple correlation coefficient might
overestimate the test-retest reliability in our sample because it does not control for the

| Trial type | Session 1 Mean | Session 1 SD | Session 2 Mean | Session 2 SD |
|------------|----------------|--------------|----------------|--------------|
| ADS | 7.71 | 2.77 | 6.96 | 2.92 |
| IDS | 8.76 | 2.84 | 7.75 | 2.75 |

Table 2

Average looking times (in seconds) for each session and condition

differences between different labs and methods (HPP, CF, and ET), we felt it was important to also conduct a Pearson correlation as it is commonly used to assess reliability. The size of the correlation coefficient was not statistically different from zero and the estimate was small, r = .09, 95% CI [-.06, .25], t(156) = 1.19, p = .237. Moreover, no significant correlations emerged in each sample considered separately (Figure 1; see Supplementary Materials S3 for a meta-analytic approach). 41.77% of the infants reversed their direction of preference for IDS versus ADS from the test to the retest session.

To investigate the test-retest reliability of each specific method, we computed Pearson 286 correlation coefficients and the same mixed-effects model described above for HPP, CF, 287 and ET separately (Table 4) in additional exploratory analyses. None of the three methods 288 showed evidence of test-retest reliability. Neither the Pearson correlation coefficients nor 289 the coefficients of the multilevel analysis were significant, all p-values > 0.12. In planned 290 secondary analyses, we found that time between test sessions, participant age, method, and 291 language background did not moderate the relationship between IDS preference in session 292 1 and session 2 (see Supplementary Materials S2). Taken together, we find no significant 293 evidence of test-retest reliability across our preregistered analyses.

Exploratory analyses with different inclusion criteria

To this point, all analyses were performed using the inclusion criteria from MB1, which required only that infants contribute at least one trial per condition for inclusion

Table 3

Coefficient estimates from a linear mixed effects model predicting IDS preference in Session 2.

| | Estimate | SE | t | р |
|--------------------------|----------|------|------|------|
| Intercept | 0.87 | 0.46 | 1.92 | 0.10 |
| IDS Preference Session 1 | 0.04 | 0.09 | 0.41 | 0.68 |

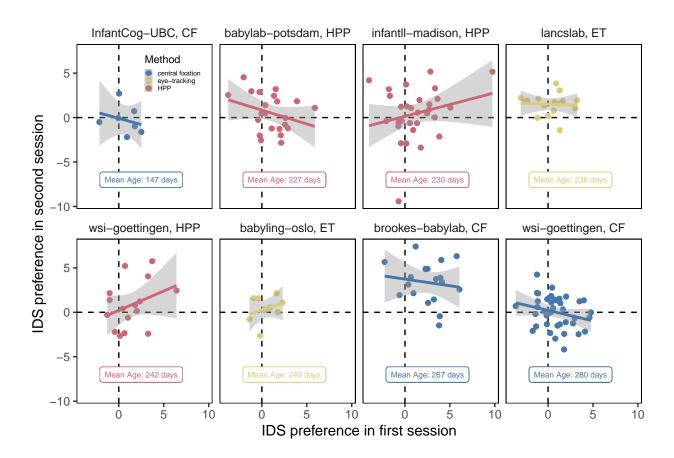


Figure 1. Correlation between IDS Preference in Session 1 and Session 2 in each lab and method. Dots indicate individual participants. Error bands represent 95 percent confidence intervals. The dashed line indicates no preference (i.e., a value of zero) for the first and second session, respectively.

Table 4

Coefficient estimates from a linear mixed effects

model predicting IDS preference in Session 2 and

Pearson correlation coefficient for each method

separately.

| Method | beta | SE | p | Pearson r |
|------------------|-------|------|------|-----------|
| central fixation | -0.20 | 0.12 | 0.12 | 0.08 |
| HPP | 0.15 | 0.14 | 0.28 | 0.13 |
| eye-tracking | 0.03 | 0.16 | 0.84 | 0.02 |

(i.e., one IDS and one ADS trial). However, more stringent inclusion criteria yielded larger 298 effect sizes in MB1. We therefore conducted exploratory analyses assessing test-retest 290 reliability after applying progressively stricter inclusion criteria, requiring two, four, six, 300 and eight valid trials per condition. Applying stricter criteria — and thereby increasing the 301 number of test trials — increased reliability numerically from r = 0.07 to r = 0.34 (Figure 302 2). In part due to a decrease in sample size, only one of these correlations was statistically 303 significant (when requiring six trial pairs): two valid trial pairs, t(152) = 0.90, p = .367; 304 four valid trial pairs, t(143) = 1.03, p = .306; six valid trial pairs, t(98) = 2.23, p = .028; eight valid trial pairs — all trials in both sessions — t(22) = 1.68, p = .108. The analyses provide tentative evidence that stricter inclusion criteria may lead to higher test-retest 307 reliability, but at the cost of substantial decreases in sample size (see Supplementary 308 Materials S4 for additional analyses, including moderator analyses using a more restricted 309 sample). 310



Figure 2. IDS preferences of both sessions plotted against each other for each inclusion criterion. n indicates the number of included infants, r is the Pearson correlation coefficient as the indicator for reliability.

Correlations between sessions for number of trials contributed and overall looking time

In exploratory analyses, we also investigated whether there were stable individual
differences in (a) the number of trials an infant contributed across the two test sessions and
(b) infants' overall looking times.

Number of trials contributed. We found a strong positive correlation between number of trials contributed during the first and the second session r = .58, 95% CI [.47, .67], t(160) = 9.00, p < .001 (Figure 3A). In other words, if infants contributed a higher number of trials in one session, compared to other infants, they were likely to contribute a higher number of trials in their next session. This finding is consistent with the hypothesis that how attentive infants are throughout an experiment (and hence how many trials they contribute) is a stable individual difference, at least for some infant looking time tasks.

Overall looking times. To what extent are participants looking times between the 323 two sessions related? To test this question, we investigated whether participants' overall 324 looking times — irrespective of condition — were correlated between the first and second 325 session. There was a robust correlation between average looking time in Session 1 and 326 Session 2: infants with longer looking times during their first session also tended to look 327 longer during their second session, r = .45, 95% CI [.31, .57], t(156) = 6.28, p < .001328 (Figure 3B). This relationship held even after controlling for number of trials (b = 0.42, 329 95% CI [0.27, 0.58], t(154) = 5.52, p < .001) and participants' average age (b = 0.44, 95% 330 CI [0.30, 0.59], t(155) = 6.16, p < .001) across the two test sessions in linear regression models. Finally, we found similar correlations in average looking time to IDS stimuli in 332 Session 1 and 2, r = .38, 95% CI [.24, .51], t(156) = 5.19, p < .001, and ADS stimuli in 333 Session 1 and 2, r = .40, 95% CI [.26, .53], t(156) = 5.49, p < .001 (Figure 3C; see 334 Supplementary Materials S9 and S10 for further details, including an investigation of 335 item-level correlations). 336

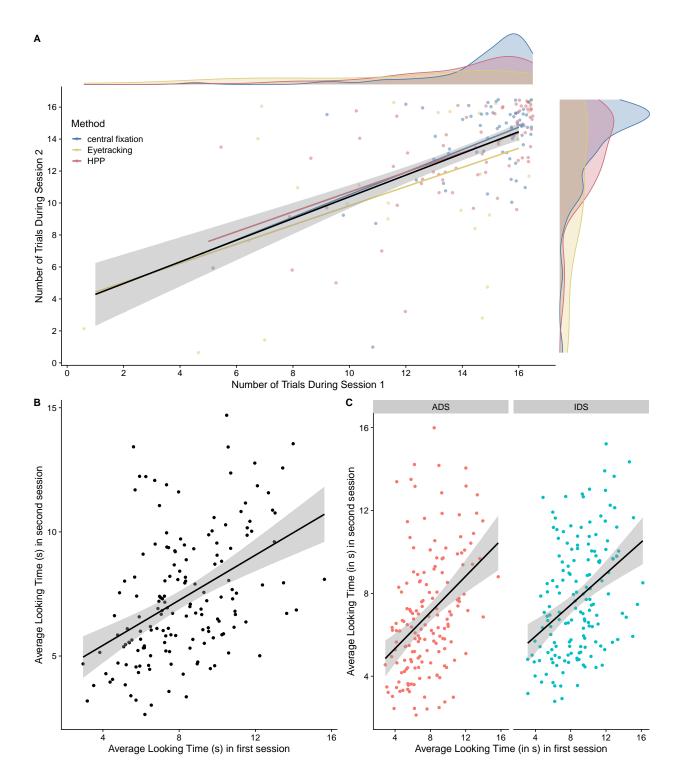


Figure 3. (A) Correlation between the number of trials contributed in Session 1 and Session 2. Each data point represents one infant. Colored lines represent linear fits for each method. (B) Overall correlations in average looking time (in s) between Session 1 and 2. (C) Correlations in average looking time (in s) between sessions, split by IDS/ADS condition.

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General Discussion

The current study investigated the test-retest reliability of infants' preference for IDS 338 over ADS. As part of the original MB1 project, we tested the IDS preference of infants in 339 two separate test sessions to assess the extent to which their pattern of preference would 340 remain consistent. While we replicated the original effect of infants' speech preference for 341 IDS over ADS for both the test and retest session on the group-level, we found that infants' speech preference measures showed no evidence of test-retest reliability. In other words, we were unable to detect stable individual differences in infants' preference for IDS. This finding is consistent with past research suggesting low test-retest reliability in other infant paradigms (Cristia et al., 2016). Given that most experimental procedures conducted in infant research are interested in the comparison of groups, individual differences between participants within a specific condition are usually minimized by the 348 experimental procedure while differences between conditions are maximized. Therefore, 349 infant preference measures may be a good approach for capturing group-level phenomena, 350 but may be less appropriate for examining individual differences in development. 351

Assessing the influence of different moderators on test-retest reliability yielded similar 352 deceiving results. While previous research suggests that HPP may be a more sensitive index 353 of infant preference and speech segmentation abilities (Junge et al., 2020; ManyBabies 354 Consortium, 2020), we find no evidence for higher test-retest reliability for HPP relative to 355 ET and CF. Similarly, our planned analyses with various other moderators such as time 356 between test sessions, infants' language background, and infants' age did not appear to exert a discernible impact on the test-retest reliability of infants' speech preference. One possible explanation may be that these analyses are largely underpowered as by far fewer 359 labs contributed to the study than we had anticipated by the time of the preregistration. 360 Alternatively, method, time between test session, infants' language background, and age 361 may simply not moderate the relationship of infant preference in the two test sessions. 362

Consistent with general psychometric theory (e.g., DeBolt, Rhemtulla, & Oakes, 363 2020), stricter inclusion criteria — and consequently a larger number of included test trials 364 per participant — tended to increase the magnitude of the correlation between test 365 sessions. However, this association was based on exploratory analyses and was in part only 366 observed descriptively, and hence should be interpreted with caution. A similar effect on 367 the group-level was found in the MB1 project, where a stricter inclusion criterion led to 368 bigger effect sizes (ManyBabies Consortium, 2020). As in MB1, higher reliability through 369 strict exclusions came at a high cost. In particular, with the strictest criterion, only a small 370 portion of the original sample size (24 out of 158 infants) could be included in the final 371 sample. In other words, applying stricter criteria leads to a higher drop-out rate and can 372 dramatically reduce the sample size. In the case of studies in the field of developmental 373 science, where there are many practical restrictions in collecting large samples of infants (e.g., birth rate in the area, restricted lab capacities, budget restrictions), a strict drop-out 375 criterion may often be difficult to implement. Note that studies in developmental science 376 already have above-average drop-out rates (Miller, 2017). In addition, drop out may not be 377 random, and so having high drop-out rates can further limit the generalizability of a study. 378 In fact, the number of trials individual infants contributed was highly correlated between 379 test sessions in the current study (see Supplementary Materials S6). Particularly in the 380 context of turning individual differences measures into diagnostic tools, high drop-out rates 381 have an additional limitation of not being broadly usable. 382

Even under best-case scenarios, reliability remained quite low. For example, when restricting the sample to infants contributing at least 6 trials in each condition in both sessions, we obtained a correlation of r = 0.22 and an intra-class correlation coefficient of α = 0.36. As Byers-Heinlein, Bergmann, et al. (2021) outline, low measurement reliability severely restricts power for detecting relationships between measures. Using the same approach as Byers-Heinlein, Bergmann, et al. (2021), we estimate that over 682 infants would be needed to have at least 80% power to observe a true correlation of r = .3 between

two measurements, assuming an intra-class correlation coefficient as large as that observed in our restricted sample ($\alpha=0.36$). Even a very large true correlation of r=.7 would require a sample size of over 120 infants. In other words, even under optimistic estimates of reliability based on strict inclusion criteria, the low reliability of IDS preference measures would severely limit the feasibility of individual difference and longitudinal research using current methods.

An alternative approach to increasing the number of valid trials is to increase the 396 number of experimental trials. This approach seeks to increase the likelihood that 397 participants will contribute sufficient trials (after trial-level exclusions) to allow for precise 398 individual-level estimates (DeBolt et al., 2020; see also Silverstein, Feng, Westermann, 390 Parise, & Twomey, 2021). While this approach is promising, it may not always be feasible, 400 because the attention span of a typical infant participant is limited. Therefore, prolonging 401 the experimental procedure to maximize the absolute number of trials is often challenging 402 in practice. Other avenues for obtaining higher numbers of valid trials may include changes 403 in the procedure (e.g., Egger, Rowland, & Bergmann, 2020) or implementing multi-day test 404 sessions (Fernald & Marchman, 2012). 405

As our results are only based on the phenomenon of IDS preference (albeit, with 406 three widely used methods: HPP, CF, ET) it is essential to further assess the underlying 407 reliability of preferential looking measures within other areas of speech perception 408 (Marimon & Höhle, 2022). While most infants prefer IDS over ADS (Dunst et al., 2012), 400 patterns of preferential looking in other tasks (e.g., speech segmentation) are often 410 inconsistent and difficult to predict (Bergmann & Cristia, 2016). These inconsistencies in looking behavior are especially important to consider in the context of relating a direction 412 of preference to later language development, and can sometimes lead to seemingly 413 contradictory findings. That is, both familiarity and novelty responses have been suggested 414 to be predictive of infants' later linguistic abilities (DePaolis, Vihman, & Keren-Portnoy, 415 2014; Newman, Ratner, Jusczyk, Jusczyk, & Dow, 2006; Newman, Rowe, & Ratner, 2016). 416

In light of our findings, researchers conducting longitudinal studies with experimental data from young infants predicting future outcomes should be cautious, as there may be large intra-individual variability affecting preference measurement.

While we observed limited evidence for test-retest reliability using preference 420 measures, we observed robust correlations for average looking times between session 1 and 421 2, both overall and for IDS and ADS stimuli considered separately (see also Supplementary Materials S9 for an investigation of item-level correlations). This finding is consistent with 423 past results in infant looking time studies finding robust correlations in average looking times across multiple sessions (Marimon & Höhle, 2022). It also raises an apparent puzzle: why are overall looking times for ADS and IDS stimuli correlated, while difference scores are not? One explanation is that infants have stable individual differences in how long they look to stimuli, but little or no stable individual differences in their preference for one 428 stimulus type over another. This only partially explains the current pattern of results, 429 however, because IDS looking time in session 1 predicted IDS looking time in session 2 430 even when controlling for ADS looking time, and vice versa (see S9). In other words, the 431 condition-specific looking time correlations are not fully explained by overall looking 432 behavior. Another long-established explanation is that difference scores tend to have poor 433 measurement reliability, because difference scores combine error from individual 434 measurements into a composite score and increasing the ratio of error relative to the 435 variance between participants (Hedge et al., 2018; Lord, 1956). Given the limitations of 436 difference scores (and composite scores in general), one goal for future research will be to 437 assess the use of trial-by-trial model-based approaches for estimating reliability (Haines et 438 al., 2020; Rouder & Haaf, 2019).

$_{\scriptscriptstyle 10}$ Limitations

While we had an above-average sample size for a study in infant research, we were unable to approach the number of participants collected within the original MB1 study. In

addition to a delayed call, the extra effort of having to schedule a second lab visit for each
participant and the fact that there were already other collaborative studies taking place
simultaneously (MB1B, Byers-Heinlein, Tsui, Bergmann, et al., 2021; MB1G,
Byers-Heinlein, Tsui, Van Renswoude, et al., 2021), might have contributed to a low
participation rate. A higher sample size and a larger number of participating labs from
different countries would have enabled us to conduct a more highly-powered test of
differences in test-retest reliability across different methods, language backgrounds, and
participant age.

A further limitation concerns the stimuli. While the order of the audio recording clips 451 presented to infants within a given trial differed between the first and second session, the 452 exact same stimulus material as in MB1 was used in both sessions. In particular, all 453 children heard the exact same voices in Session 1 and in Session 2. From a practical point 454 of view, this was the most straightforward solution for coordinating the experiment within 455 the larger MB1 project. However, familiarity effects might have influenced infants' looking 456 behavior. Infants with longer looking times in their first session might have had more 457 opportunity to recognize familiar audio clips in their second session. For infants with short 458 looking times, familiar audio clips would only occur towards the end of second-session 459 trials, thus offering infants less opportunity to recognize voices from their first session. Therefore, inconsistent familiarity with the stimulus material in the second session across infants might have artificially lowered test-retest reliability. On the other hand, one factor that mitigates this concern is that infants' looking times generally declined in session 2 compared to session 1 (consistent with past work, e.g. Marimon & Höhle, 2022; Santolin, Garcia-Castro, Zettersten, Sebastian-Galles, & Saffran, 2021), limiting opportunities for 465 infants to encounter previously experienced stimulus material.

467 Conclusion

Following the MB1 protocol, the current study could not detect test-retest reliability
in measures of infants' preference for IDS over ADS. Subsequent analyses provided
tentative evidence that stricter criteria for the inclusion of participants may enhance
test-retest reliability at the cost of high drop-out rates. Developmental studies relying on
stable individual differences between their participants need to consider the underlying
reliability of their measures, and we recommend a broader assessment of test-retest
reliability in infant research.

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