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

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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 allowed 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 found 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

Word count: 5229

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, 2022). 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. A necessary precondition for this link to be observable is that individual 116

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 131 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). In sum, there remains limited evidence and 135 considerable uncertainty about the test-retest reliability of infant looking time measures. 136

Against this background, the current study investigated test-retest reliability of infants' performance in a speech preference task. Within MB1, a multi-lab collaboration, we examined whether infants' preferential listening behavior to IDS and ADS is reliable across two different test sessions. By collecting data from multiple labs, we were able to conduct a preregistered, large-scale analysis of test-retest reliability within a standardized looking-time task that yields reliable condition effects in infants. In addition to assessing overall test-retest reliability, we also planned to investigate the influence of several

potential moderators on the reliability of IDS preference: the experimental method, infants' age and linguistic background, and the time between test sessions.

One main moderator analysis of interest was whether there were any differences in 146 test-retest reliability between three widely used methods: central fixation (CF), 147 eye-tracking (ET), and the head-turn preference procedure (HPP). Exploring differences in 148 CF, ET, and HPP, Junge et al. (2020) provided experimental and meta-analytic evidence 149 in favor of using the HPP in speech segmentation tasks. Similarly, the MB1 project 150 reported an increase in the effect size for HPP compared to CF and ET (ManyBabies 151 Consortium, 2020). HPP requires gross motor movements relative to other methods, such 152 as CF and ET paradigms, for which subtle eye movements towards a monitor located in 153 front of the child are sufficient. One possible explanation for the stronger effects with HPP 154 may be a higher sensitivity to the contingency of the presentation of auditory stimuli and 155 infants' head turns away from the typical forward-facing position. While these findings 156 suggest that HPP may be a more sensitive index of infant preference, they do not 157 necessarily imply higher reliability for individual infants' performance using HPP. For 158 example, Marimon and Höhle (2022) found no evidence for test-retest reliability when 159 testing infants' prosodic preferences using the HPP method across three testing sessions, each 7-8 days apart on average. 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 162 individual infants (Byers-Heinlein et al., 2022). In the current study, we therefore assessed 163 whether HPP yields higher test-retest reliability compared to CF and ET in looking-time 164 measures of IDS preference. 165

In our second set of moderator analyses, we aimed to address whether characteristics
of the infant, specifically their age and linguistic background, were associated with
differences in test-retest reliability. In MB1, older infants yielded larger effect sizes. Given
that older infants have had more linguistic experience, we predicted that their preference
may be more stable than that of younger infants. Infants also varied with respect to their

linguistic background. All infants were tested using a North American English (hereafter,
NAE) stimulus set, which was either their native or non-native language. We predicted
that infants for whom NAE was not their native language — and who therefore had little
or no experience with NAE — would demonstrate more variable and less reliable looking
behavior than English-learning infants. Finally, we assessed whether time between test and
retest influenced the reliability of the preference measure. Specifically, we investigated
whether test-retest reliability decreased for participants with longer durations between
their first and second test session.

179 Method

180 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.

184 Data Collection

A call was issued to all labs participating in the original MB1 study on January 29th,
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.

91 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 (approximately 8.06 months;

range: 108 – 373 days).

206 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.

Auditory stimuli. Our study was faced with a critical design choice: what stimuli 211 to use to assess test-retest reliability. One constraint on our study was that, since it was a 212 follow-on to MB1, any stimulus we used would always be presented after the MB1 stimuli. 213 One option would be simply to bring back infants and have them hear exactly the same stimulus materials. A weakness of this design would be the potential for stimulus 215 familiarity effects, however, since infants would have heard the materials before. Further complicating matters, infants might show a preference for or against a familiar stimulus 217 depending on their age (Hunter & Ames, 1988). The ideal solution then would be to create 218 a brand new stimulus set with the same characteristics. Unfortunately, because of the 219

process of how MB1 stimuli were created, we did not have enough normed raw recordings 220 available to make brand new stimulus items that conformed to the same standards as the 221 MB1 stimuli. We therefore chose an intermediate path: we reversed the ordering of MB1 222 stimuli. A second set of naturalistic IDS and ADS recordings of mothers either talking to 223 their infant or to an experimenter was created for the retest session by reversing the order 224 of clips within each sequence of the original study. This resulted in eight reordered 225 sequences of natural IDS and eight reordered sequences of natural ADS with a length of 18 226 seconds each. Average looking times in MB1 were always lower than 9s per trial, even for 227 the youngest children on the earliest trials (the group who looked the longest on average), 228 so most children in MB1 did not hear the second half of most trials. Thus, by reversing the 220 order, we had a perfectly matched stimulus set that was relatively unfamiliar to most 230 infants. The disadvantage of this design was that infants who looked longer might be more likely to hear a familiar clip heard in the previous session. If infants then showed a 232 familiarity preference — an assumption which might not be true — the end result could be to inflate 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 235 relatively low, but do note that it is a limitation of our design. In addition to the 16 236 reversed-order IDS and ADS speech stimuli, we used the identical training stimuli of piano 237 music from MB1. 238

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. 4 252 participants were excluded for being preterm (defined as a gestation time of less than 37 253 weeks). 6 participants were excluded due to session errors involving an experimenter error 254 (e.g., inaccurate coding or presentation of retest stimuli on the first test session). 255 Individual trials were excluded if they were marked as trial errors (5.45% of remaining 256 trials), i.e., if the infant was reported as fussy, an experimental or equipment error 257 occurred, or there was parental interference during the task (e.g., if the parent spoke with 258 the infant during the trial). Trials were also excluded if the minimum looking time of 2 s 259 was not met (12.60% of the remaining trials). If a participant was unable to contribute at 260 least one IDS and one ADS trial for either test or retest after trial-level exclusions, all data 261 of that participant was excluded from the test-retest analyses (12 additional participants). 262

263 Results

264 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; see Supplementary Materials S5 for robustness analyses using alternative dependent measures). 68.35% of infants in Session 1 and 63.29% of

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

infants in Session 2 showed a preference for IDS. In order to test whether there was a difference in 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.

277 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

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

the test-retest reliability in our sample because it does not control for the differences 283 between different labs and methods (HPP, CF, and ET), we felt it was important to also 284 conduct a Pearson correlation as it is commonly used to assess reliability. The size of the 285 correlation coefficient was not statistically different from zero and the estimate was small. 286 r = .09, 95% CI [-.06, .25], t(156) = 1.19, p = .237. Moreover, no significant correlations 287 emerged in each sample considered separately (Figure 1; see Supplementary Materials S3 288 for a meta-analytic approach). 41.77% of the infants reversed their direction of preference 280 for IDS versus ADS from the test to the retest session (see Supplementary Materials S7 for 290 additional analyses of infants' patterns of preferential looking across sessions). 291

To investigate the test-retest reliability of each specific method, we computed Pearson 292 correlation coefficients and the same mixed-effects model described above for HPP, CF, 293 and ET separately (Table 4) in additional exploratory analyses. None of the three methods 294 showed evidence of test-retest reliability. Neither the Pearson correlation coefficients nor the coefficients of the multilevel analysis were significant, all p-values > 0.12. In planned secondary analyses, we found that time between test sessions, participant age, method, and 297 language background did not moderate the relationship between IDS preference in Session 298 1 and Session 2 (see Supplementary Materials S2). Taken together, we find no significant 299 evidence of test-retest reliability across our preregistered analyses. 300

Table 3

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

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

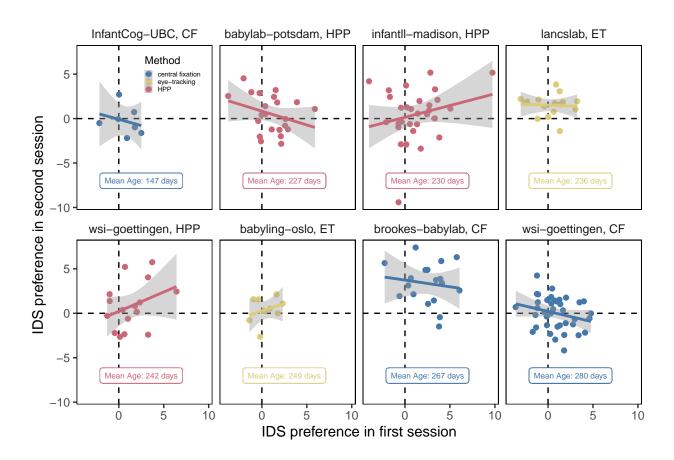


Figure 1. Correlation between IDS Preference in Session 1 and Session 2 in each lab and method. Dots indicate individual participants. Error bands represent 95 percent confidence intervals. The dashed line indicates no preference (i.e., a value of zero) for the first and second session, respectively.

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

Exploratory analyses with different inclusion criteria

To this point, all analyses were performed using the inclusion criteria from MB1, 302 which required only that infants contribute at least one trial per condition for inclusion 303 (i.e., one IDS and one ADS trial). However, more stringent inclusion criteria yielded larger 304 effect sizes in MB1. We therefore conducted exploratory analyses assessing test-retest 305 reliability after applying progressively stricter inclusion criteria, requiring two, four, six, 306 and eight valid trials per condition. Applying stricter criteria — and thereby increasing the 307 number of test trials — increased reliability numerically from r = 0.07 to r = 0.34 (Figure 308 2). In part due to a decrease in sample size, only one of these correlations was statistically 309 significant (when requiring six trial pairs): two valid trial pairs, t(152) = 0.90, p = .367; 310 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 313 reliability, but at the cost of substantial decreases in sample size (see Supplementary 314 Materials S4 and S6 for additional analyses, including moderator analyses using a more 315 restricted sample). 316



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 329 two sessions related? To test this question, we investigated whether participants' overall 330 looking times — irrespective of condition — were correlated between the first and second 331 session. There was a robust correlation between average looking time in Session 1 and 332 Session 2: infants with longer looking times during their first session also tended to look 333 longer during their second session, r = .45, 95% CI [.31, .57], t(156) = 6.28, p < .001334 (Figure 3B). This relationship held even after controlling for number of trials, b = 0.42, 335 95% CI [0.27, 0.58], t(154) = 5.52, p < .001, and participants' average age, b = 0.44, 95% 336 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 338 Session 1 and 2, r = .38, 95% CI [.24, .51], t(156) = 5.19, p < .001, and ADS stimuli in 339 Session 1 and 2, r = .40, 95% CI [.26, .53], t(156) = 5.49, p < .001 (Figure 3C; see Supplementary Materials S8 and S9 for further details, including an investigation of 341 item-level correlations).

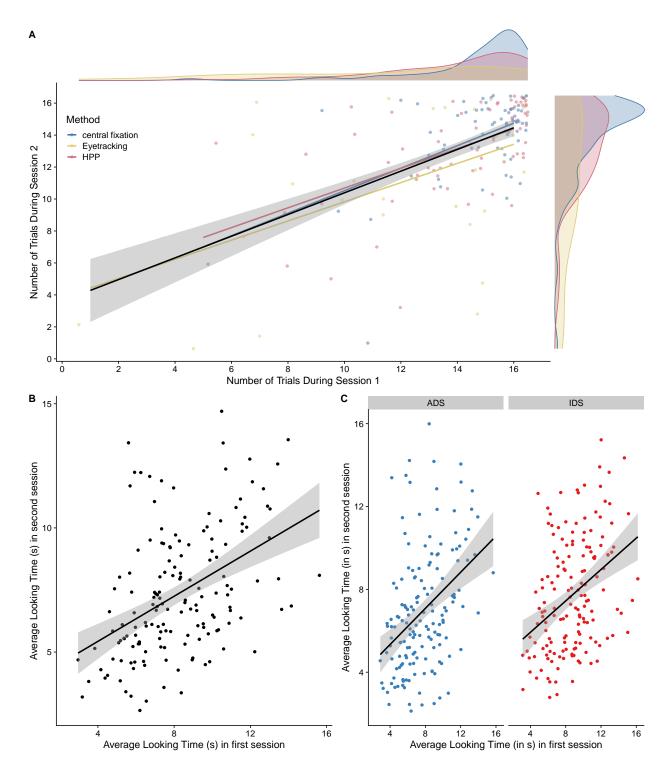


Figure 3. (A) Correlation between the number of trials contributed in Session 1 and Session 2. Each data point represents one infant. Colored lines represent linear fits for each method. (B) Overall correlations in average looking time (in s) between Session 1 and 2. (C) Correlations in average looking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

343

General Discussion

The current study investigated the test-retest reliability of infants' preference for IDS 344 over ADS. As a spin-off of the original MB1 project, we tested the IDS preference of infants 345 in two separate test sessions to assess the extent to which their pattern of preference would 346 remain consistent. While we replicated the original effect of infants' speech preference for 347 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 across all preregistered analyses. In other words, we were unable to find confirmatory evidence of stable individual differences in infants' preference for IDS. This finding is consistent with 351 past research suggesting low test-retest reliability in other infant paradigms (Cristia et al., 2016; Marimon & Höhle, 2022). Given that most experimental procedures conducted in infant research are interested in the comparison of groups, individual differences between 354 participants within a specific condition are usually minimized by the experimental 355 procedure while differences between conditions are maximized. As a consequence, infant 356 preference measures may be a good approach for capturing group-level phenomena, but 357 may be less appropriate for examining individual differences in development. 358

We also found no robust evidence that several hypothesized moderators influenced 359 test-retest reliability. While previous research suggests that HPP may be a more sensitive 360 index of infant preference and speech segmentation abilities (Junge et al., 2020; 361 ManyBabies Consortium, 2020), we find no evidence for higher test-retest reliability for 362 HPP relative to ET and CF. Similarly, our planned analyses found no evidence that the test-retest reliability of infants' infant-directed speech preference varied as a function of the time between test sessions, infants' language background, or infants' age. The absence of evidence for moderating effects should be treated with caution, given that fewer labs 366 contributed to the study than we had anticipated by the time of the preregistration, 367 resulting in limited power to detect interaction effects (see S2.1 for more detail on power 368

considerations). However, these analyses also suggest that the lack of test-retest reliability was not be due to variation across any of our hypothesized moderators.

Consistent with general psychometric theory (e.g., DeBolt, Rhemtulla, & Oakes, 371 2020), stricter inclusion criteria — and consequently a larger number of included test trials 372 per participant — tended to increase the magnitude of the correlation between test 373 sessions. However, this association was based on exploratory analyses and was in part only 374 observed descriptively, and hence should be interpreted with caution. A similar effect on 375 the group-level was found in the MB1 project, where a stricter inclusion criterion led to 376 bigger effect sizes (ManyBabies Consortium, 2020). As in MB1, higher reliability through 377 strict exclusions came at a high cost. In particular, with the strictest criterion, only a small 378 portion of the original sample size (24 out of 158 infants) could be included in the final 379 sample. In other words, applying stricter criteria leads to a higher drop-out rate and can 380 dramatically reduce the sample size. In the case of studies in the field of developmental 381 science, where there are many practical constraints on collecting large samples of infants 382 (e.g., birth rate in the area, restricted lab capacities, budget restrictions), a strict drop-out 383 criterion may often be difficult to implement. Note that studies in developmental science 384 already have above-average drop-out rates (Miller, 2017). In addition, drop out is likely not random: the number of trials individual infants contributed was highly correlated between test sessions in the current study. Therefore, implementing strict inclusion criteria that 387 result in high drop-out rates can further limit the generalizability of a study. Particularly 388 in the context of turning individual differences measures into diagnostic tools, high 380 drop-out rates have an additional limitation of not being broadly usable.

Even under best-case scenarios, reliability remained 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 et al. (2022) outline, low measurement reliability severely restricts power for detecting relationships between measures. Using the same approach as

Byers-Heinlein et al. (2022), 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 403 number of experimental trials. This approach seeks to increase the likelihood that 404 participants will contribute sufficient trials (after trial-level exclusions) to allow for precise 405 individual-level estimates (DeBolt et al., 2020; see also Silverstein, Feng, Westermann, 406 Parise, & Twomey, 2021). While this approach is promising, it may not always be feasible, 407 because the attention span of a typical infant participant is limited. Therefore, prolonging 408 the experimental procedure to maximize the absolute number of trials is often challenging 409 in practice. Other avenues for obtaining higher numbers of valid trials may include changes 410 in the procedure (e.g., Egger, Rowland, & Bergmann, 2020) or implementing multi-day test 411 sessions (Fernald & Marchman, 2012), which could become easier with the adoption of online testing methods (Lo, Hermes, Kartushina, Mayor, & Mani, 2023; Scott & Schulz, 413 2017; Weaver, Zettersten, & Saffran, 2022). 414

As our results are only based on the phenomenon of IDS preference (albeit, with
three widely used methods: HPP, CF, ET), it is essential to further assess the underlying
reliability of preferential looking measures within other areas of speech perception
(Marimon & Höhle, 2022). While most infants prefer IDS over ADS (Dunst et al., 2012),
patterns of preferential looking in other tasks (e.g., speech segmentation, artificial grammar
learning) are often inconsistent and can vary based on factors such as infants' experience
with the testing paradigm (Bergmann & Cristia, 2016; Santolin, Garcia-Castro, Zettersten,
Sebastian-Galles, & Saffran, 2021). These inconsistencies in looking behavior are especially

important to consider in the context of relating a direction of preference to later language
development, and can sometimes lead to seemingly contradictory findings. That is, both
familiarity and novelty responses have been suggested to be predictive of infants' later
linguistic abilities (DePaolis, Vihman, & Keren-Portnoy, 2014; Newman, Ratner, Jusczyk,
Jusczyk, & Dow, 2006; Newman, Rowe, & Ratner, 2016). 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 431 measures, we observed robust correlations for average looking times between session 1 and 432 2, both overall and for IDS and ADS stimuli considered separately (see also Supplementary 433 Materials S8 for an investigation of item-level correlations). This finding is consistent with 434 past results in infant looking time studies finding robust correlations in average looking 435 times across multiple sessions (Marimon & Höhle, 2022). It also raises an apparent puzzle: 436 why are overall looking times for ADS and IDS stimuli correlated, while difference scores 437 are not? One explanation is that infants have stable individual differences in how long they 438 look to stimuli, but little or no stable individual differences in their preference for one 430 stimulus type over another. This only partially explains the current pattern of results, however, because IDS looking time in Session 1 predicted IDS looking time in Session 2 441 even when controlling for ADS looking time, and vice versa (see Supplementary Materials 442 S8). In other words, the condition-specific looking time correlations are not fully explained 443 by overall looking behavior. Another long-established explanation is that difference scores tend to have poor measurement reliability, because difference scores combine error from individual measurements into a composite score and increasing the ratio of error relative to the variance between participants (Hedge et al., 2018; Lord, 1956). Given the limitations of difference scores (and composite scores in general), one goal for future research will be to 448 assess the use of trial-by-trial model-based approaches for estimating reliability (Haines et

450 al., 2020; Rouder & Haaf, 2019).

Limitations

While we had an above-average sample size for a study in infant research, we were 452 unable to approach the number of participants collected within the original MB1 study. 453 Several factors likely contributed to the lower participation rate. The call to participate in 454 the test-retest spin-off was delayed relative to the MB1 call, scheduling a second lab visit 455 for each participant involved a significant additional effort for participating labs, and there 456 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). A higher sample size and a larger number of participating labs from different 459 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. 461 A further limitation concerns the stimuli. While the order of the audio recording clips 462 presented to infants within a given trial differed between the first and second session, the 463 exact same stimulus material as in MB1 was used in both sessions. In particular, all 464 children heard the exact same voices in Session 1 and in Session 2. From a practical point 465 of view, this was the most straightforward solution for coordinating the experiment within the larger MB1 project. However, familiarity effects might have influenced infants' looking 467 behavior. Infants with longer looking times in their first session might have had more 468 opportunity to recognize familiar audio clips in their second session. For infants with short looking times, familiar audio clips would only occur towards the end of second-session trials, thus offering infants less opportunity to recognize voices from their first session. 471 Therefore, inconsistent familiarity with the stimulus material in the second session across infants might have artificially lowered test-retest reliability. However, in supplementary analyses, we found that test-retest reliability was not significantly moderated by infants' 474 overall looking time during their first testing session (see Supplementary Materials S10).

Moreover, infants' looking times generally declined in Session 2 compared to Session 1 (consistent with past work, e.g., Marimon & Höhle, 2022), limiting opportunities for infants to encounter previously experienced stimulus material.

479 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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