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## Research Highlights

- 2     • This study demonstrates a reliable infant-directed speech preference among African  
3       infants aged 3–15 months using a large-scale, multisite experimental design.
- 4     • Our findings showed no significant difference in IDS preference magnitude between  
5       African infants and method-matched samples from prior North American and  
6       European MB1 studies.
- 7     • Our study provides evidence for the cross-cultural generalizability of IDS preference  
8       while identifying practical challenges of conducting international multisite  
9       developmental research.

<sup>10</sup> Exploring variation in infants' preference for infant-directed speech: Evidence from a  
<sup>11</sup> multi-site study in Africa

12

## Abstract

13 Infants show a preference for infant-directed speech (IDS) over adult-directed speech  
14 (ADS). This preference has been linked to infants' language processing and word learning  
15 in experimental settings, and also correlates with later language outcomes. Recently, the  
16 cross-cultural consistency of infants' IDS preference has been confirmed by large-scale,  
17 multisite replication studies, but conclusions from these studies were primarily based on  
18 participants from North America and Europe. The current study addressed this sampling  
19 bias via a large-scale, multisite study of infants (3-15 months) across communities in  
20 Africa. We investigated whether participants showed a preference for IDS over ADS, and if  
21 so, whether the magnitude of their preference differs from effects documented in other  
22 populations of infants. Across six sites (total N = 200), we observed a preference for IDS  
23 over ADS ( $\hat{\beta}_{IDS \text{ vs. } ADS} = 0.06$ ), suggesting that infants look on average 6% longer on the  
24 IDS trials than the ADS trials. There was no significant difference between African infants  
25 in this study and a method-matched subsample of infants from prior studies of IDS  
26 preference. This study provides new evidence on the generalizability of IDS preference and  
27 looking-time methods more broadly, while also highlighting some of the challenges of global  
28 big team science.

29

*Keywords:* infant-directed speech; reproducibility; Africa; infants; generalizability

30

Word count: 9412

31 Exploring variation in infants' preference for infant-directed speech: Evidence from a  
32 multi-site study in Africa

33 Adults often speak to infants differently than to other adults, using a speech register  
34 known as infant-directed speech (IDS). Infant-directed speech tends to have exaggerated  
35 prosodic characteristics, including higher pitch, greater pitch variation, longer pauses,  
36 simplified grammatical structure, and shorter and slower utterances as compared to  
37 adult-directed speech (ADS; e.g. Fernald et al., 1989; Trainor & Desjardins, 2002). Even  
38 very young infants from a variety of language backgrounds have a preference for listening  
39 to IDS over ADS (Cooper, Abraham, Berman, & Staska, 1997; Cooper & Aslin, 1994;  
40 Fernald, 1985; Hayashi, Tamekawa, & Kiritani, 2001; Kitamura & Lam, 2009; Newman &  
41 Hussain, 2006; Pegg, Werker, & McLeod, 1992; Santesso, Schmidt, & Trainor, 2007; Singh,  
42 Morgan, & Best, 2002; Werker & McLeod, 1989). Infants' preference for IDS over ADS has  
43 also been demonstrated in a meta-analysis; across 34 studies, IDS preference had a fairly  
44 large average effect size with a value of Cohen's d 0.72 (Dunst, Gorman, & Hamby, 2012).

45 Why do infants prefer IDS? Perhaps IDS is intrinsically salient to infants because of  
46 its perceptual characteristics (e.g., higher pitch, greater pitch variability). Or perhaps, as  
47 infants are exposed to IDS, familiarity leads to preference. These explanations have  
48 different developmental predictions: while the intrinsic view would suggest an early  
49 preference (e.g., Cooper & Aslin, 1990), the exposure account would predict developmental  
50 increases in preference. Further, these explanations are not mutually exclusive: infants'  
51 early preference for IDS may motivate their parents to use more IDS, which in turn could  
52 lead infants to show a stronger IDS preference. Regardless of its origins, infants' preference  
53 for IDS may benefit their early language development. For example, in experimental  
54 studies, infants can segment words better in fluent speech produced in IDS than ADS  
55 (Thiessen, Hill, & Saffran, 2005), show better recognition of words introduced in IDS after  
56 a 24-hour delay (Singh, Nestor, Parikh, & Yull, 2009), and more successfully learn words

57 from IDS than ADS (Graf Estes & Hurley, 2013).

58 Further evidence comes from correlational studies, which have found that the amount  
59 of IDS in the language environment is positively related to children's language outcomes,  
60 such as vocabulary size (e.g., Ramírez-Esparza, García-Sierra, & Kuhl, 2014; L. A.  
61 Shneidman & Goldin-Meadow, 2012; L. Shneidman, Arroyo, Levine, & Goldin-Meadow,  
62 2013; but cf., Casillas, Brown, & Levinson, 2020; Casillas, Brown, & Levinson, 2021, who  
63 found similar timing of language development milestones even in a population that hears  
64 very limited IDS). Together, this work suggests that infants' preference for IDS over ADS  
65 may support their language development, which explains why infants' IDS preference  
66 continues to be an important topic in the literature on early childhood.

67 However, it is important to note that almost all prior studies, including the  
68 meta-analysis by Dunst et al. (2012) have included mainly infants learning English in  
69 Western, educated, industrialized, rich, and democratic (WEIRD) societies (Henrich,  
70 Heine, & Norenzayan, 2010), with only a few studies extended to non-Western infant  
71 populations learning languages other than English (Hayashi et al., 2001; Werker, Pegg, &  
72 McLeod, 1994). As such, there is a large sampling bias in the existing data about infants'  
73 preference for IDS, as in many other research topics in developmental psychology (see  
74 Nielsen, Haun, Kärtner, & Legare, 2017). This sampling bias is a problem for generalizing  
75 findings about infants' IDS preference to infants growing up in different cultures and  
76 learning different languages. In light of this generalizability issue – as well as the recent  
77 replication crisis in psychology (e.g., Open Science Collaboration, 2015) – infant  
78 researchers have begun to collaborate on large-scale, multi-site studies to replicate key  
79 developmental findings (Frank et al., 2017).

80 One of these multi-site projects investigated infants' preference for IDS over ADS: the  
81 ManyBabies1 study (MB1; ManyBabies Consortium, 2020). MB1 collected monolingual  
82 data from 67 laboratories, with a total sample of 2329 monolingual infants 3 – 15 months

83 old. The protocol for this experiment was simple: infants listened to alternating audio clips  
84 of IDS and ADS while viewing an uninformative visual stimulus (a colored checkerboard).  
85 Their looking time was measured over the course of up to 16 trials, 18s each in length (8  
86 IDS and 8 ADS). Notably, all participants in the study listened to stimuli that were  
87 constructed from naturalistic speech by North American mothers (speaking either to  
88 another adult or to their own infant). The mismatch between the stimuli and the native  
89 language of many infants in the study allowed inferences about native language effects and  
90 also minimized variability due to differences in the stimuli (a follow-up project now in  
91 progress seeks to measure native-language preferences in a subset of MB1 labs). Overall,  
92 older infants showed a stronger preference for IDS than younger infants. There was also an  
93 effect of infants' language backgrounds: North American infants exhibited a stronger IDS  
94 preference than infants who were not exposed to North American English (NAE).  
95 Although infants' ages and language backgrounds affected the magnitude of IDS  
96 preference, essentially all groups of infants preferred NAE IDS over ADS.

97 Despite the breadth of its sample relative to previous work, the MB1 study still  
98 constitutes a biased sample of infant populations in the world. Most of the data in MB1  
99 were contributed by laboratories in economically-advantaged areas, accessing relatively  
100 high socio-economic status participant populations. Further, although this large-scale  
101 study had a diverse sample from 17 countries, 60 out of the 67 participating laboratories  
102 were from Europe and North America, only a handful of laboratories were from Australia  
103 and Asia, and none were from Africa or South America. Thus, the sample studied in MB1  
104 came almost exclusively from Western, educated, affluent populations who heard  
105 Indo-European languages, limiting the generalizability of the findings to infants growing up  
106 in other cultural and linguistic contexts. This lack of evidence on generalizability of a key  
107 finding about infants' preference restricts our ability to build robust developmental theories  
108 of language learning across cultural contexts. Our current study takes a step towards  
109 addressing this gap.

We investigate whether infants growing up in a variety of African cultures show an IDS preference, using the paradigm developed by the MB1 study. Our study has both a theoretical goal and a practical goal. Theoretically, we are interested in whether IDS preference is a culturally and linguistically invariant developmental pattern (Nielsen et al., 2017). The inclusion of infants across many African cultures (who are acquiring many different languages, see Table 1) provides an important test of generalizability of the IDS preference. Practically, increasing sample diversity also promotes diversity among researchers engaged in developmental science and hopefully increasing exchanges between researchers across cultures. Thus, one goal of our study is building research networks to facilitate further studies with the communities represented in the current study.

Our study builds on a foundation of prior descriptive work investigating the generality of IDS across cultures. Although this work has investigated a variety of different cultures and languages, it can be (and often is) crudely summarized via the distinction between WEIRD and non-WEIRD cultures discussed above. We follow this convention here without endorsing this distinction as necessarily being meaningful in the context of our study, as IDS in WEIRD and non-WEIRD cultures shares similar prosodic properties. For example, Broesch and Bryant (2015) reported that IDS produced by North-American mothers, as well as by Kenyan and Fijian mothers, is produced with higher pitch, greater pitch variation, and is spoken at a slower rate than ADS. This finding is consistent with past work reporting that IDS shares some common exaggerated prosodic features (e.g., higher pitch, larger pitch variation) across diverse languages, which include French, Italian, German, Japanese, British English, American English (Fernald et al., 1989), Mandarin Chinese (Grieser & Kuhl, 1988), Thai, Australian English (Kitamura, Thanavishuth, Burnham, & Luksameeyanawin, 2001), Arabic (Farran, Lee, Yoo, & Oller, 2016).

IDS can also be recognized as being infant-directed by listeners from non-WEIRD cultures. Bryant, Liénard, and Clark Barrett (2012) reported that Turkana adults in Kenya can discriminate between NAE IDS and ADS (see similar results in Bryant &

137 Barrett, 2007 for Shuar hunter horticulturists from Amazonian Ecuador). These studies are  
138 consistent with findings from the MB1 studies showing that children who are not learning  
139 NAE, including children from Singapore and Korea, nonetheless show a preference for NAE  
140 IDS over ADS. Taken together, the common acoustic properties of IDS across different  
141 languages, and how NAE IDS can be recognized by non-native participants, raise the  
142 possibility of infants' IDS preference over ADS being quite consistent across different  
143 cultures and languages. However, it is possible that the strength of this preference would  
144 nonetheless be influenced by similarity between the test language (English) and the  
145 language(s) that each infant is learning, which could bolster the measured preferences to  
146 the extent that test and native language are similar (as in the case of infants learning other  
147 Indo-European languages with similar phonetic and acoustic properties). If this is the case,  
148 we expect that phylogenetic similarity between Indo-European languages and our stimuli  
149 would lead to comparable or stronger observed IDS preferences in samples of infants  
150 learning Indo-European languages than those learning languages in other families (e.g.,  
151 Bantu, the language family we expect to be most prevalent in our sample).

152 Despite evidence for general recognition of and preference for IDS across cultures, the  
153 strength of IDS preferences is likely modulated by exposure. Exposure to IDS in the home  
154 environment varies widely both within and between cultures (Casillas et al., 2020, 2021;  
155 Cristia, Dupoux, Gurven, & Stieglitz, 2019; LeVine, 1994; L. A. Shneidman &  
156 Goldin-Meadow, 2012; Vogt, Mastin, & Schots, 2015). Differences in IDS quantity have  
157 also been hypothesized to reflect differences in child-rearing practices across cultures. For  
158 example, direct verbal interaction between parents and infants can be rare in some societies  
159 (Heath, 1983; LeVine, 1994; LeVine & LeVine, 2016; Schneidman & Goldin-Meadow, 2012;  
160 Weber, Fernald, & Diop, 2017). Children in these societies – which are typically  
161 non-WEIRD, though certainly not all non-WEIRD societies can be characterized this way  
162 – are often expected to learn through observation and participation according to their skill  
163 levels (see Legare, 2019, for a review). Thus, infants and young children in such societies

164 may hear less IDS directly from their caregivers than those in WEIRD societies in which  
165 the norm involves a greater degree of direct address to parents. Of course, variation is also  
166 present within as well as across cultures. Within-culture variation has primarily been  
167 studied in North American contexts, where children from higher socioeconomic status  
168 (SES) families tend to hear more IDS than children from lower SES families (e.g., Hart &  
169 Risley, 1995; Hoff, 2006a; Huttenlocher, Waterfall, Vasilyeva, Vevea, & Hedges, 2010;  
170 Rowe, 2012; Schneidman & Goldin-Meadow, 2012; Weisleder & Fernald, 2013).

171 By virtue of our broad sample of African cultures, we expect that our study can  
172 capture substantial cultural variation in the average amount of IDS in children's  
173 environments. The African sites we sample vary widely in their degree of urbanization,  
174 their culture, their parenting values, and the average resources available in children's home  
175 environments – all of which have been argued to be meaningful dimensions governing  
176 children's early linguistic environment. For example, Keller (2012) suggested three  
177 prototypical cultural environments for children based on the degree of urbanization of the  
178 families in Western and non-Western societies. In this framework, in Western middle-class  
179 urban societies, highly educated parents generally aim to help children develop individual  
180 psychological autonomy. In contrast, in non-Western rural subsistence-based societies,  
181 parents generally aim to help children develop communal action autonomy, so that children  
182 have a strong sense of social responsibility and can contribute to the economic functioning  
183 of the family (e.g., farming). Importantly, non-Western middle-class urban societies are a  
184 hybrid of non-Western, rural and Western, urban societies, where parents generally want  
185 their children to develop more individual autonomy but also emphasize the importance of  
186 social responsibility in a large family. Broadly speaking, African families are from the  
187 non-Western, urban and non-Western, rural groups in this taxonomy (see Table 1).

188 The confirmatory analyses of our study are designed to test whether there are  
189 differences in the magnitude of IDS preferences measured in this sample and in the prior  
190 samples of MB1. Although the average IDS production in the African sites we examine is

191 unknown, consistent differences along this dimension might plausibly lead to variation in  
192 the magnitude of IDS preferences between our current study and MB1. In addition, our  
193 exploratory analyses attempt to understand whether variation in IDS preference among  
194 infants in our sample of African cultures is explained by demographic proxies related to  
195 this taxonomy (e.g., urbanization and/or socioeconomic status).

196 Since multilingualism is common in Africa (e.g., Posel & Zeller, 2016; Rosenhouse &  
197 Goral, 2004), many African children begin learning two or more different languages during  
198 infancy. Does early multilingualism alter infants' preferences for IDS? The  
199 ManyBabies1-Bilingual (MB1B) study provides some evidence that bilingual infants  
200 showed a similar preference for NAE IDS when compared to monolingual infants  
201 (Byers-Heinlein et al., 2021). MB1B examined bilingual infants' preference for NAE IDS at  
202 6 to 9 months and 12 to 15 months and found that bilingual and monolingual infants did  
203 not differ in terms of the magnitude of their IDS preferences. MB1B also found similar  
204 results to MB1, that older bilingual infants and those bilinguals with higher exposure to  
205 NAE show stronger IDS preference. However, as in the MB1 study, data collected in MB1B  
206 mainly came from laboratories in WEIRD areas, such as North America and Europe, with  
207 no laboratories from Africa, so the same caveats of generalizability apply to MB1B as to  
208 MB1. Thus, in the current study, we included both monolingual and multilingual infants,  
209 allowing us to assess the generalizability of MB1B's conclusions to our samples in Africa.

210 In sum, there are three primary (confirmatory) goals for the current study. First, we  
211 aim to measure infants' preference for North-American English IDS across a range of  
212 cultural and linguistic contexts in Africa. Second, we seek to measure developmental  
213 changes in this preference. As we found that older infants show stronger IDS preferences  
214 than younger infants in both MB1 and MB1B, we evaluate whether participants in our  
215 study show the same developmental increases in IDS preference. Finally, we investigate  
216 whether there are differences in IDS preferences between infants in Africa in our study and  
217 those in Europe and Asia in MB1 and MB1B. As an exploratory aim, we also examine

Table 1

*Test locations, participant ages, and languages by country.*

Country	Test location	Target age	Test language(s)
Ghana	University of Ghana, Accra	3–9 months	Akan, Ga, Ewe
Kenya	Nanyuki	9–15 months	Kikuyu, Kiswahili, English
Malawi	University of Malawi – Chancellor College, Zomba	6–15 months	Chichewa, English, or both
Nigeria	University of Jos, Plateau	3–15 months	Hausa, English, Birom, Ngas
Rwanda	Kigali	6–15 months	Kinyarwanda
Senegal	Dakar	3–15 months	French, Wolof
South Africa	Pretoria	3–15 months	Setswana, Xitsonga
Uganda	Makerere University, Kampala	3–15 months	Luganda, English

<sup>218</sup> relationships between parents' demographics, their responses to survey items regarding  
<sup>219</sup> subjective use of IDS, and their child's IDS preference.

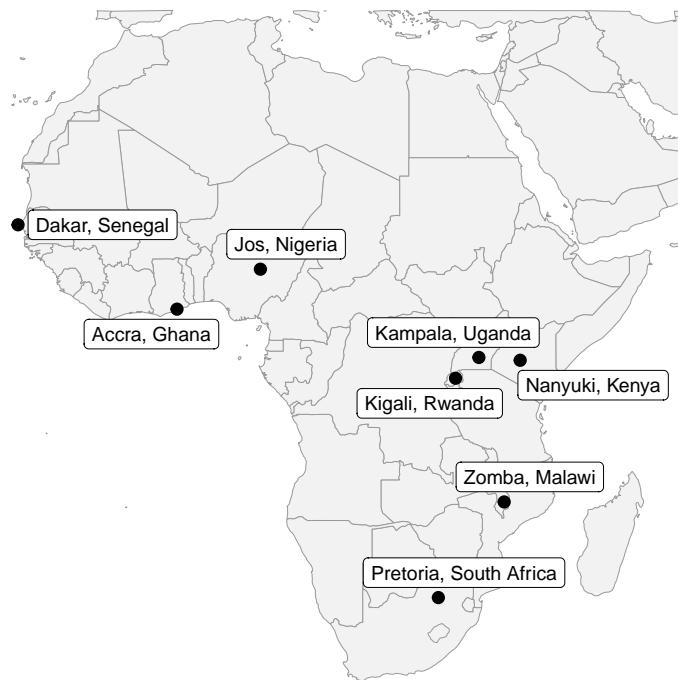
<sup>220</sup>

## Methods

<sup>221</sup> All deviations from the preregistration in the Stage 1 Registered Report are given in  
<sup>222</sup> Appendix A.

### <sup>223</sup> Participation Details

<sup>224</sup> **Time-frame.** On July 23, 2018, we issued an open call for participation by African  
<sup>225</sup> researchers via listservs and social/professional networks. Prior to submission, 11 labs  
<sup>226</sup> committed to data collection, but some labs were unable to collect data, in part due to  
<sup>227</sup> changing professional obligations and delays due to Covid-19 closures. We received data



*Figure 1.* Map shows the location of the eight participating labs.

from 8 labs (See Table 1 for target sample characteristics of each site; Figure 1 for the location of the lab). Our participating laboratories would recruit infants living in eastern (e.g., Kenya), western (e.g., Senegal) and southern (e.g., South Africa) regions of Africa. Because many of our participating laboratories are located in East Africa, thus East African participants are disproportionately represented in our sample. Data collection began September 21, 2021. We initially anticipated finishing data collection a year later, but labs encountered a wide variety of unforeseen circumstances due to the COVID-19 pandemic, challenges with receiving IRB approval, and equipment and staffing issues. Thus, data collection was extended through 2023. Unfortunately, due to experimental setup issues, data from two labs was not analyzable, leading to a final sample of 6 datasets. In both excluded datasets, looking times were not recorded: this was due to a software bug in one lab and experimenter error in the other.

**Age distribution.** Each participating laboratory was asked to recruit participants in two age bins: 3;0 – 9;0 and 9;1 – 15;0 months. Similar to MB1, each laboratory was

242 asked to collect data spanning the age bin window, but aiming for the mean of the age bin.

243       **Sample size determination.** We estimated the effect size of infants' IDS  
244 preference on the basis of the data from MB1. We used data from laboratories in MB1 that  
245 used the single-screen central visual-fixation preference procedure (which we also use here:  
246 see below) and that tested infants with no exposure to North American English (similar to  
247 our population of interest). In a mixed-effects model, we examined the effect of test trial  
248 type (IDS vs. ADS) on infants' looking time (log-transformed seconds), while controlling  
249 for normally-distributed random intercepts by infant and laboratory. The intercept,  
250 representing infants' average log-looking time across ADS trials, was 1.91; the variances of  
251 the random intercepts were 0.074 and 0.022 at the infant and laboratory levels respectively.  
252 The fixed-effect coefficient representing infants' preference for IDS over ADS was 0.080 and  
253 the residual variance was 0.33.

254       In the first power analysis, we simulated datasets based on the above coefficient  
255 estimates and variances. Using the simr package in R (Green & MacLeod, 2016), we ran a  
256 power analysis for a mixed-effect analysis with the above-mentioned simulated datasets  
257 (number of simulations = 1000). We were uncertain exactly how many labs to assume but  
258 settled on 10, given the likelihood of some later signups as well as some lab attrition.  
259 Assuming that we had 240 infants across 10 laboratories in each simulated dataset and an  
260 alpha level of 0.05, we found that the average power was 99.40% [95% confidence interval:  
261 98.70% – 99.78%] to detect the fixed ADS vs. IDS coefficient of 0.08. This first power  
262 analysis was based on very small random-effect variances estimated from MB1 and MB1B  
263 datasets. Given that most of the laboratories that participated in MB1 and MB1B had  
264 more resources and more extensive experience in running infancy studies in comparison to  
265 the participating laboratories in Africa, we planned for potentially higher variances in the  
266 data collected in the current project. Thus, we ran a conservative second power analysis by  
267 doubling the values of the random intercept and residual variances reported in the datasets  
268 from MB1 and MB1B, while holding constant the intercept and the fixed-effect coefficient

representing infants' preference for IDS over ADS. With larger variances, the average power estimate dropped to 87.20% [95% confidence interval: 84.97% – 89.21%] for a total sample of 240 infants. The power analysis can be found at <https://osf.io/jgr79>.

Given that MB1 reported around 15% data excluded in the final analysis, we expect the exclusion rate for our project is around 15% to 20%. Thus, each laboratory agreed to contribute a minimum of 32 infants (16 infants in each age bin), including infants tested but excluded for reasons not related to the demographic and age inclusion criteria (e.g., fussiness). Further, we encouraged each laboratory to contribute additional data beyond that minimum. We propose that our projected sample size of 352 would have sufficient power, as 80% of this sample size exceeds our targeted final sample size ( $n = 240$ ) based on the power analysis described above.

**Ethics.** All laboratories collected data under their own independent IRB protocol. Videos of individual infant participants during the experiment were recorded and stored at each laboratory. However, these videos were not shared with the central data analysis team. Laboratories were instead asked to only submit de-identified data for central data analyses.

## Exclusion Criteria

All data collected for the study (i.e., every infant for whom a data file was generated, regardless of how many trials were completed) were uploaded to a central database for data analysis. Every laboratory followed the protocol to report any infants who were tested in this study, including those who were excluded from the analysis. Furthermore, each laboratory followed the protocol to make note of the reasons that infants were excluded from the study. A total number of 274 infants were tested in this study, and 74 infants were excluded from the final analysis.

Typically, participants were only included in the analysis if they met all of the criteria below. However, we allowed parents to choose not to answer some of the questions

294 (e.g., about full-term gestation and developmental disorders) because disclosures might  
295 violate cultural norms in some areas of Africa. Thus, participating laboratories may have  
296 included infants who did not fully meet the inclusion criteria defined here

297       **Full-term.** We defined full term as gestation times greater than or equal to 37  
298 weeks. 11 (4.01%) of infants tested did not meet this criterion, and were excluded from  
299 further analysis. To maximize parents' comfort in participating in the experiment, they  
300 were given the option of not responding to questions about gestation.

301       **No developmental disorders or hearing loss.** We excluded infants with  
302 parent-reported developmental disorders (e.g., chromosomal abnormalities, etc.) or  
303 diagnosed hearing impairments. Developmental disorders and delays are stigmatized in  
304 some cultures in Africa (e.g., negative attitudes towards children with disorders or delays),  
305 therefore some parents may decline to answer the question about children's developmental  
306 disorders. In this case, we still tested the infants and included the infants' data in the  
307 analysis. This inclusion criterion was chosen to allow us to retain as much data as possible  
308 while ensuring our questionnaire accommodates cultural norms. Further, we noted that  
309 only 2 participants (i.e. less than 0.1%) in MB1 were excluded based on parents' report of  
310 developmental disorders. Accordingly, we do not expect that including children whose  
311 parents decline to answer this question would lead to an inclusion of large numbers of  
312 children with developmental disorders that could potentially skew the results in the study.  
313 4 (1.46%) of the infants tested did not meet this criterion. (We did not plan exclusions  
314 based on self-reported ear infections unless parents reported medically-confirmed hearing  
315 loss.)

316       **Trial-level and session-level errors.** Following MB1 and MB1B, we adopted a  
317 relatively liberal inclusion criterion for this study. To be included in the study, a child must  
318 have contributed non-zero looking time on at least one pair of test trials (i.e., one trial each  
319 of IDS and ADS from a particular stimulus pair). We asked laboratories to identify two  
320 different types of errors when uploading their data: trial-level errors and session-level

321 errors. Trial-level exclusions were based on whether we could use infants' data from a  
322 particular test trial. For example, if an infant only completed the first six test trials of the  
323 experiment, we entered this infant's data from the first six trials and discarded data from  
324 all other trials. In this case, laboratories would identify this infant's data from the first to  
325 sixth trials as "no trial errors" and any trials from the seventh trial onwards would be  
326 identified as "trial errors". In contrast, session-level errors were errors that occurred when  
327 running a particular participant. This type of error is different from the trial-level error  
328 exclusions because it indicates that errors occurred which affected an entire session (e.g.,  
329 failure to save data in the experiment). If a laboratory indicated a session-level error for a  
330 particular infant, all data from this infant was excluded from the analysis. In sum, infants  
331 who can contribute at least one pair of test trials (i.e., one IDS trial and one ADS trial)  
332 would have some data excluded at the trial level whereas infants who cannot contribute  
333 one pair of test trials would be excluded at the session level. In general, errors included the  
334 following: equipment error (e.g., no sound or visuals on the first pair of trials),  
335 experimenter error (e.g., an experimenter was unblinded in setups where infant looking was  
336 measured by live button press), or evidence of parent interference or other types of  
337 interference (e.g., talking or pointing by parents, construction noise, sibling pounding on  
338 door), and infants being uncooperative or fussy (e.g., crying, not willing to do the  
339 experiment).

340 Overall, at the trial level, 18 trials (0.39% of all trials) were excluded; relatively few  
341 trials were marked by labs as having errors. Due to experimental setup errors, data from  
342 two sites were unusable, leading to the exclusion of 60 infants (21.58% of all tested  
343 participants). No additional test sessions were excluded.

## 344 Participants

345 **Final sample.** Our final sample included 200 infants (see Table 2 for more specific  
346 sample demographic information) from 6 laboratories (mean sample size per laboratory:

Table 2

*Final sample's demographics and language background by country. Mean age in months.*

Country	N	Mean Age (SD)	Sex	Language background
Ghana	32	8.66 (3.72)	F: 17; M: 15	Monolingual: 3; Bilingual: 8; Other: 21
Kenya	27	11.14 (2.88)	F: 15; M: 12	Monolingual: 17; Bilingual: 5; Other: 5
Malawi	38	8.82 (4.10)	Unknown	Monolingual: 36; Other: 2
Rwanda	31	9.83 (4.23)	F: 16; M: 15	Monolingual: 31
South Africa	31	9.00 (3.71)	F: 16; M: 14; Unknown: 1	Monolingual: 23; Other: 3; Unknown: 5
Uganda	41	12.27 (2.17)	F: 21; M: 20	Unknown

<sup>347</sup> 33.33, SD: 5.16, range: 27 – 41). The mean age of infants included in the study was 296.49  
<sup>348</sup> days (range: 71 – 606). Similar to MB1, each laboratory was asked to collect data  
<sup>349</sup> spanning the two target age ranges (3;0 – 9;0 and 9;1 – 15;0 months); however, in practice,  
<sup>350</sup> many laboratories recruited participants outside the intended windows (younger than  
<sup>351</sup> 3-month-old:  $N = 4$ ; older than 15-month-old:  $N = 16$ ). We did not exclude these infants  
<sup>352</sup> from our analyses. See Figure 2 for a distribution of the age of the included participants in  
<sup>353</sup> each site. An additional 60 infants were tested but excluded (see the full details on  
<sup>354</sup> exclusions above).

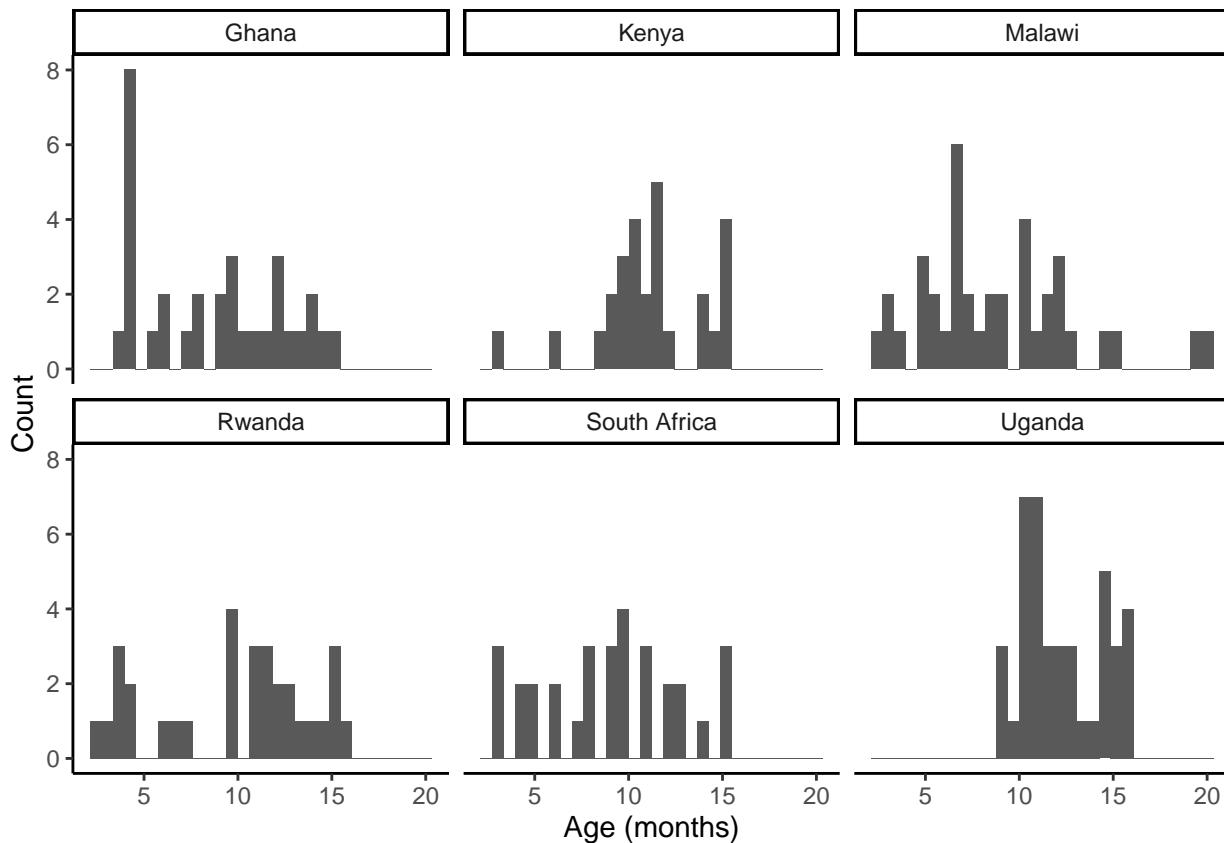


Figure 2. Histograms of the age distribution in each lab. X-axis represents the age in months.

As mentioned in the Introduction, multilingualism is common in Africa. Thus, many infants in the final sample are likely to have been exposed to more than one language. To assess infants' language backgrounds, each laboratory completed a family questionnaire with the participating parents (see materials in linked repository: [https://osf.io/jgr79/?view\\_only=5ee43f58762742daaa2caa21b85e3780](https://osf.io/jgr79/?view_only=5ee43f58762742daaa2caa21b85e3780)). Our family language background questionnaire was created based on the family language background questionnaire in the MB1 and MB1B studies, and included questions asking parents to estimate the number of hours that their infants heard different languages. We calculated the percentage of time that infants were exposed to a given language as the number of hours they hear that language (per day) divided by the total number of hours the infant hears any language each day. This method is simpler than the traditional interview method

366 used in assessing bilingual infants' language exposure (Byers-Heinlein et al., 2019), but in  
367 order to minimize the burden on participating laboratories and families, we decided to use  
368 a short questionnaire method to assess infants' language backgrounds.

369 In this paper, we define bilingualism following the criteria established in MB1B  
370 (Byers-Heinlein et al., 2021). Monolingual infants are defined as those who have a minimum  
371 of 90% exposure to one language. Simultaneous bilingual infants are defined using the  
372 following criteria: (i) infants are regularly exposed to two or more languages beginning  
373 within the first month of life; (ii) they have a minimum of 25% exposure to each of their  
374 languages. In other words, bilingual infants are exposed to two languages between 25% to  
375 75% of their time. Based on these criteria, it is possible that bilingual infants in our paper  
376 were exposed to multiple languages. For example, an infant with 45% English, 45% French,  
377 and 10% Spanish exposure would be regarded as a bilingual infant. Infants who did not  
378 meet the bilingual or monolingual criteria were designated as "other language background."  
379 All infants were included in the main, confirmatory analyses regardless of language  
380 background. Language background groupings were treated as a covariate in the analyses.

381 Based on the above-mentioned criteria, 87 infants were classified as monolingual  
382 infants, 13 infants were classified as bilingual infants, and 28 infants were classified as  
383 other. The remaining 72 infants' language background was unknown.

## 384 Materials

385 Visual stimuli. All visual stimuli were the same as those used in the MB1 study. We  
386 used a brightly colored static checkerboard as the fixation stimulus, and an animation with  
387 shrinking concentric multi-colored circles to ensure infants were attending to the screen at  
388 the start of each trial. All of the stimuli can be found at <https://osf.io/wh7md/>.

389 Auditory stimuli. All auditory stimuli were identical to those used in the MB1 study.  
390 The stimuli were recordings of North-American English mothers either speaking with

391 experimenters (ADS) or with their infants whose ages ranged from 122 to 250 days in a  
392 laboratory setting. Mothers were provided with a set of objects and were asked to talk  
393 about the objects with the experimenters and their infants in separate recording sessions.  
394 In total, two sets of auditory stimuli were created: one set consisted of 8 IDS stimuli and  
395 the other set consisted of 8 ADS stimuli. Each stimulus lasted for 18 seconds. The details  
396 of stimulus creation can be found in the report of MB1 (ManyBabies Consortium, 2020).

397 Volume. Each laboratory measured stimulus volume level using a smartphone app  
398 (e.g., the Android app “Sound Meter”). Labs kept the stimulus volume close to 63 – 65 dB  
399 SPL. According to the protocol, labs would measure and report the background noise level  
400 and the stimulus level. However, this information was not collected.

## 401 Procedure

402 Apparatus. Each laboratory used a laptop computer that had the experiment  
403 programmed in Habit 2.26 (Oakes, Sperka, DeBolt & Cantrell, 2019). Moreover, each  
404 laboratory used a computer monitor to present the visual stimuli, a speaker for audio  
405 stimuli, a webcam for the experimenter to observe and record infants’ performance,  
406 curtains/room dividers that separated the experimenter from the infant and parent during  
407 the experiment, and two sets of headphones: one for the experimenter and one for the  
408 parent.

409 Experimental procedure. The procedure was identical to the single-screen central  
410 visual fixation preference procedure reported in the MB1 study (ManyBabies Consortium,  
411 2020). Using the single-screen central fixation method, researchers measured in real time  
412 the duration of infants’ looking time to the computer monitor while they listened to the  
413 audio recordings. Infants’ looking time to the computer monitor indicated their preference  
414 for the audio recordings (i.e., IDS/ADS). Each laboratory followed procedural instructions  
415 closely (based on pre-recorded videos illustrating the procedures, which were shared with

416 all participating laboratories) to maintain the consistency of the experimental procedure  
417 across laboratories.

418 The experimenter explained the study to the parent and obtained consent from the  
419 parent before running the experiment. After completing the consent form, the  
420 experimenter led the participant to the testing room. To minimize distraction, the  
421 experimenter was separated from the infant and parent by curtains or a room divider.  
422 During the experiment, the infant sat on the parent's lap. To minimize any bias introduced  
423 by the experimenter or parent hearing the stimuli, each of them wore headphones and  
424 heard masking music during the experiment.

425 Parents were instructed not to speak to the infant during the experiment and not to  
426 point to the screen. Infants' performance was recorded by a webcam that was placed in  
427 front of and below the computer monitor. Infants' looking time to each trial was measured  
428 online by the experimenter, who observed the infant's behavior via the webcam. At the  
429 beginning of each trial, a short video of a colorful circle was presented to orient the infant's  
430 attention to the screen. Once the infant fixated on the screen, the experimenter started the  
431 trial. The first two trials of the session were warm-up trials that accustomed infants with  
432 the procedure of the experiment, so the infant's looking time during warm-up trials was  
433 not analyzed. The auditory stimuli for the warm-up trials was piano music that lasted 18  
434 seconds on each trial and the visual stimulus was the same as in the test trials (i.e., a  
435 colorful checkerboard). After the first two warm-up trials, the infant was tested with 16  
436 trials presenting the IDS and ADS stimuli. Each infant was randomly assigned to one of  
437 four pseudo-random orders to counterbalance the order of presentation of IDS and ADS  
438 stimuli. Within each order, there were four blocks and each block presented 2 IDS and 2  
439 ADS trials in alternating order. The presentation of the trials within each block were  
440 counterbalanced such that two blocks started with an IDS trial, and the other two blocks  
441 started with an ADS trial. On each trial, the auditory stimulus would continue to play  
442 until the infant looked away for 2 consecutive seconds or reached the maximum length of

443 the auditory stimulus (18 seconds). Experimenters used the Habit program to record all  
444 looking time for every trial. There was no minimum looking time per trial that was  
445 required for continuation of the experiment. However, as in the MB1 study, any looking  
446 time that was less than 2 seconds was not analyzed. We excluded 335 (11.46%) trials that  
447 had less than 2 seconds looking time in total.

448 After the main looking-time task, the parents answered questions from the  
449 experimenter about participant and family demographic information, such as infant sex,  
450 date of birth, language exposure, and preterm/full term status. The questionnaire was  
451 translated into the appropriate language(s) for participants from each data collection site.  
452 See supplementary materials for the English template and adaptations.

453 **General Lab Practices**

454 **Training of the experimenters.** Three of the authors conducted a 2-day training  
455 workshop in Nairobi, Kenya on January 28 – 29, 2020, which was attended by lead  
456 researchers from 8 of the participating laboratories. The training session provided an  
457 overview of the experimental procedure, advice on setting up the apparatus at the  
458 researcher's institution, and training, instructions and guidelines for running the  
459 experiment. Further, the first author sent instructions for experiment set-up and the  
460 workshop materials to all participating laboratories, and kept close contact with all lead  
461 researchers in the participating laboratories to provide technical support for the  
462 experiment.

463 **Training of research assistants.** Each laboratory was responsible for maintaining  
464 good experimenter training practices. We extended an invitation for the training workshop  
465 to one research assistant in each laboratory, so that the researcher primarily responsible for  
466 data collection could receive training directly as well. Following the MB1 study, each  
467 laboratory reported on which research assistant ran each infant using pseudonyms or  
468 numerical codes. After data collection, each laboratory completed a questionnaire

469 regarding their training practices, the experience and academic status of each  
470 experimenter, and their basic participant greeting practices.

471 **Results**

472 **Confirmatory Analyses**

473 Data processing and analytic framework. Our primary dependent variable of interest  
474 was infants' looking time (LT). Infants' looking time was defined as time spent fixating on  
475 the computer screen during test trials. We did not count LT when infants looked away  
476 from the screen, though the trial was discontinued if an infant looked away and did not  
477 look back to the screen within 2 seconds. Following MB1 and MB1B, we log-transformed  
478 looking times prior to statistical analysis (Csibra, Hernik, Mascaro, Tatone, & Lengyel,  
479 2016). We made this decision because we wanted to compare the data of the current study  
480 with those in MB1 and MB1B.

481 We tested our research questions via general linear mixed effects models. We fit all  
482 models using a maximal random effects structure (Barr, Levy, Scheepers, & Tily, 2013).  
483 Under this approach, we first specified all random effects that are appropriate for the  
484 experimental design (e.g., IDS/ADS trial type varied within subjects in our experimental  
485 design, thus it can be specified as a random effect by subject; see below for the full list of  
486 effects considered). If any of these mixed-effects models failed to converge, we used an  
487 iterative pruning strategy: first removing random slopes nested within subjects, next  
488 removing random slopes nested within labs, and finally removing random intercepts from  
489 groupings in the same order, retaining effects of trial type as these were of greatest  
490 theoretical interest. Following MB1 and MB1B, we fit all models using the lme4 package  
491 with the bobyqa optimizer, version 1.1-35.3 (Bates, Mächler, Bolker, & Walker, 2015) and  
492 computed confidence intervals and p values using the lmerTest package (Kuznetsova,  
493 Brockhoff, & Christensen, 2017).

In addition to the mixed-effect models, we assessed the reliability of measurement in our study by reporting the reliability of the infants' looking time difference to the IDS vs ADS stimuli across different trials. We reported the intraclass correlation coefficient (ICC) as our reliability measure. The ICC was computed using the psych package in R (Revelle, 2017). We reported an ICC3k measure, on the basis of a two-way random effects model, a mean-rating of 8 (i.e., we had 8 pairs of IDS and ADS trials) and consistency agreement (Koo & Li, 2016; Parsons, Kruijt, & Fox, 2019). The estimated ICC was 0.18, 95% CI [0.00, 0.34]. This relatively low value is consistent with previous work showing limited test-retest reliability in IDS studies (Schreiner et al., 2024).

Below is a description of variables in our mixed-effect models:

- Log\_lt: Dependent variable. Log-transformed looking time in seconds.
- Trial\_type: a dummy coded variable with two levels: ADS (reference) and IDS. A positive coefficient means that infants look longer to IDS trials compared to ADS trials.
- Age\_months: a continuous variable measuring the infant's age in months (centered).
- Trial\_num: An index for the current trial (1-16 for infants who completed the experiment). Excluded trials were reflected as missing trial numbers.
- Language\_background: this consisted of two dummy coded variables that represented infants from three different language backgrounds: monolinguals ( $\geq 90\%$  exposure to one's native language); bilinguals ( $\geq 25\%$  to each of their languages); other (any infants who were not categorized as monolinguals or bilinguals). Using monolinguals as the reference level, the two dummy-coded variables are: (i) bilingual – infants who were categorized as bilinguals would be coded as 1 and all other infants would be coded as 0; (ii) Other (any infants who are not monolinguals or bilinguals) – infants who were categorized as other would be coded as 1 and all other infants would be coded as 0. In this case, monolingual

520 infants would be coded as 0 in the above-mentioned dummy-coded variables.

- 521 • Infant Type: a dummy coded variable with two levels, representing infants living in  
522 Africa in our current study (coded as 1) and infants living in Europe, Australia and  
523 Asia who were not hearing North American English, with data from MB1(B) (coded  
524 as 0).

525 As a reminder, we examined the following research questions in our paper: (1) IDS

526 preference: whether infants in our multi-site African sample showed a preference for IDS

527 and what is the corresponding effect size of this preference; (2) Age effect: whether there

528 were changes in the infants' IDS preference across different ages; (3) Population

529 comparison: examine whether the magnitude of infants' IDS preference in our study

530 differed from infants in MB1 and in MB1B (comparing only infants in these three samples

531 who were not exposed to North American English).

532 Research questions 1 and 2: Infants' IDS preference and age effect. We addressed our

533 first two research questions using only the data collected in the current paper from

534 laboratories in Africa. We specified the following model: `log_lt ~ trial_type +`

535 `trial_num + age_months + trial_type * trial_num + age_months * trial_num +`

536 `age_months * trial_type + (trial_type * trial_num | subid) + (trial_type |`

537 `lab).`

538 The fixed-effects structure of this model included main effects of trial type (IDS vs

539 ADS), age, and trial number. This structure controls for the effects of each independent

540 variable on infants' looking time (e.g., longer looking times for IDS, shorter looking times

541 on later trials). In addition, we included several two-way interaction terms: trial type

542 interacting with trial number to model the possibility of infants' faster habituation to ADS,

543 age interacting with the trial type to model the developmental trajectory of infants' IDS

544 preference, and age interacting with trial number to model faster habituation for older

545 children. The random effects structure of the model controlled for subject-level and

546 lab-level grouping. For subject-level grouping, we added random intercepts and random  
 547 effects of trial type, trial number, and their interaction to model the possibility that each  
 548 infant may have different rates of habituation for IDS and ADS trials. For lab-level  
 549 grouping, we added a random effect trial type to model differences in IDS preferences  
 550 across labs.

551 After pruning for non-convergence, our final model specification was: `log_lt ~`  
 552 `trial_type + trial_num + age_months + trial_type * trial_num + age_months *`  
 553 `trial_num + age_months * trial_type + (1 | subid) + (1 | lab)`.

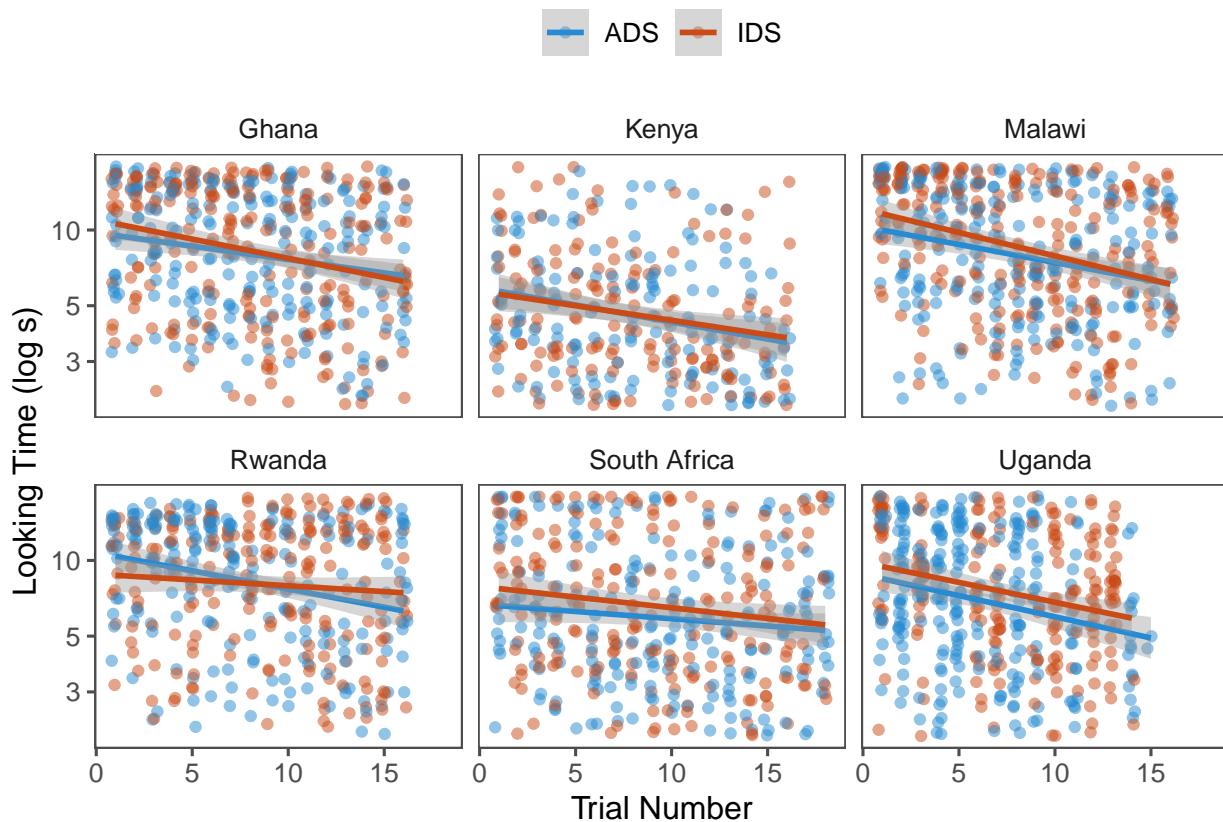
554 As in MB1 and MB1B, the fixed effect estimate for trial type corresponds to the  
 555 predicted infant-directed speech preference effect in units of log looking time (research  
 556 question 1). The fixed effect estimate for the interaction of trial type and age indicates the  
 557 estimated age-related change in infant-directed speech preference in log seconds per month  
 558 (research question 2).

Table 3

*Model estimates for Research Questions 1 and 2. The baseline for trial type is adult-directed speech. Trial number is mean-centered. Age is measured in months and standardized.*

Term	Estimate [95% CI]	std.error	t	p
(Intercept)	1.89 [1.66, 2.12]	0.09	21.34	< .01
Trial Type	0.06 [0.02, 0.10]	0.02	2.76	0.01
Trial Number	-0.03 [-0.04, -0.02]	0.00	-9.72	< .01
Age	-0.09 [-0.14, -0.04]	0.03	-3.40	< .01
Trial Type * Trial Number	0.00 [-0.01, 0.01]	0.00	-0.22	0.83
Trial Number * Age	0.00 [0.00, 0.01]	0.00	1.60	0.11
Trial Type * Age	0.02 [-0.02, 0.06]	0.02	0.94	0.35

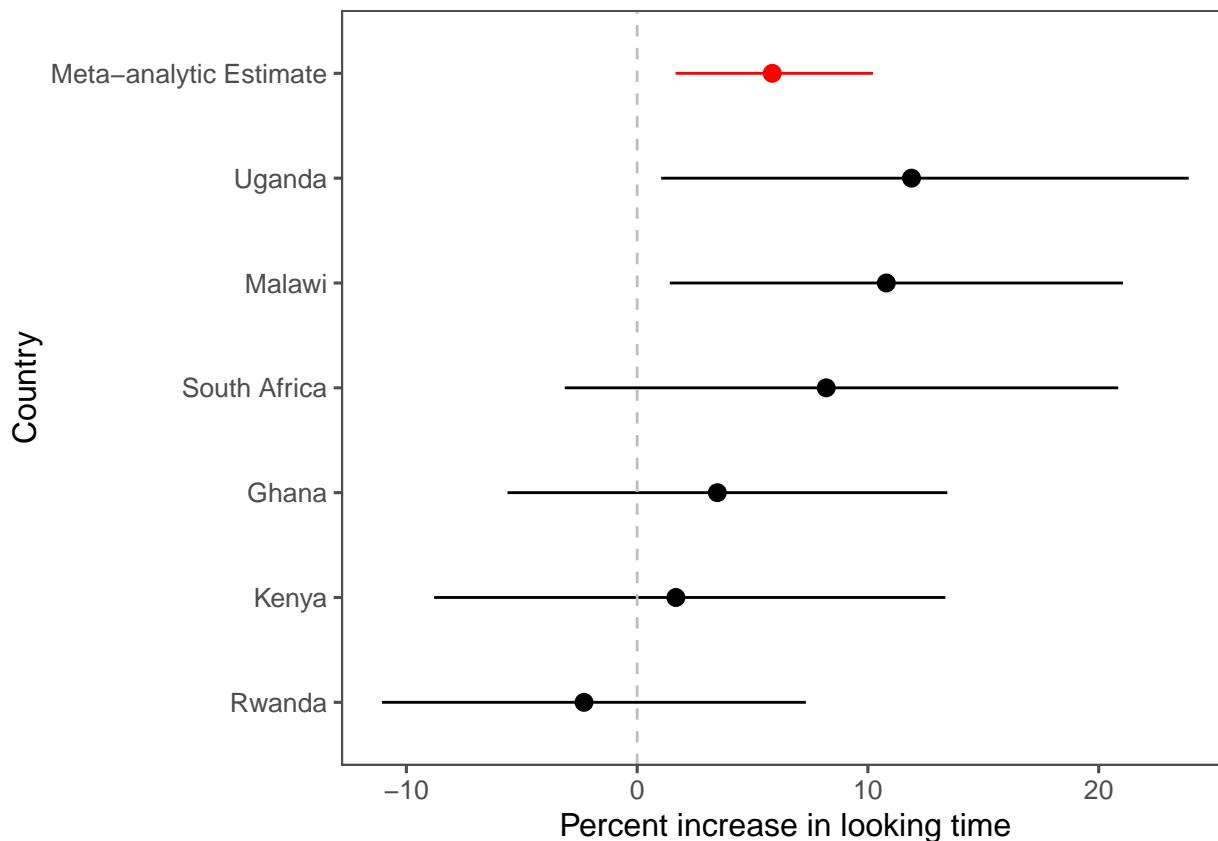
559 The model revealed a significant main effect of trial type, such that infants looked



*Figure 3.* By lab results. Each dot represents one trial of looking time. X-axis represents the trial number. Y-axis represents the log looking time in seconds. Red represents IDS trials. Blue represents ADS trials.

longer at IDS trials than ADS trials (Figure 3, Figure 4; Table 3;  $\beta = 0.06 [0.02, 0.10]$ ;  $SE = 0.02$ ;  $t = 2.76$ ;  $p = 0.01$ ). There was also a significant negative effect of trial number, indicating that looking times decreased over the course of the session ( $\beta = -0.03 [-0.04, -0.02]$ ;  $SE = 0$ ;  $t = -9.72$ ;  $p < 0.01$ ). Age in months was also a significant predictor, with older infants showing shorter looking times overall ( $\beta = -0.09 [-0.14, -0.04]$ ;  $SE = 0.03$ ;  $t = -3.40$ ;  $p < 0.01$ ). None of the interaction terms reached statistical significance, including the interaction between trial type and age, suggesting that the magnitude of IDS preference did not change reliably with age.

Research question 3: Population comparison. In this analysis, we compare the data



*Figure 4.* Percent increase in infants' looking time to infant-directed speech (IDS) relative to adult-directed speech (ADS) across six African sites and the meta-analytic estimate. While the meta-analytic estimate (red point) reflects the fixed-effect coefficient from the preregistered model, individual site estimates (black points) were obtained by fitting site-specific mixed-effects models predicting looking time using the same specification without the by-site random effect. Percent changes were computed from the fixed effect estimate, and horizontal lines represent 95% confidence intervals. Positive values indicate longer looking to IDS.

<sup>569</sup> collected from the laboratories in Africa to data collected in MB1 and MB1B in Germany,

<sup>570</sup> Italy, New Zealand, Turkey, United Kingdom. We selected the subset of data from MB1

<sup>571</sup> and MB1B that was collected using central fixation procedures (to match methods across

<sup>572</sup> studies) and from infants who were not exposed to North American English (non NAE) (to

573 match stimulus un-familiarity due to language background). While we could have  
574 controlled the methodological and demographic variables statistically (and hence included  
575 all data from MB1 and MB1B in the full model), we believed that the increase in model  
576 complexity – and comparable decrease in interpretability – outweighed the benefits of this  
577 strategy.

578 We examine whether our sample of infants' IDS preference is different from those in  
579 MB1 and MB1B with the following model: `log_lt ~ trial_type + trial_num +`  
580 `age_months + infant_type + language_background + trial_type * trial_num +`  
581 `age_months * trial_num + age_months * trial_type + trial_type * infant_type`  
582 `+ trial_num * infant_type + trial_type * language_background + (trial_type *`  
583 `trial_num | subid) + (trial_type | lab)`

584 In this mixed-effects model, the fixed-effects included main effects of trial type,  
585 language background, age, trial number, infants in our study/non NAE infants in MB1(B)  
586 and language background. In addition, we included several two-way interaction terms in  
587 the fixed effects structure: (i) trial type interacted with trial number, modeling the  
588 possibility of infants' faster habituation to ADS, (ii) age interacted with trial number,  
589 modeling faster habituation for older children, (iii) age interacted with trial type, modeling  
590 the developmental trajectory of infants' IDS preference, (iv) trial type interacted with  
591 infants in our sample, modeling the possible difference in IDS preference between infants in  
592 Africa and infants tested in MB1 and MB1B, (v) trial num interacted with infants in our  
593 sample, modeling the possible difference in habituation between our sample of infants and  
594 infants tested in MB1 and MB1B, and (vi) trial type interacted with language background,  
595 modeling the possible difference in IDS preference from infants with different language  
596 backgrounds. We adopted the same baseline random effects as in the previous model.

597 After pruning for non-convergence, our final model specification was: `log_lt ~`  
598 `trial_type + trial_num + age_months + infant_type + language_background +`

599 trial\_type \* trial\_num + age\_months \* trial\_num + age\_months \* trial\_type +  
600 trial\_type \* infant\_type + trial\_num \* infant\_type + trial\_type \*  
601 language\_background + (1 | subid) + (1 | lab). The fixed effect estimate  
602 corresponding to our research question is the trial\_type \* infant\_type, which captures  
603 differences in measured IDS preference between the current data and data from  
604 MB1/MB1B in units of log seconds of looking time.

605 The model revealed no significant difference in IDS preference between infants tested  
606 in our sample and those tested in MB1/MB1B, as indicated by the non-significant  
607 interaction between trial type and infant type (Table 4;  $\beta = 0 [-0.06, 0.06]$ ;  $SE = 0.03$ ;  $t =$   
608 0.03;  $p = 0.97$ ). However, there was a significant main effect of trial type ( $\beta = 0.02 [-0.04,$   
609 0.07];  $SE = 0.03$ ;  $t = 0.62$ ;  $p = 0.53$ ), suggesting that infants showed a reliable preference  
610 to IDS over ADS.

611 Consistent with expectations, looking times decreased over the course of the session  
612 ( $\beta = -0.05 [-0.05, -0.05]$ ;  $SE = 0$ ;  $t = -31.08$ ;  $p < 0.01$ ), and older infants looked for less  
613 time overall ( $\beta = -0.12 [-0.15, -0.09]$ ;  $SE = 0.01$ ;  $t = -8.26$ ;  $p < .001$ ). We also found that  
614 older infants habituated more quickly, as indicated by a significant negative interaction  
615 between age and trial number ( $\beta = 0 [-0.01, 0]$ ;  $SE = 0$ ;  $t = -3.79$ ;  $p < .001$ ). The  
616 interaction between trial number and infant type was also significant ( $\beta = 0.02 [0.01, 0.03]$ ;  
617  $SE = 0$ ;  $t = 6.64$ ;  $p < .001$ ), indicating that looking times declined more slowly across  
618 trials for infants in our sample compared to those in MB1/MB1B. No other interactions  
619 reached significance.

## 620 Exploratory Analyses

621 SES. Previous research in North America (e.g., Hart & Risley, 1995; Hoff, 2006b;  
622 Weisleder & Fernald, 2013) has shown that the quantity and quality of child-directed  
623 speech vary across families with different SES backgrounds. These differences in language

Table 4

*Model estimates for Research Questions 3. The baseline for trial type is adult-directed speech (ADS), trial number is mean-centered, age is measured in months and standardized, the baseline for infant type is infants from MB1/MB1B (non-NAE sample), and the baseline for language background is bilingual infants.*

Term	Estimate [95% CI]	SE	t	p
(Intercept)	1.76 [1.62, 1.91]	0.07	24.16	< .01
Trial Type	0.02 [-0.04, 0.07]	0.03	0.66	0.51
Trial Number	-0.05 [-0.05, -0.05]	0.00	-30.82	< .01
Age	-0.12 [-0.15, -0.09]	0.02	-8.38	< .01
Infant Type	0.13 [-0.09, 0.36]	0.12	1.32	0.21
Language Background (Monolingual)	-0.02 [-0.12, 0.07]	0.05	-0.48	0.63
Language Background (Other)	0.00 [-0.14, 0.14]	0.07	0.24	0.81
Trial Type * Trial Number	0.00 [-0.01, 0]	0.00	-0.81	0.42
Trial Number * Age	0.00 [-0.01, 0]	0.00	-4.39	< .01
Trial Type * Age	0.02 [0, 0.04]	0.01	1.86	0.06
Trial Type * Infant Type	0.00 [-0.06, 0.06]	0.03	-0.26	0.80
Trial Number * Infant Type	0.02 [0.01, 0.03]	0.00	4.96	<.01
Trial Type * Language Background (Monolingual)	0.04 [-0.02, 0.11]	0.03	1.42	0.16
Trial Type * Language Background (Other)	0.01 [-0.08, 0.10]	0.05	0.07	0.95

<sup>624</sup> input may drive differences in infants' preference for IDS. Thus, we explored how SES  
<sup>625</sup> affects infants' preference for IDS. SES was measured by primary caregiver's formal  
<sup>626</sup> education (number of years). We entered primary caregiver's formal education (in years) as  
<sup>627</sup> a predictor in the regression model specified for RQ1 and RQ2, along with its interaction  
<sup>628</sup> with trial type.

629        The interaction between trial type and primary caregiver education was not  
630 significant ( $\beta = -0.01 [-0.05, 0.03]$ ;  $SE = 0.02$ ;  $t = -0.46$ ;  $p = 0.65$ ), not allowing rejection  
631 of the null hypothesis that SES does not moderate infants' preference for IDS. In other  
632 words, the magnitude of IDS preference was similar regardless of caregivers' years of formal  
633 education. At the same time, we did observe a significant main effect of trial type, with  
634 infants looking longer at IDS than ADS trials overall ( $\beta = 0.06 [0.01, 0.10]$ ;  $SE = 0.02$ ;  $t =$   
635  $2.55$ ;  $p = 0.01$ ). Looking times decreased significantly across trials ( $\beta = -0.04 [-0.04, -0.03]$ ;  
636  $SE = 0$ ;  $t = -9.92$ ;  $p < 0.01$ ), and older infants looked for less time overall ( $\beta = -0.10$   
637  $[-0.15, -0.05]$ ;  $SE = 0.03$ ;  $t = -3.62$ ;  $p < 0.01$ ). No other interactions were significant. As a  
638 robustness check, we re-ran the model on the subset of infants with female primary  
639 caregivers (80.52% infants). The pattern of results was qualitatively unchanged.

640        Meta-analysis. For comparison with ManyBabies 1 and 1B, we computed  
641 standardized effect sizes for each lab using looking time, following the method used in  
642 ManyBabies Consortium (2020). We also confirmed that the effect size was similar when  
643 computing using log-transformed looking time. The resulting meta-analytic plot is shown  
644 in Appendix B: Figure B1. The meta-analytic effect size is  $0.17 [-0.03, 0.37]$  (log  
645 transformed:  $0.14 [-0.06, 0.34]$ ), which is numerically smaller than the  $.35 [0.29, 0.42]$   
646 reported in MB1 (Manybabies Consortium, 2020) and the  $.26 [0.09, 0.43]$  for bilingual  
647 infants reported in MB1B (Byers-Heinlein et al., 2021). But these estimates are not  
648 directly comparable due to the different method and age distribution in the current project  
649 (cf., Zettersten et al., 2024).

## 650                      General Discussion

651        Infants' preference for IDS is both an important phenomenon for understanding  
652 language learning and a case study for infant methods more broadly. The MB1 study  
653 investigated variation in IDS preference across laboratories and across countries and found  
654 a small but reliable effect such that infants preferred IDS over ADS. Although there was

655 substantial variation across labs, our analyses suggest that much of it reflected random  
656 sampling error, as the moderators we tested accounted for little of the between-lab  
657 variance. Nevertheless, the effect was moderated by infant age, language background, and  
658 experimental method. Though MB1 and its sister project MB1B investigated children from  
659 a wide variety of language backgrounds, no sites from Africa were included in this initial  
660 group of participating laboratories. The current study was designed to fill this gap.

661 We summarize our findings with respect to three research questions in the paper.  
662 First, consistent with MB1 and MB1B, we found evidence for a significant IDS preference  
663 in a sample of 200 African infants. Second, and unlike MB1, we did not find significant  
664 age-related variation in IDS preference, but given the relatively small magnitude of the  
665 overall effect, we may not have had sufficient power to detect an interaction with age in our  
666 primary analytic model. In addition, the variability in age across sites was limited, with  
667 four of the six labs having mean ages clustered around 8 to 9 months, leaving us little age  
668 variation to detect such an effect.

669 Finally, we were interested in comparing the magnitude of the IDS preference in the  
670 current study to the estimates obtained in MB1 and MB1B (with multilingual data  
671 providing an important comparison because of the diverse language backgrounds in our  
672 current sample). We did not find significant study effects in a model comparing data from  
673 the current study to a method- and language-background matched sample of infants from  
674 MB1 and MB1B. The magnitude of the IDS preference in African infants in the current  
675 study ( $d = 0.17$ ) was numerically smaller than the overall estimate reported by MB1  
676 ( $d = 0.35$ ) and the estimates for bilingual infants reported by MB1B ( $d = 0.26$ ). But the  
677 two estimates are not directly comparable; among other things, the current study was  
678 conducted with infants who were not growing up learning North American English and it  
679 used the central fixation method, both of which were associated with overall lower effect  
680 sizes in MB1 compared with other groups (Zettersten et al., 2024).

681 The current study provides important evidence on the generalizability of the IDS  
682 preference and of looking-time methods in infancy more broadly. Despite the diversity of  
683 MB1, as noted above, the vast majority of labs were from Western countries where  
684 Indo-European languages are spoken. Thus, the findings of the current study provide  
685 evidence that the IDS preference observed in MB1 is present in infants growing up in a  
686 diverse array of non-WEIRD environments. More broadly, there is limited work using  
687 looking-time methods for infancy research in the African context (cf. Pyykkö et al., 2019).  
688 Recent studies in Ghana have begun to build this foundation, demonstrating that Ghanaian  
689 infants show vowel-harmony preferences and can use harmony cues for speech segmentation  
690 using the same central-fixation paradigm (Omane, Benders, & Boll-Avetisyan, 2024;  
691 Omane, Boll-Avetisyan, & Benders, 2025). The current work complements these studies by  
692 extending the approach across multiple African sites and testing a different domain,  
693 thereby providing a broader demonstration of methodological feasibility across cultures.

694 Importantly, our findings also suggest that the IDS preference extends to infants who  
695 grow up in highly multilingual environments. Many infants in our study were likely  
696 exposed to more than one language from birth, a common characteristic in African  
697 contexts. For instance, infants in Ghana may receive input from two to six different  
698 languages within a single household (Omane, Benders, & Boll-Avetisyan, 2025). The  
699 persistence of the IDS preference at the group level despite such linguistic diversity  
700 underscores the robustness of this effect.

701 A second goal of the current study was to build a team of laboratories in Africa  
702 collecting data with infants. We were successful in accomplishing this goal and we believe  
703 that the current study represents the largest experimental study of African infants to date.  
704 Despite this success, we encountered a number of substantial challenges. Some of these  
705 were unique to the African context (e.g. variable internet connectivity) and to the specific  
706 time-period of our study (e.g., spanning the Covid-19 pandemic) while others were more  
707 general to the project of conducting “big team science” investigations across institutions

708 with a wide range of resources.

709        Although we received commitments for data collection from 11 teams, three of these  
710 teams were unable to collect data due to a variety of challenges relating to personnel and  
711 resources. One source of these challenges was the initiation of Covid-19 lockdowns soon  
712 after our initial project training in the winter of 2020. In some cases, personnel left the  
713 project or priorities changed, leading teams to lose the ability to participate. These  
714 lockdowns were also difficult even for teams who stayed involved in the project. Due to  
715 turnover and the long delay between initial training and setup for data collection after  
716 restrictions were eased, a number of procedural deviations were introduced. In two cases,  
717 these deviations were so severe that we could not analyze data from the site. In an ideal  
718 world, our group would have been able to conduct additional site visits and training after  
719 sites began collecting data; unfortunately this was not possible due to budget and  
720 personnel limitations.

721        In addition to logistical and training challenges, it is important to acknowledge the  
722 broader structural differences between the participating laboratories in this project and the  
723 Western laboratories that contributed to MB1 and MB1B. Most Western labs had access  
724 to established infrastructure for infant research, such as participant databases, mailing  
725 lists, as well as stable funding and institutional traditions of developmental  
726 experimentation. In contrast, many of the African sites in our study were building these  
727 capacities for the first time. Recruiting families without existing participant pools required  
728 extensive community engagement and word-of-mouth networks. The success of this effort  
729 therefore reflects not only careful coordination and training, but also the commitment and  
730 creativity of local researchers who developed sustainable pathways for infant recruitment  
731 and testing in settings with limited prior infrastructure.

732        These efforts resonate with broader calls to diversify and globalize infant research.  
733 Recent commentaries have highlighted that developmental science continues to draw

734 disproportionately from Western contexts, and have urged the field to address the  
735 structural barriers that prevent broader participation (Kidd & Garcia, 2022; Singh, Cristia,  
736 Karasik, Rajendra, & Oakes, 2023). Equitable global collaboration requires attention not  
737 only to sampling diversity but also to the power dynamics and infrastructural inequities  
738 that shape how research is conducted and credited. Encouragingly, recent initiatives such  
739 as the /L+/ Global School on Language Acquisition (Aravena-Bravo et al., 2024), co-led  
740 by several members of the MB1-Africa team, demonstrate how capacity building can foster  
741 sustainable participation from underrepresented regions.

742 The current study has a number of further scientific limitations, including some  
743 shared with prior studies including MB1 and MB1B. First, although we attempted to  
744 estimate IDS preference, we did so using a specific set of speech stimuli and a specific  
745 paradigm. It is likely that the stimuli used here are less extreme than many used in prior  
746 studies, and further, they are produced in North American English, making them  
747 linguistically unfamiliar to one degree or another to all of the infants in our study.  
748 Followups using native language stimuli are needed to measure the importance of this  
749 choice to the IDS preference (i.e., MB1N, an in-progress follow up study to MB1 examining  
750 native language IDS preferences). Second, although we invited broad participation, our  
751 samples are convenience samples in at least two ways: both of the sites who participated  
752 and the infants who participated at each site. Thus our effect size estimates cannot be  
753 treated as population effects but rather “proof of concept” that an IDS preference can be  
754 observed in African infants across a diverse set of sites. Finally, although we did not  
755 observe major demographic variation, we caution against over-interpretation of any  
756 demographic differences in IDS preference given that IDS preference has not been shown to  
757 be individually predictive of any later outcomes (Soderstrom et al., 2024).

758 In sum, our study offers a case study of “big team science” (Coles, Hamlin, Sullivan,  
759 Parker, & Altschul, 2022) carried out via a collaboration between African researchers and  
760 the ManyBabies Consortium. Although it faced a variety of logistical challenges (many

761 shared with other grass roots efforts; Baumgartner et al., 2023), it nevertheless yields  
762 important evidence on generalizability, of a key phenomenon in early language learning, of  
763 looking time methods, and finally of a broad-based collaborative model for studying infant  
764 development.

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## Appendix A

### Discrepancies between pre-registration and final analyses

#### 1. ICC comparison

*Pre-registration:* We planned to compare the estimated ICC in our study with the ICC reported in MB1(B).

*Final:* This comparison was not conducted because MB1(B) did not report ICC values.

#### 2. Metrics of effect heterogeneity

*Pre-registration:* We planned to report distributional metrics describing heterogeneity across labs (Mathur & VanderWeele, 2020) using the *MetaUtility* package.

*Specifically,* we proposed to estimate: (1) the percentage of effects greater than 0, (2) the percentage greater than Cohen's  $d = 0.2$ , and (3) the percentage less than  $d = -0.2$ . These metrics were to be reported only if random slopes of trial type by lab were included in the final model, their variance was estimated as greater than 0, and at least 10 labs contributed data.

*Final:* Random slopes of trial type by lab were not included in the final model, and only six labs contributed data. As a result, we did not estimate these metrics.

#### 3. Subset of MB1 and MB1(B) data

*Pre-registration:* We planned to subset MB1 and MB1(B) data to include only infants tested with central fixation procedures and infants not exposed to North American English (NAE), to maximize comparability with the present study.

*Final:* In MB1 and MB1(B) data, information about English exposure often did not distinguish between American and British English. Therefore, we excluded infants based on country of residence (United States/Canada) rather than reported language exposure.

#### 4. Urban–rural exploratory analysis

1004       *Pre-registration:* We planned an exploratory analysis testing whether infants' IDS  
1005       preference differed between urban and rural areas, motivated by prior findings of  
1006       differences in parental speech input across these contexts (e.g., Keller, 2012; Vogt et  
1007       al., 2015). *Final:* We did not collect information on urban versus rural residence, so  
1008       this analysis was not conducted.

1009       **5. Socioeconomic status (SES) analyses**

1010       *Pre-registration:* We planned to measure SES using both mothers' years of formal  
1011       education and the MacArthur Scale of Subjective Social Status (MacSSS), including  
1012       both variables in regression models.

1013       *Final:* We did not have MacSSS data for most participants, and information on  
1014       mothers' education was incomplete. We therefore used primary caregiver education  
1015       as the main SES variable and conducted a robustness check using the subset of infant  
1016       whose primary caregiver's sex is female.

## Appendix B

### By-lab Meta-analysis

1017 We computed a single effect size per lab and fit an intercept-only mixed-effect  
1018 meta-regression to estimate the overall IDS preference across sites. This approach provides  
1019 a comparable summary of results across sites. Even with a standardized protocol, sites  
1020 differ in their cultures, recruitment pools, equipment, and experimenter behavior. A  
1021 random-effects meta-analysis treats those differences as legitimate heterogeneity rather  
1022 than noise, yielding a conservative estimate of the cross-lab mean and its uncertainty.

1023 To do so, we calculated each infant's mean IDS–ADS difference score <sup>1</sup>, standardized  
1024 these within lab to obtain effect sizes, and estimated their sampling variances. These  
1025 lab-level estimates were then entered into a REML random-effects model to produce the  
1026 pooled effect size and 95% confidence interval (Figure 1). The meta-analytic effect size is  
1027 0.17[-0.03, 0.37], which is numerically smaller than the .35 [0.29, 0.42] reported in MB1  
1028 (Manybabies Consortium, 2020) but these estimates are not directly comparable due to the  
1029 different method and age distribution in the current project (cf. Mathur et al., 2024).

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<sup>1</sup> Due to an experimental procedure error, infants at the South Africa site were not always presented with complete IDS–ADS stimulus pairs; in some cases, the same stimulus was played multiple times. While this issue did not affect the random-effects model in the main analysis, it does impact the present meta-analysis. We therefore trimmed the data by (1) retaining only the first presentation of each trial and (2) including only trials in which both the IDS and ADS versions were presented. The trimmed data includes 176 trials from 28 infants.

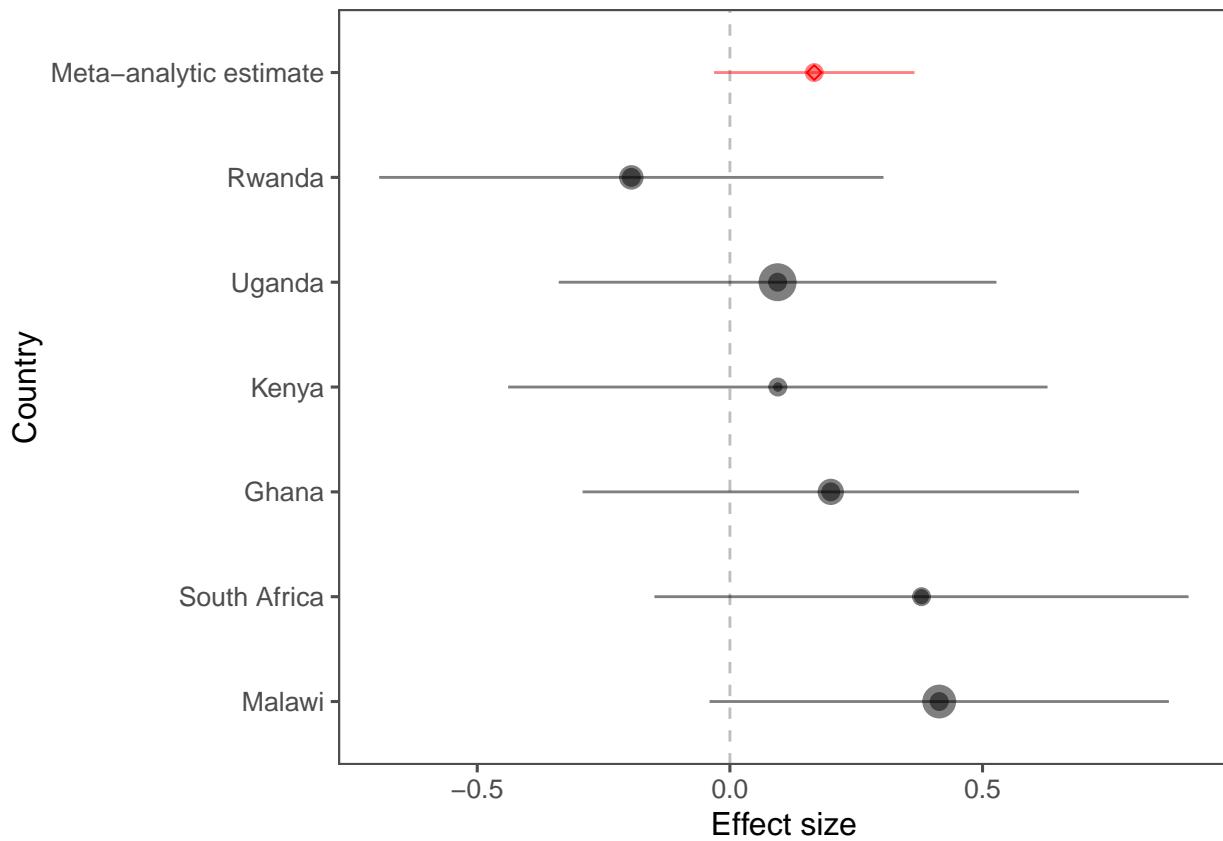


Figure B1. Forest plot of lab-level standardized effect sizes (for the IDS–ADS preference). Points represent individual country estimates, with size proportional to the inverse of their sampling variance; horizontal bars show 95% confidence intervals. The meta-analytic aggregate (top, red) is from an intercept-only random-effects model.