

Manybabies1 Test-Retest Supplementary Information

1

2

3

#### 4 S1: Deviations from the preregistration

5 Below, we document all deviations from the preregistered methods and analyses  
6 <https://osf.io/v5f8t>.

- 7 • All infants with usable data for both test and retest session were included in the  
8 analyses, regardless of the number of total of infants a lab was able to contribute after  
9 exclusion. This decision is consistent with past decisions in ManyBabies projects to  
10 be as inclusive about data inclusion as possible (ManyBabies Consortium, 2020).
- 11 • A small number of infants with a time between sessions above 31 days were also  
12 included in the analyses (n=2).
- 13 • Consistent with analytic decisions in ManyBabies 1 (ManyBabies Consortium, 2020),  
14 total looking times were truncated at 18 seconds (the maximum trial time) in the  
15 small number of cases where recorded looking times were slightly greater than 18s  
16 (presumably due to small measurement error in recording infant looking times). -The  
17 preregistration included a series of secondary planned analyses that ultimately did  
18 run because we collected fewer participants than originally expected.
- 19 • In assessing differences in IDS preference between test and retest sessions, we  
20 preregistered an additional linear mixed-effects model including a by-lab random  
21 slope for session. This model yielded qualitatively equivalent results (see R  
22 markdown analysis script for the main manuscript). However, the model resulted in a  
23 singular fit, suggesting that the model specification may be overly complex and that  
24 its estimates should be interpreted with caution. We therefore focused only on the  
25 first preregistered model (including only by-lab and by-participant random  
26 intercepts) in reporting the analyses in the main manuscript.

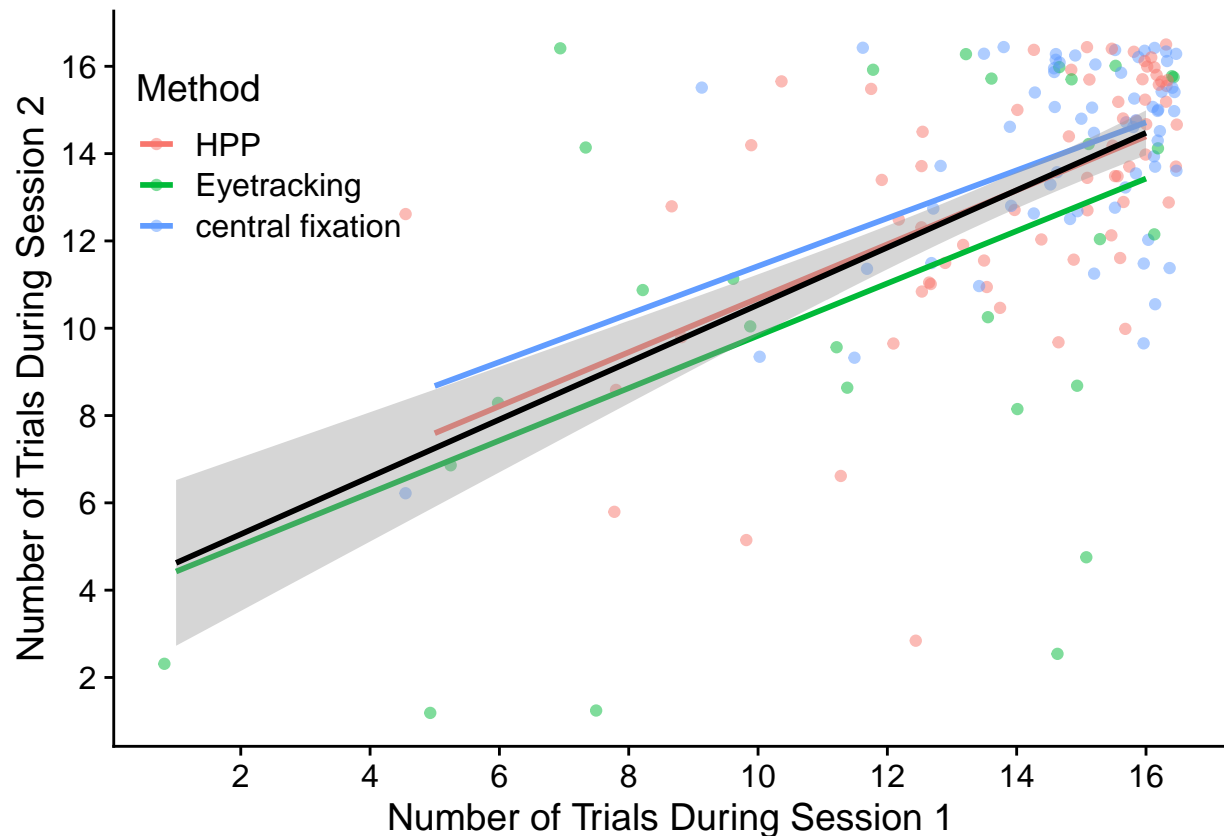


Figure 1. Correlation between the number of trials contributed in session 1 and session 2. Each data point represents one infant. Colored lines represent linear fits for each method.

## S2: Relationship between the number of trials infants contribute in each session

Are there stable individual differences in how likely an infant is to contribute a high number of trials? To answer this question, we conducted an exploratory analysis investigating whether there is a relationship between the number of trials an infant contributed in session 1 and session 2. Do infants who contribute a higher number of trials during their first testing session also tend to contribute more trials during their second testing session? A positive correlation between trial numbers during the first and second session would indicate that there is some stability in a given infant's likelihood of remaining attentive throughout the experiment. On the other hand, the absence of a correlation would indicate that the number of trials a given infant contributes is not predictive of how

many trials they might contribute during their next session.

We found a strong positive correlation between number of trials contributed during the first and the second session  $r = .58$ , 95% CI  $[.47, .68]$ ,  $t(159) = 9.05$ ,  $p < .001$  (see Figure 1). This result suggests that if infants contribute a higher number of trials in one session, compared to other infants, they are 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. Researchers should therefore be mindful of the fact that decisions about including or excluding infants based on trials contributed may selectively sample a specific sub-set of the infant population they are studying (Byers-Heinlein, Bergmann, & Savalei, 2021; DeBolt, Rhemtulla, & Oakes, 2020).

### **S3: Patterns of preference across sessions**

We also conducted analyses to explore whether there were any patterns of preference reversal across test sessions. While there was no strong correlation in the magnitude of IDS preference between test session 1 and test session 2, here we asked whether infants consistently expressed the same preference across test sessions. Overall, 58.20% of the infants had a consistent preference from test to retest session, indicating that infants were not more likely than chance to maintain their preference from test session 1 to test session 2 (exact binomial test;  $p = 0.05$ ). Of the 158 total infants, 44.90% of infants showed a consistent infant-directed speech preference and 13.30% showed a consistent adult-directed speech preference. 23.40% of infants switched from an infant-directed speech preference at test session 1 to an adult-directed speech preference at test session 2 and 18.40% switched from an adult-directed speech preference to an infant-directed speech preference.

Next, we explored whether we could detect any systematic clustering of infants with distinct patterns of preference across the test and retest session. We took a bottom-up

approach and conducted a  $k$ -means clustering of the test-retest difference data. We found little evidence of distinct clusters emerging from these groupings: the clusterings ranging from  $k=2$  (2 clusters) to  $k=4$  (4 clusters) appear to simply track whether participants are approximately above or below the mean looking time difference for test session 1 and test session 2, and the diagnostic elbow plot shows little evidence of a qualitative improvement as the number of clusters is increased.

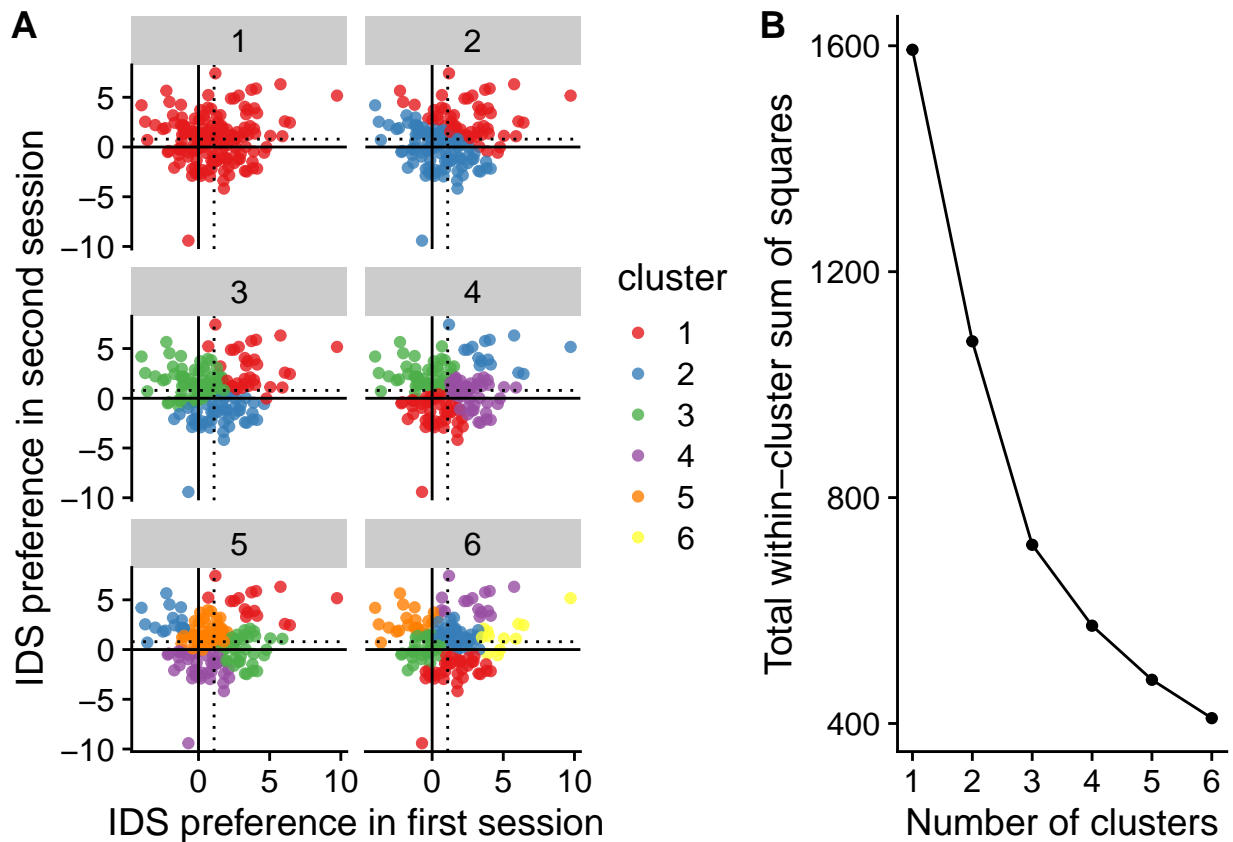


Figure 2. (A) Results from the  $k$ -means clustering analysis of IDS preference in session 1 and 2 for different numbers of  $k$  and (B) the corresponding elbow plot of the total within-cluster sum of squares. In (A), points represent individual participants' magnitude of looking time difference at test sessions 1 (x-axis) and 2 (y-axis). The solid line indicates no preference for IDS vs. ADS, the dotted lines indicate mean IDS preference at test session 1 and 2, respectively. Colors indicate clusters from the  $k$ -means clustering for different values of  $k$ .

68 **S4: Correlations in average looking times between sessions**

69 **S4.1: Relations between overall looking time in session 1 and 2.** There is a  
70 strong relationship between average overall looking in the first test session and the second  
71 test session, even after controlling for number of trials in the first and second session.

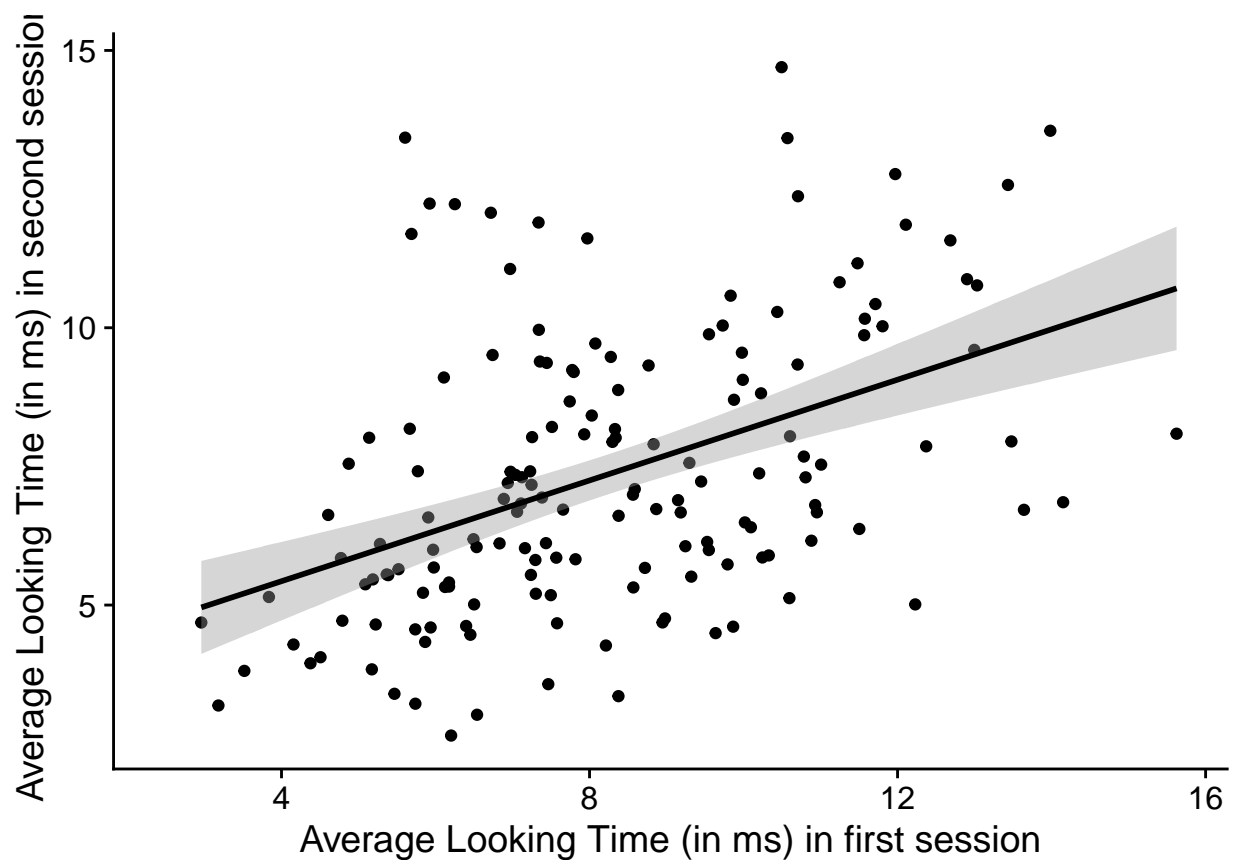
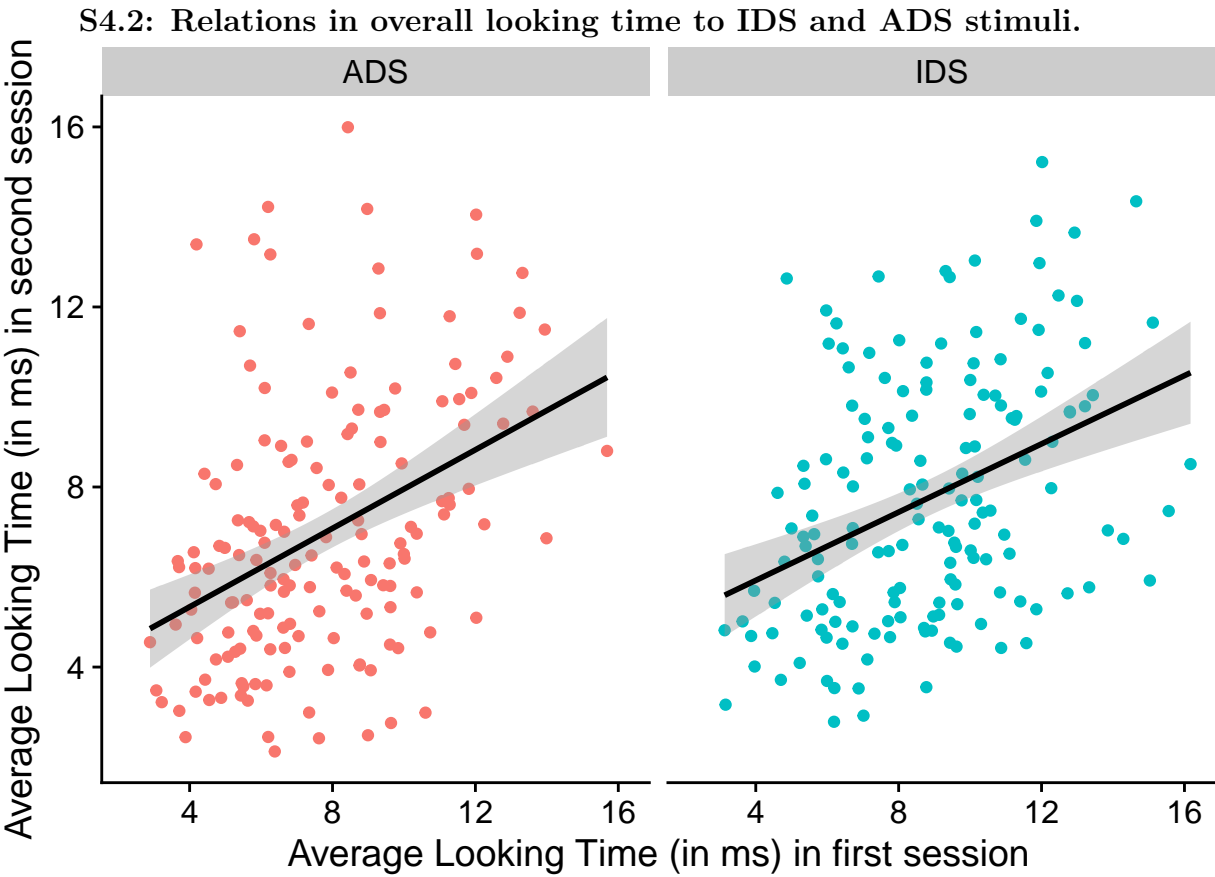


Table 1

*Average Looking during session 1 predicted from average looking at session 2, controlling for trial number for each session.*

Predictor	<i>b</i>	95% CI	<i>t</i>	<i>df</i>	<i>p</i>
Intercept	2.55	[0.38, 4.73]	2.32	154	.022
Mean lt 1	0.42	[0.27, 0.58]	5.52	154	< .001
N 1	-0.08	[-0.24, 0.08]	-0.96	154	.338
N 2	0.18	[0.04, 0.32]	2.52	154	.013



```

76 ## Call:
77 ## lm(formula = LT_Retest_IDS ~ LT_Test_IDS + LT_Test_ADS, data = agg_by_subj_condition_
78 ##
79 ## Residuals:
80 ##      Min       1Q   Median       3Q      Max
81 ## -4.2721 -1.7567 -0.2799  1.4822  6.4805
82 ##
83 ## Coefficients:
84 ##              Estimate Std. Error t value Pr(>|t|)
85 ## (Intercept)   3.9749     0.6902   5.759 4.41e-08 ***
86 ## LT_Test_IDS   0.2123     0.1008   2.105  0.0369 *
87 ## LT_Test_ADS   0.2467     0.1044   2.362  0.0194 *
88 ## ---
89 ## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
90 ##
91 ## Residual standard error: 2.52 on 155 degrees of freedom
92 ## (7 observations deleted due to missingness)
93 ## Multiple R-squared:  0.1771, Adjusted R-squared:  0.1665
94 ## F-statistic: 16.68 on 2 and 155 DF,  p-value: 2.751e-07
95 ##
96 ## Call:
97 ## lm(formula = LT_Retest_ADS ~ LT_Test_IDS + LT_Test_ADS, data = agg_by_subj_condition_
98 ##
99 ## Residuals:
100 ##      Min       1Q   Median       3Q      Max
101 ## -5.556 -1.771 -0.489  1.254  8.901
102 ##

```



```

103 ## Coefficients:
104 ##           Estimate Std. Error t value Pr(>|t|)
105 ## (Intercept)   3.2374     0.7356   4.401   2e-05 ***
106 ## LT_Test_IDS   0.1103     0.1075   1.026   0.30641
107 ## LT_Test_ADS   0.3563     0.1113   3.201   0.00166 **
108 ## ---
109 ## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
110 ##
111 ## Residual standard error: 2.686 on 155 degrees of freedom
112 ##   (7 observations deleted due to missingness)
113 ## Multiple R-squared:  0.1677, Adjusted R-squared:  0.157
114 ## F-statistic: 15.62 on 2 and 155 DF,  p-value: 6.619e-07

```

115 **S4.3: Relations for specific ADS and IDS stimuli between the first and**  
 116 **second session.**

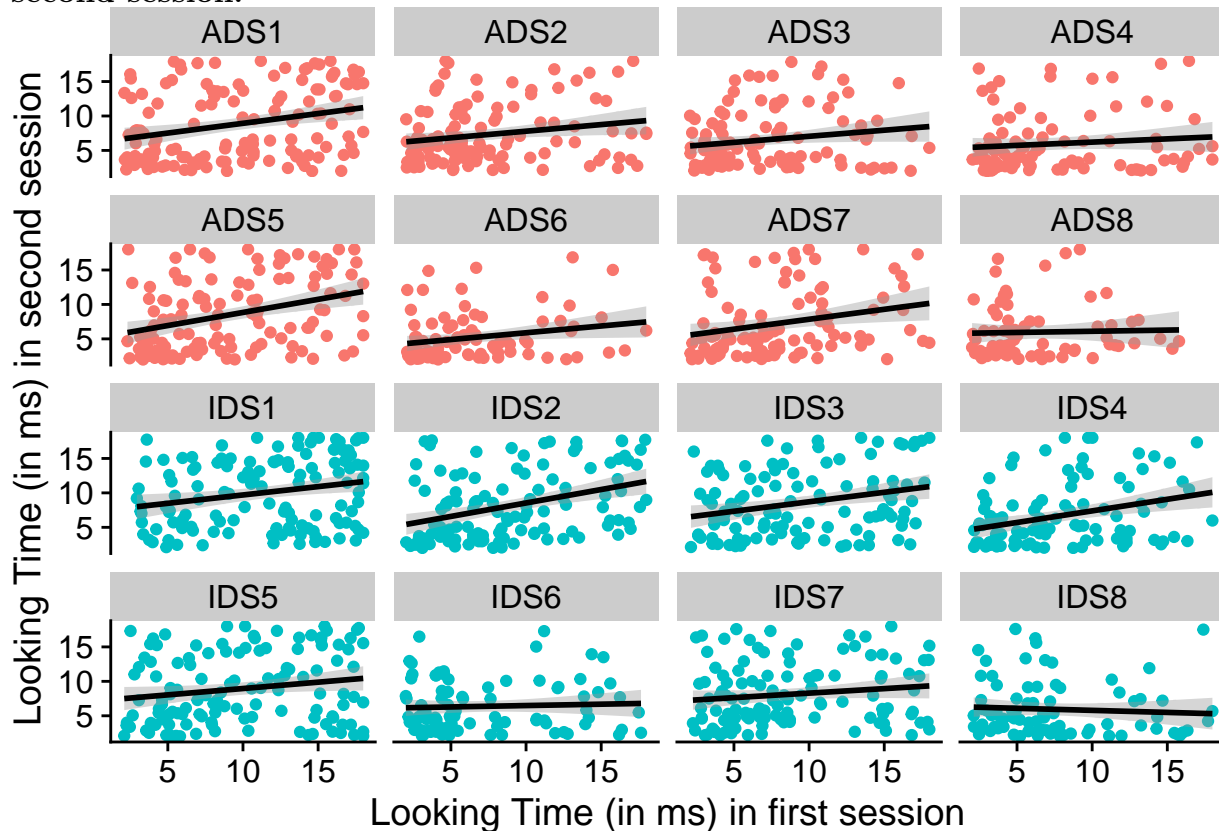
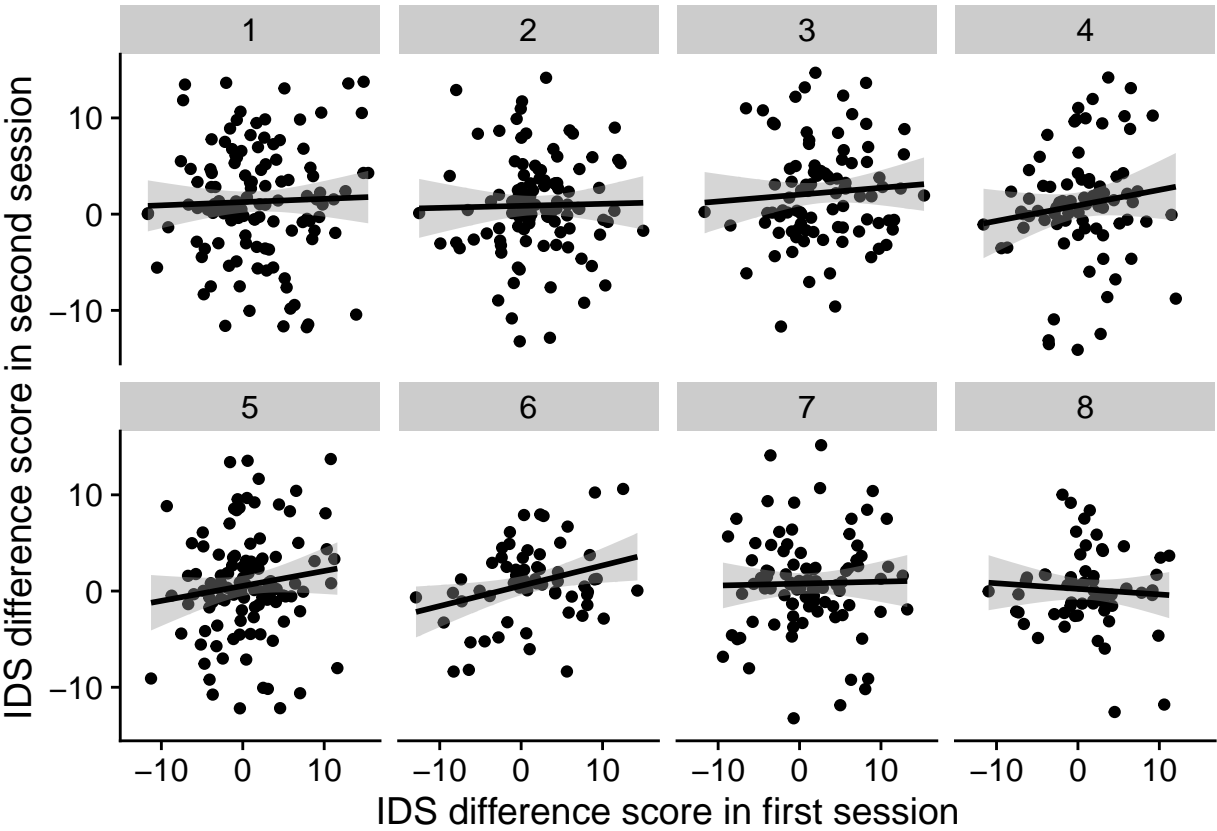


Table 2  
*Mixed-effects model results predicting looking time during session 1 from looking time at session 2 at the stimulus level.*

Term	$\hat{\beta}$	95% CI	$t$	$df$	$p$
Intercept	6.04	[4.99, 7.08]	11.35	6.88	< .001
LT Test	0.13	[0.05, 0.20]	3.46	25.38	.002

118 S5: By-item-pair preference scores across sessions



119

120

Table 3  
*Mixed-effects model results predicting IDS preference during session 1 from IDS preference at session 2 at the stimulus level.*

Term	$\hat{\beta}$	95% CI	$t$	$df$	$p$
Intercept	0.87	[0.45, 1.30]	4.04	122.79	< .001
Diff 1	0.10	[-0.02, 0.22]	1.63	6.31	.151

References

Byers-Heinlein, K., Bergmann, C., & Savalei, V. (2021). Six solutions for more reliable infant research. *Infant and Child Development*, e2296.

DeBolt, M. C., Rhemtulla, M., & Oakes, L. M. (2020). Robust data and power in infant research: A case study of the effect of number of infants and number of trials in visual preference procedures. *Infancy*, 25(4), 393–419.

ManyBabies Consortium. (2020). Quantifying sources of variability in infancy research using the infant-directed-speech preference. *Advances in Methods and Practices in Psychological Science*, 3(1), 24–52.