

Research Articles: Behavioral/Cognitive

# Language Exposure Relates to Structural Neural Connectivity in Childhood

Rachel R. Romeo<sup>1,2,3</sup>, Joshua Segaran<sup>2</sup>, Julia A. Leonard<sup>2</sup>, Sydney T. Robinson<sup>2</sup>, Martin R. West<sup>4</sup>, Allyson P. Mackey<sup>2,5</sup>, Anastasia Yendiki<sup>6</sup>, Meredith L. Rowe<sup>4</sup> and John D. E. Gabrieli<sup>2,4</sup>

DOI: 10.1523/JNEUROSCI.0484-18.2018

Received: 21 February 2018

Revised: 20 July 2018 Accepted: 22 July 2018

Published: 13 August 2018

**Author contributions:** R.R.R., J.A.L., M.R.W., A.P.M., M.L.R., and J.G. designed research; R.R.R., J.A.L., and S.T.R. performed research; R.R.R., J.S., A.Y., M.L.R., and J.G. analyzed data; R.R.R. wrote the first draft of the paper; R.R.R., J.A.L., M.R.W., A.P.M., A.Y., M.L.R., and J.G. edited the paper; R.R.R. wrote the paper.

Conflict of Interest: The authors declare no competing financial interests.

Research was funded by the Walton Family Foundation (to M.R.W.), National Institute of Child Health and Human Development (F31HD086957 to R.R.R.), Harvard Mind Brain Behavior Grant (to R.R.R.), and a gift from David Pun Chan (to J.D.E.G). We thank the Athinoula A. Martinos Imaging Center at the McGovern Institute for Brain Research (MIT); Atshusi Takahashi, Steve Shannon, and Sheeba Arnold for data collection support; Kelly Halverson, Emilia Motroni, Lauren Pesta, Veronica Wheaton, and Christina Yu for assistance in administering behavioral assessments; Megumi Takada for help with data collection/organization; Hannah Grotzinger for MRI quality assurance; Matthias Goncalves for data processing assistance; and Transforming Education, John Connolly and Glennys Sanchez from 1647 Families plus Ethan Scherer from the Boston Charter Research Collaborative for extensive recruitment support.

Correspondence should be addressed to Author to whom correspondence should be addressed: Rachel Romeo, MIT (Office 46-4037), 43 Vassar St, Cambridge, MA 02139, P: 617-324-6515, E: rromeo@mit.edu

Cite as: J. Neurosci; 10.1523/JNEUROSCI.0484-18.2018

Alerts: Sign up at www.jneurosci.org/cgi/alerts to receive customized email alerts when the fully formatted version of this article is published.

Accepted manuscripts are peer-reviewed but have not been through the copyediting, formatting, or proofreading process.

<sup>&</sup>lt;sup>1</sup>Division of Medical Sciences, Harvard University, 02115

<sup>&</sup>lt;sup>2</sup>Department of Brain and Cognitive Sciences and McGovern Institute for Brain Research, Massachusetts Institute of Technology, 02139

<sup>&</sup>lt;sup>3</sup>Boston Children's Hospital, Division of Developmental Medicine, 02215

<sup>&</sup>lt;sup>4</sup>Harvard University Graduate School of Education, 02138

<sup>&</sup>lt;sup>5</sup>Department of Psychology, University of Pennsylvania, 19104

<sup>&</sup>lt;sup>6</sup>Athinoula A. Martinos Center for Biomedical Imaging, Massachusetts General Hospital and Harvard Medical School, 02129

1						
2	Language Exposure Relates to Structural Neural Connectivity in Childhood					
3						
4	Rachel R. Romeo <sup>1,2,3</sup> *, Joshua Segaran, <sup>2</sup> Julia A. Leonard <sup>2</sup> , Sydney T. Robinson <sup>2</sup> ,					
5	Martin R. West <sup>4</sup> , Allyson P. Mackey <sup>2,5</sup> , Anastasia Yendiki <sup>6</sup> , Meredith L. Rowe <sup>4</sup> ,					
6	John D. E. Gabrieli <sup>2,4</sup>					
7						
8	<sup>1</sup> Division of Medical Sciences, Harvard University, 02115					
9	<sup>2</sup> Department of Brain and Cognitive Sciences and McGovern Institute for Brain					
10	Research, Massachusetts Institute of Technology, 02139					
11	<sup>3</sup> Boston Children's Hospital, Division of Developmental Medicine, 02215					
12	<sup>4</sup> Harvard University Graduate School of Education, 02138					
13	<sup>5</sup> Department of Psychology, University of Pennsylvania, 19104					
14	<sup>6</sup> Athinoula A. Martinos Center for Biomedical Imaging, Massachusetts General Hospital					
15	and Harvard Medical School, 02129					
16						
17	*Author to whom correspondence should be addressed:					
18	Rachel Romeo, MIT (Office 46-4037), 43 Vassar St, Cambridge, MA 02139					
19	P: 617-324-6515, E: <u>rromeo@mit.edu</u>					
20						
21	Number of Pages: 37 24 Abstract Word Count: 231					
22	Number of Figures: 4 25 Introduction Word Count: 543					
23	Number of Tables: 0 26 Discussion Word Count: 1498					

27 28	Conflict of Interest
29 30 31 32 33	The authors declare no competing financial interests.
34	Acknowledgements
35	Research was funded by the Walton Family Foundation (to M.R.W.), National Institute
36	of Child Health and Human Development (F31HD086957 to R.R.R.), Harvard Mind
37	Brain Behavior Grant (to R.R.R.), and a gift from David Pun Chan (to J.D.E.G). We
38	thank the Athinoula A. Martinos Imaging Center at the McGovern Institute for Brain
39	Research (MIT); Atshusi Takahashi, Steve Shannon, and Sheeba Arnold for data
40	collection support; Kelly Halverson, Emilia Motroni, Lauren Pesta, Veronica Wheaton,
41	and Christina Yu for assistance in administering behavioral assessments; Megumi
42	Takada for help with data collection/organization; Hannah Grotzinger for MRI quality
43	assurance; Matthias Goncalves for data processing assistance; and Transforming
44	Education, John Connolly and Glennys Sanchez from 1647 Families plus Ethan Scherer
45	from the Boston Charter Research Collaborative for extensive recruitment support.
46	

### 47 Abstract

18	Neuroscience research has elucidated broad relationships between socioeconomic
19	status (SES) and young children's brain structure, but there is little mechanistic
50	knowledge about specific environmental factors that are associated with specific
51	variation in brain structure. One environmental factor, early language exposure, predicts
52	children's linguistic and cognitive skills and later academic achievement, but how
53	language exposure relates to neuroanatomy is unknown. By measuring the real-world
54	language exposure of young children (ages 4-6 years, 27 male/13 female), we
55	confirmed the preregistered hypothesis that greater adult-child conversational
56	experience, independent of SES and the sheer amount of adult speech, is related to
57	stronger, more coherent white matter connectivity in the left arcuate and superior
58	longitudinal fasciculi on average, and specifically near their anterior termination at
59	Broca's area in left inferior frontal cortex. Fractional anisotropy of significant tract sub-
60	regions mediated the relationship between conversational turns and children's
61	language skills and indicated a neuroanatomical mechanism underlying the SES
62	"language gap." Post-hoc whole-brain analyses revealed that language exposure was
63	not related to any other white matter tracts, indicating the specificity of this
64	relationship. Results suggest that the development of dorsal language tracts is
65	environmentally influenced, specifically by early, dialogic interaction. Furthermore,
66	these findings raise the possibility that early intervention programs aiming to ameliorate
67	disadvantages in development due to family SES may focus on increasing children's
68	conversational exposure in order to capitalize on the early neural plasticity underlying
59	cognitive development.

### Significance Statement

Over the last decade, cognitive neuroscience has highlighted the detrimental impact of disadvantaged backgrounds on young children's brain structure. However, to intervene effectively, we must know which proximal aspects of the environmental aspects are most strongly related to neural development. The present study finds that young children's real-world language exposure, and specifically the amount of adult-child conversation, correlates with the strength of connectivity in the left hemisphere white matter pathway connecting two canonical language regions, *independent* of SES and the sheer volume of adult speech. These findings suggest that early intervention programs aiming to close the achievement gap may focus on increasing children's conversational exposure in order to capitalize on the early neural plasticity underlying cognitive development.

	II	n	tr	O.	d	u	С	ti	O	n
--	----	---	----	----	---	---	---	----	---	---

85

Socioeconomic status (SES) is a multifaceted index of one's financial resources, 86 87 educational capital, and relative social status. Neuroimaging studies have found 88 relatively consistent evidence that variation in SES is associated with variation in brain 89 development, including gray matter volume (Raizada et al., 2008; Jednorog et al., 2012; 90 Noble et al., 2012; Hanson et al., 2013; Luby et al., 2013), thickness (Lawson et al., 91 2013; Mackey et al., 2015; Romeo et al., 2017), and surface area (Noble et al., 2015), in 92 addition to white matter macrostructure (Raizada et al., 2008; Luby et al., 2013) and 93 microstructure (Gianaros et al., 2013; Ursache et al., 2016). Presumably these neural 94 disparities arise because of systematic differences in certain immediate environmental 95 factors during early childhood. There is, however, a paucity of evidence as to which 96 specific aspects of children's experiences are associated with individual variation in 97 specific neuroanatomical developments. 98 99 Behaviorally, it is well known that the quantity and quality of the language young 100 children are exposed to early in life predicts their later linguistic and cognitive skills 101 (Huttenlocher et al., 1991; Rodriguez and Tamis-LeMonda, 2011; Rowe, 2012; 102 Weisleder and Fernald, 2013; Hirsh-Pasek et al., 2015). Furthermore, children from 103 lower SES backgrounds are exposed to, on average, fewer utterances of lower 104 complexity than their higher-SES peers (Hoff et al., 2002; Rowe et al., 2005; 105 Huttenlocher et al., 2007). A seminal study estimated that by the time children reach 106 school age, children growing up in higher-SES families were, on average, exposed to

30 million more words than children growing up in lower-SES families (Hart and Risley, 1995).

Subsequent research has found that more important than the simple *quantity* of words heard is the *quality* of language exposure, including linguistic features such as vocabulary diversity and sophistication, grammatical complexity, and narrative use (Rowe, 2012), as well as interactional features such as contiguous (time-locked), contingent (topically similar), back-and-forth conversation (Hirsh-Pasek et al., 2015). Conversational turn-taking involves a rich experience of high quality linguistic, attentional, and social features. There is now some evidence that certain aspects of children's language environments relate to *functional* brain responses in prefrontal cortical regions (Sheridan et al., 2012; Garcia-Sierra et al., 2016; Romeo et al., 2018). However, there is no evidence as yet relating children's language exposure to their brain *structure*, including the white matter tracts that connect brain regions into networks.

The white matter tract most associated with language is the left arcuate fasciculus, a component of the superior longitudinal fasciculus (SLF) that connects two cortical regions critical for language: the left inferior frontal gyrus ("Broca's area") and the left posterior superior temporal gyrus ("Wernicke's area"). Microstructure of this tract has been associated with scores on language and literacy measures in children (Yeatman et al., 2011; Saygin et al., 2013; Skeide et al., 2016), and is often altered in both children and adults with disorders of speech, language, and/or literacy (Catani and Mesulam,

2008; Vandermosten et al., 2012a). Given the importance of this tract for language development, we tested the pre-registered hypothesis that early language experience—independent of SES—might be related to the microstructure of the left arcuate/superior longitudinal fasciculi; if true, this would suggest that these dorsal language tracts may be a neuroanatomical mechanism by which children's language environments affect their linguistic and cognitive skills.

### **Methods**

### **Experimental Design**

A priori hypotheses and exploratory analyses were pre-registered at <a href="https://osf.io/fes4i/register/564d31db8c5e4a7c9694b2be">https://osf.io/fes4i/register/564d31db8c5e4a7c9694b2be</a>. Specifically, the present study was designed to confirm or refute the hypothesis that young children's language exposure, and particularly the number of conversational turns with adults, would be positively correlated with the fractional anisotropy of the left arcuate/superior longitudinal fasciculi (and/or a portion thereof), independent of SES and the sheer quantity of adult and child speech alone. As such, this experiment aimed to recruit a socioeconomically diverse sample of young children and their parents to complete diffusion magnetic resonance imaging (dMRI), standardized cognitive assessments, and two full days of real-word auditory language recordings. All analyses were within-group correlations with specific covariates (nuisance and interest) as described below.

### **Participants**

Forty children (27 male) aged 4 years, 2 months to 6 years, 10 months (M = 5.78 years, SD = 0.72 years) and their parents completed this study. Children were in either pre-Kindergarten or Kindergarten grades, and were required to be native English speakers with no history of premature birth (< 37 weeks), neurological disorders, developmental delay, speech/language therapy, or grade repetition. Nineteen additional children were initially assessed and excluded for not meeting these inclusion criteria.

Twenty-three other children participated but did not have complete data sets, either because they did not complete the home recordings (n = 6), did not participate in the DTI scan (n = 7), or exhibited excessive movement during the DTI scan (n = 10, details below). Excluded participants did not differ from the included sample on age, SES, behavioral scores, or language exposure measures. However, the groups did differ on child gender; unintentionally, all home-recording non-completions occurred with female participants, so that girls were more likely to be excluded. Thus, all analyses control for gender. Additionally, half of the final sample additionally participated in a larger randomized controlled intervention study on parenting practices; only their baseline data (before learning of group assignment) was used here. Furthermore, task-based functional MRI results were previously reported for a partially overlapping subset of this sample (Romeo et al., 2018). Forty-four participants had either/both useable fMRI and DTI data; of these, 32 had both useable fMRI and DTI data, 4 had useable fMRI data only (for a final fMRI sample of 36), and 8 had useable DTI data only (for a final DTI sample of 40 for all analyses reported here). All procedures were approved by the

Institutional Review Board at the Massachusetts Institute of Technology, and written informed consent was obtained from parents.

#### Socioeconomic Measures

Participants were from a wide SES range. Parent(s) filled out a short questionnaire about total gross annual household income and the highest level of education obtained by each parent and/or primary caregiver (0 = less than high school, 1 = high school, 2 = some college/associate's degree, 3 = bachelor's degree, 4 = advanced degree). When two parents were present in the home, maternal and paternal years of education were averaged to create a parental education metric. For the final sample, parental education ranged from 0.5 to 4 (M = 2.81, Mdn = 3.50, SD = 1.17, Fig. 1), and gross household income ranged from \$6,000 to \$250,000 (M = \$108,728, Mdn = \$93,000, SD = \$69,064, Fig. 1), which is equivalent to the median family income of the Metro region from which participants were sampled (American Community Survey, 2016). For mediation analyses, education and income metrics were z-scored and averaged.

### **Standardized Behavioral Assessments**

Children completed standardized behavioral assessments to characterize verbal and nonverbal cognitive skills. A nonverbal composite score comprised the average of the age-normed standard scores from the Matrix Reasoning, Picture Memory, and Bug Search subtests of the Wechsler Preschool and Primary Scale of Intelligence, 4<sup>th</sup> edition (WPPSI-IV) (Wechsler, 2012). A verbal composite score comprised the average age-normed standard scores of the Peabody Picture Vocabulary Test (PPVT-4) (Dunn and

Dunn, 2007) and the Core Language Score of the Clinical Evaluation of Language Fundamentals, 5<sup>th</sup> edition (CELF-5; Wiig et al., 2013). To be included in the final sample, participants were required to score scores greater than or equal to one standard deviation below the mean (16th percentile) on both composite scores.

201

202

203

204

205

206

207

208

209

210

211

212

213

214

215

216

217

218

197

198

199

200

### **Neuroimaging Data Acquisition**

Neuroimaging sessions occurred at the Athinoula A. Martinos Imaging Center at the McGovern Institute for Brain Research, at the Massachusetts Institute of Technology. Children were acclimated to the MRI environment and practiced lying still in a mock MRI scanner before data acquisition on a 3 Tesla Siemens MAGNETOM Trio Tim scanner equipped for echo planar imaging (EPI; Siemens, Erlangen, Germany) with a 32channel phased array head coil. First, an automated scout image was acquired, and shimming procedures were performed to optimize field homogeneity. Then a wholehead, high-resolution T1-weighted multi-echo MPRAGE structural image was acquired using a protocol optimized for movement-prone pediatric populations (TR = 2530 ms, TE = 1.64 ms/3.5 ms/5.36 ms/7.22 ms, TI = 1400 ms, flip angle = 7°, resolution = 1 mm isotropic). Whole brain diffusion-weighted images were acquired in 74 axial interleaved slices of thickness 2mm and axial in-plane isotropic resolution 2mm (128×128×74 image matrix, TR = 9.3 s, TE = 84 ms, and GRAPPA acceleration factor 2). The series included 10 non-diffusion weighted reference volumes (b = 0) and 30 diffusion-weighted volumes ( $b = 700 \text{ s/mm}^2$ ). Resting state and one task-based functional scans were also collected in the same session, but are not reported here.

219

220	Code Accessibility
221	All code necessary to replicate results, along with links to necessary software packages
222	are freely available at <a href="https://github.com/rromeo2/openmindMIT">https://github.com/rromeo2/openmindMIT</a> .
223	
224	Neuroimaging Processing and Analysis
225	First, all diffusion data underwent quality control via visual inspection of all volumes
226	followed by the fully automated DTIPrep pipeline (Oguz et al., 2014), which corrects
227	artifacts caused by Eddy currents, head motion, bed vibration/pulsation, and slice-wise,
228	interlace-wise, and gradient-wise intensity inconsistencies. Participants with more than
229	5 unusable volumes (12.5%) were excluded (n = 10), leaving the final sample of 40
230	participants.
231	
232	All preprocessing was implemented via a custom script in Nipype version 0.13.0
233	(Gorgolewski et al., 2011). All images in the diffusion series were aligned to the first
234	non-diffusion-weighted image using affine registration, and corresponding diffusion-
235	weighting gradient vectors were reoriented accordingly, in order to reduce
236	misalignment. A per-subject total head motion index (TMI) was computed from volume-
237	by-volume translation and rotation, percentage of slices with signal dropout, and signal
238	drop-out severity (Yendiki et al., 2014). All analyses statistically control for the TMI.
239	
240	Eighteen major white matter fascicles were automatically reconstructed using
241	TRACULA implemented in FreeSurfer version 6.0 (Yendiki et al., 2011), which uses
242	global probabilistic tractography and the ball-and-stick model of diffusion to estimate the

243

244

245

246

247

248

249

250

251

252

253

254

255

256

257

258

259

260

261

262

263

264

265

### LANGUAGE EXPOSURE AND CONNECTIVITY

posterior probability distribution of each pathway. This distribution includes the prior probabilities of the pathway given the cortical parcellation and subcortical segmentation of the anatomical image, which had been processed and manually edited as necessary in FreeSurfer (version 5.3.0; Fischl, 2012) to ensure correct gray and white matter boundaries. Each pathway distribution was thresholded at 20% of the maximum value, and the values at each voxel in the pathway were weighted by the pathway probability at that voxel in order to obtain whole-tract average measures of microstructure. Of interest were three measures of water diffusion within tracts: axial diffusivity (AD), which measures the rate of diffusion parallel to the tract; radial diffusivity (RD), which measures the rate of diffusion perpendicular to the tract; and fractional anisotropy (FA), a summary measure of microstructural organization that indexes the overall strength and directionality of diffusion (Lebel et al., 2017). These measures were analyzed within two a priori components of the left SLF: the arcuate fasciculus, which runs between inferior frontal and superior posterior temporal regions (roughly corresponding to SLF II), and SLF III, which runs between inferior frontal and inferior parietal regions<sup>25</sup> (henceforth referred to as SLF). TRACULA was also used to calculate FA at successive cross-sections as a function of position along the trajectory of both tracts in an anterior-to-posterior direction. Correspondence of nodes across subjects was based on the Euclidean distance in MNI

space. Because tracts were reconstructed in each subject's native space and not in a

template space, individual participants' tracts were of varying length. For participants

with shorter tracts, tail FA values were extrapolated by calculating moving averages of the previous 3 points in order to ensure uniform length (35 points along the superior longitudinal fasciculus and 48 points along the arcuate fasciculus). The presented results do not change if instead no extrapolations were made.

Finally, whole-brain voxel-wise statistical analysis was conducted with Tract-Based Spatial Statistics (TBSS)(Smith et al., 2006), as implemented in FSL version 5.0.9 (Jenkinson et al., 2012). Diffusion space FA images were aligned to each participant's anatomical image using boundary-based registration (BBR)(Greve and Fischl, 2009), which was then affine aligned to MNI space. Each subject's MNI-space image was eroded to remove the highly variable lateral regions of the FA map. The images were averaged to generate an inter-subject FA skeleton, and each voxel from participants' FA volumes were projected onto the FA skeleton. Voxel-wise regression analyses were conducted with FSL's randomise tool with 5,000 permutations, and threshold free-cluster enhancement (TFCE) was used to correct for multiple comparisons with p < 0.05 (Smith and Nichols, 2009). Significant voxels were then back-projected from skeleton positions to the position at the center of the nearest tract in the subject's FA image in standard space. These points were then inversely warped to each subject's native diffusion space for localization within the probabilistic tractography.

### **Home Audio Recordings**

Specific details of the home audio recordings have been previously reported (Romeo et al., 2018). Briefly, parents recorded two consecutive weekend days of audio from the

child's perspective via the Language Environmental Analysis (LENA) Pro system (Gilkerson et al., 2017). LENA software automatically processes the recordings and estimates the number of words spoken by an adult in the child's vicinity ("adult words), the number of utterances the key child made ("child utterances"), and the number of dyadic conversational turns, defined as a discrete pair of consecutive adult and child utterances in any order, with no more than 5 seconds of separation ("conversational turns"). As such, conversational turns measure the contiguous, linguistic interaction between children and adults. Running totals for each speech category were calculated for each consecutive 60 minutes across the two days in 5-minute increments (e.g., 7:00 AM – 8:00 AM, 7:05 AM – 8:05 AM, etc.), and the per-participant highest hourly total of adult words, child utterances, and conversational turns were separately extracted for statistical analysis. This metric helped minimize differences in language measures due solely to different recording lengths and/or loud activities that may have masked speech and misrepresented language input.

#### Statistical Analysis

Statistical analysis of behavioral and summary diffusion measures was executed in SPSS Statistics version 24 (IBM Corp., 2016). Given that all participants constituted a single group and all independent and dependent variables were continuous, all relational analyses were two-tailed regressions, reporting Pearson's r (if no covariates) or partial r (with covariates listed in results). For the node analysis within tracts, independent regressions with listed covariates were conducted with FA at each node as

the dependent variable, and p-values were FDR corrected for the total number of nodes in both tracts (n = 83).

Mean FA was extracted from the significant TBSS cluster and entered into two bootstrapped mediation analyses (controlling for age, gender, and motion) with 5000 repetitions, as executed in the PROCESS macro (Hayes, 2018; Preacher & Hayes, 2004). In the first model, the number of conversational turns was the independent variable, composite verbal score was the dependent variable, and cluster FA was the mediator. In the second model, composite SES was the independent variable, composite verbal score was the dependent variable, and both conversational turns and cluster FA were entered as mediators. The bootstrapped 95% confidence intervals for the direct (*c*) and indirect (*ab*) effects are reported; the mediation was considered "significant" if the 95% confidence interval for the indirect effect did not contain 0. Effect sizes were determined by the mediation ratio, which is the ratio of the indirect effect coefficient to the total effect coefficient; this measure indicates the proportion of the total effect that is mediated.

### Results

Replicating prior studies, higher SES was strongly correlated with higher composite verbal scores (education: r(38) = 0.65,  $p = 5 \times 10^{-6}$ ; income: r(38) = 0.46, p = 0.003) and to a lesser extent, with higher composite nonverbal scores (education: r(38) = 0.35, p = 0.03; income: r(38) = 0.16, p = n.s.). SES was also positively correlated with measures of language exposure, including adult words (education: r(38) = 0.41, p = 0.008;

334	income: $r(38) = 0.28$ , $p = 0.08$ ) and conversational turns (education: $r(38) = 0.38$ , $p = 0.08$ )
335	0.02; income: $r(38) = 0.40$ , $p = 0.01$ ), but not child utterances alone (both $r(38) < 0.27$ ,
336	both $p > 0.10$ ). After controlling for SES (parental education and income), the number of
337	conversational turns was the only exposure measure that correlated with children's
338	composite verbal scores (partial $r(36) = 0.51$ , $p = 0.001$ ; adult words: partial $r(36) =$
339	0.08, $p = 0.65$ ; partial $r(36) = 0.10$ , $p = 0.57$ ), indicating that differences in
340	conversational exposure relate to variance in children's language skills over and above
341	socioeconomic disparities. Nonverbal scores were not related to any of the language
342	exposure measures (all $r(38) <  0.18 $ , all $p > 0.2$ ).
343	
344	Controlling for age, gender, and head motion, neither the number of adult words nor the
345	number of child utterances were correlated with any diffusion measure in either the
346	arcuate or SLF (all partial $r(35) < 0.17$ , all $p > 0.32$ ). However, the number of
347	conversational turns correlated positively with FA (arcuate: partial $r(35) = 0.46$ , $p =$
348	0.004; SLF: partial $r(35) = 0.45$ , $p = 0.005$ ; Fig. 2) and negatively with RD (arcuate:
349	partial $r(35) = -0.34$ , $p = 0.04$ ; SLF: partial $r(35) = -0.37$ , $p = 0.02$ ), but did not correlate
350	with AD (both partial $r(35)$ < abs(0.07), both $p > 0.70$ ). Combined, these measures
351	indicate that greater conversational turns correspond with greater coherence of diffusion
352	parallel to the tract, which may be a marker of greater axonal myelination (Lebel et al.,
353	2017). Importantly, the relationships between conversational turns and FA/RD remained
354	significant when controlling for potential confounding variables of SES, the two other
355	LENA measures, or composite language scores. Specifically: controlling for SES
356	(arcuate FA partial $r(33) = 0.48$ , $p = 0.003$ ; SLF FA partial $r(33) = 0.45$ , $p = 0.007$ ;

357	arcuate RD partial $r(33) = -0.37 p = 0.03$ ; SLF RD partial $r(33) = -0.36$ , $p = 0.04$ );
358	controlling for the two other LENA measures (arcuate FA partial $r(33) = 0.46$ , $p = 0.005$ ;
359	SLF FA partial $r(33) = 0.42$ , $p = 0.01$ ; arcuate RD partial $r(33) = -0.35$ $p = 0.04$ ; SLF RD
360	partial $r(33) = -0.37$ , $p = 0.03$ ); and controlling for children's composite language scores
361	(arcuate FA partial $r(34) = 0.46$ , $p = 0.005$ ; SLF FA partial $r(34) = 0.35$ , $p = 0.038$ ;
362	arcuate RD partial $r(34) = -0.349$ , $p = 0.037$ ; SLF RD partial $r(34) = -0.268$ , $p = 0.114$ ).
363	These findings indicate that the relations between conversational turns and SLF
364	microstructure cannot be explained by these other child-level or environmental
365	variables.
366	
367	A node analysis was conducted to explore whether a specific sub-location within these
368	tracts was driving observed relationships. Controlling for age, gender, motion, and SES,
369	25 (of 83) nodes exhibited significant correlations (FDR-corrected $p < 0.05$ ) between
370	conversational turns and local FA; these nodes occurred in four clusters located toward
371	both the anterior and posterior ends of the left arcuate and SLF (Fig. 3), suggesting that
372	the strong correlations in these regions drive the relation between conversational turns
373	and whole tract averages.
374	
375	Finally, post-hoc analyses aimed to ascertain the anatomical specificity of these
376	correlations across all white matter tracts. Correlations between conversational turns
377	and all 18 TRACULA-defined tracts revealed no significant correlations with any tracts
378	other than left arcuate and left SLF (all FDR-corrected $p > 0.2$ ). Additionally, a whole-
379	brain, voxel-wise analysis with TBSS (controlling for age, gender, and motion) revealed

that, convergent with the node analysis, the number of conversational turns was positively correlated (TFCE corrected p < 0.05) with FA in a cluster of 513 voxels at the anterior end of the left arcuate/SLF where these tracts terminate with Broca's area in the left inferior frontal gyrus (Fig. 4). To confirm localization in each participant's native space, back-projection revealed that the maximally significant voxel of this cluster occurred within the TRACULA-defined bounds of the intertwining arcuate/SLF near the anterior termination.

The average FA from this cluster was extracted for mediation analyses so as to better characterize the relationship between early language experience, white matter microstructure, and language skill. Controlling for age, gender, and motion, FA in the left anterior arcuate/SLF significantly mediated the relationship between conversational turns and the composite language score (direct effect = 0.095 [95% CI = 0.022-0.169], indirect effect = 0.043 [95% CI = 0.002-0.100], indirect/total effect = 0.311), such that variation in regional FA accounted for 31% of the total relationship between language experience and language skill. Furthermore, conversational turns and FA jointly mediated the relationship between SES and language scores, (direct effect = 7.134 [95% CI = 3.057-11.210], indirect effect = 3.007 [95% CI = 0.680-5.829], indirect/total effect = 3.007, indicating that combined behavioral and neural mechanisms explained nearly a third (30%) of the socioeconomic "language gap." FA in this region was not significantly related to nonverbal scores (r(35) = 0.117, p = 0.49).

### **Discussion**

These results provide the first evidence of direct association between a specific aspect of children's language experience, namely adult-child conversational turns, and particular neuroanatomical structural properties, specifically the connectivity of the left arcuate and the left superior longitudinal fasciculi. The number of adult-child conversational turns young children experienced, independent of SES, was positively correlated with the strength of coherence of two dorsal white matter tracts: the left arcuate fasciculus and the left superior longitudinal fasciculus. This relationship appeared to be driven by anisotropy in a sub-region near where these tracts terminate in the left inferior frontal lobe at a known hub for expressive and receptive language processing (Friederici, 2012). Mediation models revealed that microstructural properties in this region provide a neural mechanism underlying the relationship between children's conversational exposure and their language skills.

This localization is consistent with functional findings that children's language exposure is related to activation specifically in left prefrontal cortical regions (Sheridan et al., 2012; Garcia-Sierra et al., 2016; Romeo et al., 2018). Together this suggests that "Broca's area" and adjacent pathways may be components of the perisylvian language network that are particularly sensitive to early linguistic input, especially dialogic conversation. Because the arcuate fasciculus bidirectionally connects Broca's area to primary receptive language regions in superior posterior temporal cortex, this uniquely human tract may be evolutionarily specialized for language (Rilling et al., 2008), as evidenced by correlations between language skill and structural properties of the left arcuate. Classically, damage to the arcuate fasciculus is associated with conduction

aphasia (Catani and Mesulam, 2008). Further, individual microstructural variation in the absence of overt damage is related to a number of linguistic skills in childhood, including phonological knowledge and literacy skills (Yeatman et al., 2011; Saygin et al., 2013), presence or risk for developmental dyslexia (Vandermosten et al., 2012a; Langer et al., 2017; Wang et al., 2017), rate of vocabulary growth (Su et al., 2018), as well as word learning (Lopez-Barroso et al., 2013), verbal memory (Catani et al., 2007), and speech perception (Vandermosten et al., 2012b) in adulthood. In all cases, greater coherence in the left arcuate fasciculus reflected better linguistic skills, suggesting that that fast, efficient connectivity between frontal and temporal areas facilitates verbal skills throughout the lifespan. The present results further suggest that variation in early childhood language experience may underlie individual differences in neuroanatomy and behavior.

The apparent environmental influence of conversational turn-taking on left arcuate and superior longitudinal microstructure is congruent with findings that dorsal language tracts (superior longitudinal and arcuate fasciculi) develop more slowly than their ventral counterparts (inferior longitudinal, inferior-frontal-occipital, and uncinate fasciculi) (Perani et al., 2011; Brauer et al., 2013). Specifically, the terminal projection of the arcuate fasciculus at the furthest anterior point near Broca's area is the latest developing component of the dorsal pathway, which is still not fully mature at age seven years (Brauer et al., 2013). As such, this period of protracted development in early and middle childhood may correspond to a sensitive period of neurodevelopment in which children's anterior dorsal language circuitry is highly susceptible to their environments.

The present finding that conversational exposure correlated positively with FA and negatively with RD in the left arcuate and superior longitudinal fasciculi indicates greater coherence of diffusion parallel to the tracts, which is often considered a marker of greater axonal myelination (Lebel et al., 2017). Considering that myelination increases throughout childhood and early adulthood (Miller et al., 2012), these findings suggest that increased conversational exposure in early childhood might advance maturation of the anterior terminations of the dorsal language pathways important for language processing. However, longitudinal studies of children are necessary to determine precise developmental trajectories in relation to language exposure.

Localization of white-matter microstructural associations with conversational turns was specific to white matter near Broca's area, but such localization is related partially to methods and limitations of neuroimaging. No other tract or region was significantly related to conversational turns in either the TRACULA or TBSS analyses. Weaker associations would not be detected if they were below the statistical thresholds employed in the present study. As in any thresholded neuroimaging study, the conclusion that white-matter microstructure near Broca's area is associated with language experience is more certain than the conclusion that no other white-matter area is more weakly associated with such exposure.

In regards to language exposure, dorsal pathway microstructure was related only to the quantity of dialogic adult-child conversational turns, and not to the sheer volume of

speech spoken in the child's presence. Conversational turns incorporate social interactional features, such as contiguity (temporal connectedness), contingency (contextual relevancy), and joint attention, beyond simple linguistic features of the spoken content. The specificity of the relation between conversational turns and white-matter microstructure further supports the idea that *qualitative* aspects of children's early language experience, as opposed to sheer *quantitative* aspects, may be the largest influence on children's language development (Zimmerman et al., 2009; Rowe, 2012; Roseberry et al., 2014; Hirsh-Pasek et al., 2015). The present findings suggest that neuroanatomical maturation and concomitant language development may critically rely on social exchanges of linguistic information rather than purely passive speech exposure or child speech production in isolation. Developmental models have argued that social interaction is a necessary precursor to language acquisition, perhaps because language may rely on evolutionarily older social neurocircuitry (e.g., Golinkoff et al., 2015; Kuhl, 2007), and the present findings contribute neuroanatomical evidence in favor of such models.

A limitation of this study is the correlational nature of the analyses, which applies to nearly all studies of SES differences as well as most neuroimaging studies comparing groups of people. There is, however, behavioral evidence that experimental manipulation of children's language environment contributes to changes in their language development (Leech, Wei, Harring, & Rowe, 2018; McGillion, Pine, Herbert, & Matthews, 2017; Suskind et al., 2016; Windsor, Moraru, Nelson, Fox, & Zeanah, 2013; Windsor et al., 2011). Further, in absence of an intervention, the relative quantity and

quality of parents' speech to children is remarkably consistent throughout early childhood (Huttenlocher, Vasilyeva, Waterfall, Vevea, & Hedges, 2007). Thus, although we measured only a limited sample of home language, it is likely that the neural and language variation across children reflected years of differential home language experience.

Several models have addressed how early cognitive stimulation, such as language exposure, may contribute to cognitive development. While some suggest that linguistic experience may uniquely contribute to language domains (e.g., Johnson, Riis, & Noble, 2016), others argue that early language interaction may contribute to other aspects of cognition more broadly (e.g., McLaughlin, Sheridan, Nelson, 2017). Although the present study did not find relationships between nonverbal cognition (operationalized as fluid reasoning, working memory, and processing speed) and either language experience or left dorsal language tracts, this does not necessarily mean that language experience solely affects verbal domains. It is possible that language exposure directly relates to other nonverbal domains such as executive functioning or spatial reasoning. It is also possible that language exposure indirectly influences nonverbal cognition at older ages via language skills at younger ages (Noble, McCandliss, & Farah, 2007; Noble, Norman, & Farah, 2005). More comprehensive, longitudinal studies are necessary to tease out the direct and indirect influences of early language experience on multiple domains of cognition throughout childhood and adolescence.

The present findings highlight the specific role that conversational turns may play in a particular aspect of brain development above and beyond SES. There are multiple studies reporting correlations between SES and brain structure and function (for review, see Farah, 2017). Crucially, in the present study, the relation between conversational turns and white-matter remained significant after SES was statistically controlled for. This implies that the critical environmental correlate was not SES per se, but rather conversational turns at any level of SES. Although higher SES was in general associated with more conversational turns, the apparent influence of conversational turns on white-matter microstructure occurred independent of SES.

The present results may also have practical implications. Community-based intervention programs designed to close the SES "word gap" have often focused on closing this gap by increasing the quantity of speech that low-SES parents direct toward children (Cartmill, 2016). However, the present results build on previous behavioral findings that the *quality* of language—specifically conversational interaction—is more strongly linked to children's behavioral outcomes by revealing that this same quality is associated with white-matter development in children's language brain circuitry. This suggests that early intervention programs should not only encourage parents to talk *to* their children, but to talk *with* their children to promote optimal brain development. Further research is needed to determine if enrichment of the language environment in at-risk children could reduce the measurable socioeconomic disparities in academic achievement and brain development (Mackey et al., 2015; Noble et al., 2015; Johnson et al., 2016). More generally, the finding that more conversational turns are associated with more coherent

540	white-matter connectivity independent of SES indicates that promoting such
541	conversational turns may enhance structural brain development and the language
542	abilities supported by that brain development in children from all backgrounds.
543	

544	Author Contributions
545	R. R. Romeo and J. D. E. Gabrieli developed the study concept. R. R. Romeo, J. A.
546	Leonard, A. P. Mackey, M. L. Rowe, and J. D. E. Gabrieli designed the study. R. R.
547	Romeo, J. A. Leonard, and S. T. Robinson collected the data. R. R. Romeo and J.
548	Segaran performed the data analysis and interpretation under the supervision of J. D. E
549	Gabrieli and M. L. Rowe. R. R. Romeo and J. D. E. Gabrieli wrote the manuscript with
550	suggestions from all other authors. All authors approved the final version of the
551	manuscript for submission.
552	

553	References
554	
555	American Community Survey (2016).
556 557	Brauer J, Anwander A, Perani D, Friederici AD (2013) Dorsal and ventral pathways in language development. Brain Lang 127:289-295.
558	Cartmill EA (2016) Mind the Gap. Policy Insights Behav Brain Sci 3:185-193.
559 560	Catani M, Mesulam M (2008) The arcuate fasciculus and the disconnection theme in language and aphasia: History and current state. Cortex 44:953-961.
561 562 563	Catani M, Allin MPG, Husain M, Pugliese L, Mesulam MM, Murray RM, Jones DK (2007) Symmetries in human brain language pathways correlate with verbal recall. Proc Natl Acad Sci USA 104:17163-17168.
564 565	Dunn LM, Dunn DM (2007) Peabody picture vocabulary test, 4th Edition. Bloomington, MN: Pearson.
566 567	Farah MJ (2017) The neuroscience of socioeconomic status: Correlates, causes, and consequences. Neuron 96:56-71.
568	Fischl B (2012) FreeSurfer. Neuroimage 62:774-781.
569 570	Friederici AD (2012) The cortical language circuit: From auditory perception to sentence comprehension. Trends Cogn Sci 16:262-268.
571 572 573	Garcia-Sierra A, Ramírez-Esparza N, Kuhl PK (2016) Relationships between quantity of language input and brain responses in bilingual and monolingual infants. Int J Psychophysiol 110:1-17.
574 575	Gianaros PJ, Marsland AL, Sheu LK, Erickson KI, Verstynen TD (2013) Inflammatory pathways link socioeconomic inequalities to white matter architecture. Cereb Cortex 23:2058-2071.
576 577 578	Gilkerson J, Richards JA, Warren SF, Montgomery JK, Greenwood CR, Kimbrough Oller D, Hansen JHL, Paul TD (2017) Mapping the early language environment using all-day recordings and automated analysis. Am J Speech Lang Pathol 26:248-265.
579 580	Golinkoff RM, Can DD, Soderstrom M, Hirsh-Pasek K (2015) (Baby)talk to me. Curr Dir Psychol Sci 24:339-344.
581 582 583	Gorgolewski K, Burns CD, Madison C, Clark D, Halchenko YO, Waskom ML, Ghosh SS (2011) Nipype: A flexible, lightweight and extensible neuroimaging data processing framework. Front Neuroinform 5:1-15.
584 585	Greve DN, Fischl B (2009) Accurate and robust brain image alignment using boundary-based registration. Neuroimage 48:63-72.

586 587	Hanson JL, Hair N, Shen DG, Shi F, Gilmore JH, Wolfe BL, Pollak SD (2013) Family poverty affects the rate of human infant brain growth. PLoS ONE 8:e80954.
588 589	Hart B, Risley T (1995) Meaningful differences in the everyday experience of young American children. Baltimore, MD: P.H. Brookes.
590 591 592	Hirsh-Pasek K, Adamson LB, Bakeman R, Owen MT, Golinkoff RM, Pace A, Yust PKS, Suma K (2015) The contribution of early communication quality to low-income children's language success. Psychol Sci 26:1071-1083.
593 594	Hayes, A. F. (2018). <i>Introduction to mediation, moderation, and conditional process analysis: A regression-based perspective</i> (2nd ed.). New York, NY: The Guilford Press.
595 596 597	Hoff E, Laursen B, Tardif T (2002) Socioeconomic status and parenting. In: Handbook of parenting, 2 Edition (Bornstein MH, ed), pp 231-252. Mahway, NJ: Lawrence Erlbaum Associates.
598 599	Huttenlocher J, Haight W, Bryk A, Seltzer M, Lyons T (1991) Early vocabulary growth: Relation to language input and gender. Dev Psychol 27:236-248.
600 601	Huttenlocher J, Vasilyeva M, Waterfall HR, Vevea JL, Hedges LV (2007) The varieties of speech to young children. Dev Psychol 43:1062-1083.
602	IBM Corp. (2016) IBM SPSS Statistics, Version 24.0. Armonk, NY: IBM Corp.
603 604 605	Jednorog K, Altarelli I, Monzalvo K, Fluss J, Dubois J, Billard C, Dehaene-Lambertz G, Ramus F (2012) The influence of socioeconomic status on children's brain structure. PLoS One 7:e42486.
606 607	Jenkinson M, Beckmann CF, Behrens TE, Woolrich MW, Smith SM (2012) FSL. Neuroimage 62:782-790.
608 609	Johnson SB, Riis JL, Noble KG (2016) State of the art review: Poverty and the developing brain Pediatrics 137:e20153075.
610	Kuhl PK (2007) Is speech learning 'gated' by the social brain? Dev Sci 10:110-120.
611 612 613	Langer N, Peysakhovich B, Zuk J, Drottar M, Sliva DD, Smith S, Becker BL, Grant PE, Gaab N (2017) White matter alterations in infants at risk for developmental dyslexia. Cereb Cortex 27:1027-1036.
614 615	Lawson GM, Duda JT, Avants BB, Wu J, Farah MJ (2013) Associations between children's socioeconomic status and prefrontal cortical thickness. Dev Sci 16:641-652.
616 617 618	Lebel C, Treit S, Beaulieu C (2017) A review of diffusion MRI of typical white matter development from early childhood to young adulthood. NMR Biomed epub ahead of print.

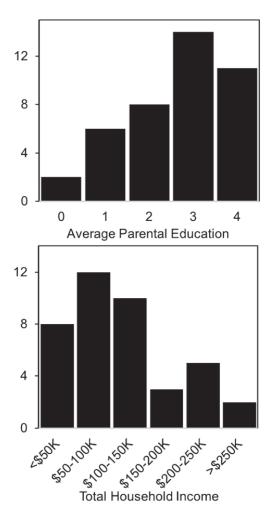
619 620	Leech K, Wei R, Harring JR, Rowe ML (2017) A brief parent-focused intervention to improve preschoolers' conversational skills and school readiness. Dev Psychol 54:15-28.
621 622 623	Lopez-Barroso D, Catani M, Ripolles P, Dell'Acqua F, Rodriguez-Fornells A, de Diego-Balaguer R (2013) Word learning is mediated by the left arcuate fasciculus. Proc Natl Acad Sci USA 110:13168-13173.
624 625 626	Luby J, Belden A, Botteron K, Marrus N, Harms MP, Babb C, Nishino T, Barch D (2013) The effects of poverty on childhood brain development: the mediating effect of caregiving and stressful life events. JAMA Pediatr 167:1135-1142.
627 628 629	Mackey AP, Finn AS, Leonard JA, Jacoby-Senghor DS, West MR, Gabrieli CF, Gabrieli JD (2015) Neuroanatomical correlates of the income-achievement gap. Psychol Sci 26:925-933.
630 631 632	McGillion M, Pine JM, Herbert JS, Matthews D (2017) A randomised controlled trial to test the effect of promoting caregiver contingent talk on language development in infants from diverse socioeconomic status backgrounds. J Child Psychol Psychiatry 58:1122–1131.
633 634	McLaughlin KA, Sheridan MA, Nelson CA (2017) Neglect as a violation of species-expectant experience: Neurodevelopmental consequences. Biol Psychiatry 82:462-471.
635 636 637 638	Miller DJ, Duka T, Stimpson CD, Schapiro SJ, Baze WB, McArthur MJ, Fobbs AJ, Sousa AMM, Šestan N, Wildman DE, Lipovich L, Kuzawa CW, Hof PR, Sherwood CC (2012) Prolonged myelination in human neocortical evolution. Proc Natl Acad Sci USA 109:16480-16485.
639 640	Noble KG et al. (2015) Family income, parental education and brain structure in children and adolescents. Nat Neurosci 18:773-778.
641 642	Noble KG, Houston SM, Kan E, Sowell ER (2012) Neural correlates of socioeconomic status in the developing human brain. Dev Sci 15:516-527.
643 644	Noble KG, McCandliss BD, Farah MJ (2007) Socioeconomic gradients predict individual differences in neurocognitive abilities. Dev Sci 10:464-480.
645 646	Noble KG, Norman MF, Farah MJ (2005) Neurocognitive correlates of socioeconomic status in kindergarten children. Dev Sci 8:74-87.
647 648	Oguz I, Farzinfar M, Matsui J, Budin F, Liu Z, Gerig G, Johnson H, Styner M (2014) DTIPrep: Quality control of diffusion-weighted images. Front Neuroinform 8.
649 650 651	Perani D, Saccuman MC, Scifo P, Anwander A, Spada D, Baldoli C, Poloniato A, Lohmann G, Friederici A, D. (2011) Neural language networks at birth. Proc Natl Acad Sci USA 108:16056–16061.

652 653 654	Preacher, K. J., & Hayes, A. F. (2004). SPSS and SAS procedures for estimating indirect effects in simple mediation models. <i>Behavior Research Methods, Instruments, &amp; Computers</i> , 36(4), 717-731.
655 656 657	Raizada RD, Richards TL, Meltzoff A, Kuhl PK (2008) Socioeconomic status predicts hemispheric specialisation of the left inferior frontal gyrus in young children. Neuroimage 40:1392-1401.
658 659	Rilling JK, Glasser MF, Preuss TM, Ma X, Zhao T, Hu X, Behrens TE (2008) The evolution of the arcuate fasciculus revealed with comparative DTI. Nat Neurosci 11:426-428.
660 661 662	Rodriguez ET, Tamis-LeMonda CS (2011) Trajectories of the home learning environment across the first 5 years: Associations with children's vocabulary and literacy skills at prekindergarten. Child Dev 82:1058-1075.
663 664 665	Romeo RR, Leonard JA, Robinson ST, West MR, Mackey AP, Rowe ML, Gabrieli JDE (2018) Beyond the "30 million word gap:" Children's conversational exposure is associated with language-related brain function. Psychol Sci 29:700-710.
666 667 668	Romeo RR, Christodoulou JA, Halverson KK, Murtagh J, Cyr AB, Schimmel C, Chang P, Hook PE, Gabrieli JDE (2017) Socioeconomic status and reading disability: Neuroanatomy and plasticity in response to intervention. Cereb Cortex 28:2297-2312.
669 670	Roseberry S, Hirsh-Pasek K, Golinkoff RM (2014) Skype me! Socially contingent interactions help toddlers learn language. Child Dev 85:956-970.
671 672	Rowe ML (2012) A longitudinal investigation of the role of quantity and quality of child-directed speech in vocabulary development. Child Dev 83:1762-1774.
673 674	Rowe ML, Pan BA, Ayoub C (2005) Predictors of variation in maternal talk to children: A longitudinal study of low-income families. Parent Sci Pract 5:285-310.
675 676 677 678	Saygin ZM, Norton ES, Osher DE, Beach SD, Cyr AB, Ozernov-Palchik O, Yendiki A, Fischl B, Gaab N, Gabrieli JD (2013) Tracking the roots of reading ability: white matter volume and integrity correlate with phonological awareness in prereading and early-reading kindergarten children. J Neurosci 33:13251-13258.
679 680	Sheridan MA, Sarsour K, Jutte D, D'Esposito M, Boyce WT (2012) The impact of social disparity on prefrontal function in childhood. PLoS One 7:e35744.
681 682	Skeide MA, Brauer J, Friederici AD (2016) Brain functional and structural predictors of language performance. Cereb Cortex 26:2127-2139.
683 684 685	Smith SM, Nichols TE (2009) Threshold-free cluster enhancement: addressing problems of smoothing, threshold dependence and localisation in cluster inference. Neuroimage 44:83-98.

686 687 688	Smith SM, Jenkinson M, Johansen-Berg H, Rueckert D, Nichols TE, Mackay CE, Watkins KE, Ciccarelli O, Cader MZ, Matthews PM, Behrens TEJ (2006) Tract-based spatial statistics Voxelwise analysis of multi-subject diffusion data. Neuroimage 31:1487-1505.
689 690 691	Su M, Thiebaut de Schotten M, Zhao J, Song S, Zhou W, Gong G, McBride C, Ramus F, Shu H (2018) Vocabulary growth rate from preschool to school-age years is reflected in the connectivity of the arcuate fasciculus in 14-year-old children. Dev Sci.
692 693 694 695	Suskind DL, Leffel KR, Graf E, Hernandez MW, Gunderson EA, Sapolich SG, Suskind E, Leininger L, Goldin-Meadow S, Levine SC (2016) A parent-directed language intervention for children of low socioeconomic status: A randomized controlled pilot study. J Child Lang 43:366-406.
696 697	Ursache A, Noble KG, PING Study (2016) Socioeconomic status, white matter, and executive function in children. Brain Behav 6:e00531.
698 699 700	Vandermosten M, Boets B, Wouters J, Ghesquiere P (2012a) A qualitative and quantitative review of diffusion tensor imaging studies in reading and dyslexia. Neurosci Biobehav Rev 36:1532-1552.
701 702 703	Vandermosten M, Boets B, Poelmans H, Sunaert S, Wouters J, Ghesquiere P (2012b) A tractography study in dyslexia: neuroanatomic correlates of orthographic, phonological and speech processing. Brain 135:935-948.
704 705 706	Wang Y, Mauer MV, Raney T, Peysakhovich B, Becker BLC, Sliva DD, Gaab N (2017) Development of tract-specific white matter pathways during early reading development in at-risk children and typical controls. Cereb Cortex 27:2469-2485.
707 708	Wechsler D (2012) Wechsler preschool and primary scale of intelligence, 4th Edition. Bloomington, MN: Pearson.
709 710	Weisleder A, Fernald A (2013) Talking to children matters: Early language experience strengthens processing and builds vocabulary. Psychol Sci 24:2143-2152.
711 712	Wiig EH, Semel EM, Secord W (2013) Clinical evaluation of language fundamentals, 5th Edition. Bloomington, MN: Pearson.
713 714	Windsor J, Benigno JP, Wing CA, Carroll PJ, Koga SF, Nelson CA, Fox NA, Zeanah CH (2011) Effect of foster care on young children's language learning. Child Dev 82:1040-1046.
715 716 717	Windsor J, Moraru A, Nelson CA, Fox NA, Zeanah CH (2013) Effect of foster care on language learning at eight years: Findings from the Bucharest Early Intervention Project. J Child Lang 40:605-627.
718 719 720	Yeatman JD, Dougherty RF, Rykhlevskaia E, Sherbondy AJ, Deutsch GK, Wandell BA, Ben-Shachar M (2011) Anatomical properties of the arcuate fasciculus predict phonological and reading skills in children. J Cogn Neurosci 23:3304-3317.

721 722	Yendiki A, Koldewyn K, Kakunoori S, Kanwisher N, Fischl B (2014) Spurious group differences due to head motion in a diffusion MRI study. Neuroimage 88:79-90.
723 724 725 726	Yendiki A, Panneck P, Srinivasan P, Stevens A, Zollei L, Augustinack J, Wang R, Salat D, Ehrlich S, Behrens T, Jbabdi S, Gollub R, Fischl B (2011) Automated probabilistic reconstruction of white-matter pathways in health and disease using an atlas of the underlying anatomy. Front Neuroinform 5:23.
727 728 729	Zimmerman FJ, Gilkerson J, Richards JA, Christakis DA, Xu D, Gray S, Yapanel U (2009) Teaching by listening: The importance of adult-child conversations to language development. Pediatrics 124:342-349.
730	

### 731 Figures



**Fig 1: Participant socioeconomic status.** Histograms representing the distribution of education and income across participants. Parental education is an average of the highest education level obtained by each parent and/or primary caregiver (0 = less than high school, 1 = high school, 2 = some college/associate's degree, 3 = bachelor's degree, 4 = advanced degree).

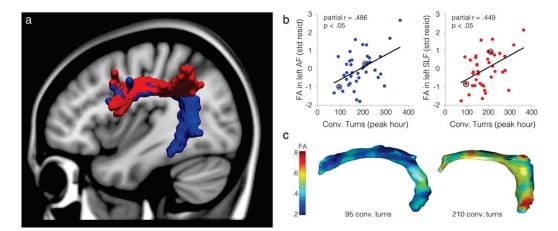


Fig 2: Conversational exposure relates to white matter microstructure. (a)

Illustration of the two left-hemisphere white matter tracts of interest. Tracts were reconstructed in each participant's native diffusion space extracted from an example participant and registered to MNI template space for visualization. Red = Superior Longitudinal Fasciculus (SLF), Blue = Arcuate Fasciculus (AF). (b) Fractional anisotropy in the left AF and left SLF as a function of the peak number of conversational turns per hour experienced by each participant, controlling for age, gender, and head motion. (c) Reconstructed left AF and SLF tracts combined for two participants matched on age, gender, and SES, but differing in the number of conversational turns experienced (open black circles in Fig. 2b). Warmer colors indicate voxels with higher FA, while cooler colors indicate voxels with lower FA.

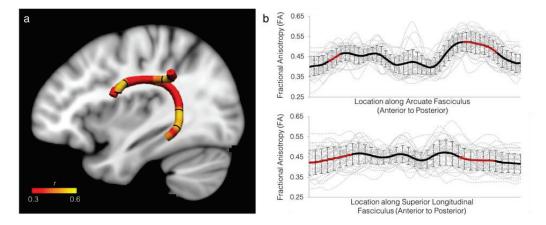


Fig. 3: Within-tract localization of the relation between conversational turns and white matter integrity. (a) Partial correlations between the number of conversational turns and FA at 35 nodes along the left Superior Longitudinal Fasciculus (SLF) and 48 points along the left Arcuate Fasciculus (AF), controlling for age, gender, motion, and SES (both parental education and family income), projected onto group average tracts in MNI space. Clusters of significant nodes are marked with black lines. (b) FA as a function of position along the AF (top) and SLF (bottom) from anterior to posterior. Gray dotted lines represent individual participants; thick dark line represents the mean of all participants; error bars represent the standard error of the mean. Regions marked in red correspond to the clusters of significant nodes marked in Fig 3a.

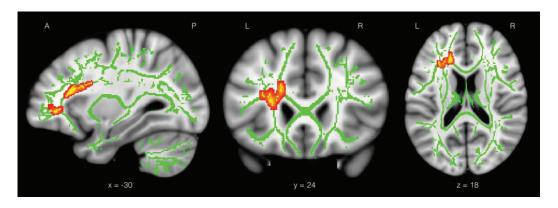


Fig 4: Whole-brain voxel-wise analysis of the relation between conversational turns and white matter integrity. The number of conversational turns was associated with FA in a cluster of voxels (p < 0.05 corrected) at the anterior end of the left SLF and left AF where these tracts terminate with Broca's area in the left inferior frontal gyrus. Analyses were computed on a skeleton (green), and thresholded values (red/yellow) were thickened and overlaid on the MNI template for visualization purposes. A = Anterior, P = Posterior, L = Left, R = Right.

