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Hemispheric dissociations in regions supporting auditory sentence comprehension in older adults

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ABSTRACT

We investigated how the aging brain copes with acoustic and syntactic challenges during spoken language comprehension. Thirty-eight healthy adults aged 54 - 80 years (M = 66 years) participated in an fMRI experiment wherein listeners indicated the gender of an agent in short spoken sentences that varied in syntactic complexity (object-relative vs subject-relative center-embedded clause structures) and acoustic richness (high vs low spectral detail, but all intelligible). We found widespread activity throughout a bilateral frontotemporal network during successful sentence comprehension. Consistent with prior reports, bilateral inferior frontal gyrus and left posterior superior temporal gyrus were more active in response to object-relative sentences than to subject-relative sentences. Moreover, several regions were significantly correlated with individual differences in task performance: Activity in right frontoparietal cortex and left cerebellum (Crus I & II) showed a negative correlation with overall comprehension. By contrast, left frontotemporal areas and right cerebellum (Lobule VII) showed a negative correlation with accuracy specifically for syntactically complex sentences. In addition, laterality analyses confirmed a lack of hemispheric lateralization in activity evoked by sentence stimuli in older adults. Importantly, we found different hemispheric roles, with a left-lateralized core language network supporting syntactic operations, and right-hemisphere regions coming into play to aid in general cognitive demands during spoken sentence processing. Together our findings support the view that high levels of language comprehension in older adults are maintained by a close interplay between a core left hemisphere language network and additional neural resources in the contralateral hemisphere.

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Introduction

Although older adults frequently experience agerelated sensory and cognitive declines [57], in many situa- 93 tions their language comprehension is remarkably good [18,67,86]. Indeed, older adults perform just as well as young adults in a number of tasks involving syntactic

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[74,81–82], semantic [6,17,93], and lexical [54,76,80] knowledge, all of which are key components in language comprehension. Nevertheless, language comprehension in healthy aging may only be stable under limited circumstances; that is, older adults may have difficulty understanding spoken sentences when made more difficult through linguistic or acoustic challenge [87,88]. The neural systems underlying preserved and fragile language processing in healthy aging are still not fully understood.

For neuroimaging studies of language in which young adults show lateralized activity, older adults frequently demonstrate recruitment of additional regions in the contralateral hemisphere [7,14,51,74,83] in addition to robust activation in the left perisylvian cortex [9,12,56,74]. This increase in bilateral activity in older adults relative to young adults is seen in many tasks and is often interpreted in the context of two classic frameworks. In the HAROLD (Hemispheric Asymmetry Reduction in Older Adults) model [8], for example, increased bilateral activity is typically seen as reflecting compensatory activity, i.e., supporting successful behavior. CRUNCH (Compensation-Related Utilization of Neural Circuits Hypothesis: [59] also posits that older adults recruit bilateral neural resources to maintain performance level. However, the involvement of the right hemisphere is only seen when the task demand is low. Critically, from both perspectives, the role of increased bilateral activity is interpreted in the context of behavioral success. By contrast, the dedifferentiation theory views age-related contralateral activity as non-specific activation, responsible for inefficient processes [33,43]. Behavioral performance is therefore key in understanding potential roles of contralateral recruitment.

In language tasks, previous studies in older adults have found both positive and negative associations between right frontal activity and behavioral performance. For example, Hoyau et al. [39] found bilateral frontal activity associated with faster response times during an object naming task, and van Ettinger-Veenstra et al. [79] showed that higher reading test scores were positively associated with the activity in right posterior temporal cortex. By contrast, a negative correlation was found between right frontal activity and performance on tasks related to picture naming [83], semantic fluency [51], or rhythm-judgment [28]. Intriguingly, Geva and colleagues [28] interpreted a negative correlation between right frontal activity and task performance as supporting a compensatory role of the right hemisphere. They based this conclusion on the fact that increased right frontal activity was only observed when error-prone subjects correctly performed the task. That is, recruitment of right frontal cortex was thought to lead to task success specifically for older adults who found the task more challenging.

In addition to the correlational findings within the older adults, group comparisons between young and older adults have indicated greater bilateral activation during various language tasks. For example, older adults activated right frontal cortex more strongly than young adults when they performed the same tasks including object naming [39], semantic categorization [45], and rhyming judgment tasks [28]. Importantly, young and old adult groups were matched in their behavioral task performance, suggesting

the importance of the right hemisphere in maintaining task demands.

It is especially difficult to ascertain the functional role of the right frontal cortex in language tasks involving both language-specific (e.g., syntactic) and domain-general (e.g., working memory) processes (see [10]. Behavioral work has established that language-specific and domain-general processes interact [24], and right frontal activity has been associated with both language-specific [48] and domain-general [89] functioning.

In order to isolate the contributions of language-specific and domain-general processes, in the present fMRI study, we adopted the experimental design used in a previous study with young adults [46]. In brief, we presented participants a series of sentences that varied in syntactic complexity depending on the type of center-embedded relative clause (e.g., object-relative vs subject-relative structures). To assess the contributions of domaingeneral processes, we acoustically degraded these sentences using noise vocoding [68] with 24 channels, thus preserving intelligibility. We draw a distinction between syntactic complexity and general sentence processing. Whereas syntactic complexity specifically taps processing operations related to word order, general sentence processing involves domain-general cognitive operations including sequence monitoring and working memory.

In the present experiment we asked the following questions: first, to what extent do older adults recruit the nondominant right hemisphere for general sentence processing? For this purpose we directly compare sentencerelated activity between older adults (current data) and young adults [46]. In addition, we compute laterality indices from the fMRI data of the two age groups. We hypothesized that older adults would engage additional resources in the right hemisphere compared to the young adults. Secondly, we ask whether the core (i.e., left frontotemporal) language network would show greater activity pertaining to increased syntactic complexity in older than young adults. To this end, we directly compare [objective-relative to subject-relative] contrast between the two age groups. In addition, we perform multiple linear regression to investigate individual differences in both general sentence processing and syntactic complexity within older adults. We hypothesized that right hemisphere activity may reflect individual differences pertaining to task demands, while left hemisphere activity would be specifically sensitive to differences in the syntactic structure of sentences. As was done in Geva et al [28], we focus on correct trials to be related to the fMRI data.

Materials and methods

Materials and behavioral task

The experimental stimuli were identical to those used in our previous study [46]. In brief, we used 24 base sentences, each containing two characters, one of which was the agent of an action. In each sentence, one of the characters was a male (king, father, uncle, etc.) and the other a female (queen, mother, aunt, etc.), and each of which

appeared in two syntactically different structures with gender of the actor being counterbalanced. The syntactic structure was manipulated by rearranging word order while equating lexical characteristics across sentences with center-embedded clauses containing either a subject-relative structure (e.g., "Brothers that aid sisters are good") or an object-relative structure ("Brothers that sisters aid are good").

In addition, half of the sentences were noise vocoded with 24 channels while the other half were unprocessed, retaining the original acoustic details. The particular sentences presented as unprocessed or noise-vocoded speech were counterbalanced across participants. Our pilot study confirmed that these vocoded sentences were still intelligible to healthy older adults. Together, these manipulations resulted in a 2×2 factorial design with varying syntactic complexity (subject-relative, object-relative) and acoustic detail (acoustically rich, acoustically less-detailed).

After each sentence was heard, the participant was asked to indicate whether the agent of the action in the sentence was male or female by pressing one of two buttons on an MRI-compatible button pad. For example, for the sentence "Brothers that help sisters are nice", the correct response would be "male". A subset of 24 experimental sentences was vocoded with a single channel to create an unintelligible control condition, for which participants were told to randomly press either the male or female button.

Pilot study

To determine the optimal range of noise vocoding, we conducted a behavioral pilot study. We presented sentences at levels of vocoding that ranged from unprocessed (clear) speech and 8, 16, 24, and 32 channel vocoded speech. Vocoding with<8 channels resulted in extremely poor signal quality as illustrated in the spectrograms in Fig. 1A. We tested 40 adults aged 62–88 years. All participants provided written informed consent according to a protocol approved by the Brandeis University Institutional Review Board.

Sentences were identical to those used in the main study. After hearing each sentence, participants were instructed to press a key, as quickly and accurately as possible, to indicate the gender of the character performing the action. The accuracy of responses as a function of acoustic clarity is shown in Fig. 1B. When<16 channels were used in noise vocoding, accuracy drastically decreased. These results indicated that noise vocoding down to 16 channels would be sufficient to ensure relatively high levels of accuracy for all conditions in the fMRI study for all participants, regardless of age. Because the pilot study was conducted in a quiet room with minimum levels of background noise, we chose 24-channel noise vocoded speech as a conservative manipulation of acoustic challenge for the noisy MR scanning environment.

fMRI study

Participants

The fMRI experiment included 46 participants. All participants reported themselves to be right-handed, in good

health, having no known neurological disorders, being native speakers of English, and having normal hearing. Among the 46 participants, eight individuals' data were later discarded due to a chance-level performance in any of the 4 functional runs (N=6), excessive head motion (translation > 10 mm) (N=1), and missing working memory scores (N=1). This resulted in a total of 38 usable subjects (54-80 years) of age; M=66 years, SD=6.7; 24 females; education M=16 years, SD=2.4; Mini-Mental Status Examination M=29, SD=0.8). All participants submitted written consent and the study was approved by the Institutional Review Board at the University of Pennsylvania.

Prior to the fMRI experiment, we measured hearing acuity and verbal working memory performance. For hearing acuity, we used standard pure-tone audiometry in which pure tones were presented at octave intervals between 250 and 8000 Hz, with each participant's pure tone threshold reflecting the softest tone they could reliably hear (Fig. 2A). We then averaged thresholds over 1000, 2000, and 4000 Hz for a pure-tone average (PTA). We used each participant's better-ear PTA as a parsimonious summary of their hearing ability.

To measure verbal working memory, we used a reading span task [13]. In this test, participants were given a set of written sentences, for each of which they were required to make a true/false judgment. Upon completion of all sentences in a set, participants were asked to recall the last words of each sentence. Participants proceeded with the next set in which the number of sentences was increased. The task was finished either when the participants correctly remembered the last word of each sentence or when they could not correctly give more than three target words.

Experimental procedure. Prior to entering the scanner, participants received instructions and performed a practice session to ensure they understood the task. Once inside the scanner, but prior to scanning, the volume of auditory stimuli was determined by asking participants to repeat each sentence of four recorded sentences (two unprocessed and two 24-channel vocoded sentences). An optimal volume level was determined upon perfect repetition of all sentences. These sentences were not included in the main stimuli set. Once set, the volume remained constant for that participant.

Following the volume determination, the participant conducted the sentence comprehension task across four fMRI runs. On each trial, either an intelligible (both clear and 24-channel vocoded speech) or unintelligible noise (1-channel vocoded speech) was presented to the participant using E-Prime 2.0 (Psychology Software Tools, Inc., Pittsburgh, PA). These sentence stimuli were presented binaurally using MRI-compatible high-fidelity insert earphones (Sensimetrics Model S14).

As indicated previously, for each trial participants were told to indicate the gender of the individual who performed the action in the sentence via a button press. These button responses were later analyzed to assess sentence comprehension. For the 1-channel vocoded speech participants were instructed to randomly press either the male or female button. Together, each run consisted of 36 trials

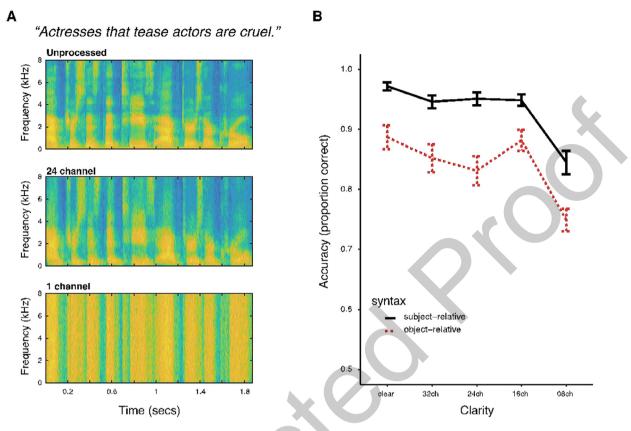


Fig. 1. A. Spectrograms of unprocessed speech, 24-channel noise vocoded speech, and 1-channel noise vocoded speech. B. Accuracy from a pilot study examining performance as a function of acoustic clarity in 40 older adults. Error bars show one standard error.

with 24 intelligible sentences (6 sentences \times 2 syntactic constructions \times 2 acoustic manipulations), 6 unintelligible speech (1-channel vocoded), and 6 trials of silence. The trials were randomized within runs and 96 trials were presented only once across four functional runs.

MRI scanning. We collected MRI data on a 3 T Siemens Trio scanner (Siemens Medical System, Erlangen, Germany) using an 8-channel head coil. The field of view was angled approximately 30° away from the AC-PC line. First, a T1weighted structural volume was acquired using a magnetization prepared rapid acquisition gradient echo (MPRAGE) protocol [axial orientation, repetition time (TR) = 1620 ms, echo time (TE) = 3 ms, flip angle = 15° , field of view (FOV) = 250×188 mm, matrix = 256×192 mm, 160 slices, voxel resolution = $0.98 \times 0.98 \times 1$ mm]. Next, four runs of blood oxygenation level-dependent (BOLD) functional MRI scanning were conducted [TR = 2000 ms, TE = 30 ms, flip angle = 78°, FOV = 192×192 mm, matrix = 64×64 mm, 32 slices, voxel resolution = $3 \times 3 \times 3$ mm with 0.75 mm gap] using an interleaved silent steady state (ISSS) protocol [66], in which five consecutive volumes were acquired in between four seconds of silence. Each ISSS trial consisted of four seconds of relative quiet period and a subsequent 10 s (5 volumes) of data collection period. Before completion of the fMRI experiment, we acquired a B0 field mapping sequence [TR = 1050 ms, TE = 4 ms, flip angle = 60° , FOV = 240×240 mm, matrix = 64×64 mm, 44 slices, slice thickness = 4 mm, voxel resolution = $3.8 \times 3.8 \times 4$ mm].

fMRI data analysis. Preprocessing was performed via SPM12 (version 6906; Wellcome Trust Centre for Human Neuroimaging) after raw images were unwarped in the prelude and flirt routines from FSL version 5.0.5 (FMRIB Software Library, University of Oxford). First, all timeseries were realigned to the first image in the first session. Next, the structural image was coregistered to the functional images before being normalized to the standard MNI T1 template using a unified segmentation approach [4]. The same transformation was then applied to normalizing all functional images to MNI space with the original voxel size preserved. Preprocessing was completed by smoothing with a Gaussian kernel of nine mm full-width half maximum (FWHM).

To model the ISSS data, we fit the functional data using a finite impulse response (FIR) function, wherein five time points were separately estimated per trial (Peelle, 2014). We separately estimated sentences associated with correct and incorrect responses. For those conditions with no errors (i.e., 100 % accuracy), the last TR was modeled as a dummy event in order to avoid a regressor of empty vector. Although we applied high-pass filtering with a 128 sec cutoff, temporal autocorrelation was disabled due to the temporal discontinuity of our data sets. Together, there were

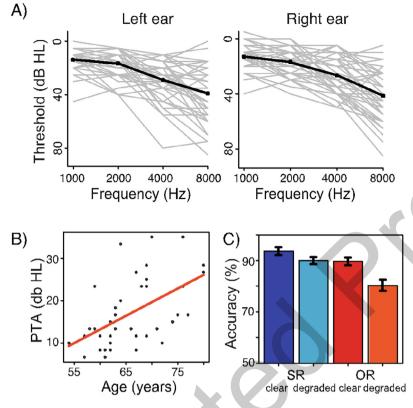


Fig. 2. A. Hearing acuity in left and right ear of 38 participants. Light gray lines indicate individual participants, black line the group average. B. Relationship between age and better-ear PTA. Age is positively associated with hearing decline [Pearson r = 0.53; p < 0.001].; C. Behavioral accuracy during the fMRI experiment. There are significant effects of syntactic complexity and acoustic detail.

20 regressors per correct and incorrect sentence conditions, 5 regressors of 1 channel vocoded speech, and 6 motion parameters, totaling 51 estimates per run. There were additional 4 run regressors in the design matrix.

After estimating the GLM, we calculated the summed positive area (SPA) using the 5 beta estimates for each condition [46]. More specifically, we integrated the area of the estimated response above zero and used this as a metric to be used for further analyses. These SPA maps were then submitted to a series of 2nd level random effects analyses including one-sample t-tests and multiple regressions. All results were obtained using a voxel-wise threshold of p < 0.001 (uncorrected) in combination with cluster-wise threshold of p < 0.05 corrected using gaussian random field theory [27,90]. Results were displayed using SPM or MRIcron [61]. Unthresholded statistical maps are available at https://neurovault.org/collections/9592/ [32].

The degree of lateralization was determined on group maps using the LI toolbox (version 1.3.2) [84]. In brief, the LI (lateralization index) was calculated by subtracting the sum of voxel values above the cut-off threshold (i.e., T = 3) of the right hemisphere from the left hemisphere, which was subsequently normalized by the total sum of voxel values of both hemispheres. The LI ranges between +1 and -1 with positive value being left lateralization. We obtained the LI separately in frontal, parietal, temporal, subcortical regions (thalamus + basal ganglia), and cerebel-

lum. In each of these regions, only voxels located>11 mm from to the midline were considered.

Results

Behavioral results

As seen in Fig. 2C, overall, participants exhibited relatively good accuracy on the comprehension task during scanning (M = 88.4 percent correct; SD = 7.9). We set up a linear mixed effects model using lmerTest package [44] in R (version 4.1.1) to predict task performance in logittransformed accuracy with a set of fixed effects including PTA, age, working memory, and PTA-by-working memory for between-group variables as well as syntax, acoustic clarity, and syntax-by-acoustic clarity as within-group variables. Participants were included as a random effect. Results are shown in Table 1. The model yielded significant main effects of syntax $[F_{(1,114)} = 43.75, p < 0.001, \eta^2 = 0.28]$ and acoustic clarity $[F_{(1,114)} = 44,07, p < 0.001, \eta^2 = 0.28];$ consistent with the behavioral pilot, accuracy of comprehension was poorer for object-relative sentences compared to subject-relative sentences, and accuracy of comprehension decreased in the presence of degraded rather than clear speech. Age, hearing acuity, and working memory had no significant impact on the task performance.

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Table 1Behavioral results on the sentence comprehension task

Variable	Estimate	Std. Error	t value	P value
syntax (SR)	0.58	0.14	4.27	4.10E-05
acoustic clarity (degraded)	-0.71	0.14	-5.1	1.35E-06
age	-0.02	0.02	-0.9	0.37
working memory	0.21	0.570	0.37	0.71
hearing acuity (pta)	-0.01	0.04	-0.29	0.78
syntax-by-acoustic	0.11	0.2	0.58	0.56
working memory-by-pta	0.004	0.03	0.13	0.9

In addition, we compared the behavioral performance between the current older adults and the young adults (N = 26) who performed the same task in our previous study [46]. An independent samples t-test using logittransformed accuracy revealed that older adults performed significantly worse than young adults $[t_{(62)} = 3.68,$ p < 0.001, Cohen's d = 0.93]. We also compared syntactic performance between older and young adults. For the comparison, we used a metric of [% accuracy of subject object-relative sentences], in which larger positive values indicate greater vulnerability to the syntactic complexity. These metrics — after confirming normal distribution using Shapiro-Wilk normality test (p > 0.05) —were submitted to the independent samples t-test. We found that older adults indeed performed significantly worse than younger adults as the syntactic structure becomes more complex $[t_{(62)} = 2.97, p < 0.001, Cohen's d = 0.66].$

fMRI results

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We first examined the activity pertaining to general sentence processing (i.e., between all sentences versus noise) using group-level random effects. As expected, widespread activity was seen bilaterally throughout both cortical and subcortical areas (Fig. 3A; Table 2). To further examine this bilateral pattern of activation, we compared the resulting map to the same group-level result obtained from young adults [46] via independent samples t-test. We found four significant clusters exclusively in the right hemisphere that were more active in the older than young adults; those include the insula, pre-supplementary motor area (SMA), and two cerebellar loci in VII and crus II (Fig. 4; Table 3). To further explore the bilateral activation in the current data, we computed LIs in 5 atlas-based regions of interest (ROIs): frontal, parietal, temporal, subcortical, cerebellum. These LIs were then compared to those obtained from the study with young adults [46]. Because these indices were drawn from group-level T maps for young and older adults' group data, no statistical comparison was made. As summarized in Table 4, frontal, parietal, temporal, and subcortical areas displayed increased bilateralization in older compared to young adults. By contrast, the cerebellum showed weaker right lateralization in the older adults.

Next, we examined the activity more specifically related to differences in syntactic structure by comparing the difference between object-relative and subject-

relative sentences. We found significant clusters predominantly in the left hemisphere including IFG (both pars triangularis and pars opercularis), posterior aspect of STG/MTG, SMA, precentral gyrus, and thalamus. In addition, bilateral clusters were seen within the insula, putamen, and cerebellum (Fig. 3B; Table 5).

As was done for the general sentence processing map above, we directly compared the group map of the [object-relative > subject-relative] contrast between the young and older adult groups via independent samples *t*-test, which did not yield any significant clusters in either [older > young] or [young > old] even at a relaxed threshold (P < 0.01 voxel-wise uncorrected and extent clustersize = 20). Nevertheless, the LI index revealed that frontal, parietal, and temporal areas were more left-lateralized in the young than older adults (Table 4).

Next, we performed a series of multiple regressions to investigate the individual differences in the degree of activations pertaining to their behavioral performance within older adults. As an independent variable, we used percent accuracy from the sentence comprehension task. As covariates, we included age, PTA, working memory, and the interaction term of PTA-by-working memory, with meancentering in the model. We found 6 significant clusters that were negatively correlated with the task accuracy. These clusters were mostly located in the right hemisphere, including IFG/insula, frontal pole, SMA, angular gyrus/supramarginal gyrus — all of which were parts of general sentence processing network described in the group level *t*-test result above. In addition, two large clusters emerged in the bilateral cerebellum (Fig. 5; Table 6).

We performed another multiple regression to identify areas that would be sensitive to individual differences in behavioral performance for syntactically more complex sentences. To this end, we used a score that reflected differences in task performance between subject-relative accuracy and object-relative condition (i.e., subjectrelative minus object-relative accuracy). As such, smaller scores indicate comparable performance between the two syntactic conditions and higher scores reflect worse performance in the object-relative condition. This score was then regressed against the syntactic contrast map (objectrelative > subject-relative). The same covariates as above were also included. We found that left insula, pre-SMA, precentral gyrus, and right cerebellum showed positive correlations with performance, indicating participants who struggled more with syntactic complexity tended to

A) Sentences > Noise

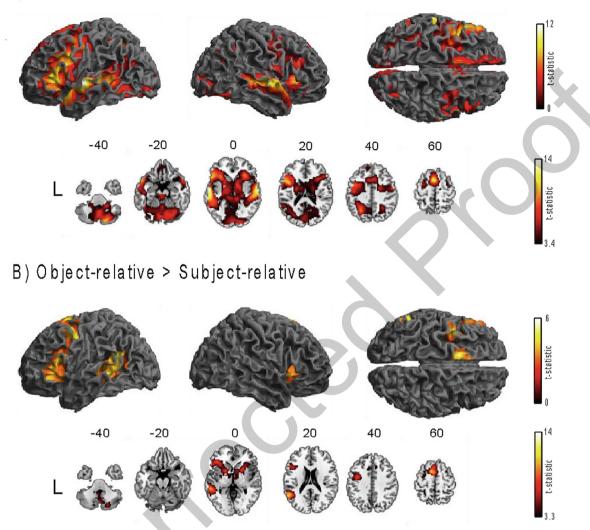


Fig. 3. A) Group-level results of sentence processing [all sentences > 1 channel vocoded speech] B) Group-level results of the main effect of syntactic complexity [Object-relative > Subject-relative]; L = Left hemisphere.

upregulate these areas during successful syntactic operation (Fig. 6, Table 7).²

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To further confirm these results derived from correct trials, we performed the same multiple regression exclusively on error trials. Because participants overall committed relatively few errors in each condition, we constructed a new model in which the error trials were collapsed across

all sentence conditions. In addition, for this model, functional runs with no errors were removed per participant in the first level analysis. The results revealed no significant voxels.

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Lastly, we evaluated the [clear > 24 ch vocoded] contrast by collapsing object-relative and subject-relative conditions and syntax-by-acoustic interactions, none of which yielded significant differences.

Discussion

In the present study, we investigated the neural underpinnings of successful sentence comprehension in healthy older adults. We found widespread activity throughout cortical and subcortical areas *bilaterally* while participants performed an auditory sentence comprehension task. Given the current study adopted the identical experimen-

² At the suggestion of a reviewer, we repeated this analysis by replacing negative scores for [subject-relative minus object-relative accuracy] with zeros, the logic being that it is difficult to interpret scores if listeners perform worse on the easier sentences. This modified analysis yielded results that were largely comparable, but slightly weaker: the left dorsolateral frontal cluster was no longer significant (see Supplementary Material for a side-by-side comparison). In addition, we replaced the [object-relative vs subject-relative] contrast with [all sentences vs 1 ch. noise] contrast to be regressed against the original score of [subject-relative minus object-relative accuracy]. This did not yield any significant clusters.

Significant regions in the [sentence > noise] group-level comparison.

	MNI Coordii	nates			
Region name	x	У	Z	z-value	volume of cluster (μl)
L anterior STG/STS	-57	-10	-1	>7.61	532,008
R anterior STG/STS	63	-13	-1	>7.61	
L MTG	-57	-25	-1	>7.61	
L MTG	-60	-28	2	>7.61	
L SMA	-3	5	62	7.61	
L IFG (Operculum)	-42	8	23	7.49	
L Orbitofrontal Cortex	-51	23	-4	7.43	
R anterior STG/STS	60	-1	-4	7.41	
R middle STG/STS	54	-22	2	7.33	
L IFG (Triangularis)	-51	26	8	7.22	
L Superior temporal pole	-54	8	-10	7.11	
Cerebellum (Crus2)	18	-79	-40	7.05	
L Superior temporal pole	-48	14	-22	7.02	
L Insula	-33	26	8	7.01	
R Cerebellum VIII	30	-64	-52	6.9	

STG: Superior Temporal Gyrus; STS: Superior Temporal Sulcus MTG: Middle Temporal Gyrus; SMA: Supplementary Motor Area IFG: Inferior Frontal Gyrus; R:Right; L: Left

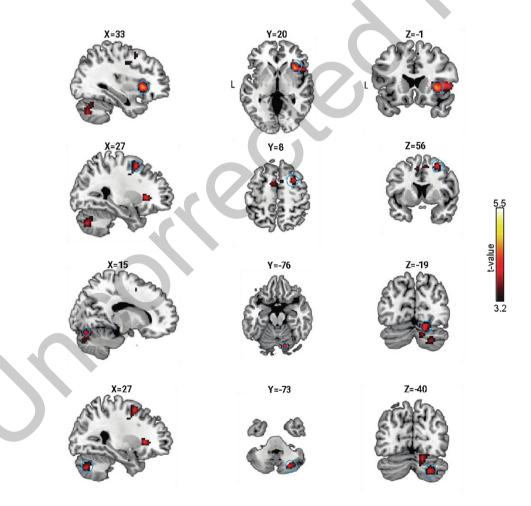


Fig. 4. Areas that were more active in the older adults than young adults during general sentence processing. These clusters were exclusively located within the right hemisphere, further confirming the bilateral activation in older adults.

Table 3Areas that were more activated in older than younger adults.

	MNI Coordi	inates			
Region name	х	У	Z	z-value	volume of cluster (μl)
R Insula	33	20 –1	-1	4.29	177
	45	20	5	4.07	
	54	17	2	3.74	
R Cerebellum VI	15	-76	-19	3.86	107
	12	-79	-31	3.56	
R PreSMA	27	8	56	3.82	178
	21	2	50	3.69	
	-9	5	53	3.57	
R Cerebellum Crus II	27	-73	-40	3.72	166
	39	-61	-28	3.66	
	36	-67	-43	3.59	

PreSMA: Pre-Supplementary Motor Area

Table 4Lateralization indices of old vs young adults data.

	Frontal		Parietal		Tempor	al	BG + Th	alamus	Cerebellu	m
	old	young	old	young	old	young	old	young	old	young
sentence > noise	0.18	0.63	0.17	0.31	0.15	0.29	0.01	0.39	-0.38	-0.86
OR > SR	0.56	0.79	0.93	N.A	0.77	0.95	0.13	0.06	-0.8	N.A

[&]quot;+": left-lateralized; "-": right-lateralized; young adults' LI are from data in Lee et al. [46].

N.A: Not Available; analyses were failed to yield an LI

BG: Basal Ganglia

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Table 5Significant regions in the [OR > SR] group-level comparison.

	MNI Coordii	nates			
Region name	х	У	z	z-value	volume of cluster (μl)
R putamen	21	8	-7	5.56	61,749
L SMA	-9	5	62	5.51	
R caudate	15	5	11	5.47	
L putamen	-18	5	8	5.29	
L precentral gyrus	-39	2	50	5.16	
L putamen	-18	8	-4	5.07	
L IFG (pars triangularis)	-36	29	5	4.78	
L IFG (orbitofrontal cortex)	-42	29	-4	4.74	
L thalamus	-12	-4	11	4.61	
L IFG (pars opercularis)	-51	17	14	4.35	
L IFG (pars triangularis)	-54	23	8	4.22	
R insula	33	17	-7	4.09	
L insula	-27	17	-4	3.97	
R insula	30	26	2	3.72	
L cerebellum VIIII	-3	-49	-37	4.24	15,633
R cerebellum VI	18	-67	-31	4.17	
cerebellum (vermis)	0	-61	-34	4.12	
R cerebellum VI	30	-61	-28	4.05	
L cerebellum VIII	-12	-64	-31	3.98	
R cerebellum (crus II)	21	-79	-40	3.93	
R cerebellum (crus I)	45	-64	-31	3.86	
R cerebellum VIII	6	-67	-46	3.76	
R cerebellum VIII	9	-70	-43	3.68	
R cerebellum (crus II)	-15	-79	-40	3.25	
R cerebellum V	15	-46	-19	3.23	
L MTG	-54	-52	20	5.49	15,363
L MTG	-57	-37	2	5.16	
L MTG	-51	-43	11	4.92	

tal design, sentence materials, and behavioral task used in our previous study [46], we were able to make some direct comparisons to the previous study, which confirmed the appearance of significant clusters exclusively within the right hemisphere that were more active in the older adults than previously shown in the young adults.

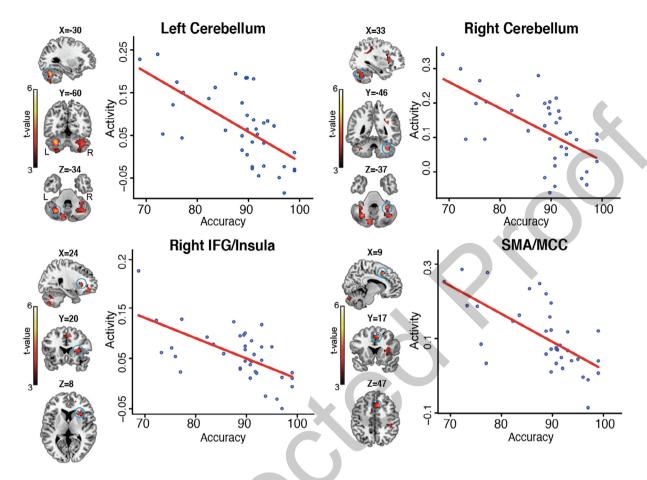


Fig. 5. Areas in which activity negatively correlates with individual differences in overall task performance. These areas are situated contralateral to the conventional language network, suggesting that compensatory neural resources support sentence comprehension. The correlation plot is for visualization purposes on significant clusters found by the whole-brain multiple regression using [sentences > 1ch. vocoded speech], in which activity was averaged across voxels within a significant cluster.

Significant regions in the multiple regression on [sentence > noise] contrast.

	MNI Coordir	iates			
Region name	х	У	Z	z-value	volume of cluster (μ <i>l</i>)
L Cerebellum (Crus I)	-30	-64	-34	4.35	14,634
L Cerebellum (Crus II)	-9	-76	-40	4.32	
L Cerebellum (Crus I)	-36	-67	-46	4.02	
R vmPFC	21	47	-10	4.31	2241
R vmPFC	18	50	2	4.04	
R vmPFC	27	41	-1	3.62	
R SMA/preSMA	9	17	47	4.11	4590
L SMA/preSMA	-15	2	53	3.97	
L SMA/preSMA	-9	-19	62	3.61	
R Cerebellum (Crus I)	33	-46	-37	4.06	8235
R Cerebellum (Crus I)	30	-70	-22	3.89	
R Cerebellum (Crus I)	24	-70	-55	3.87	
R Supramarginal gyrus	36	-49	29	3.86	2187
R Postcentral gyrus	39	-34	47	3.53	
R Supramarginal gyrus	36	-37	35	3.43	
R Insular cortex	24	20	8	3.86	2376
R Insular cortex	36	17	14	3.63	
R Frontal operculum	30	14	23	3.32	

vmPFC: ventromedial prefrontal cortex; SMA: suppelmentary motor cortex

L: Left; R: Right

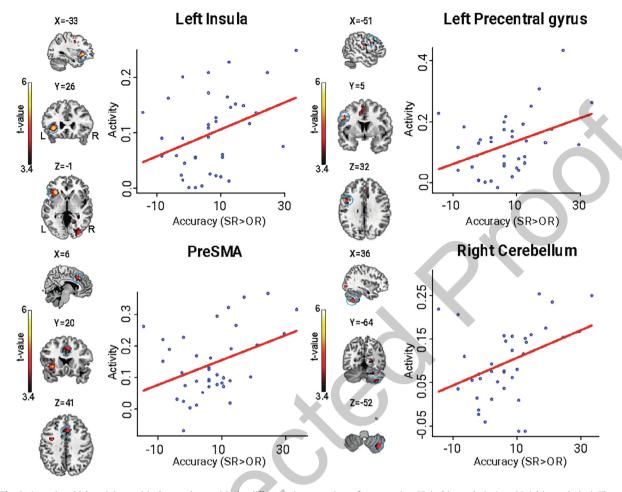


Fig. 6. Areas in which activity positively correlates with the difference in syntactic performance, i.e., SR (subject-relative) vs OR (object-relative). These areas are situated in the left-lateralized frontal network and right cerebellum. The correlation plot is for visualization of the significant clusters found by the whole-brain multiple regression, in which activity was averaged across voxels within a significant cluster.

Table 7Significant regions in the multiple regression on [OR > SR] contrast.

	MNI Coordii	nates			
Region name	х	У	Z	z-value	volume of cluster (μ <i>l</i>)
L Insula	-33	26	-1	4.64	4590
L Precentral Gyrus	-51	5	32	4.22	3024
L Middle Frontal Gyrus	-33	-1	41	4.13	
L Postcentral Gyrus	-51	-16	23	3.96	
R Lateral Occipital Cortex	36	-85	-1	4.1	4617
R Intracalcarine Cortex	18	-76	14	3.73	
R Intracalcarine Cortex	33	-76	2	3.66	
R preSMA	6	20	41	4.06	5427
L preSMA	-9	11	47	3.94	
L preSMA	-3	8	53	3.79	
R Cerebellum VIIb	36	-64	-52	4.01	5211
R Cerebellum VI	27	-52	-31	3.89	
R Cerebellum VI	15	-55	-28	3.75	

SMA: Supplementaryt Motor Area

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As noted in the introduction, for tasks in which young adults show strongly lateralized activity, bilateral activity in older adults has been ascribed to either compensation or dedifferentiation, depending on the relationship with behavioral performance. That is, from a compensatory perspective, the contralateral hemisphere is recruited in older 522

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adults to maintain the level of behavioral performance that is supported solely by the dominant hemisphere in young adults. By contrast, a dedifferentiation perspective suggests that increased bilateral activity in older adults relative to young adults reflects non-specific and inefficient processing, hampering task performance. Indeed, when behavioral performance on the same auditory comprehension task was compared between the two age groups, older adults performed significantly worse than young adults in the current data.

To better understand the role of the bilateral activity in older adults, we performed multiple regression analyses within older adults. Although we only included correct trials in our main imaging analyses, participants varied in their overall performance, which allowed us to relate overall task accuracy to neural activity pertaining to general sentence processing using whole-brain multiple regressions. We indeed found a set of right cortical areas (mostly in frontal cortex) that showed greater activity in participants with poorer performance. Similar to the present finding, Wierenga et al. [83] reported a negative correlation between right frontal activity and picture naming accuracy. Meinzer et al. [51] also found the same relationship between right frontal activity and semantic fluency.

Such a negative correlation may, at first glance, suggest that the right frontal activity is non-beneficial or even harmful to language processing. However, it is important to consider that we found the negative association in the correct trials only, ruling out a potential contribution of the stronger right activity toward failure. Similarly, Geva and colleagues [28] found a negative correlation between right frontal activity associated with correct trials and behavioral performance on a rhyme-judgment task in healthy older adults. Although further research is warranted, our data suggest that the right prefrontal cortex may aid in language performance of older adults who experience more difficulty in the task than others.

What compensatory roles do these right cortical areas play during sentence comprehension? Frontoparietal regions (typically in the left hemisphere) are often implicated in various processes intrinsic to sentence processing, as briefly described below. First, the ventromedial portion of prefrontal cortex is thought to monitor upcoming events from the ongoing action sequences [62]. As such, upregulation of this prefrontal region might help facilitate successive word analyses while listening to speech. The insula, especially the anterior portion, has been implicated in studies examining degraded speech perception [77]. For example, Erb et al [20] reported greater activity of the right insula in degraded (i.e., 4 channel vocoded sentences) than clear speech. They also showed that the right insula's activity was negatively associated with older adults' hearing acuity.

In contrast to Erb et al. [20], we did not find a significant correlation between the right insula and hearing acuity. This may be due to our use of 24 channel vocoded speech, which was intended to present a minimal acoustic challenge. Moreover, sound intensity was calibrated for each participant prior to MRI scanning, which could have further obscured differences in acoustic sensitivity among the older adults. Instead, we found that bilateral insula

showed a significant negative correlation with sentence comprehension accuracy. The SMA/pre-SMA may be recruited for procedural and predictive coding which is important for syntactic construction [38,65,78]. In addition, the pre-SMA plays a role in semantic and lexical processing during language comprehension, as evidenced by studies reporting increased activity during semantic priming [75] and semantic/lexical decision tasks [52].

Outside of the prefrontal cortex, we also observed activity in the right supramarginal gyrus (SMG), presumably due to its role in phonological processing. Activity in SMG has been reported in studies that required explicit phonological or rhyme judgment tasks [58,92]. In support of this notion, an inhibitory TMS (transcranial magnetic stimulation) resulted in slower responses in phonological, but not semantic processing [70]. Together, these right frontoparietal regions may begin to play a compensatory role in linguistic or speech processes and are more leveraged by older adults who are more prone to age-related declines in language operations.

Alternatively, however, right frontoparietal activity also reflect domain-general processing, language-specific processing per se [55]. Previous studies have suggested that requiring listeners to perform an explicit task resulted in a lateralization shift from left to right, whereas task-free sentence comprehension evoked activities only in the left-lateralized frontotemporal system [14,16]. In the present study, we used a gender judgment task on every sentence except occasional unintelligible stimuli (i.e., 1 channel vocoded speech), for which participants were allowed to press the male or female button at their discretion. As such, trials with sentence stimuli were presumably more demanding than trials with unintelligible noise stimuli, indicating that the right hemisphere may play an important role in keeping up with general task demands.

In stark contrast to the right-lateralized activity associated with general sentence processing (sentences vs noise), robust left-lateralization was found to be sensitive to the syntactic complexity (object-relative vs subject-relative) in older adults as is typically found in young adults. For example, consistent with extant literature [26,35,36,49,73], we observed the left dorsolateral prefrontal cortex and the posterior aspect of superior temporal gyrus to exhibit stronger activity in response to object-relative than subject-relative sentences. Hence, the conventional language system in the left frontotemporal network is still robustly sensitive to the syntactic structure of sentences, independent of aging. However, it is worth correlation noting that the of the [objectrelative > Subject-relative | with the behavioral scores of [subject-relative minus object-relative] did not implicate the core syntactic regions including pars triangularis, pars opercularis in the left IFG, or posterior STG/MTG. Instead, the analysis yielded areas outside of the core language network [22,21,23]. Indeed, these frontal clusters (left insula, precentral gyrus, and pre-SMA) are parts of the multiple demand network [19], which are active across different cognitive domains. As such, it appears that those older adults who experience greater difficulty in syntactically complex sentences (i.e., object-relative sentences) leverage

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domain-general constructs such as verbal working memory [11,31,41,47,50,60]. Future research using systematic manipulation of the syntactic structure [53,63] is warranted to better understand the compensatory role of these domain-general regions on syntactic analysis.

In the present case, when the same group map of [object-relative > subject-relative] was compared between older and young adults, the degree of activation was comparable between the two groups. Importantly, however, upregulation of the left frontotemporal language system was observed in older adults who were more vulnerable to the syntactic complexity. Indeed, object-relative sentences are known to be more demanding than subjectrelative sentences, as evident here by lower behavioral scores (see also [88,87]). Despite the differences in task demands, syntactic comparisons — in both group-level ttest and multiple regression - yielded activity mostly in the left hemisphere, in contrast to the results regarding general sentence processing. This finding is in line with the notion that the core language function is stable across lifespan, and that it continues to rely on the left-lateralized frontotemporal system [9.67.74.93].

Stable language function in healthy aging has been supported by previous neuroimaging work showing that the left frontotemporal system is consistently activated during language comprehension across a large cohort of young and older participants, and with no intrinsic functional connectivity difference in syntactic processing between young and older adults [9]. Relatedly, decreased restingstate functional connectivity within the left frontotemporal system in older adults is associated with agrammatism in early stage of primary progressive aphasia (i.e., nonfluent/agrammatic variant PPA or naPPA) even before atrophy becomes noticeable [5]. In a clinical-pathological study of naPPA, non-fluent speech errors and grammatical simplifications were associated with increased burden of tau pathology in left mid-frontal cortex [30] and associated frontal white matter regions [29]. Contrary to these findings, a recent study using transcranial Direct Current Stimulation study reported that increase of language score was associated with decrease of functional connectivity between left IFG and posterior MTG [25]. Future research is required to provide more insights into the relationship between functional connectivity and grammar processing in both healthy older adults and those with naPPA. For example, it needs to be determined whether or not an effective behavioral therapy for agrammatism [37] is by re-stabilizing the aberrant functional connectivity independent of compensatory activation in the deteriorating left frontotemporal system of patients with naPPA [34,85].

Although we found robust activation in the left frontotemporal network associated with syntactic processing in our healthy older adults, the laterality index pertaining to the syntactic manipulation indicated less left-lateralization in the older adults compared to young adults. Such bilateral activations in cortical areas are paralleled by bilateral activation in the cerebellum which typically shows contralateral activity (i.e., more robust right-lateralized) during language processing. Although the cerebellum has been traditionally viewed as a motoric processor, there is a growing body of evidence that the

cerebellum plays a key role in language comprehension [15,72], especially syntactic processing [1,3,2;42]. Typically, right cerebellar activity is seen along with left-lateralized frontotemporal network activity during language comprehension [40]. Early case reports with right cerebellar lesions revealed that both expressive and receptive syntactic operations were impaired [69,91]. Schwartze and Kotz [64] theorized that the cerebellum mediates, in concert with basal ganglia, analyses of temporal structures (e.g., boundaries across words and phrases) during speech comprehension. In particular, lobule VII in the right cerebellum is thought to play a key role in syntactic analysis with its robust connection to the prefrontal cortex [71].

In the present study, we found a large expanse of the right cerebellum spanning across lobule V to VIII respond to syntactically complex speech. Moreover, lobule VII from the right cerebellar cluster was found to be associated with age-group difference (i.e., stronger in older adults) for general sentence processing as well as individual differences within older adults in syntactic perfromance. Although less common, both bilateral and left-lateralized cerebellum activities have been reported in studies involving language perception and production tasks [42,71]. Indeed, the left cerebellum, along with the right frontoparietal areas, was most robustly (in both cluster-size and t-statistic) associated with individual differences in general sentence processing. Together, we found different hemispheric roles in the cerebellum that correspond to the opposite hemisphere of the core language network.

In summary, in the present study we investigated the neural underpinnings of successful sentence comprehension in healthy older adults. Our main findings within older adults suggest distinct functional roles in supporting spoken sentence processing: whereas the right frontal cortex and the left cerebellum were newly recruited to maintain general sentence processing [14], the left frontal cortex and the right cerebellum - the conventional language network – continued to play a key role in core syntactic function despite aging (see also [9]. Furthermore, these regions were leveraged to a greater extent by worse performers. In addition, we found that, compared to young adults, older adults recruited a much wider brain network that included the traditional left sylvian cortex as well as right frontoparietal and cerebellar regions. We note that the direct comparison between the two age groups was made in a post-hoc and supplementary manner, which posed some issues including unbalanced sample sizes. Future study should employ larger and matched sample size as well as more planned statistical analyses in delineating neuroanatomical differences between young and older adults in sentence processing. Together, our findings are broadly compatible with a neural compensation perspective and the observed resilience of the core language system in healthy aging.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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References

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- Ackermann H, Mathiak K, Riecker A. The contribution of the cerebellum to speech production and speech perception: Clinical and functional imaging data. Cerebellum 2007;6:202–13. doi: https://doi.org/10.1080/14734220701266742.
- [2] Adamaszek M, Kirkby KC. Chapter 5 Cerebellum and Grammar Processing. In: Mariën P, Manto M, editors. The Linguistic Cerebellum. San Diego: Academic Press; 2016. p. 81–105. doi: https://doi.org/10.1016/B978-0-12-801608-4.00005-0.
- [3] Adamaszek M, Strecker K, Kessler C. Impact of cerebellar lesion on syntactic processing evidenced by event-related potentials. Neurosci Lett 2012;512:78–82. doi: https://doi.org/10.1016/j.neulet.2012.01.020.
- [4] Ashburner J, Friston KJ. Unified segmentation. Neuroimage 2005;26:839–51. doi: https://doi.org/10.1016/j.neuroimage.2005.02.018.
- [5] Bonakdarpour B, Rogalski EJ, Wang A, Sridhar J, Mesulam MM, Hurley RS, Functional Connectivity is Reduced in Early-stage Primary Progressive Aphasia When Atrophy is not Prominent. Alzheimer Dis Assoc Disord 2017;31:101–6. doi: https://doi.org/10.1097/WAD.000000000000193.
- [6] Bowles NL. Age and Semantic Inhibition in Word Retrieval. J Gerontol 1989;44:P88–90. doi: https://doi.org/10.1093/geronj/44.3.P88.
- [7] Bozic M, Tyler LK, Ives DT, Randall B, Marslen-Wilson WD. Bihemispheric foundations for human speech comprehension. PNAS 2010;107:17439-44. doi: https://doi.org/10.1073/ pnas.1000531107.
- [8] Cabeza R. Hemispheric asymmetry reduction in older adults: the HAROLD model. Psychol Aging 2002;17:85–100. doi: https://doi.org/10.1037//0882-7974.17.1.85.
- [9] Campbell KL, Samu D, Davis SW, Geerligs L, Mustafa A, Tyler LK, et al. Robust Resilience of the Frontotemporal Syntax System to Aging. J Neurosci 2016;36:5214–27. doi: https://doi.org/10.1523/JNEUROSCI.4561-15.2016.
- [10] Campbell KL, Tyler LK. Language-related domain-specific and domain-general systems in the human brain. Curr Opin Behav Sci, Evol Language 2018;21:132–7. doi: https://doi.org/10.1016/ i.cobeha.2018.04.008.
- [11] Caplan D, Waters GS. Verbal working memory and sentence comprehension. Behav Brain Sci 1999;22:77–94. doi: https://doi.org/10.1017/S0140525X99001788.
- [12] Cooke A, Zurif EB, DeVita C, Alsop D, Koenig P, Detre J, et al. Neural basis for sentence comprehension: Grammatical and short-term memory components. Hum Brain Mapp 2002;15:80–94. doi: https://doi.org/10.1002/hbm.10006.
- [13] Daneman M, Carpenter PA. Individual differences in working memory and reading. J Verbal Learn Verbal Behav 1980;19:450-66. doi: https://doi.org/10.1016/S0022-5371(80)
- [14] Davis SW, Zhuang J, Wright P, Tyler LK. Age-related sensitivity to task-related modulation of language-processing networks. Neuropsychologia 2014;63:107-15. doi: https://doi.org/10.1016/j.neuropsychologia.2014.08.017.
- [15] Desmond JE, Fiez JA. Neuroimaging studies of the cerebellum: language, learning and memory. Trends Cognit Sci 1998;2:355–62. doi: https://doi.org/10.1016/S1364-6613(98)01211-X.
- [16] Diachek E, Blank I, Siegelman M, Affourtit J, Fedorenko E. The Domain-General Multiple Demand (MD) Network Does Not Support Core Aspects of Language Comprehension: A Large-Scale fMRI Investigation. J Neurosci 2020;40:4536–50. doi: https://doi.org/ 10.1523/INEUROSCI.2036-19.2020.
- [17] Diaz MT, Johnson MA, Burke DM, Madden DJ. Age-related Differences in the Neural Bases of Phonological and Semantic Processes. J Cognit Neurosci 2014;26:2798–811. doi: https://doi. org/10.1162/jocn a 00665.

- [18] Diaz MT, Rizio AA, Zhuang J. The Neural Language Systems That Support Healthy Aging: Integrating Function, Structure, and Behavior. Language Linguistics Compass 2016;10:314–34. doi: https://doi.org/10.1111/lnc3.12199.
- [19] Duncan J. The multiple-demand (MD) system of the primate brain: mental programs for intelligent behaviour. Trends Cognit Sci 2010;14:172-9. doi: https://doi.org/10.1016/i.tics.2010.01.004.
- [20] Erb J, Henry MJ, Eisner F, Obleser J. The Brain Dynamics of Rapid Perceptual Adaptation to Adverse Listening Conditions. J Neurosci 2013;33:10688–97. doi: https://doi.org/10.1523/JNEUROSCI.4596-12.2013.
- [21] Fedorenko E, Behr MK, Kanwisher N. Functional specificity for high-level linguistic processing in the human brain. Proc Natl Acad Sci 2011;108:16428–33. doi: https://doi.org/10.1073/pnas.1112937108.
- [22] Fedorenko E, Duncan J, Kanwisher N. Language-Selective and Domain-General Regions Lie Side by Side within Broca's Area. Curr Biol 2012;22:2059–62. doi: https://doi.org/10.1016/j.cub.2012.09.011.
- [23] Fedorenko E, Hsieh P-J, Nieto-Castañón A, Whitfield-Gabrieli S, Kanwisher N. New Method for fMRI Investigations of Language: Defining ROIs Functionally in Individual Subjects. J Neurophysiol 2010;104:1177–94. doi: https://doi.org/10.1152/jn.00032.2010.
- [24] Ferreira F, Patson ND. The 'Good Enough' Approach to Language Comprehension. Language Linguistics Compass 2007;1:71–83. doi: https://doi.org/10.1111/j.1749-818X.2007.00007.x.
- [25] Ficek BN, Wang Z, Zhao Y, Webster KT, Desmond JE, Hillis AE, et al. The effect of tDCS on functional connectivity in primary progressive aphasia, NeuroImage: Clinical 2018;19:703–15. doi: https://doi.org/10.1016/j.nicl.2018.05.023.
- [26] Friederici AD. Evolution of the neural language network. Psychon Bull Rev 2017;24:41–7. doi: https://doi.org/10.3758/s13423-016-1090-x.
- [27] Friston KJ, Worsley KJ, Frackowiak RSJ, Mazziotta JC, Evans AC. Assessing the significance of focal activations using their spatial extent. Hum Brain Mapp 1994;1:210–20. doi: https://doi.org/10.1002/hbm.460010306.
- [28] Geva S, Jones PS, Crinion JT, Price CJ, Baron J-C, Warburton EA. The Effect of Aging on the Neural Correlates of Phonological Word Retrieval. J Cognit Neurosci 2012;24:2135–46. doi: https://doi.org/10.1162/jocn_a_00278.
- [29] Giannini LAA, Peterson C, Ohm D, Xie SX, McMillan CT, Raskovsky K, et al. Frontotemporal lobar degeneration proteinopathies have disparate microscopic patterns of white and grey matter pathology. Acta Neuropathol Commun 2021;9:30. doi: https://doi.org/10.1186/s40478-021-01129-2.
- [30] Giannini LAA, Xie SX, McMillan CT, Liang M, Williams A, Jester C, et al. Divergent patterns of TDP-43 and tau pathologies in primary progressive aphasia. Ann Neurol 2019;85:630-43. doi: https://doi.org/10.1002/ana.25465.
- [31] Gibson E. Linguistic complexity: locality of syntactic dependencies. Cognition 1998;68:1–76. doi: https://doi.org/10.1016/S0010-0277 (98)00034-1.
- [32] Gorgolewski KJ, Varoquaux G, Rivera G, Schwarz Y, Ghosh SS, Maumet C, et al. NeuroVault.org: a web-based repository for collecting and sharing unthresholded statistical maps of the human brain. Front Neuroinform 2015;9:8. doi: https://doi.org/ 10.3389/fninf.2015.00008.
- [33] Grady C. The cognitive neuroscience of ageing. Nat Rev Neurosci 2012;13:491–505. doi: https://doi.org/10.1038/nrn3256.
- [34] Grossman M. The non-fluent/agrammatic variant of primary progressive aphasia. Lancet Neurol 2012;11:545–55. doi: https://doi.org/10.1016/S1474-4422(12)70099-6.
- [35] Grossman M, Cooke A, DeVita C, Chen W, Moore P, Detre J, et al. Sentence Processing Strategies in Healthy Seniors with Poor Comprehension: An fMRI Study. Brain Lang 2002;80:296–313. doi: https://doi.org/10.1006/brln.2001.2581.
- [36] Hagoort P. On Broca, brain, and binding: a new framework. Trends Cognit Sci 2005;9:416–23. doi: https://doi.org/10.1016/j.tics.2005.07.004.
- [37] Henry ML, Hubbard HI, Grasso SM, Mandelli ML, Wilson SM, Sathishkumar MT, et al. Retraining speech production and fluency in non-fluent/agrammatic primary progressive aphasia. Brain 2018;141:1799–814. doi: https://doi.org/10.1093/brain/awy101.
- [38] Hertrich I, Dietrich S, Ackermann H. The role of the supplementary motor area for speech and language processing. Neurosci Biobehav Rev 2016;68:602–10. doi: https://doi.org/10.1016/ji.neubiorev.2016.06.030.

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- [39] Hoyau E, Boudiaf N, Cousin E, Pichat C, Fournet N, Krainik A, et al. Aging Modulates the Hemispheric Specialization during Word Production. Front Aging Neurosci 2017;9. doi: https://doi.org/10.3389/fnagi.2017.00125.
- [40] Jansen A, Flöel A, Randenborgh JV, Konrad C, Rotte M, Förster A-F, et al. Crossed cerebro-cerebellar language dominance. Hum Brain Mapp 2005;24:165–72. doi: https://doi.org/10.1002/hbm.20077.
- [41] Just MA, Carpenter PA. A capacity theory of comprehension: Individual differences in working memory. Psychol Rev 1992;99:122–49. doi: https://doi.org/10.1037/0033-295X.99.1.122.
- [42] Justus T. The Cerebellum and English Grammatical Morphology: Evidence from Production, Comprehension, and Grammaticality Judgments. J Cognit Neurosci 2004;16:1115–30. doi: https://doi.org/10.1162/0898929041920513.
- [43] Koen JD, Rugg MD. Neural Dedifferentiation in the Aging Brain. Trends Cognit Sci 2019;23:547–59. doi: https://doi.org/10.1016/j.tics.2019.04.012.
- [44] Kuznetsova A, Brockhoff PB, Christensen RHB. ImerTest Package: Tests in Linear Mixed Effects Models. J Stat Softw 2017;82:1–26. doi: https://doi.org/10.18637/jss.v082.i13.
- [45] Lacombe J, Jolicoeur P, Grimault S, Pineault J, Joubert S. Neural changes associated with semantic processing in healthy aging despite intact behavioral performance. Brain Lang 2015;149:118–27. doi: https://doi.org/10.1016/j.bandl.2015.07.003.
- [46] Lee Y-S, Min NE, Wingfield A, Grossman M, Peelle JE. Acoustic richness modulates the neural networks supporting intelligible speech processing. Hear Res 2016;333:108–17. doi: https://doi.org/10.1016/j.heares.2015.12.008.
- [47] Lewis RL, Vasishth S, Van Dyke JA. Computational principles of working memory in sentence comprehension. Trends Cognit Sci 2006;10:447–54. doi: https://doi.org/10.1016/j.tics.2006.08.007.
- [48] Matar S, Pylkkänen L, Marantz A. Left occipital and right frontal involvement in syntactic category prediction: MEG evidence from Standard Arabic. Neuropsychologia 2019;135:. doi: https://doi.org/10.1016/j.neuropsychologia.2019.107230107230.
- [49] Matchin W, Hickok G. The Cortical Organization of Syntax. Cereb Cortex 2020;30:1481–98. doi: https://doi.org/10.1093/cercor/bhz180.
- [50] Matchin WG. A neuronal retuning hypothesis of sentence-specificity in Broca's area. Psychon Bull Rev 2018;25:1682–94. doi: https://doi.org/10.3758/s13423-017-1377-6.
- [51] Meinzer M, Flaisch T, Wilser L, Eulitz C, Rockstroh B, Conway T, et al. Neural Signatures of Semantic and Phonemic Fluency in Young and Old Adults. J Cognit Neurosci 2009;21:2007–18. doi: https://doi.org/10.1162/jocn.2009.21219.
- [52] Moore-Parks EN, Burns EL, Bazzill R, Levy S, Posada V, Müller R-A. An fMRI study of sentence-embedded lexical-semantic decision in children and adults. Brain Lang, Special Issue Lang Dev 2010;114:90-100. doi: https://doi.org/10.1016/j.bandl.2010.03.009.
- [53] Pallier C, Devauchelle A-D, Dehaene S. Cortical representation of the constituent structure of sentences. Proc Natl Acad Sci 2011:108:2522-7. doi: https://doi.org/10.1073/pnas.1018711108.
- [54] Park DC, Lautenschlager G, Hedden T, Davidson NS, Smith AD, Smith PK. Models of visuospatial and verbal memory across the adult life span. Psychol Aging 2002;17:299–320.
- [55] Peelle JE. Listening Effort: How the Cognitive Consequences of Acoustic Challenge Are Reflected in Brain and Behavior. Ear Hear 2018;39:204–14. AUD.00000000000000494.
- [56] Peelle JE, Troiani V, Wingfield A, Grossman M. Neural processing during older adults' comprehension of spoken sentences: age differences in resource allocation and connectivity. Cereb Cortex 2010;20:773–82. doi: https://doi.org/10.1093/cercor/bhp142.
- [57] Peelle JE, Wingfield A. The neural consequences of age-related hearing loss. Trends Neurosci 2016;39:486–97. doi: https://doi.org/10.1016/j.tins.2016.05.001.
- [58] Raizada RDS, Poldrack RA. Selective Amplification of Stimulus Differences during Categorical Processing of Speech. Neuron 2007;56:726–40. doi: https://doi.org/10.1016/j.neuron.2007.11.001.
- [59] Reuter-Lorenz PA, Lustig C. Brain aging: reorganizing discoveries about the aging mind. Curr Opin Neurobiol, Cognit Neurosci 2005;15:245–51. doi: https://doi.org/10.1016/j.conb.2005.03.016.
- [60] Rogalsky C, Matchin W, Hickok G. Broca's area, sentence comprehension, and working memory: an fMRI study. Frontiers in Human Neuroscience 2, 2008.
- [61] Rorden C, Brett M. Stereotaxic display of brain lesions. Behav Neurol 2000;12:191–200.

- [62] Sakai K, Passingham RE. Prefrontal interactions reflect future task operations. Nat Neurosci 2003;6:75–81. doi: https://doi.org/10.1038/nn987.
- [63] Santi A, Grodzinsky Y. Working memory and syntax interact in Broca's area. NeuroImage 2007;37:8–17. doi: https://doi.org/10.1016/i.neuroimage.2007.04.047.
- [64] Schwartze M, Kotz SA. Contributions of cerebellar event-based temporal processing and preparatory function to speech perception. Brain Lang, Contrib Cereb Lang Funct 2016;161:28–32. doi: https://doi.org/10.1016/i.bandl.2015.08.005.
- [65] Schwartze M, Rothermich K, Kotz SA. Functional dissociation of pre-SMA and SMA-proper in temporal processing. NeuroImage 2012;60:290-8. doi: https://doi.org/10.1016/j.neuroimage.2011.11.089.
- [66] Schwarzbauer C, Davis MH, Rodd JM, Johnsrude I. Interleaved silent steady state (ISSS) imaging: A new sparse imaging method applied to auditory fMRI. NeuroImage 2006;29:774–82. doi: https://doi.org/ 10.1016/j.neuroimage.2005.08.025.
- [67] Shafto MA, Tyler LK. Language in the aging brain: The network dynamics of cognitive decline and preservation. Science 2014;346:583-7. doi: https://doi.org/10.1126/science.1254404.
- [68] Shannon RV, Zeng F-G, Kamath V, Wygonski J, Ekelid M. Speech Recognition with Primarily Temporal Cues. Science 1995;270:303-4. doi: https://doi.org/10.1126/science.270.5234.303.
- [69] Silveri MC, Leggio MG, Molinari M. The cerebellum contributes to linguistic production: A case of agrammatic speech following a right cerebellar lesion. *Neurology* 44, 1994, 2047–2047. doi:10.1212/ WNL.44.11.2047.
- [70] Sliwinska MWW, Khadilkar M, Campbell-Ratcliffe J, Quevenco F, Devlin JT. Early and Sustained Supramarginal Gyrus Contributions to Phonological Processing. Front Psychol 2012;3. doi: https://doi.org/10.3389/fpsyg.2012.00161.
- [71] Stoodley CJ, Schmahmann JD. Functional topography in the human cerebellum: A meta-analysis of neuroimaging studies. NeuroImage 2009;44:489–501. doi: https://doi.org/10.1016/j.neuroimage.2008.08.039.
- [72] Strick PL, Dum RP, Fiez JA. Cerebellum and Nonmotor Function. Annu Rev Neurosci 2009;32:413–34. doi: https://doi.org/10.1146/annurev.neuro.31.060407.125606.
- [73] Tyler LK, Marslen-Wilson W. Fronto-temporal brain systems supporting spoken language comprehension. Philos Trans R Society B: Biol Sci 2008;363:1037-54. doi: https://doi.org/10.1098/stb.2007.2158.
- [74] Tyler LK, Shafto MA, Randall B, Wright P, Marslen-Wilson WD, Stamatakis EA. Preserving Syntactic Processing across the Adult Life Span: The Modulation of the Frontotemporal Language System in the Context of Age-Related Atrophy. Cereb Cortex 2010;20:352–64. doi: https://doi.org/10.1093/cercor/bhp105.
- [75] Ulrich M, Hoenig K, Grön G, Kiefer M. Brain Activation during Masked and Unmasked Semantic Priming: Commonalities and Differences. J Cognit Neurosci 2013;25:2216–29. doi: https://doi. org/10.1162/jocn_a_00449.
- [76] Uttl B. North American Adult Reading Test: Age Norms, Reliability, and Validity. J Clin Exp Neuropsychol 2002;24:1123–37. doi: https://doi.org/10.1076/icen.24.8.1123.8375.
- [77] Vaden KI, Kuchinsky SE, Ahlstrom JB, Teubner-Rhodes SE, Dubno JR, Eckert MA. Cingulo-Opercular Function during Word Recognition in Noise for Older Adults with Hearing Loss. Exp Aging Res 2016;42:67–82. doi: https://doi.org/10.1080/0382.016.1108784.
- [78] Vagharchakian L, Dehaene-Lambertz G, Pallier C, Dehaene S. A Temporal Bottleneck in the Language Comprehension Network. J Neurosci 2012;32:9089–102. doi: https://doi.org/10.1523/JNEUROSCI.5685-11.2012.
- [79] van Ettinger-Veenstra HM, Ragnehed M, Hällgren M, Karlsson T, Landtblom A-M, Lundberg P, et al. Right-hemispheric brain activation correlates to language performance. NeuroImage 2010;49:3481–8. doi: https://doi.org/10.1016/ji.neuroimage.2009.10.041.
- [80] Verhaeghen P. Aging and vocabulary score: A meta-analysis. Psychol Aging 2003;18:332–9. doi: https://doi.org/10.1037/0882-7974.18.2.332.
- [81] Waters G, Caplan D. The relationship between age, processing speed, working memory capacity, and language comprehension. Memory 2005;13:403–13. doi: https://doi.org/10.1080/09658210344000459.
- [82] Waters GS, Caplan D. Age, working memory, and on-line syntactic processing in sentence comprehension. Psychol Aging 2001;16:128-44. doi: https://doi.org/10.1037/0882-7974.16.1.128.

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83] Wierenga CE, Benjamin M, Gopinath K, Perlstein WM, Leonard CM, Rothi LJG, et al. Age-related changes in word retrieval: Role of bilateral frontal and subcortical networks. Neurobiol Aging 2008;29:436–51. doi: https://doi.org/10.1016/j.neurobiolaging.2006.10.024.

[84] Wilke M, Lidzba K. Ll-tool: a new toolbox to assess lateralization in functional MR-data. J Neurosci Methods 2007;163:128–36. doi: https://doi.org/10.1016/j.jneumeth.2007.01.026.

- [85] Wilson SM, Dronkers NF, Ogar JM, Jang J, Growdon ME, Agosta F, et al. Neural Correlates of Syntactic Processing in the Nonfluent Variant of Primary Progressive Aphasia. J Neurosci 2010;30:16845–54. doi: https://doi.org/10.1523/JNEUROSCI.2547-10.2010.
- [86] Wingfield A, Grossman M. Language and the Aging Brain: Patterns of Neural Compensation Revealed by Functional Brain Imaging. J Neurophysiol 2006;96:2830-9. doi: https://doi.org/10.1152/in.00628.2006.
- [87] Wingfield A, McCoy SL, Peelle JE, Tun PA, Cox CL. Effects of Adult Aging and Hearing Loss on Comprehension of Rapid Speech Varying in Syntactic Complexity. J Am Acad Audiol 2006;17:487–97. doi: https://doi.org/10.3766/jaaa.17.7.4.
- [88] Wingfield A, Peelle JE, Grossman M. Speech Rate and Syntactic Complexity as Multiplicative Factors in Speech Comprehension by

- Young and Older Adults. Aging, Neuropsychol, Cognit 2003;10:310–22. doi: https://doi.org/10.1076/anec.10.4.310.28974.
- [89] Woolgar A, Hampshire A, Thompson R, Duncan J. Adaptive Coding of Task-Relevant Information in Human Frontoparietal Cortex. J Neurosci 2011;31:14592-9. doi: https://doi.org/10.1523/ INEUROSCI.2616-11.2011.
- [90] Worsley KJ, Evans AC, Marrett S, Neelin P. A three-dimensional statistical analysis for CBF activation studies in human brain. J Cereb Blood Flow Metab 1992;12:900–18. doi: https://doi.org/10.1038/icbfm.1992.127.
- [91] Zettin M, Cappa SF, D'amico A, Rago R, Perino C, Perani D, et al. Agrammatic speech production after a right cerebellar haemorrhage. Neurocase 1997;3:375–80. doi: https://doi.org/10.1080/13554799708411976.
- [92] Zevin JD, McCandliss BD. Dishabituation of the BOLD response to speech sounds. Behav Brain Funct 2005;1:4. doi: https://doi.org/10.1186/1744-9081-1-4.
- [93] Zhuang J, Johnson MA, Madden DJ, Burke DM, Diaz MT. Age-related differences in resolving semantic and phonological competition during receptive language tasks. Neuropsychologia 2016;93:189–99. doi: neuropsychologia.2016.10.016.

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