

Neural Correlates of Word Class Processing: An fMRI Study

Cynthia K. Thompson*^{†‡}, Borna Bornakdarpour*, Henrike K. Blumenfeld*, Stephen C. Fix*[#], Todd B. Parrish^{^‡}, Darren R. Gitelman^{†^‡}, M.-Marsel Mesulam^{†‡}

Departments of *Communication Sciences & Disorders, #Linguistics, Northwestern University, and [†]Neurology, [^]Radiology, & [‡]Cognitive Neurology and Alzheimer's Disease Center, Northwestern University Feinberg School of Medicine



INTRODUCTION

Aphasic individuals often show impairments that are specific to word class. Most notable are (double) dissociations between noun and verb production (Miceli et al., 1984, 1988; Zingeser & Berndt, 1990).

- Anomic aphasia: nouns more difficult than verbs
- Agrammatic aphasia: verbs more difficult than nouns

VERBS AND ARGUMENT STRUCTURE

In fact, the verb production impairment in agrammatic aphasia is more than an across-the-board impairment for all types of verbs. It is sensitive to several factors, such as lexical frequency, imageability, and semantic factors. Most notably, agrammatic verb production is a function of a verb's *argument structure*.

VERB ARGUMENT STRUCTURE

argument = participant in verb action

Possible argument structures [arguments in blue]:

- 1-place (intransitive) verb: John slept
- 2-place (transitive) verb: John punched Alex
- 3-place (ditransitive) verb: John gave money to Mary

For agrammatic speakers, verb production becomes more difficult as verb argument structures become more complex. That is, 3-place verbs are more difficult than 2-place verbs, which are more difficult than 1-place verbs (Figure 1). This finding has been reported in English (Thompson et al., 1997; Kim & Thompson, 2000, 2004; Thompson, 2003) as well as in other languages, such as Dutch (Jonkers & Bastiaanse, 1996, 1998), Italian (Luzzatti & Chierchia, in press), Hungarian (Kiss, 2000), and German (de Bleser & Kauschke, 2000).

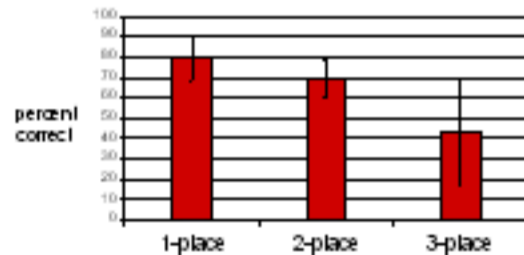


Figure 1. Aphasic production of verbs by argument structure. (Kim & Thompson, 2000; cf. Thompson et al., 1997; Kim & Thompson, 2004)

This verb processing impairment seen in agrammatism suggests that anterior left-hemisphere regions are crucial for processing verbs and verb argument structure. Some data support this notion:

- Petersen et al. (1988) reported left frontal activation in a PET verb production task.
- Shapiro et al. (2001) reported left frontal disruption of verb production but not noun production in a repetitive transcranial magnetic stimulation (rTMS) study.

However, posterior areas may participate in verb processing as well:

- Perani et al. (1999) compared, with fMRI, lexical decision of concrete and abstract nouns and verbs.

Verbs compared to nouns = frontal and posterior perisylvian activation

- Grossman et al. (2002), using fMRI, examined:

- motion verbs (*run, walk*) – observable events
- cognition verbs (*think, know*) – non-observable events; complex argument structure

Cognition minus motion verbs = posterior activation, including Wernicke's area

- Hadar & Palti (in press) reported greater Wernicke's area activation for verbs with more complex argument structures, in an fMRI study using a semantic judgment task.

HYPOTHESIS/EXPERIMENTAL DESIGN

QUESTION

Given these argument structure effects in aphasic populations as well as results from neuroimaging, what parts of the language network are recruited for verb argument structure processing?

HYPOTHESIS

We anticipated that increases in posterior perisylvian activation (e.g., Wernicke's area) would coincide with increases in verb argument structure complexity.

EXPERIMENTAL DESIGN

PARTICIPANTS

N = 15; 6 males, 9 females
Age range 20-30; mean = 25.2
Monolingual English speakers
Right-handed (Edinburgh Handedness Inventory); no familial left-handedness
No history of neurological/psychological problems

MATERIALS

Visually presented words/pseudowords: displayed for 1200 ms followed by 300 ms ITI

Table 1. Experimental stimuli.

	Total number	Mean frequency*
Animal names	40	9.3
Tool names	40	9.1
1-place verbs	40	9.5
2-place verbs	40	9.3
3-place verbs	40	9.65
Pseudowords	50	---

* per million, from the CELEX database (Rayson, Coxhead, & van Rijn, 1993)

Null events = 1700 or 3400 msec

65% of stimuli were nulls

Stimuli appeared in two runs of 6 min 4 sec (including 6 initial dummy scans in each)

Order pseudorandomized using OPTSEQ (<http://surfer.nmr.mgh.harvard.edu/optseq>)

SCANNING PARAMETERS

3T scanner (Trio, Siemens)

T2-weighted functional scan

Single-shot echo-planar imaging

24 slices parallel to AC-PC line

TR=2 sec

TE = 30 msec

flip angle = 90°

FOV = 220 mm

matrix = 64 x 64

slice thickness = 3 mm

T1-weighted anatomical scan

3D FLASH sequence with saturation

band inferior to image volume

TR= 2.1 sec

TE = 4.38 sec

flip angle = 20°

FOV = 220 mm

matrix = 256 x 256

slice thickness = 1 mm

DATA ANALYSIS

All analysis used SPM2 (Wellcome Dept. of Imaging Neuroscience, Inst. for Neurology, University College London) running in a Matlab 6.5 environment (Mathworks Inc., Natick, MA). Functional scans were corrected for slice-acquisition timing and realigned to a mean functional image. The anatomical volume was coregistered to the mean image, and normalized to the MNI 152-subject template brain (ICBM, NIH P-20 project). The functional volumes were then normalized using the same transformation and were smoothed using a 10mm (FWHM) isotropic kernel. Resulting stereotactic coordinates, in MNI-space, were transformed into Talairach space using a script written by Matthew Brett (<http://www.nrc-cbu.cam.ac.uk/Imaging/Common/mnispace.shtm>).

RESULTS

Scan data from five of the 15 subjects were discarded because of excess noise or motion artifact. The remaining subjects showed a clear pattern of perisylvian activation under verb processing conditions. These subjects were analyzed as a group.

VERB AND NOUN ACTIVATION

For all verbs taken together, activation was seen in the inferior frontal gyrus (Brodmann's Area [BA] 44, 47) and in the superior temporal gyrus (BA 22) and superior temporal sulcus (BA 21/22). Additional activation was evident in the posterior-superior regions of the parietal lobe (see Figures 2 and 3, and Table 2). Nouns, however, elicited more widespread, and little perisylvian, activation (see Figure 2).

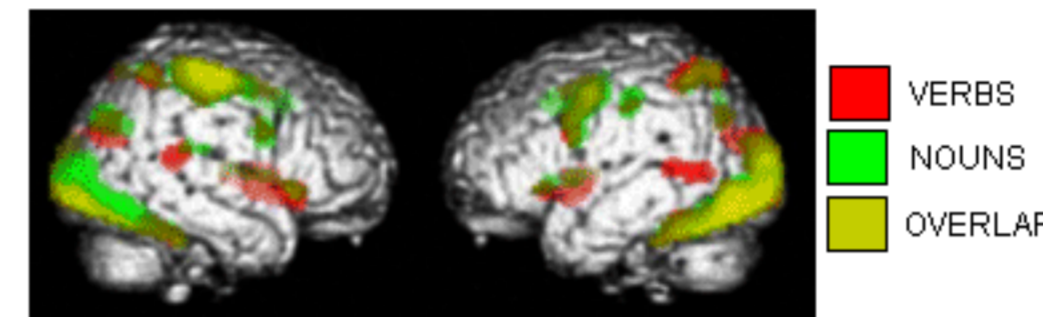


Figure 2. Lateral images showing activation for verb conditions and noun conditions. Verb conditions are red, noun conditions are green, overlap is yellow. Activations are significant $p < .001$ (uncorrected) at a 3 voxel extent threshold.

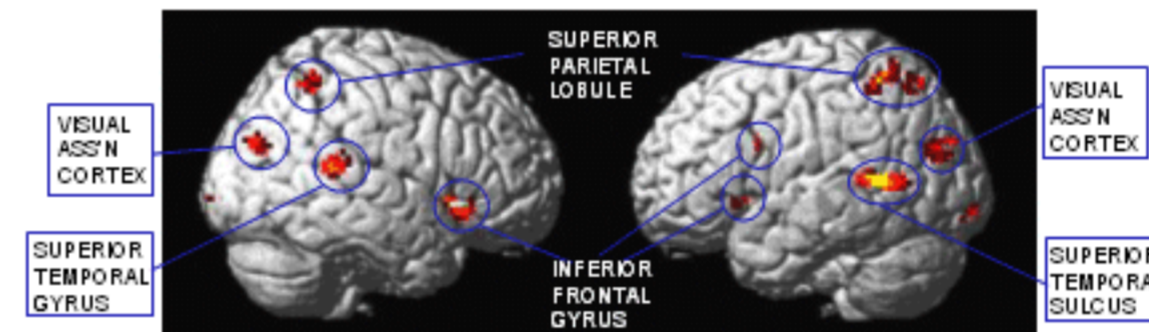


Figure 3. Lateral images showing activation for verb conditions as compared to activation for noun conditions. Activations are significant $p < .001$ (uncorrected) at a 3 voxel extent threshold.

Table 2. Regions of statistically significant activation under verb processing conditions as compared to noun conditions.

Brodmann Area	Hemisphere	Side	MNI Coordinates		Z Value	Brodmann Area	Hemisphere	Side	MNI Coordinates		Z Value	
			x	y					x	y		
Inferior Frontal Gyrus (Brodmann's Area)	44	Left	-36	10	3.32	Visual Association Cortex	19	Left	-21	-71	2.6	
Inferior Frontal Gyrus	47	Right	30	14	2.9	Superior Temporal Gyrus	19	Right	30	14	2.9	
Inferior Frontal Gyrus	47	Left	-30	21	4	3.31	Parietal	17	Left	-55	66	3
Inferior Frontal Gyrus	47	Right	33	21	4	3.33	Superior Temporal Gyrus	17	Right	6	69	2
Superior Temporal Sulcus (Wernicke's Area)	22	Left	-59	42	8	4.36	Superior Parietal Lobule	7	Left	-24	47	3.2
Superior Temporal Gyrus (Wernicke's Area)	22	Right	65	37	13	4.01	Superior Parietal Lobule	7	Left	-24	47	3.2
Posterior Gyrus	31	Left	-36	40	10	3.88	Superior Temporal Gyrus	17	Right	21	62	3.0
							Superior Temporal Gyrus	17	Right	42	51	3.12
							Superior Temporal Gyrus	17	Right	31	40	3.2

Note: BA = Brodmann's area. Voxel-wise (uncorrected) $p < .001$, cluster size = 3.

ARGUMENT STRUCTURE ACTIVATION

Comparison of activation for 3-place verbs to activation for both 1-place and 2-place verbs revealed activation in Wernicke's area (BA 22) bilaterally (Figure 4; Table 3). Similar activation was evident when 3-place verbs were compared to 1-place verbs, and when 3-place verbs were compared to 2-place verbs.

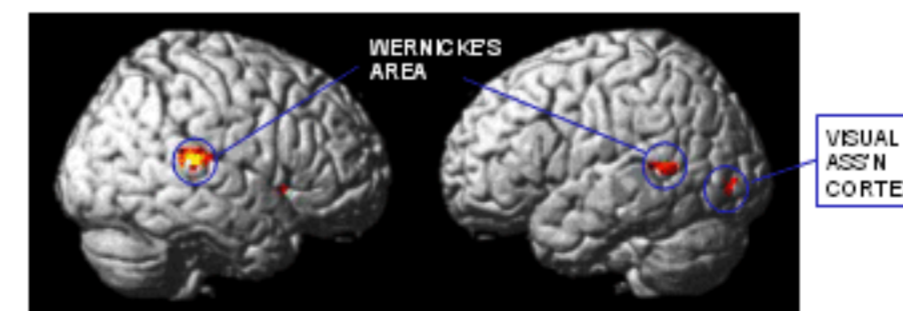


Figure 4. Lateral images showing activation for the 3-place verb condition compared to activation during 1- and 2-place verb conditions. Activations are significant $p < .001$ (uncorrected) at a 3 voxel extent threshold.

Table 3. Regions of statistically significant activation under 3-place verb condition as compared to 1-place and 2-place verb conditions.

Brodmann Area	Hemisphere	Side	MNI Coordinates		Z Value
			x	y	
Midle Temporal Gyrus (Wernicke's Area)	22	Left	-53	49	3.31
Superior Temporal Gyrus (Wernicke's Area)	22	Right	62	34	3.87
Posterior Gyrus	31	Left	-36	44	3.36
Posterior Gyrus	31	Left	-27	42	3.38
Posterior Gyrus	31	Right	30	42	4.11
Visual Association Cortex	19	Left	-21	62	3.44

Note: BA = Brodmann's area. Voxel-wise (uncorrected) $p < .001$, cluster size = 3.

DISCUSSION

Anterior perisylvian cortex (including Broca's area) and posterior perisylvian cortex (including Wernicke's area) were active in verb conditions but not in noun conditions. This role of Broca's area is consistent with results indicating processing of syntactic information associated with verbs (Friederici et al., 2000; Newman et al., in press; Perani et al., 1999; Warburton et al., 1996).

Activation of posterior regions, including Wernicke's area, is consistent with previous data suggesting that Wernicke's area is involved in verb processing (the aforementioned studies of Perani et al., Grossman et al., and Hadar & Palti). Most notably, the Hadar & Palti and Grossman et al. studies correlated Wernicke's area activation with increased verb argument structure complexity.

Patient studies provide another line of evidence. In several studies, Wernicke's aphasic subjects demonstrated inability to process aspects of argument structure information. McCann & Edwards (2000) found an inability to detect argument structure violations (e.g., *John gives a car*) in Wernicke's aphasic patients. Shapiro & Levine (1990) reported results of an online sentence processing task in which verb argument structure was manipulated. Wernicke's patients (but not Broca's patients) showed a lack of sensitivity to argument structure, suggesting that they were unable to access argument structure information. This in turn suggests that Wernicke's area is necessary for verb argument structure processing.

Together with these findings, our data indicate that posterior portions of the language network are essential for processing verb argument structure.

REFERENCES

Beynon, R. H., Piepenbrock, R., & van Rijn, H. (1993). *The CELEX Lexical Database (CD-ROM)*. Philadelphia, PA: University of Pennsylvania Linguistics Data Consortium.

De Bleser, R., & Havelka, C. (2000). Acquisition and loss of nouns and verbs: Parallel or divergent patterns? Paper presented at the British Psychological Society, Cognitive Psychology Section XVII Annual Conference, University of Essex.

Friederici, A., Meyer, M., von Cramon, D.Y. (2000). Auditory language comprehension: An event-related fMRI study on the processing of syntactic and lexical information. *Brain and Language*, 75, 289-300.

Grossman, M., Koenig, P., DeVita, C., et al. (2002). The neural representation of verb meaning: An fMRI study. *Human Brain Mapping*, 13(2), 124-134.

Hadar, U., & Palti, D. (in press). The divergence of grammatical category and grammatical gender in lexical processing: An fMRI investigation.

Jonkers, R., & Bastiaanse, R. (1994). The influence of instrumentality and transitivity on action naming in Broca's and anomic aphasia. *Brain and Language*, 55, 37-39.

Jonkers, R., & Bastiaanse, R. (1998). How selective are selective word class deficits? Two case studies of action and object naming. *Aphasiology*, 3, 245-254.

Kim, M., & Thompson, C.K. (2004). Verb deficit in Alzheimer's disease and agrammatism: Implications for lexical organization. *Brain and Language*, 88, 1-20.

Kim, M., & Thompson, C.K. (2000). Patterns of comprehension and production of nouns and verbs in agrammatism: Implications for lexical organization. *Brain and Language*, 74, 1-25.

Kir, K. (1999). Effects of verb complexity on agrammatic aphasic sentence production. In R. Bastiaanse & Y. Grodzinsky (eds.), *Grammatical disorders in aphasia* (pp. 152-170). London: Wm. & W.

Luzzatti, C., & Chierchia, G. (in press). On the nature of selective deficit involving nouns and verbs. *Italian Journal of Linguistics*.

McCann, C., & Edwards, J. (2002). Verb problems in frontal aphasia. *Brain and Language*, 85, 1-20.

Mitch, G., Silvari, M., Vilh, G., & Caramazza, A. (1984). On the basis for the agrammatic's difficulty in producing main verbs. *Cortex*, 20, 207-220.

Mitch, G., Silvari, M., Nocentini, U., & Caramazza, A. (1988). Patterns of dissociation in comprehension and production of nouns and verbs. *Aphasiology*, 2, 351-358.

Newman, A., Paulsen, R., Olson, K., et al. (in press). An event-related fMRI study of syntactic and semantic violations. *Journal of Psycholinguistic Research*, 31.

Peterson, S., Fox, P., Perner, M., et al. (1988). Perimenstrual studies of the cortical anatomy of single word processing. *Nature*, 331, 585-589.

Perani, D., Cappa, S., Schnur, I., et al. (1999). The neural correlates of verb and noun processing: A PET study. *Brain*, 122, 2337-2344.

Shapiro, K., Pascual-Leone, A., Mottaghy, F., et al. (2001). Grammatical distinctions in the left frontal cortex. *J Cognitive Neuroscience*, 13, 713-720.

Shapiro, L., & Levine, B. (1990). Verb processing during sentence comprehension in aphasia. *Brain and Language*, 38, 21-47.

Thompson, C.K., Lange, K., Schneider, J., & Shapiro, L. (1997). A grammatical and non-brain-damaged subject's verb and argument structure production. *Aphasiology*, 11, 473-490.

Thompson, C.K. (2003). Unaccusative verb production in agrammatic aphasia: The argument structure complexity hypothesis. *J Neurolinguistics*, 16, 151-147.

Warburton, E., Wise, R., Price, C., et al. (1994). Noun and verb retrieval by normal subjects: Studies with PET. *Brain*, 119, 179-179.

Zingeser, L., & Berndt, R. (1990). Retrieval of nouns and verbs in agrammatism and anomia. *Brain and Language*, 39, 14-32.

ACKNOWLEDGMENT

Supported by NIH grant DC01948-1 (CKT).