

Lesion mapping of syntactic and lexical features derived from Natural Language Processing of narrative speech elicited by patients with chronic post-stroke aphasia

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Background

- Language assessment in patients with aphasia poststroke is usually conducted with batteries of standardized tests aimed at identifying residual language capacities. Many of these tests rely on patients' responses to isolated stimuli, sometimes threatening their ecological validity in certain conditions.
- Narrative/connected speech, however, can provide a rich source of response samples to obtain insightful information about patients' language skills. The analysis of such data, however, is highly timeconsuming and requires specific training.
- We sought to analyze connected speech in patients with chronic stroke using Natural Language Processing to identify measures sensitive to aphasia and to understand their neurobiological bases with VLSM.

Methods

- 65 participants (66.2 % M) who had a lefthemisphere stroke at least 6 months ago (19 without residual aphasia, 18 Broca's, 14 anomic, 9 conduction, 3 Wernicke's, and 1 global). Mean age was 58.5 (SD = 10.3) and mean months post-stroke was 37.2 (SD = 40.2) months.
- 10 age- and gender- comparable health controls with no history of neurological or psychiatric disease
- All participants completed the WAB-R and were asked to describe, for 2 minutes each, three scenes:
 - 1. Cookie Theft scene from the BDAE
 - 2. Picnic scene from the WAB
 - 3. Circus scene from the Apraxia Battery for Adults
- Discourse was transcribed verbatim

By means of sentence boundary disambiguation, parsing, and parts-of-speech tagging, we derived 37 lexical and syntactic features

We applied Principal Components Analysis (PCA) on these features and applied voxel-based lesionsymptom mapping (VLSM) to identify the neural basis associated with principal components that accounted for > 50% of the variance.

Principal Component Analysis		 Factor 1 was strongly
1	2	composed of mainly syntactic
44.8%	11.5%	
.954	110	features and VLSM was thus
.949	084	reatures and vesivi was thus
.926	.063	
.923	.154	controlled for WAB fluency
.919	133	(in decreasing order of loading scores: claus
.917	.258	
.917	.258	width, width/height and number of verbal
.907	.123	phrases, number of words, speech rate,
.903	010	•
.897	110	number of clauses and their height, distance
	Compo 1 44.8% .954 .949 .926 .923 .919 .917 .917 .907 .903	Components 1 2 44.8% 11.5% .954 110 .949 084 .926 .063 .923 .154 .919 133 .917 .258 .917 .258 .907 .123 .903 010

Numb Diff Words

VP to VP distance

Length of Sentence

Clause Per Sentence

oken Type Ratio

Number of Preposition

Dept Clause Per Clause

Noun Phrases

Mean Imagery

Lexical Density

Numb Nouns

Numb Adverbs

Adverb Variation

Noun Variation

Verb Variation

Word Length

Mean AoA

Lexical Variation

Mean Length Clause

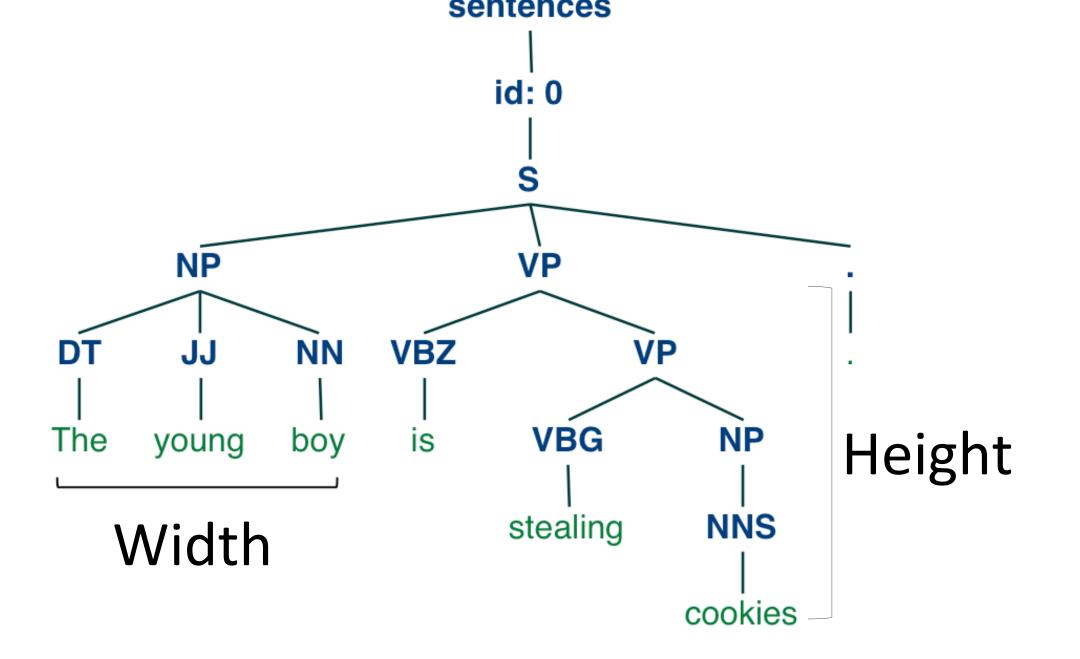
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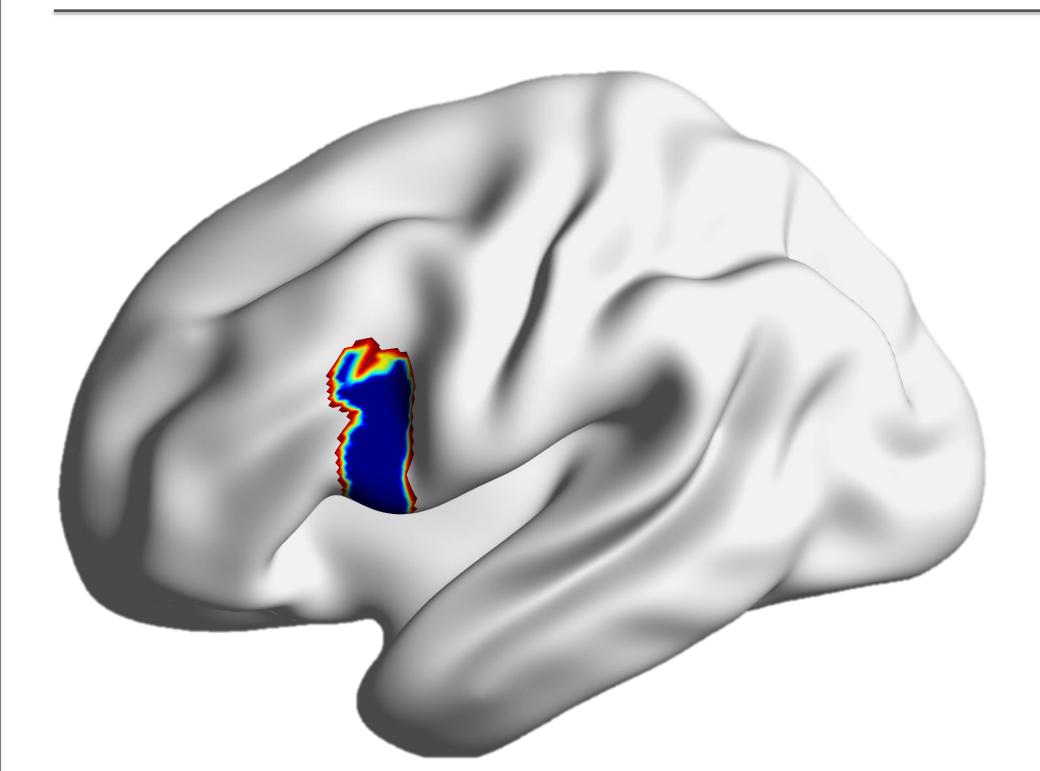
Factor 2 was strongly composed of mainly lexical features and VLSM was thus controlled for WAB naming (in decreasing order of loading scores: frequency of all words, number of nouns, adverbs, and adjectives, as well as adverb, noun, verb and overall lexical variation)

between noun phrase and verb phrases, etc.)

Methods & Results

Sample parsing of the sentence "The boy is stealing cookies" demonstrating some features of syntactic complexity such as width (in this case, the NP has a width of 3) and height (in this case, the VP has a height of 2)

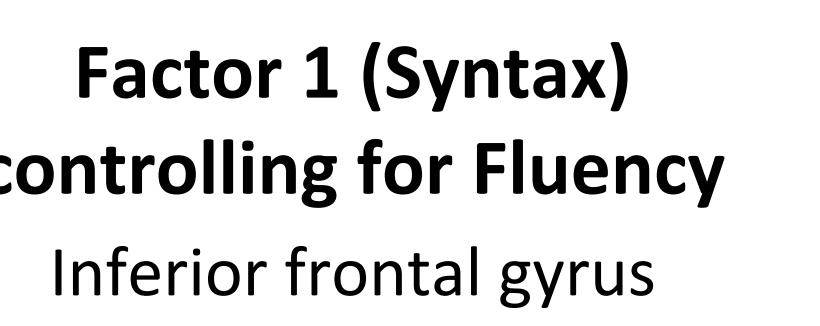




controlling for Fluency

Factor 2 (Lexical) controlling for Naming

> Precentral gyrus Postcentral gyrus Pos superior temporal gyrus Supramarginal gyrus



Conclusions

We showed that syntactic performance beyond speech fluency might rely on the inferior frontal gyrus (Broca's area), while lexical performance beyond fluency may depend on areas in the precentral and post-central gyri as well as the postero-superior temporo and inferio-parietal regions. Our findings show that NLP applied to connected speech elicited by patients with post-stroke aphasia can shed light on the organization of language in brains with vascular damage.