## **Supplementary Information**

# Neural encoding and production of functional morphemes in the posterior temporal lobe

Daniel K. Lee<sup>1\*</sup>, Evelina Fedorenko<sup>2,3,4\*</sup>, Mirela V Simon<sup>5</sup>, William T Curry<sup>1</sup>, Brian Nahed<sup>1</sup>, Dan P Cahill<sup>1</sup>, Ziv M Williams<sup>1,6,7</sup>

- 1. Department of Neurosurgery, Harvard Medical School, Massachusetts General Hospital, Boston MA
- 2. Department of Psychiatry, Harvard Medical School, Massachusetts General Hospital, Boston MA
- 3. Harvard Program in Speech and Hearing Bioscience, Cambridge MA
- 4. Massachusetts Institute of Technology, McGovern Institute for Brain Research, Cambridge MA
- 5. Department of Neurology, Harvard Medical School, Massachusetts General Hospital, , Boston MA
- 6. Harvard-MIT Division of Health Sciences and Technology, Boston MA
- 7. Harvard Medical School, Program in Neuroscience, Boston MA

\* These authors contributed equally to the work

### SUPPLEMENTARY TABLES

Patient	Testing	Lesion Location	cm <sup>2</sup>	Pathology	
1	Stimulation	Anterior frontal	121	Oligoastrocytoma	
2	Stimulation	Superior temporal	61	Oligoastrocytoma	
3	Stimulation	Temporal, parietal	28	Amyloid Angiopathy	
4	Stimulation	Frontal, temporal	293	Astrocytoma	
5	Stimulation	Temporal	141	Glioblastoma	
6	Stimulation	Temporal	138	Astrocytoma	
7	Stimulation	Temporal	183	Oligodendroglioma	
8	Stimulation	Temporal	83	Astrocytoma	
9	Stimulation	Parietal	57	Astrocytoma	
10	Stimulation	Temporal	27	Ganglioglioma	
11	Stimulation	Temporal	184	Epidermoid Cyst	
12	Stimulation	Temporal	50	Oligoastrocytoma	
13	Stimulation	Temporal, parietal	14	Glioblastoma	
14	Stimulation	Temporal	27	Metastatic Melanoma	
15	Recording	Temporal	5	Metastatic Melanoma	
16	Recording	Frontal, temporal	46	Glioblastoma	
17	Recording	Temporal, parietal	30	Glioblastoma	
18	Recording	Temporal, parietal	54	Glioblastoma	
19	Recording	Temporal	19	Glioblastoma	
20	Lesion	Temporal*	2	Glioblastoma	
21	Lesion	Temporal*	2	Glioblastoma	
22	Lesion	Temporal	3	Glioblastoma	
23	Lesion	Temporal	9	Astrocytoma	
24	Lesion	Temporal	17	Metastatic Carcinoma	
25	Lesion	Temporal	n/a	Mesial temporal sclerosis	
26	Lesion	Temporal	n/a	Mesial temporal sclerosis	
27	Lesion	Temporal	n/a	Mesial temporal sclerosis	

**Supplementary Table 1. Testing modality, lesion location and size, and underlying pathology.** Participants are color-coded by testing modality: stimulation (light green), recording (light blue), or post-operative lesion evaluation (\*the first two patients had P-STG lesions). Lesion size is represented in cubic centimeters, and approximate anatomical locations are provided. All lesions involved the left dominant hemisphere. As noted in the Supplement text, there was no difference in the main neurophysiological findings based on patient demographics or pathology.

Condition	Target	Reg/Irreg	Number	Sentence	Input	Correct
			Tense	preamble	word	production
Control	Noun	Regular	Singular	This is the	apple	apple
Syntactic	Noun	Regular	Plural	These are the	apple	apples
Syntactic	Noun	Regular	Singular	This is the	apples	apple
Control	Noun	Regular	Plural	These are the	apples	apples
Control	Noun	Irregular	Singular	This is the	scarf	scarf
Syntactic	Noun	Irregular	Plural	These are the	scarf	scarves
Syntactic	Noun	Irregular	Singular	This is the	scarves	scarf
Control	Noun	Irregular	Plural	These are the	scarves	scarves
Control	Verb	Regular	Future	Today we will	talk	talk
Syntactic	Verb	Regular	Past	Yesterday, we	talk	talked
Syntactic	Verb	Regular	Future	Today we will	talked	talk
Control	Verb	Regular	Past	Yesterday, we	talked	talked
Control	Verb	Irregular	Future	Today we will	run	run
Syntactic	Verb	Irregular	Past	Yesterday, we	run	ran
Syntactic	Verb	Irregular	Future	Today we will	ran	run
Control	Verb	Irregular	Past	Yesterday, we	ran	ran

**Supplementary Table 2**. **Syntactic manipulations.** Examples of the different types of trials in the critical syntactic condition, where participants had to morpho-syntactically manipulate the target word prior to production (targets shown in red) and the control condition, where participants had to simply read the target word out loud (targets shown in green). The trials varied in whether the target word was a) a noun or a verb, b) morphologically regular or irregular, and c) singular or plural for nouns, or in the past vs. future tense for verbs. The latter manipulation resulted in some cases where a morpheme had to be *added* and other cases where it had to be *subtracted* to produce the correct word.

#### SUPPLEMENTARY FIGURES



**Supplementary Figure 1. Pre-operative baseline performance for participants in the stimulation and recording components.** =Performance was near ceiling across conditions across patients (*n*=19 total). To test for potential effects of task demands on performance, we additionally compared pre-operative baseline performance to performance during stimulation based on (i) suffix addition vs. subtraction comparison, and (ii) evaluation of between-condition differences in pre-operative performance (i.e., 2-4% error vs. 0-2% error). As detailed in the main text, the results suggested that there was little variation in effect in the P-STG based on task demands. Error bars indicate +/- 95% CI s.e.m..



#### Supplementary Figure 2. Stimulated sites and elicited deficits on a patient-by-

**patient level.** 54 sites were tested across 14 participants, transformed here to a standard MNI152 cortical atlas. Stimulation sites by participants are color-coded, as indicated in the legend. Sites eliciting a selective morpho-syntactic deficit in functional morpheme production vs. a global linguistic deficit are circled in red and green, respectively. Here, 9 of the 14 patients had specific access to the P-STG.



**Supplementary Figure 3. Participants with anterior stimulation sites.** Four of the 14 participants had craniotomies which covered a small portion of the inferior frontal gyrus. The sites tested in these patients are colored in red. Sites that involved frontal cortex are circled.



**Supplementary Figure 4. Spatial distribution of deficits during lexical and semantic conditions.** Stimulation sites that produced a deficit during the both the lexical and semantic conditions are displayed in light green. Sites that produced a deficit during the lexical but not semantic condition are shown in dark green. The shaded regions are calculated based on an average 1 cubic centimeter area estimated to be affected by focal stimulation.

ECOG Processing Workflow



**Supplementary Figure 5. ECOG processing workflow.** Electrocorticography data were notch-filtered and normalized for each channel separately. The raw signals (blue) were notch-filtered at 60 Hz (orange) to remove land-line noise, and, in some patients, at 105 Hz and 20 Hz, due to instruments in the operative field. To maximize the signal-to-noise ratio, data were normalized by subtraction of the average baseline (i.e. time periods between trials) (green) for each channel.



-1500 to -500ms

-500 to 500 ms

500 to 1500 ms

Supplementary Figure 6. The spatial-temporal activity dynamic of syntacticsensitive sites. The times are based on a 1 second window centered before, during, and after speech onset (t=0). Any site with significant differentiation between syntactic and control conditions (permutation test,  $n_{perm}$ =3000, p<0.05) within the 1 second window are drawn.



-1500 to -500 ms

-500 to 500 ms

500 to 1500 ms

Supplementary Figure 7. Time-course and location of functional interactions. The results from a sample participant of the functional interactions observed between a site within the P-STG (red) and other neighboring sites (blue) over time. Only significant interactions are shown (time-series analysis,  $n_{pair}=15$ , p<0.05). The times are based on a 1 second window centered before, during, and after speech onset (t=0).