Faster hemodynamic response latencies to speech in posterior superior temporal gyrus

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Summary

Recent electrocorticography (ECoG) recordings of speech processing in superior temporal gyrus (STG) have found evidence of neural selectivity to speech onsets in posterior STG (pSTG), with response latencies like those in Heschl's gyrus (HG) [1]. The high spatiotemporal resolution of ECoG enables precise determination of the latency and spatial encoding of fast and transient speech features, such as speech onsets. However, the invasive nature of ECoG limits its applicability to a broader population. In this study, we investigate the hemodynamic response to speech onsets using functional magnetic resonance imaging (fMRI) across

areas in STG. Here we explore the location and timing of brain activation to speech onsets. Models of speech onsets during a passive listening task show a significant response to speech onsets in pSTG, with predicted hemodynamic responses peaking faster in pSTG and HG than aSTG.



Methods

Participants: 19 adult native English-speaking controls, with normal speech, language, hearing, and reading abilities. Language-Localizer Task : Participants listened passively to audio recordings of natural speech (podcasts, interviews) in 18-s blocks of intact and acoustically degraded conditions. **MRI Data Acquisition:** Whole-brain structural and functional data were acquired using a 3T Siemens Trio Scanner. EPI fMRI parameters: TR=750ms, 45 axial slices, 5-slice SMS, 3mm³ voxels, 484 TRs.

Data Analysis: Speech onsets from the "speech" blocks of the language localizer were automatically defined by periods of 750 ms or more of preceding silence. The onsets were convolved with a canonical hemodynamic response function (HRF) and its temporal derivative, which accounts for the rate of change in the hemodynamic response. These speech onsets were incorporated into a mass univariate linear model. Fitted responses with higher temporal derivative beta values indicate hemodynamic responses that peak faster.





We focused on 3 principal anatomical regions of interest: HG; pSTG; and aSTG. [2]

Fast hemodynamic response latencies to speech onsets in pSTG

Subject 1: The profile of strong model significance suggests speech onsets are good predictors of BOLD responses in pSTG. The temporal derivative parameter of the HRF was generally higher in pSTG than aSTG, and the time to peak tended to be earlier in **pSTG** and **HG** than in **aSTG**.



Subject 3 & Subject 4: Speech onset models continue to explain activation in pSTG well; however, the pattern of fitted HRFs differs from those of Subjects 1 and 2. Subject 3 showed HRFs with negative responses, and Subject 4 showed no clear relationship between the HRF coefficients and faster peak times in pSTG. For these subjects, the fastest times to peak were in HG with no overall difference between aSTG and pSTG. In future work we will consider alternative HRF basis sets to better characterize neural responses to fast speech events [3].

References: [1] Hamilton et al. (2021). *Cell*; [2] Tourville et al. (2019). J. Speech Lang. Hear. Res. [3] Polimeni and Lewis (2021). Prog. Neurobiol. **Contact:** aamoore@bu.edu

Funding: This project was supported by NIH grants T32 DC01301 (C. Stepp); R03 HD096098 and R03 DC014045 (T. Perrachione); F32 DC019531 (T. Scott); and an NSF GRFP (to A. Moore)

Subject 2: The model again showed high significance in pSTG, suggesting responsiveness to speech onsets. The temporal derivative parameter of the HRF tended to have higher values in pSTG than aSTG. In this subject, time to peak tended to be earlier in HG than pSTG, but both of these were faster than aSTG.



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