

Do motor symptoms and antipsychotic medications influence the digital speech assessment of negative symptoms in schizophrenia spectrum disorders?

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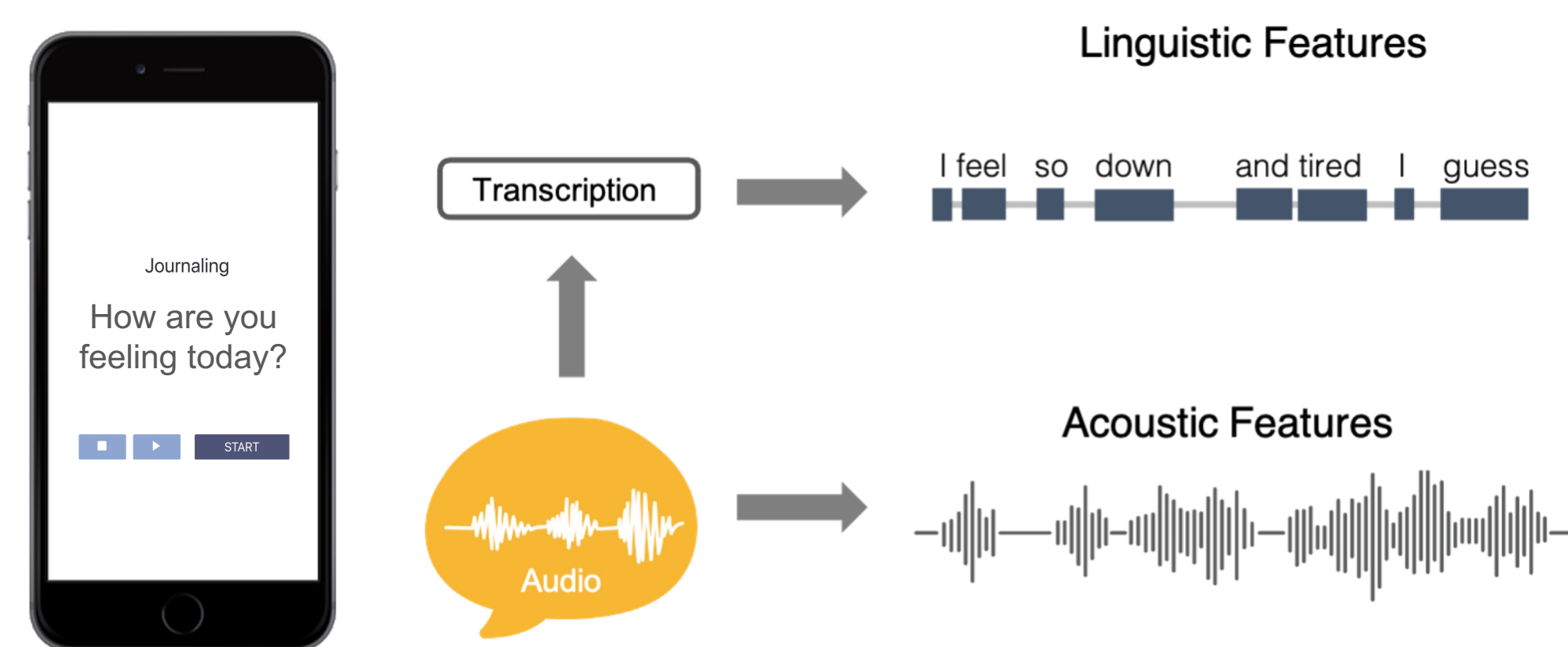
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Background

- Computational speech analysis may help provide a quantitative assessment of negative symptoms in schizophrenia and overcome limitations of traditional symptom scales.
- However, the influence of potential confounding clinical variables (e.g., comorbid motor symptoms) has not yet been systematically ruled out, which is necessary for the clinical validation of computational speech metrics.
- This study examined whether motor symptoms and antipsychotic medication were associated with speech markers of negative symptoms in participants with schizophrenia spectrum disorders (SSD).**

Methods

- Participants:** 43 inpatients with SSD
- Clinical Assessments:** Negative symptoms (Scale for the Assessment of Negative Symptoms; SANS); motor symptoms (Extrapyramidal Symptom Rating Scale; ESRs).
- Speech tasks (from the Winterlight assessment app):** Journaling (x2), Picture Description (x3), Phonemic Fluency, Semantic Fluency, Paragraph Reading.
- Quantitative speech features:** 8 acoustic and 10 timing variables extracted for each participant from transcribed speech recordings.

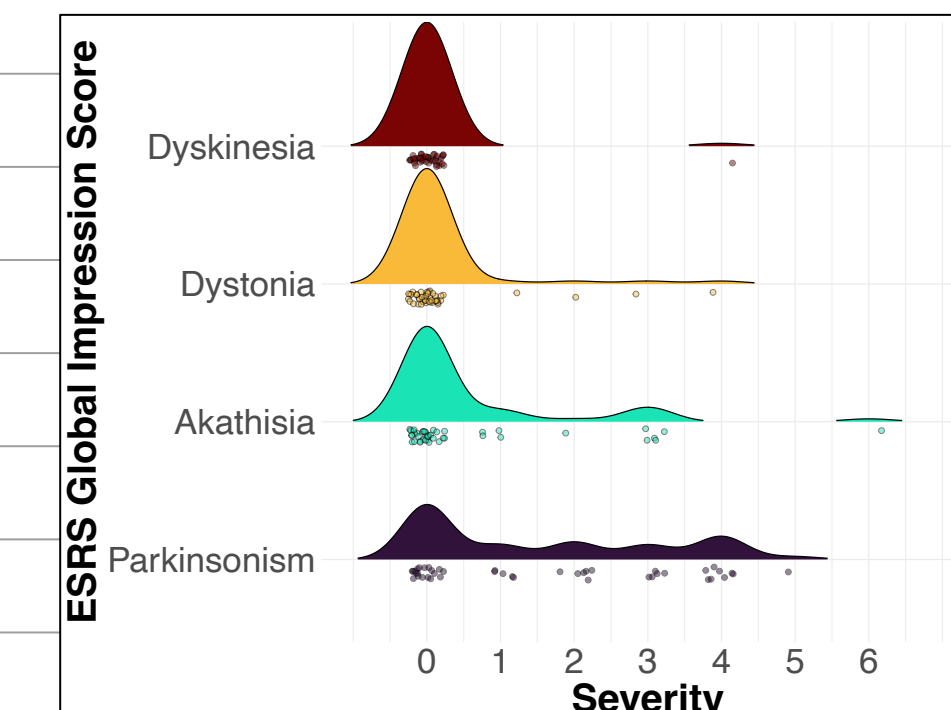


- Analyses:** Associations (age- and sex-adjusted partial Kendall rank correlations) were evaluated between quantitative speech features and the following: negative symptoms (SANS), motor symptoms (ESRS global impression scores), and antipsychotic medication dose (chlorpromazine equivalent; CPZE).
- Statistical significance: set at $p < .05$, FDR-corrected within task.
- Bayesian analyses were used to further evaluate evidence for the absence of associations between speech and motor symptoms or antipsychotic medication dose.

Participant characteristics

Age (years)	26.16 (5.08)
Sex	14 female, 29 male
Race/Ethnicity	21% Asian, 47% Black, 16% White-not Hispanic, 9% Multiple, 7% Other
Education (years)	14.19 (1.97)
Diagnosis	3 bipolar disorder w/ psychosis, 11 schizoaffective, 20 schizophrenia, 1 schizophreniform, 8 unspecified psychotic disorder
BPRS Total	48.47 (11.58)
SANS Total	26.58 (9.98)
ESRS Parkinsonism	1.53 (1.67)
ESRS Akathisia	0.56 (1.26)
ESRS Dystonia	0.23 (0.81)
ESRS Dyskinesia	0.09 (0.61)
CPZE dose	289.66 (220.54)

Note. Means and standard deviations are reported where relevant.

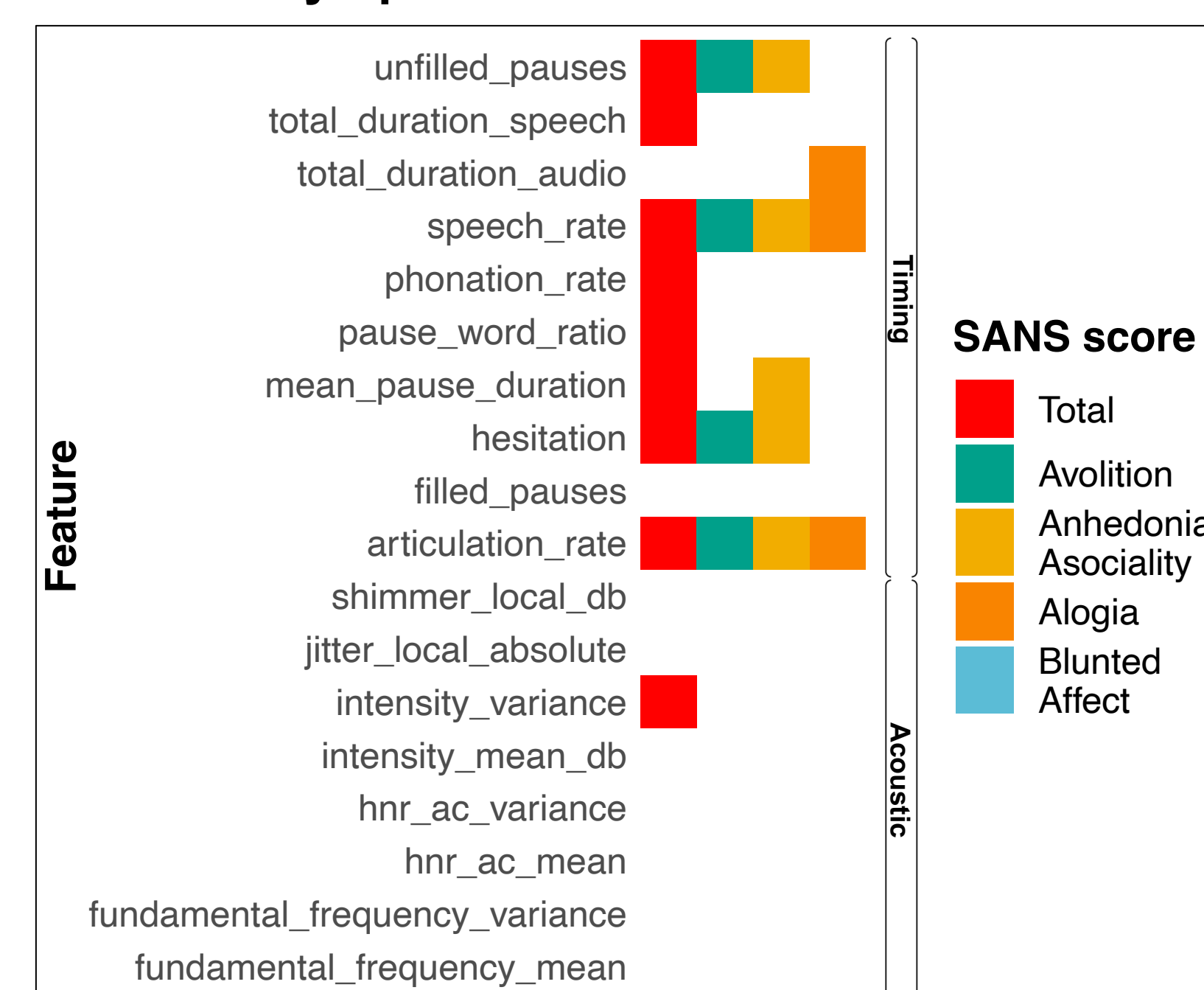


Highlights

Quantitative speech features are sensitive to negative symptom severity in schizophrenia spectrum disorders and do not appear to be confounded by motor symptoms or antipsychotic medications.

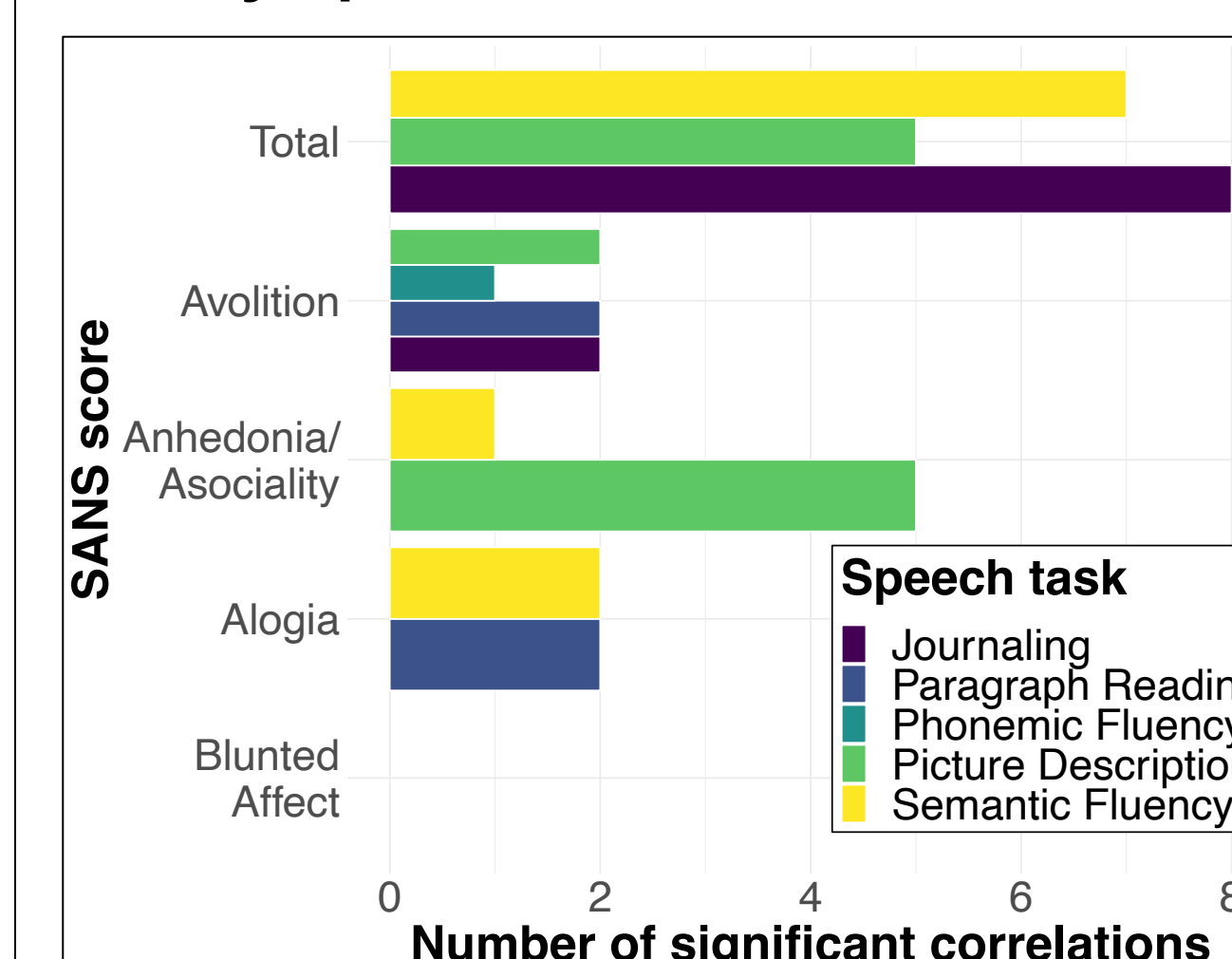
Results: Speech and negative symptoms

Features significantly associated with SANS symptoms on at least one task



- The majority of significant feature correlations were for timing features, with most associations present across multiple tasks.
- Association strength was in the small-to-medium range for positive (Kendall's tau = 0.21 to 0.31) and negative (Kendall's tau = -0.37 to -0.20) correlations.

Number of significant correlations between speech features and negative symptom scores in each task



Results: Speech, motor sx, and antipsychotics

- No significant correlations between speech features and parkinsonism, akathisia, or CPZE dose after FDR correction.
- Bayesian analyses indicated no more than anecdotal support for the alternative hypothesis for all but one feature: mean fundamental frequency (with akathisia; moderate support; $BF_{10} = 3.56$), which was not associated with negative symptoms.
- Bayes factors otherwise indicated moderate support ($BF_{01} > 3$) for the absence of an association between speech and motor symptoms or CPZE dose for most features.

Bayesian tests for correlations with relevant speech features

Feature	ESRS Akathisia	ESRS Parkinsonism	CPZE dose
Unfilled pauses^a (picture description)	$BF_{10} = 0.2$ $BF_{01} = 5.01$	$BF_{10} = 0.205$ $BF_{01} = 4.876$	$BF_{10} = 0.291$ $BF_{01} = 3.434$
Total duration speech^a (semantic fluency)	$BF_{10} = 0.407$ $BF_{01} = 2.456$	$BF_{10} = 0.201$ $BF_{01} = 4.979$	$BF_{10} = 0.162$ $BF_{01} = 6.154$
Total duration audio^a (paragraph reading)	$BF_{10} = 0.459$ $BF_{01} = 2.176$	$BF_{10} = 0.272$ $BF_{01} = 3.676$	$BF_{10} = 0.268$ $BF_{01} = 3.737$
Speech rate (semantic fluency)	$BF_{10} = 0.234$ $BF_{01} = 4.273$	$BF_{10} = 0.21$ $BF_{01} = 4.76$	$BF_{10} = 0.335$ $BF_{01} = 2.988$
Phonation rate^a (semantic fluency)	$BF_{10} = 0.407$ $BF_{01} = 2.456$	$BF_{10} = 0.201$ $BF_{01} = 4.979$	$BF_{10} = 0.162$ $BF_{01} = 6.154$
Pause word ratio (journaling)	$BF_{10} = 0.206$ $BF_{01} = 4.856$	$BF_{10} = 0.206$ $BF_{01} = 4.856$	$BF_{10} = 0.177$ $BF_{01} = 5.648$
Mean pause duration^a (journaling)	$BF_{10} = 0.562$ $BF_{01} = 1.778$	$BF_{10} = 0.684$ $BF_{01} = 1.462$	$BF_{10} = 0.307$ $BF_{01} = 3.255$
Hesitation^a (semantic fluency)	$BF_{10} = 0.338$ $BF_{01} = 2.96$	$BF_{10} = 1.86$ $BF_{01} = 0.538$	$BF_{10} = 0.264$ $BF_{01} = 3.787$
Articulation rate^a (picture description)	$BF_{10} = 0.198$ $BF_{01} = 5.06$	$BF_{10} = 0.198$ $BF_{01} = 5.06$	$BF_{10} = 0.161$ $BF_{01} = 6.22$
Intensity variance (semantic fluency)	$BF_{10} = 0.303$ $BF_{01} = 3.3$	$BF_{10} = 0.203$ $BF_{01} = 4.93$	$BF_{10} = 0.173$ $BF_{01} = 5.793$

Note. Bayesian tests are reported for speech from the task demonstrating the strongest association with SANS Total score (or subscore when the correlation with Total score was not significant).

BF_{01} = Bayes factor for the null hypothesis; BF_{10} = Bayes factor for the alternative hypothesis.

Bayes Factor (BF) interpretation: 1-3 = anecdotal support; 3-10 = moderate support; > 10 = strong support.

^aFeatures that showed a significant ($pFDR < .05$) correlation with negative symptoms and a significant (uncorrected $p < .05$) correlation with motor symptoms or CPZE dose.

Conclusions

- Speech features are sensitive to negative symptom severity in SSD and do not appear to be confounded by motor symptoms or antipsychotic dose.
- Additional research in samples with more severe motor symptoms and that examines other factors influencing speech (e.g., culture) will help to further validate computational speech-based assessment of negative symptoms.