

TWO-MINUTE VOCAL TEST AND ACOUSTIC ANALYSIS REVEAL VOICE AND SPEECH DISORDERS IN EARLY UNTREATED PARKINSON'S DISEASE

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BACKGROUND and AIMS

Disorders of voice and speech in Parkinson's disease (PD) affect all subsystems including respiration, phonation, articulation, and prosody [1-3]. Quick vocal test consisting of sustained phonation, fast syllable repetition, and running speech was designed in order to be gender independent [4]. Main aim of this study was to separate early untreated PD from healthy control (HC) participants based upon automated acoustic analysis [5].

PATIENTS and DATA

24 PD speakers (20 men & 4 women)

- examined before the symptomatic treatment was started
- age $60.9 \pm SD 11.2$ years
- duration of PD symptoms 31.3 ± 22.3 months
- H&Y stage 2.2 ± 0.5
- UPDRS III motor score 17.4 ± 7.1

22 HC speakers (15 men & 7 women)

- no history of neurological or communication disorders
- age 58.7 ± 14.6 years

Table I: Summary of the speech data. For reproducibility of data, each task was repeated at least 2 times.

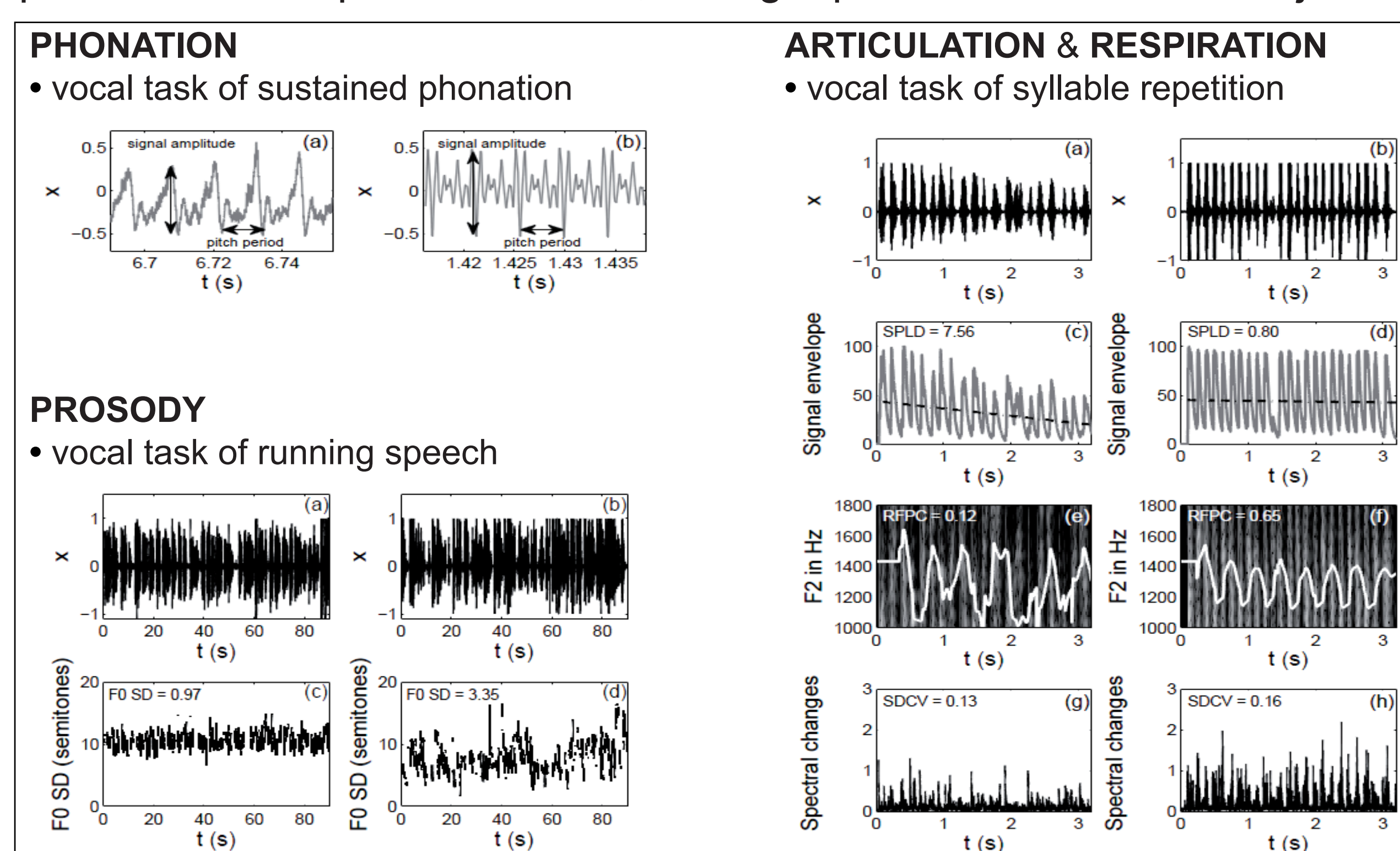
Task no	Speech data
[VT1]	Sustained phonation on one breath at a comfortable pitch and loudness as constant and long as possible, at least 5-sec.
[VT2]	Rapid steady /pa/-/ta/-/ka/ syllables repetition, called diadochokinetic (DDK) task, on one breath as constant and long as possible, repeated at least 5-times.
[VT3]	Running speech, at least approx. 80-sec.

METHODS

Table II: Overview of the measurement methods used.

Determined from task	Acoustic measurement	Acoustic measurement description
[VT1]	Jitter	Regular or irregular variations of glottal cycle duration, the average absolute difference between a period and the average of it and its four closest neighbours, divided by the average period.
[VT1]	Shimmer	Regular or irregular variations of amplitude maxima in subsequent glottal cycles, the average absolute difference between the amplitudes of consecutive periods, divided by the average amplitude.
[VT1]	NHR	Noise-to-Harmonics-Ratio, the amplitude of noise relative to tonal components.
[VT1]	HNR	Harmonics-to-Noise-Ratio, the amplitude of tonal relative to noise components.
[VT2]	SPLD	Sound Pressure Level Decline, the robust linear regression of the signal envelope.
[VT2]	RFPC	Robust Formant Periodicity Correlations, the first autocorrelation coefficient of F2 contour.
[VT2]	SDCV	Spectral Distance Change Variations, the variations of spectral distance changes in spectrum.
[VT3]	F0 SD	Variations of fundamental frequency (F0) in semitones, the variations of vibration rate of vocal folds.

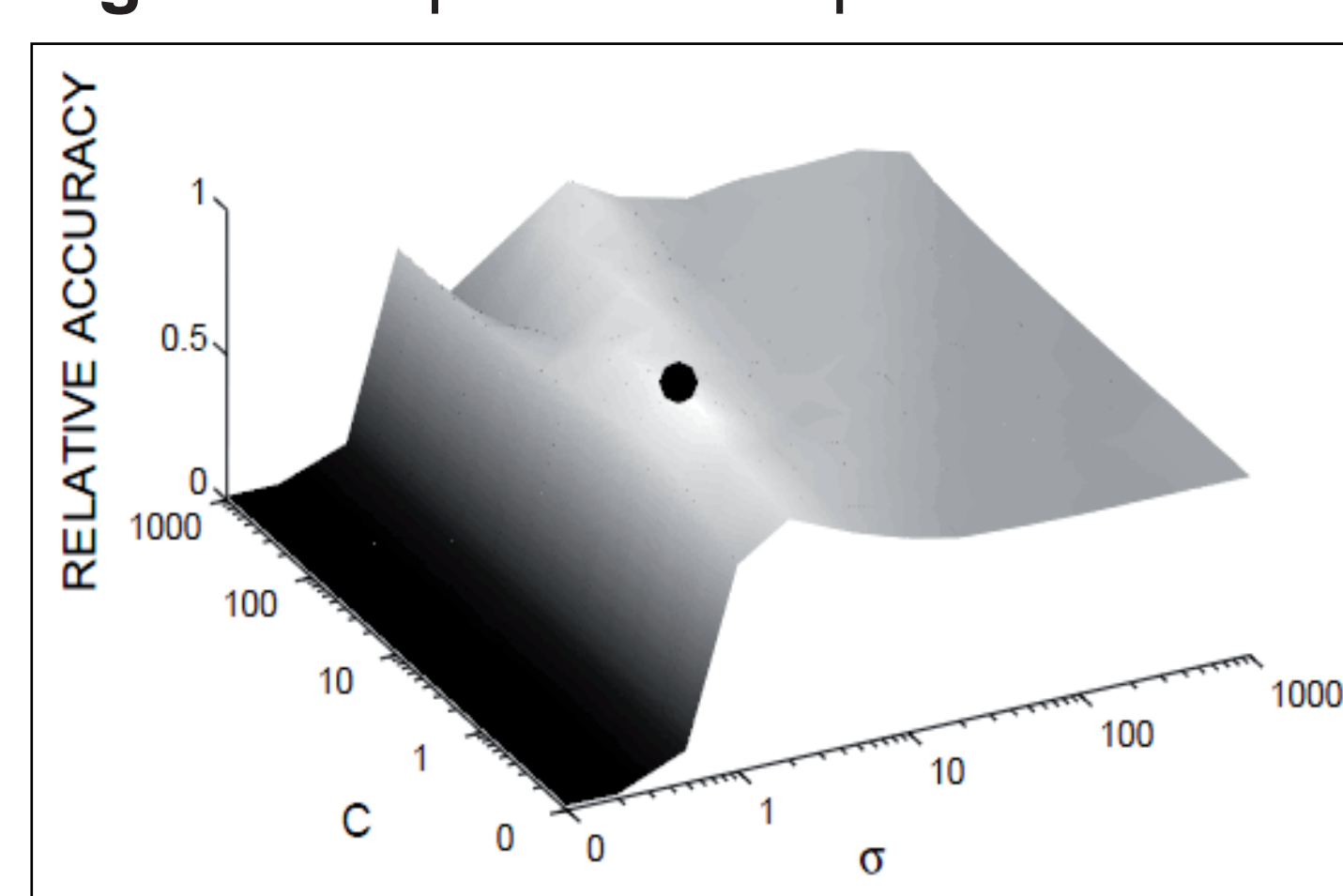
Figure 1: Details of measures used for subsequent analysis. The left panels are for a person with PD, the right panels are for a HC subject.



Statistics

- predictive model was built using a kernel support vector machine (SVM)
- exhaustive search of all possible measure combinations and optimal SVM parameters (C, σ)
- cross-validation with the leave-one-out method was used to validate reproducibility
- best combination of measurements was found to differentiate PD from HC subjects

Figure 2: Optimal SVM parameters.



RESULTS

- 116 vocal recordings were used for classification (56 for PD/60 for HC)
- relationships between measures of articulation & phonation and subscores of bradykinesia & rigidity

- best performance of 85.0 ± 6.1% in combination of four measures that represent all PD-related affected speech subsystems
- 81.3 ± 6.9% for running speech
- 75.6 ± 8.3% for sustained phonation
- 71.4 ± 8.3% for DDK task

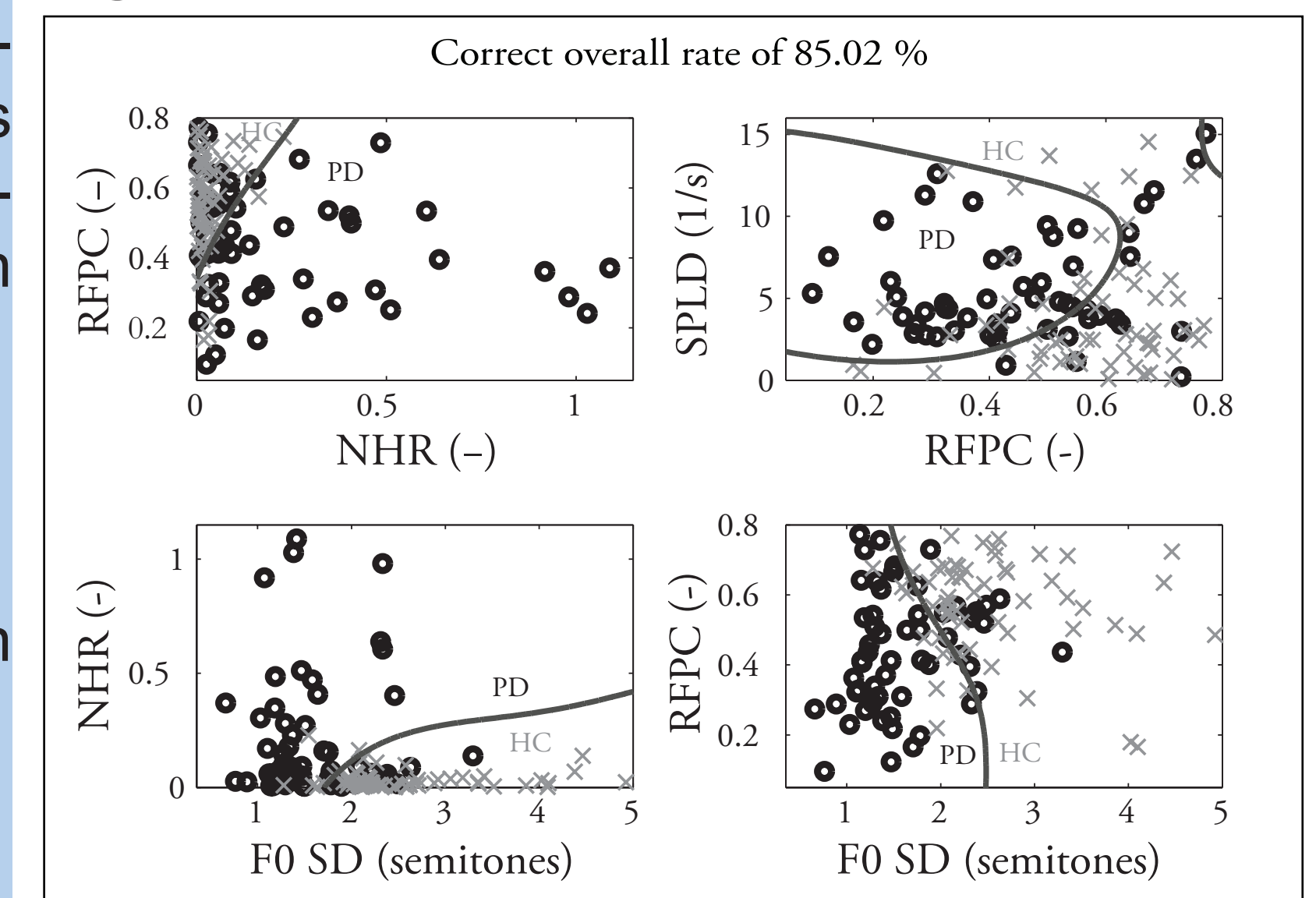
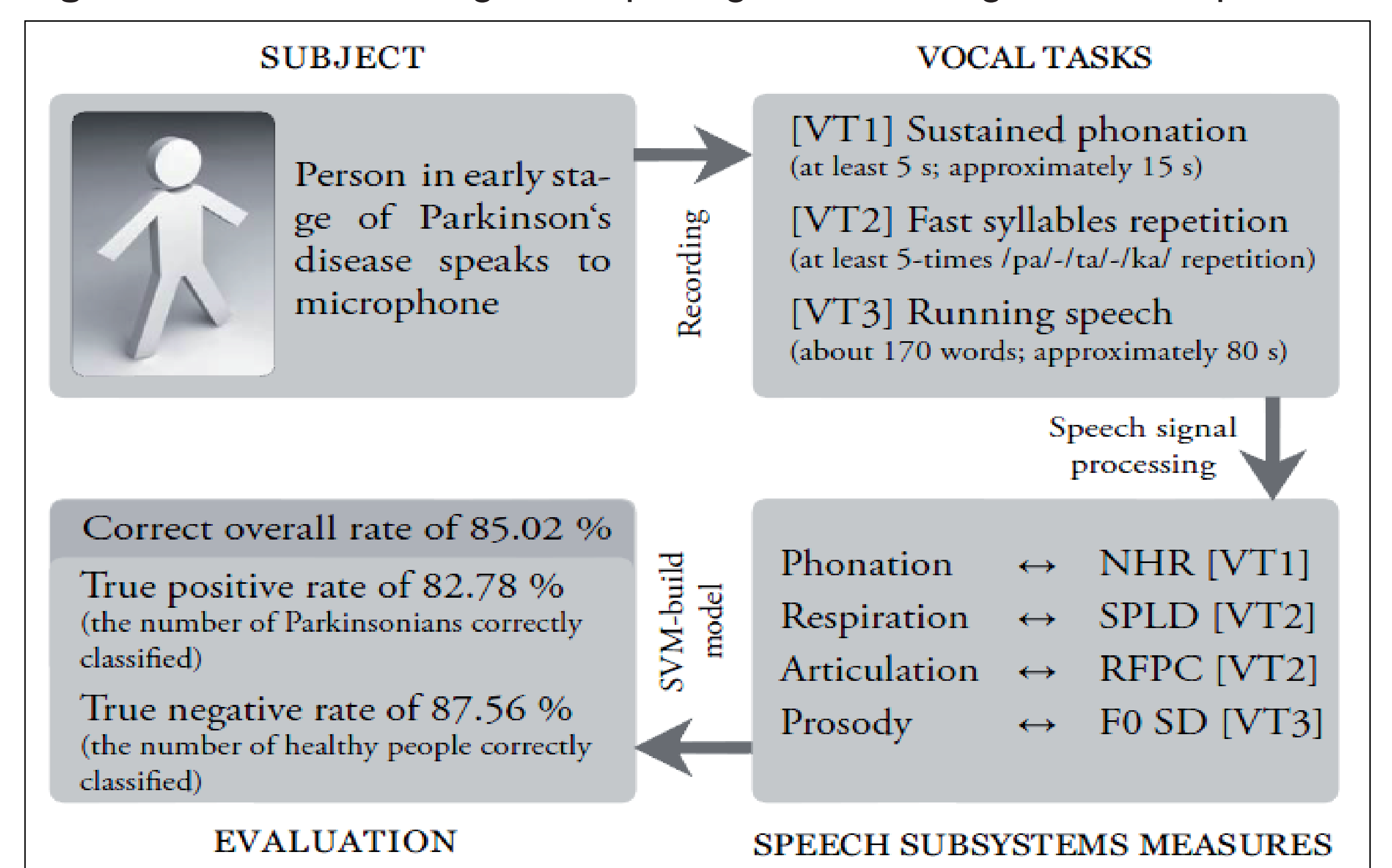


Table III: Result of the speech examination.

Measurement	Subjects		Differences between groups	Correlation between PD patients' speech performances and UPDRS III									
	PD Mean (SD)	HC Mean (SD)		A composite UPDRS III subscore of			Rigidity						
				total score	Bradykinesia	PIGD							
[VT1] Sustained phon.													
<i>Phonation</i>													
Jitter (%)	0.91	0.68	0.33	0.21	$P < .001$	$R = .31$	$P = .14$	$R = .42$	$P < .05$	$R = .21$	$P = .32$	$R = .24$	$P = .25$
Shimmer (%)	8.57	4.60	3.25	1.57	$P < .001$	$R = .28$	$P = .18$	$R = .33$	$P = .12$	$R = .01$	$P = .95$	$R = .35$	$P = .09$
NHR (-)	0.22	0.25	0.04	0.03	$P < .001$	$R = .36$	$P = .08$	$R = .43$	$P < .05$	$R = .21$	$P = .33$	$R = .39$	$P = .06$
HNR (dB)	14.05	6.01	22.55	4.28	$P < .001$	$R = -.40$	$P = .05$	$R = -.44$	$P < .05$	$R = -.14$	$P = .52$	$R = -.42$	$P < .05$
[VT2] DDK task													
<i>Respiration</i>													
SPLD (1/s)	5.68	2.99	3.85	3.01	$P < .05$	$R = -.12$	$P = .59$	$R = -.19$	$P = .38$	$R = -.37$	$P = .08$	$R = .16$	$P = .46$
<i>Articulation</i>													
RFPC (-)	0.43	0.14	0.58	0.10	$P < .001$	$R = -.33$	$P = .12$	$R = -.18$	$P = .40$	$R = -.13$	$P = .55$	$R = -.23$	$P = .28$
SDCV (-)	0.14	0.03	0.17	0.03	$P < .01$	$R = -.40$	$P = .05$	$R = -.44$	$P < .05$	$R = -.28$	$P = .19$	$R = -.28$	$P = .18$
[VT3] Running speech													
<i>Prosody</i>													
F0 SD (semitones)	1.52	0.43	2.62	0.75	$P < .001$	$R = .12$	$P = .56$	$R = .14$	$P = .53$	$R = .26$	$P = .22$	$R = .19$	$P = .37$

Figure 4: Schematic diagram depicting the recording of the PD speakers.



CONCLUSIONS

- method demonstrated that it can accurately differentiate PD patients from HCs
- subtle abnormalities such as reduced melody in running speech were detectable from the early stage of PD
- acoustic analysis may serve as a simple screening test in view of the expected advent of neuroprotective treatment
- acoustic vocal tests can be used for clinical monitoring of speech progression, effect of medication on speech production, and feedback in voice treatment

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