Title: The quantitative assessment of inter-utterance stability: application to dysarthria

Authors: Fred Cummins¹, Anja Lowit², and Frits van Brenk²

School of Computer Science & Informatics

Computer Science Building

Belfield

Dublin 4

Ireland

Strathclyde University

40 George Street

Glasgow G1 1QE

UK

Corresponding Author:

Anja Lowit, PhD

School of Psychological Sciences and Health

Strathclyde University

40 George Street

UK - Glasgow G1 1QE

a.lowit@strath.ac.uk

Tel. +44 141 548 3102

Fax +44 141 548 4001

¹ University College Dublin

² School of Psychological Sciences and Health

Abstract

Purpose: Following recent attempts to quantify articulatory impairment in speech, we evaluate the usefulness of a novel measure of motor stability to characterise dysarthria.

Method: We investigated 8 speakers with ataxic dysarthria (AD), 16 speakers with hypokinetic dysarthria (HD) as a result of Parkinson's Disease and 24 unimpaired control participants. Each participant performed a series of sentence repetitions under habitual, fast and slow speaking rate conditions. We used an algorithm to measure utterance-to-utterance spectro-temporal variation (UUV; Cummins, 2009). Speech rate and intelligibility were also measured.

Results: UUV scores were significantly correlated with perceptually based intelligibility scores. There were significant differences in UUV between control speakers and the ataxic, but not the HD groups, presumably due to differences in intelligibility in the samples employed, and not to differences in pathology. Habitual speaking rate did not correlate with UUV scores. All speaker groups had greater UUV levels in the slow conditions compared to habitual and fast speaking rates.

Conclusion: UUV results are consistent with those of other variability indices and thus appear to capture motor control issues in a similar way. The results suggest that the UUV could be developed into an easy to use clinical tool that could function as a valid and reliable assessment and outcome measure.

Introduction

The last few decades have seen a surge in the development of more and more refined techniques and measurements to investigate movement disorders. Such methods are useful, not only to shed more light on the symptoms of these disorders and their causes, but they can also be of significant clinical value in that they can support differential diagnosis of different types of motor speech problems, function as outcomes measures for treatment approaches, and allow clinicians to detect sub-clinical problems, thus enabling them to time treatment more effectively.

These methods are based on a wide variety of methodological paradigms. Noteworthy developments are papers by Liss and co-workers, who have published a number of methodologies over recent years that are successful in discriminating different types of dysarthric speech from each other and healthy control speakers (Liss, LeGendre, & Lotto, 2010; Liss et al., 2009; Utianski, Liss, Lotto, & Lansford, 2012). A further technique worth scrutinising is the variability measurement pioneered by Smith and co-workers (Smith, Goffman, Zelaznik, Ying, & McGillem, 1995) and subsequently expanded on by a number of other research groups. These have been used less to discriminate between different types of dysarthria, but more to highlight differences in motor control across healthy and disordered speakers as well as different elicitation paradigms.

Liss et al.'s (2009, 2010) and Utianski et al.'s (2012) research focused on a group of 70 speakers with four different types of dysarthria (ataxic, mixed spastic-flaccid, hypokinetic, and hyperkinetic) and healthy control speakers. They investigated the discriminating possibilities of a range of acoustic measures which focus both on the segmental and suprasegmental aspects of speech, namely rhythm (Liss, et al., 2009), amplitude envelope

modulations (EMS) (Liss, et al., 2010) and long-term average spectra (LTAS) (Utianski, et al., 2012). In each case, they were able to demonstrate their measures' ability to distinguish between healthy and impaired speakers, as well as between different types of dysarthric speakers with relatively good success. Their measures were thus able to reflect particular qualitative aspects of the speech signal that might be present in one but not another type of dysarthria. It became evident that some measures were more suitable to detect impairments in particular disorders, for example, some were distinguished better by rhythm measures if this was a particular feature of the disorder, whereas other dysarthria types were better discriminated with the amplitude measure. The authors thus advocate that the complete protocol of measures should be employed rather than just one measure in isolation. Such an application would be facilitated by the fact that they are all based on the same speech data, which makes the analysis more clinically viable by reducing assessment time. One of the drawbacks of Liss et al.'s (2009, 2010) and Utianski et al.'s (2012) methodology, however, is that their measures are time consuming to evaluate and require a good knowledge of acoustics (e.g. in order to identify consonant and vowel boundaries in the spectrographic signal for the rhythm measure) as well as data processing skills by the examiner, probably to a level that cannot be expected of the average clinician.

Another technique that has been used increasingly to research motor control aspects of speech and that has the potential for being applicable in the clinical context is the spatio-temporal variability index (STI, Smith, et al., 1995). The STI is a composite measure of articulatory variability that examines the stability and patterning of speech movement sequences. Its value reflects the degree to which movement patterns are consistent over a number of repetitions. Variability is thought to reflect movement control as the production of fluent and intelligible speech requires the coordinated action of the relatively autonomous acting pulmonary, laryngeal and vocal tract articulation components. The ability of speakers

to consistently and accurately execute a specific pattern of speech motor movements during a sequence of identical speech productions is reflected in the degree of token-to-token variability in the resulting speech output. Any abnormalities in strength, speed, steadiness or accuracy of speech movements will likely be reflected in the amount of variability present in speech production tasks with reiterated utterances.

The STI was initially used to investigate healthy speakers to gain insight into the nature of speech motor control processes. These studies highlighted that the stability of speech movements could be reduced under certain experimental conditions such as increased or slowed speech rate (Smith et al. 1995), the performance of concurrent tasks (Dromey & Bates, 2005; Dromey & Benson, 2003; Dromey & Shim, 2008), or the complexity of the carrier phrase (Kleinow & Smith, 2000). Wohlert and Smith (1998) furthermore noted that older speakers tended to be more variable than the younger participants. The STI has also been able to highlight differences between disordered and unimpaired speakers as demonstrated by Kleinow and Smith (2000) as well as Smith & Kleinow (2000) in people who stutter, and Kleinow, Smith, and Ramig (2001) in people with HD. In addition, the STI has been used to demonstrate which therapy techniques were likely to be most beneficial for speakers. Kleinow et al. (2001) found that loud voice resulted in more stable articulation, suggesting that the Lee Silverman Voice Treatment (LSVT®, Ramig, Countryman, Thompson, & Horii, 1995) is an effective treatment to improve loudness as well as articulation. In another study by McHenry (2003), the STI highlighted differences across speakers in disease severity and, in addition, indicated which treatment strategy for slowed speech (stretched speech versus inserting pauses) was most appropriate for the speakers. McHenry (2003) thus argued that the STI has an important role to play diagnostically, as well as in establishing optimum treatment strategies.

The above studies have established differences between disordered and unimpaired speakers, and provided a range of experimental setups that serve to bring to light such differences. They furthermore highlighted the value of the STI for increasing our understanding of motor control processes. However, similarly to Liss et al.'s (2009, 2010) and Utianski et al.'s (2012) methods, the STI is insufficiently developed to function as an effective research or clinical tool. One of the biggest drawbacks is the invasive and complex nature of the associated technology, which generally involves lip tracking. This requires specialist equipment and the procedures might not be easy to tolerate for participants with motor and sensory impairments. Participant numbers in previous studies have therefore been low, impacting on the statistical power and predictions made by these experiments. In order to address these problems, methods involving acoustic data collection and analysis have been successfully validated in replicating findings of the kinematic STI (Howell, Anderson, Bartrip, & Bailey, 2009). Acoustic recordings are a well established method of data collection in speech disorders and allow capturing of performance from a large number and wider variety of speakers. In addition, they allow greater freedom in experimental set-up, such as the choice of speech task and recording environments. A further methodological development has been to use a nonlinear data processing approach (Functional Data Analysis (FDA), Lucero, 2005; Lucero & Koenig, 2000; Ramsay, Munhall, Gracco, & Ostry, 1996). This approach differs from the STI in that it produces separate information on spatial and temporal variability. Studies by Anderson, Lowit, and Howell (2008) and van Brenk and Lowit (2012) applied FDA to acoustic data in the form of formant, intensity and pitch tracks and demonstrated that the technique has good potential of distinguishing between different types of dysarthria.

Although the technique requires less time to segment the signal than e.g. the rhythm measure evaluated by Liss et al. (2009) by only marking the beginnings and end of an utterance, a good knowledge of acoustics and a significant amount of time is still necessary to check the

signal for any aberrant data points, particularly for the formant and pitch analyses, thus precluding it from being a readily useable clinical tool in its current form. For any acoustic or other instrumental measure to have any clinical value, it is thus necessary to develop an evaluation procedure that is quick and easy to use, yet at the same time provides valid data that reflect the severity and type of speech impairment experienced by an individual.

In what follows we pursue another variant on measuring variability that might achieve this goal. A common insight underlying several of the above studies is the observation that utterance-to-utterance invariance is a characteristic of highly skilled coordinated action, and that impaired speech is commonly associated with an increase in variability, or, equivalently, a decrease in stability (Anderson et al., 2008; McHenry, 2003; Kleinow, et al., 2001, van Brenk & Lowit, 2012). Given that it is relatively easy to obtain audio recordings of repeated productions of a standard phrase, we present a computational method that quantifies the utterance-to-utterance variability. Measuring variability is not entirely straightforward for a rich multi-dimensional signal such as speech. We make use of a parametric representation of speech, the Mel Frequency-scaled Cepstral Coefficient, which is known to capture numerically many of the spectral and temporal properties of speech that are important for speech perception. These representations are used as a standard in both speech recognition and speech synthesis applications. We combine this representation of speech with the Dynamic Time Warping algorithm, that allows one sequence to be mapped onto another, and we use a quantitative measure of the amount of warping to arrive at an index of utterance-toutterance variability. We wish to see whether the quantification of utterance-to-utterance variability (hereafter, UUV) can provide a sensitive index of dysarthria suited to evaluating severity either as an outcome measure, or as a diagnostic tool. The tool would thus provide a significant contribution to evidence based practice, allowing clinicians to demonstrate

effectiveness of their interventions and researchers to evaluate different treatment methods against each other.

Methods

Participants

The speakers and tasks reported in this study form a subset of a larger study (see van Brenk & Lowit, 2012). All data collection and analysis procedures adhered to current ethical guidelines and had been approved by Strathclyde University as well as relevant health boards. We examined data from 8 speakers with ataxic dysarthria (AD) and from 16 speakers with Hypokinetic Dysarthria (HD) as a consequence of Parkinson's disease, along with age (+/- 5 years) and gender matched healthy control speakers. For one HD participant, matching was only possible by age but not gender (Table 1). All speakers were native speakers of British English. Suitability for participation in the experiment was assessed through informal evaluation of the speakers' hearing and visual skills, a cognitive screen (ACE-R, Mioshi, Dawson, Mitchell, Arnold, & Hodges, 2006) and a short medical history to exclude any previous speech and language problems or other health issues that could have affected task performance. The severity and type of the dysarthria were judged by the referring SLT, and confirmed by two of the current authors. In order to quantify the level of severity for experimental purposes, intelligibility scores were derived from three tasks, a monologue about a holiday, passage reading (Grandfather passage, Darley, Aronson, & Brown, 1975) and the Unpredictable Sentence Intelligibility Test (McHenry & Parle, 2006). The monologue and reading passage were scored on a 9-point scale (Dobinson, 2007), and the sentences on a percentage scale (% of words correctly transcribed). Fifteen trained listeners (SLP students) evaluated the data. To assess the overall consistency amongst the 15 listeners, the intraclass

correlation coefficient (ICC) was calculated (Sheard, Adams & Davis, 1991). The ICC was .868 (95% CI: .778 to .933) and thus showed good agreement between listeners. Results of all three assessments correlated strongly with each other. For ease of reference, only the data of the sentence intelligibility test, expressed as percentage of correctly transcribed words, are reported here (Table 1). The data show that there was a difference in intelligibility between the HD and AD speakers, with the AD speakers generally showing a lower intelligibility level (mean intelligibility HD: 84%, AD: 68%), although there was some overlap between the two groups.

Insert table 1 around here

Recording and Analysis Procedures

Participants were recorded in a quiet environment in their homes, at the university or at the local clinic they were attending. Audio recordings were taken using a wave recorder (Edirol R-09HR) connected to a head-mounted condenser microphone (AKG C-420). The data recorder supplied Phantom Power to the microphone. Data were sampled at 44.1 kHz at 16 bits. The head-mounted device allowed for a constant distance of approximately 4 cm between the speaker's mouth and microphone during recording.

Experimental Tasks

The experimental task consisted of a simple utterance repetition paradigm. Participants were instructed to repeat the phrase "Tony knew you were lying in bed" with as little variability as possible from one utterance to the next. In the larger study which focused on spatio-temporal

variability, they repeated this task under six different conditions. For the purposes of the current analysis, three tasks commonly used in previous research on variability were chosen, i.e. habitual, fast and slow speaking rates. Coordinative stability is frequently rate-dependent, which motivates the examination of multiple speaking rates. It also allowed the comparison of the current results with those from other previously published measures such as STI and FDA, which detected higher levels of variability in altered rate conditions (e.g. Smith et al. 1995), in order to evaluate the validity of our results. Participants were allowed to self-select rate in the habitual condition and were instructed to speak half as fast or twice as fast as their usual rate for the slow and fast conditions respectively. The aim was to achieve 20 to 25 repetitions of the phrase, however, not all speakers could achieve this target.

Measurement Parameters

The method under investigation in this paper is the utterance-to-utterance variability (UUV) measured across the repetitions of the test utterance. The variability measured is not simply durational, as would be the case in for example the various forms of the Pairwise Variability Index (Low, Grabe, & Nolan, 2000). Rather, we employ a full spectral representation, the Mel Frequency-scaled Cepstral Coefficient (MFCC), estimated in overlapping sliding windows of approximately 23 ms, and we use the similarity of the entire spectral representation in evaluating similarity among successive utterances. The method was introduced in Cummins (2009) for the assessment of the degree of synchrony in parallel utterances of the same text, but we re-purpose it here to identify utterance to utterance variability in successive phrases. Full details of the computational procedure are provided in Cummins (2009). An estimate of this variability was obtained for each speaker and condition in the following manner.

First, an automated Praat script was used to divide the recording up into individual utterances. This script used a sliding window to look for pauses based on waveform intensity. To facilitate the analysis of a lot of data, the resulting segmentation into individual utterances was visually checked rapidly by the experimenter, omitting data where the algorithm either failed to place a boundary correctly, or placed inappropriate boundaries. For each sequence of utterances, then, utterance i was compared to utterance i+1, by aligning the two utterances and estimating the amount of warping that would be required to map one onto the other. For highly similar utterances, this is a low number, and for disparate utterances, it increases. For n utterances, this provides n-1 comparisons, and these estimates can then be used for intraand inter-group comparisons.

Two utterances to be compared are first represented as sequences of MFCCs, yielding a sequence of vectors for each utterance. Dynamic Time Warping (DTW) is used to map one utterance on to the other. The estimate of variability is derived from the mapping process, and is proportional to the amount of compression and stretching required for the optimal mapping of one utterance onto the other. Technical details of the algorithm are as follows: We used the first 12 MFCC parameters, as is standard in speech technological applications, omitting the zero-th, or energy, component. We estimated MFCCs using Hamming windows of 1024 samples, or approximately 23 ms, with an overlap of half a window between successive frames. DTW was then used to map one sequence (the comparator) onto the other (the referent), yielding a warp path, that can be represented as a function stretching above and below a horizontal line, as explained in Cummins (2009). The unsigned area under this curve, estimated during voiced portions of the referent only, provides the quantitative assessment of dissimilarity for the two utterances. Restriction to the voiced only portions has been found to improve the stability of the method when used on many speakers and texts. The entire comparison procedure is automated within Matlab.

Within-trial variability in the inter-utterance variability score was rather high for both disordered speakers and control participants. This presumably arose because of the difficulty in maintaining an invariant repetition style for a prolonged speaking time. We therefore removed from a trial all inter-utterance variability scores that were more than two times the interquartile range from the median. All subsequent analyses were performed on the trimmed data. Trimming removed between 2.0% and 7.0% of data points. Comparable amounts were removed for both control participants and those with dysarthria.

A further measure taken from the data for control purposes was rate. Because participants can respond to the experimental imperative to speak more or less rapidly in various ways, we initially evaluated both articulation and speech rate behaviour. Subsequent statistical comparisons with other speech parameters revealed the same results for both measures and we therefore restrict the discussion to articulation rate. As some of the disordered speakers had a tendency to omit individual phonemes from their production, we decided to use the reciprocal of articulation rate by measuring the average interval duration within a trial from the onset of voicing in "Tony" to the end of voicing in the word "in". To validate the measure we calculated the actual articulation rate for the control speakers based on their interval durations, which provided a mean of 4 syllables per second, which is within the normal range for a reading task. For simplicity of language, we will use the term articulation rate in the following discussion.

Statistical Analysis

We employed a range of parametric tests to investigate the relationship between the various measurement parameters. For within group comparisons, all control participants were pooled

into one group. For across group comparisons, the division between AD and HD controls was maintained in order to ascertain a close match with the disordered groups in terms of age and gender. Bonferroni corrections were applied to all statistical analyses.

Results

Articulation Rate and UUV

We first examined articulation rate to see whether participants did in fact modulate rate as a function of experimental condition. Within each condition, we conducted paired t-tests for rate, with the conservative Bonferroni correction of p-values for repeated tests. The results are provided in Table 2, from which it can be seen that the control group and the HD group successfully modulated rate, while the ataxic group (AD) did not display substantial modulations.

Insert Table 2 about here.

We then compared the utterance to utterance variability (UUV) exhibited by each patient group to their respective controls in each of the rate conditions, habitual, fast and slow. Figure 1 shows the distribution of the median UUV scores for both patient groups and their respective control participants. Bonferroni-protected paired t-tests are reported in Table 3. It is evident that UUV differentiates between the AD group and their controls, but not between the HD group and their respective controls, despite the fact that participant numbers are considerably smaller in the AD comparison.

Insert Figure 1 about here

Insert Table 3 about here

To further examine the relationship between the UUV scores and rate we performed a repeated measures analysis for each group separately across the three speaking conditions,

habitual, fast, and slow rate (the AD and HD control participants were considered as one group for this analysis). We applied the Greenhouse-Geisser correction for departure from sphericity. The medians are shown in box plots in Figure 2. There was a significant effect of condition for control participants (F(1.4, 32.6)=12.4, p < .001), for the speakers with Parkinson's Disease (F(1.0, 15.7)=8.1, p < .05), as well as for the AD group (F(1.3, 9.3)=5.2, p < .05). Tukey HSD post hoc tests revealed significant differences in UUV between slow and habitual rates for controls and HD participants, and between slow and fast conditions for all three groups, with slow rates having the highest UUV value in each case. These results suggest that although speakers' performances were affected by the speaking task, the direction of between task differences was similar across all participant groups.

Insert figure 2 about here

Correlations between UUV, rate and intelligibility

In summary, the results showed that UUV changed across rate conditions in all three participant groups, but that the AD group were more affected in the UUV than the HD group compared to their respective control participants. There are a number of potential reasons for this difference in results between the HD and AD groups. One factor neould have been rate, which was modulated differently in the two groups. The other could have been dysarthria severity, as the HD group had higher intelligibility levels than the AD group. Finally, the results could reflect inherent differences in the underlying pathology and how this affects the speaker's ability to repeat the same utterance in a consistent manner.

In order to investigate the above possibilities a number of correlational analyses were performed. We first examined the relationship between articulation rate and UUV for AD and HD participants separately. For both groups, correlations in the habitual and fast rate conditions were non-significant, but in the slow condition all participants displayed strong

correlations between UUV and rate (AD: r(6)=0.78, p<.05, HD: r(14)=0.74,p<.01). The fact that both HD and AD speakers were affected in similar ways suggests that rate differences between the two groups were not the cause of the varying results observed for the UUV. However, these results lead to some further interesting questions. First, the fact that UUV performance at habitual and fast rates was independent of actual articulation rate, but that there was some relationship at slow rates requires further investigation. Second, the similarity in results between HD and AD participants is interesting given the absence of any significant rate changes across the conditions displayed by the AD group. This suggests that although rate itself was not a factor, task demands might have been, i.e. AD speakers appeared to have greater difficulties slowing down, affecting their stability in motor control while executing this task.

Sentence intelligibility (table 1) was negatively correlated with UUV for the HD group only in both habitual and fast rates, but not the slow rate (habitual: r(14)=-0.65, p<.01; fast: r(14)=-0.62, p<.05). Figure 3 plots UUV against intelligibility for both disordered groups in the habitual rate condition. A single regression line based on the combined data has been fitted (r(22)=-0.68, p<0.001). These data suggest at least a tendency for the relationship between intelligibility and UUV. It was particularly interesting to see changes in UUV even at the mild end of the severity range where speakers did not necessarily fall outside the normal range for intelligibility yet. This might suggest that the UUV is able to detect early, subclinical signs of speech disorders, however, this would require further investigation with larger participant numbers.

Insert figure 3 about here

Discussion

This paper investigated whether the UUV has the potential to act as a valid and clinically viable tool to quantify the severity of dysarthric speech, and perhaps to differentiate among speech pathologies.

Validity

In terms of its validity, the across task comparisons for healthy as well as disordered speakers showed the same differences as have previously been reported for STI, i.e. higher variability for slow speech rates compared to habitual and fast rates (McHenry 2003, Smith et al. 1995). In addition, we also found that the disordered groups followed the same pattern of across task differences (cf. Smith & Kleinow, 2000; Kleinow & Smith 2000). The UUV thus appears to capture similar aspects of motor control as these other measures.

In addition, there was a strong correlation with the perceptual intelligibility scores for HD patients. A larger participant number might have resulted in a similar result for the AD speakers, as the current data showed at least a trend towards such a relationship in this group. The results thus suggest that the UUV metric may exhibit good validity for representing the severity of a speech disorder. It is thus again comparable to previously published variability measures such as the STI or the FDA in this respect. McHenry (2003) reported correlations between severity and the STI in an HD population. Similarly, van Brenk and Lowit (2012) identified a relationship between a number of FDA variables and intelligibility, albeit for the AD rather than the HD group. However, their results were based on a larger participant sample, and when the analyses are re-run on the current speaker selection, both their HD and AD groups show significant relationships between intelligibility scores and variability in the fast speaking condition. The suggested sensitivity of the UUV across the range of severities present in the current sample would have significant clinical advantages in that the measure should be applicable as an outcome measure. More research will have to be performed to

establish the minimum detectable change in a speaker's intelligibility as represented by the UUV; however, these preliminary results are highly encouraging in this respect.

The current results thus suggest that the UUV has the same potential to reflect the severity of a person's speech disorder and capture task dependent variations in motor control demands as other measures of variability. What sets is apart from them is that it has a practical advantage in terms of processing time and skills required by the examiner, which makes it a more viable option for clinical application.

Technical Implementation

Measurements are based on the same, easy to capture data as the acoustic STI/FDA paradigms, i.e. audio data of simple sentence repetitions. The measurement process includes two steps, each of which is straightforward and requires no technical knowledge of signal processing. The first step is segmentation of a string of utterances into individual utterances. This requires application of a relatively simple Praat script which identifies phrase boundaries, and visual inspection of their correct placement, a task which even those not particularly skilled at acoustic analysis can perform, and which can be done relatively quickly compared to some other methods. The second step, quantifying the utterance to utterance variability, is implemented in a series of Matlab functions. This step would require some further development in order to make it applicable in a stand-alone version without the Matlab platform. A reference implementation and some supporting Praat scripts that can help in the segmentation of long strings of utterances are provided as supplementary materials. These are intended as a starting point for researchers interested in developing their own full implementation. Discussion of the many possible variants of the basic time warping algorithm are provided in Senin (2008).

As a note of caution, our experience with the application of this method suggests that it will be of most utility in cases in which UUV is mild to moderate, offering the possibility of sensitively indexing changes in articulatory stability over time. In repeated productions of the sort employed here, a single highly aberrant token will generate values that appear as outliers in the series. We found such outliers in both the control data and the patients, and to comparable extent in each group, suggesting that it is a reliable characteristic of such sequential productions. This suggests that a certain degree of utterance to utterance stability should be considered a prerequisite for the application of the method, and alternatives might have to be sought for more severe cases. However, the current participant sample did not include patients at the more severe end of the spectrum and further studies need to be conducted to assess the method's suitability across the full range of client presentations.

Future Directions

There are evidently a number of limitations in the current research, most prominently the small participant numbers, particularly in the AD group. Having said that, our study does not compare unfavourable with many other studies trying to develop new technologies; for example, although the Liss et al. (2009, 2010) and Utianski et al. (2012) studies have 70 speakers in total, these are distributed across five speaker groups, resulting in an average of 14 speakers per group. In addition, the fact that our correlation results are strong even with our small group sizes lends further support to the viability of the UUV as a valid measure of intelligibility and motor control. Another factor that significantly limited our ability to comment on the diagnostic value of the UUV is the fact that the HD and AD groups were not comparable in terms of their severity. No judgements can thus be made in terms of how our measure compares to those proposed by Liss et al. (2009, 2010) in its discriminating capacity, and this thus needs to be the focus of further research.

In conclusion, we have introduced a new quantitative measure of utterance to utterance variability that can be employed in the quantitative assessment of dysarthria. Preliminary results show good comparability to existing validated measures, highlighting the potential to develop this measure into a clinically viable assessment tool and outcome measure in addition to its application to fundamental research on motor control in both healthy and disordered speakers. These findings now need to be confirmed by conducting further research with larger and more varied participant groups, further investigating the diagnostic value of the UUV, its sensitivity to change, as well as establishing a firm normative range for this measure.

Acknowledgments

We would like to express our thanks to all our participants who dedicated their time to this project. We would also like to acknowledge the support of the Scottish Funding Council for the PhD studentship which provided the speech data for the current investigation.

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Table 1: Participant information, stating gender, age, medical diagnosis, intelligibility level and medication (HD participants only).

ADI M 40 CA 79 AD2 F 38 FA 58 AD3 F 63 MS 85 AD4 M 58 SCA8 47 AD5 F 44 MS 72 AD6 M 70 MS 55 AD7 M 37 MS 69 AD8 M 46 MS 79 AD8 M 46 MS 79 AD8 Mean (SD) 49.8 (14.3) HD01 M 40 IPD 98 A 1x16mg; K 1x10mg HD02 F 55 IPD 93 A 4x6mg; E 6xXmg HD03 M 48 IPD 89 A 1x4mg HD04 M 56 HD05 F 76 IPD 89 D 4x25/100mg; D 4x12.5/50mg; F 4x200mg HD07 M 76 AD8 M 76 BD9 B 1x150mg; E 2x25mg; G 8x50/12.5mg; G 8x50/12.5mg; G 8x50/12.5mg; G 8x50/12.5mg; G 8x50/12.5mg; G 8x50/12.5mg BD0 M 54 BD0		gender	age	diagnosis	intell	medication
AD3	AD1				79	
AD5	AD2	F	38	FA	58	
AD5	AD3	F	63	MS	85	
AD6	AD4	M	58	SCA8	47	
AD7	AD5	F	44	MS	72	
AD8	AD6	M	70	MS	55	
Mean (SD) 49.5 (12.5) 68 (13.4)	AD7	M	37	MS	69	
AD controls 3 F, 5 M Mean (SD) 49.8 (14.3)	AD8	M	46	MS	79	
Mean (SD) 49.8 (14.3)		Mean (SD)	49.5 (12.5)		68 (13.4)	
Mean (SD) 49.8 (14.3)						
HD01 M	AD controls	3 F, 5 M				
HD02		Mean (SD)	49.8 (14.3)			
HD02	11001	7.6	40	IDD	0.0	A 1 1 6 W 1 1 0
HD03 M						
HD04 M 56 PD A 6x2mg; B 4x25/100mg; D						· ·
HD04 M	HD03	M	48		89	Č
HD05 F				IPD		
HD07 M 76 63 1x25mg E 2x25mg; G 3x25/100/200mg; H 1x40mg; L 1x25mg C 8x50/12.5mg; G 8-10x50/200/200mg HD10 M 64 IPD 88 A 3x7mg HD11 M 75 IPD 72 C 3x50/12.5mg; D 6x25/100mg HD12 M 73 IPD 76 B 1x50/200mg; D 6x25/100mg HD13 M 78 IPD 70 B 3x25/100mg; C 2x100/25mg IPD A 3x8mg; B 1x50/200mg; E HD14 F 75 88 3x62.5mg C 2x100/25mg; D 3x25/100mg; HD15 M 78 IPD A 3x4mg; B 1x50/200mg; D HD15 M 78 IPD A 3x4mg; B 1x50/200mg; D HD16 M 67 F 67 A 3x4mg; B 1x50/200mg; D 3x25/100mg; D 3x25/100mg IPD A 3x8mg; B 3x25/100mg; D 3						
HD07 M	HD05	F	76		89	
HD07 M				IPD		
HD09 M 54 P94 10x50/200/200mg HD10 M 64 IPD 88 A 3x7mg	110.5	3.6				
HD09 M 54 94 10x50/200/200mg HD10 M 64 IPD 88 A 3x7mg HD11 M 75 IPD 72 C 3x50/12.5mg; D 6x25/100mg HD12 M 73 IPD 76 B 1x50/200mg; D 6x25/100mg HD13 M 78 IPD 70 B 3x25/100mg; C 2x100/25mg HD14 F 75 88 3x62.5mg HD14 F 75 88 3x62.5mg HD15 M 78 88 I 3x0.7mg and 3x0.18mg HD15 M 78 88 I 3x0.7mg and 3x0.18mg HD16 M 67 77 3x25/100mg HD16 M 67 77 3x25/100mg HD17 M 60 98 1x100/25mg; D 4x50/200mg; I HD20 F 65 IPD 75 D 4x25/100mg HD20 F 65 IPD 75 D 4x25/100mg HD20	HD07	M	76		63	
HD10	HD00	3.4	5.4	IPD	0.4	
HD11 M 75 IPD 72 C 3x50/12.5mg; D 6x25/100mg HD12 M 73 IPD 76 B 1x50/200mg; D 6x25/100mg HD13 M 78 IPD 70 B 3x25/100mg; C 2x100/25mg HD14 F 75 88 3x62.5mg HD15 M 78 IPD C 2x100/25mg; D 3x25/100mg; I 3x25/100mg; I 3x0.7mg and 3x0.18mg HD16 M 67 77 3x25/100mg HD16 M 67 77 3x25/100mg HD17 M 60 98 1x100/25mg; D 4x50/200mg; J 1x100mg; K 1x1mg HD20 F 65 IPD 75 D 4x25/100mg HD controls 5 F, 11 M 84 (10.5) HD 4x25/100mg				IDD		<u> </u>
HD12 M						ŭ
HD13 M 78 IPD 70 B 3x25/100mg; C 2x100/25mg HD14 F 75 88 3x62.5mg HD15 IPD C 2x100/25mg; D 3x25/100mg; D 3x25/100mg; D 3x25/100mg; I 3x0.7mg and 3x0.18mg HD16 M 67 77 A 3x4mg; B 1x50/200mg; D 3x25/100mg; D 3x25/100mg; D 3x25/100mg; D 3x25/100mg; D 3x25/100mg HD16 M 67 77 A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J 1x100/25mg; D 4x50/200mg; J 1x100mg; K 1x1mg HD17 M 60 98 1x100mg; K 1x1mg HD20 F 65 IPD 75 D 4x25/100mg HD controls 5 F, 11 M 84 (10.5) IPD IPD						
HD14 F 75 88 3x62.5mg C 2x100/25mg; D 3x25/100mg; E HD15 M 78 88 I 3x0.7mg and 3x0.18mg I 3x0.7mg and 3x0.18mg I 3x0.7mg and 3x0.18mg I 3x25/100mg; D A 3x4mg; B 1x50/200mg; D A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J I 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		+				
HD14 F 75 88 3x62.5mg HD15 IPD C 2x100/25mg; D 3x25/100mg; I 3x0.7mg and 3x0.18mg HD16 IPD A 3x4mg; B 1x50/200mg; D 3x25/100mg HD16 M 67 77 3x25/100mg IPD A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J 1x100/25mg; D 4x50/200mg; J 1x100mg; K 1x1mg IPD 75 D 4x25/100mg HD20 F 65 IPD 75 D 4x25/100mg HD controls 5 F, 11 M 84 (10.5) IPD	HD13	M	/8		70	
HD15 M 78 B8 I 3x0.7mg and 3x0.18mg IPD A 3x4mg; B 1x50/200mg; D 3x25/100mg; D A 3x8mg; B 3x25/100mg; D A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J HD17 M 60 98 1x100mg; K 1x1mg HD20 F 65 IPD 75 D 4x25/100mg Mean (SD) 65 (11.8) 84 (10.5) HD controls 5 F, 11 M HD controls 5 F, 11 M HD HD C 2x100/25mg; D 3x25/100mg; D A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J A 3x8mg; B 3x25/100mg; D	IID14	E	75	IPD	00	[
HD15 M 78 88 I 3x0.7mg and 3x0.18mg HD16 IPD A 3x4mg; B 1x50/200mg; D 3x25/100mg HD16 IPD A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J 4x50/200mg; J 4x50/200mg; J 4x50/200mg; J 4x50/200mg; J 4x50/200mg HD17 M 60 98 1x100mg; K 1x1mg HD20 F 65 IPD 75 D 4x25/100mg Mean (SD) 65 (11.8) 84 (10.5) HD controls 5 F, 11 M	HD14	Г	/3	IDD	88	C
HD16 M 67 F 65 IPD 75 D 4x25/100mg Mean (SD) 65 (11.8) HD controls 5 F, 11 M IPD A 3x4mg; B 1x50/200mg; D 3x25/100mg A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J 1x100mg; K 1x1mg D 4x25/100mg D 4x25/100mg	IID15	M	70	IPD	00	
HD16 M 67 77 3x25/100mg IPD A 3x8mg; B 3x25/100mg; C 1x100/25mg; D 4x50/200mg; J 1x100mg; K 1x1mg HD17 M 60 98 1x100mg; K 1x1mg HD20 F 65 IPD 75 D 4x25/100mg Mean (SD) 65 (11.8) 84 (10.5) HD controls 5 F, 11 M	מומח	IVI	78	IDD	00	
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HD17 M 60 98 1x100mg; K 1x1mg HD20 F 65 IPD 75 D 4x25/100mg Mean (SD) 65 (11.8) 84 (10.5) HD controls 5 F, 11 M Image: Control of the				עוו		
HD20 F 65 IPD 75 D 4x25/100mg Mean (SD) 65 (11.8) 84 (10.5) HD controls 5 F, 11 M	HD17	M	60		98	_
Mean (SD) 65 (11.8) 84 (10.5) HD controls 5 F, 11 M				IPD		
HD controls 5 F, 11 M	11020	<u> </u>		11.0		- mailtoing
· · · · · · · · · · · · · · · · · · ·		1,10011 (51)	35 (11.0)		3. (10.5)	
· · · · · · · · · · · · · · · · · · ·	HD controls	5 F. 11 M				
		Mean (SD)	63.6 (11.3)			

Abbreviations: AD – ataxic dysarthria, HD – hypokinetic dysarthria, CA – cerebellar ataxia, FA – Friedreich's ataxia, SCA – spinocerebellar ataxia, MS – Multiple Sclerosis, IPD – idiopathic Parkinson's Disease

Key to medication: A: Ropinirole, B: Sinemet Controlled Release (carbidopa/levodopa), C: Madopar Controlled Release (levodopa/benserazide), D: Sinemet-Plus (carbidopa/levodopa), E: Sinemet (levodopa), F: Entacapone, G: Stalevo (carbidopa/levodopa/entacapone), H: Stimvastatin, I: Pramipexole, J: Amantadine, K: Rasagiline, L: Atenolol.

Table 2: Bonferroni-protected t-tests of the differences observed in a measure of articulation rate.

	Ataxic Dysarthria		Hypokinetic Dysarthria		Control Group	
	t	p	t	p	t	p
Habitual vs. Fast	t(7) = 2.3	n.s.	t(15) = 5.2	***	t(23) = 8.6	***
Habitual vs. Slow	t(7) = -1.7	n.s.	t(15) = -3.3	*	t(23) = -4.4	***
Fast vs. Slow	t(7) = -2.1	n.s.	t(15) = -4.8	***	t(23) = -5.5	***

NB: Due to the Bonferroni correction, only significance levels rather than exact p values are reported: *** = p<.001, ** = p<.01, * = 0<.05

Table 3: Bonferroni-protected t-tests of UUV differences between participant groups at each rate.

	Habitual		Fast		Slow	
	t	p	t	p	t	p
AD vs AD Control	t(7) = 2.5	n.s.	t(7) = 3.5	*	t(7) = 2.8	*
HD vs. HD Control	t(15) = 0.52	n.s.	t(15) = -0.66	n.s.	t(15) = -0.28	n.s.

NB: Due to the Bonferroni correction, only significance levels rather than exact p values are reported: *** = p < .001, ** = p < .01, * = 0 < .05

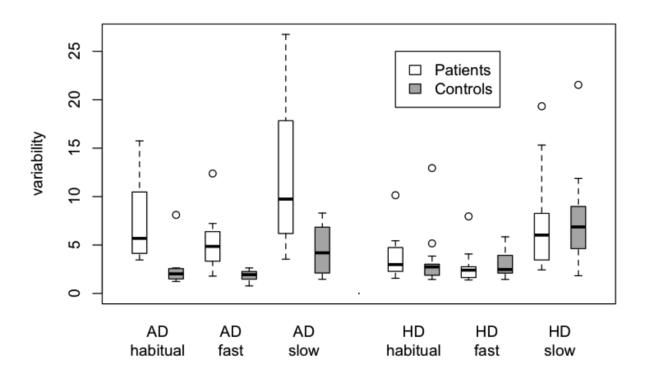


Figure 1: Boxplot of utterance-to-utterance variability scores for disordered speakers and their relevant controls in the three rate conditions.

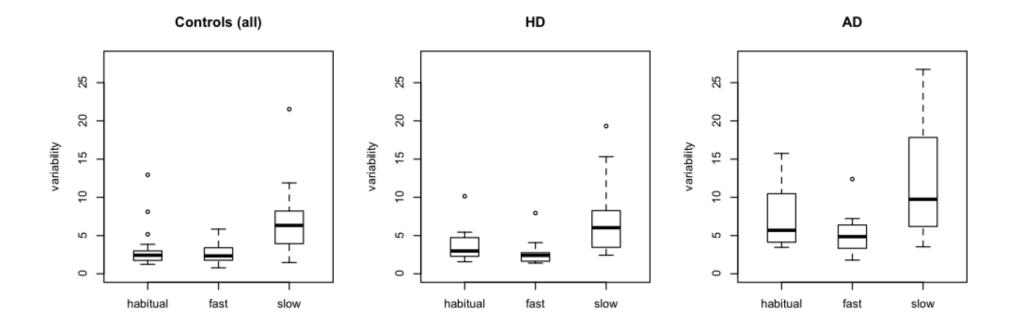


Figure 2: Boxplot of utterance-to-utterance variability scores for each group across speaking rate conditions.

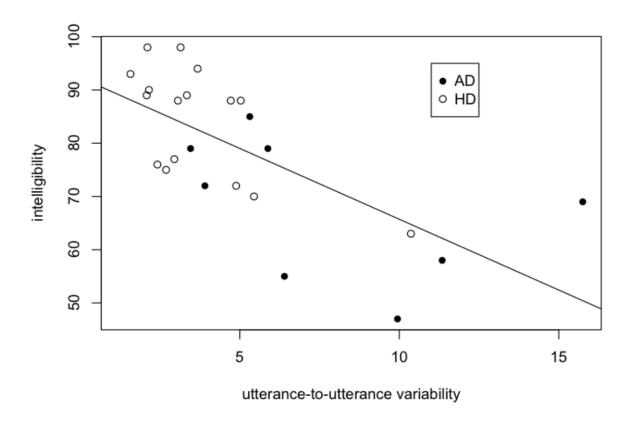


Figure 3: Scatter plot of utterance-to-utterance variability scores for the habitual condition and intelligibility results for HD and AD speakers.