Anatomical Directional Dissimilarities in Tri-axial Swallowing Accelerometry Signals

Faezeh Movahedi, Atsuko Kurosu, James L. Coyle, Subashan Perera, Ervin Sejdić

Abstract—Swallowing accelerometry is a non-invasive approach currently under consideration as an instrumental screening test for swallowing difficulties, with most current studies focusing on the swallowing vibrations in anterior-posterior (A-P) and superior-interior (S-I) directions. However, the displacement of the hyolaryngeal structure during the act of swallowing in patients with dysphagia involve declination of the medial-lateral (M-L), which suggests that the swallowing vibrations in the M-L direction have the ability to reveal additional details about the swallowing function. With this motivation, we performed a broad comparison of the swallowing vibrations in all three anatomical directions. Tri-axial swallowing accelerometry signals were concurrently collected from 72 patients suspected of dysphagia undergoing videofluoroscopic evaluation of swallowing (mean age: 63.94 ± 12.58 years). Participants swallowed one or more thickened liquids with different consistencies including thin-thick liquids, nectar-thick liquids, and pudding-thick liquids with either a comfortable self-selected volume from a cup or a controlled volume by the examiner from a 5ml spoon. Swallows were grouped based on the viscosity of swallows and the participant's stroke history. Then, a comprehensive set of features was extracted in multiple signal domains from 881 swallows. The results highlighted inter-axis dissimilarities among tri-axial swallowing vibrations including the extent of variability in amplitude of signals, the degree of predictability of signals, and the extent of disordered behavior of signals in time-frequency domain. First, the upward movement of the hyolaryngeal structure, representing the S-I signals, were actually more variable in amplitude and showed less predictable behavior than the sideways and forward movements, representing the A-P and M-L signals, during swallowing. Second, the S-I signals, which represent the upward movement of the hyolaryngeal structure, behaved more disordered in the time-frequency domain than the sideways movement, M-L signals, in all groups of study except for the pudding swallows in the stroke group. Third, considering the viscosity and the participant's pathology, thin liquid swallows in the non-stroke group presented the most directional differences among all groups of study. Despite of few directional dissimilarities, M-L axis characteristics were nearly similar to the two other axes characteristics, indicating we might be able to interpret M-L axis characteristics, which are not observable in videofluoroscopic images, based on the A-P and

Index Terms—Keywords: Tri-axial swallowing accelerometry, cervical auscultation, dysphagia, signal processing.

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I. INTRODUCTION

S WALLOING is a complex neuromuscular process requiring a series of biomechanical events, which result in the safe passage of the liquids and solids from mouth to the stomach without entering the airway [1], [2]. Due to the complexity of the swallowing process, many health complications, such as head/neck cancers, and neurological disorders, like strokes, can affect the swallowing function. Dysphagia, swallowing disorders, can affect any of the four stages of swallowing. Serious and potentially fatal health consequences associated with dysphagia negatively impact the health, safety, and quality of life of patients [3], [4]. Aspiration, dehydration, and malnourishment are probable complications of dysphagia [5], [6], [7], [8]. Approximately 1 in 25 adults in the United States will have a swallowing problem annually [9]. Moreover, studies have shown that dysphagia is prevalent among the elderly population due to age-related effects on swallowing physiology in addition to age-related diseases [3], [4], [7], [10], [11], [12].

There are two primary imaging instrumental examinations used for diagnosis and management of dysphagia: videofluoroscopic swallow studies (VFSS) and fiberoptic endoscopic swallowing tests (FEES) [13], [14], [15], [16]. VFSS, which has been considered the gold standard of dysphagia test for over 30 years [13], [14], consists of radiographic imaging of the structure and the biomechanical functions of the upper aerodigestive tract during the ingestion of different consistencies of barium coated materials, and how the swallowed material (bolus) flows through the oral and pharyngeal cavities [14], [17]. However, VFSS is not always available or feasible, and asymptomatic patients are routinely not referred for a VFSS because they do not display overt signs of dangerous or disordered swallowing function. VFSS also requires an examiner with specialized clinical expertise and the use of a radiology suite, which may be unavailable in some areas. [13], [18]. Early identification of dysphagia through screening tests has received much attention in the past 10 years. These efforts have attempted to accomplish two goals. First, the screening is designed to preemptively identify dysphagia before patients with impaired protection of the airway are put in harm's way and are allowed to orally ingest food and liquids. Second, the screening identifies the patients in need of gold-standard imaging testing with VFSS or FEES and identifies from those who do not need the expensive testing. When the screening test are used precisely according to protocol, they possess reasonable sensitivity for identifying dysphagia or its absence in symptomatic patients. However, since they rely on human

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observation of the overt signs of dysphagia (coughing or choking while eating or drinking), they completely fail to identify patients without overt signs of dysphagia who are at risk for dysphagia. Moreover, adherence to the published screening protocols has not been universally good due to clinical expediency. Therefore, there has been a growing interest in complimentary dysphagia screening methods that provide high accuracy, are non-invasive and easy to administer by minimally trained individuals, and consume few resources in health care institutions.

Many studies have recently considered the use of swallowing accelerometry for dysphagia screening. Swallowing accelerometry is based on the translation of the vibrations recorded from the upper aerodigestive tract structure during the act of swallowing into a voltage signal [19], [20], [21], [22]. Numerous investigation have been performed in attempts to find the physiological sources. Some of the studies have considered the displacement of the hyoid bone and larynx as a physiological source of swallow signals [17], [19]. The displacement of hyoid bone is an essential component of the swallowing function that contributes to the protection of the airway during the swallow, as well as, the traction forces necessary to distend the upper esophageal sphincter and allow the flow of swallowed material into the digestive portion of the mechanism [1], [19], [23], [24], [25].

The final goal of the swallowing accelerometry method is to classify healthy swallows and unhealthy swallows based on swallowing vibrations recorded by accelerometers. To reach this goal, at the first step, we should be bale to characterize swallowing vibrations by extracting various features which provide us crucial information about swallowing. Some studies investigated simple features from swallow signals in timedomain such as onset of swallows, duration of swallows, timing of the different physiological stages of swallow, and magnitude of signals [19], [26], [27], [28], [29], [30]. On the other hand, some studies have extracted advanced features in time, frequency, and time-frequency domains from swallow vibrations [26], [31], [32], [33], [34]. The results from past studies have shown the validity of swallowing accelerometry method using one or two directions of anteriorposterior (A-P) and superior-inferior (S-I), representing the forward and upward movements of hyolaryngeal structures during swallowing, in swallow classification and capturing swallowing difficulties [26], [22], [31], [35], [32], [36]. While the focus of most studies were on the assessing performance of different classification methods based on the computed features, few studies have directly compared extracted features in two directions of A-P and S-I to find out about the dissimilarities between the swallowing vibrations in different anatomical directions [22], [36], [37], [38]. The results have showed that two axes of A-P and S-I contain different temporal and spectral information about swallowing. Their findings indicated that the employment of two axes of dualaccelerometer in swallowing accelerometry analysis would be beneficial as each axis provides unique information about swallowing. However, it should be mentioned that focus of previous researches were on the healthy subjects [22], [36], [37], [38] rather than unhealthy subjects who suffer dysphagia, even though the ultimate goal of swallowing accelerometry is to catch any swallowing difficulties in patients who have or are at the risk of dysphagia using swallowing vibrations. No study has comprehensively compared features of swallowing signals in the A-P and S-I directions under various conditions such as swallowing different volumes and viscosities in subjects who suffer from swallowing difficulties. It should be also mentioned that the selection of features in previous studies were subjective as there is no clear answer to which features probably characterize two directions of swallow vibrations. In addition, previous studies utilized different procedural methods, which limits the ability to compare their results. Therefore, there is still a need of further investigation about swallowing vibrations in two axes of A-P and S-I to have a better understanding of swallow in different anatomical directions. While most attention to swallowing-related movements of hyolaryngeal structures has been to their S-I and A-P directions, both of which are clearly observed with VFSS, little to no attention has been directed to medial-lateral (M-L) movements. It is plausible that hyolaryngeal movement in the M-L plane occurs due to the origins and the insertions of paired muscles responsible for displacing the hyolaryngeal complex and the selective, asymmetrical weakening of these muscles in disease states such as certain types of stroke, cranial nerve injury, maxillofacial trauma, head and neck resections, and others. Therefore, due to the interesting observations that arose when a second axis of recording was added to singleaxis accelerometry, the addition of the final, third axis (M-L) may provide intriguing information about the relationship of swallowing accelerometry signals to actual biomechanical swallowing activity, particularly in patients with diseases affecting symmetry of hyolaryngeal muscle contractions.

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The purpose of this study is to comprehensively compare amongst the three anatomical directions of tri-axial swallowing accelerometry signals in both stroke and non-stroke subjects while examining various viscosities. The main question in this study is to examine whether accelerometry signals derived from different anatomical planes provide discriminate information about changes in swallowing function. To answer this question, nine features in time, information-theoretic, frequency, and time-frequency domains have been extracted from

tri-axial swallowing accelerometry signals from dysphagic patients. The outcome of this study provides insights into how using tri-axial accelerometers can be helpful to identify swallowing difficulties. In addition, the possible effect of age and gender on the differences between the values of the extracted features in the three directions has been investigated in this study.

II. METHODOLOGY

A. Data Acquisition

72 patients (mean age 63.94 ± 12.58 years, 42 male, 30 female), who underwent videofluoroscopic examinations at the Presbyterian University Hospital of the University of Pittsburgh Medical Center (Pittsburgh, Pennsylvania), participated in this study. There were 20 patients who had a previous history of stroke as a reason behind their swallowing difficulties (neuromuscular issue) while other 52 stroke-free patients had other various reasons such as head/neck cancer (anatomical issue) or esophageal spasm. All participants signed informed consent and the data collection protocol was approved by the University of Pittsburgh Institutional Research Board. As a part of their examinations, the participants swallowed one or more thickened Liquids with different consistencies including thin-thick liquid (Varibar Thin Liquid with < 5 cPs viscosity), nectar-thick liquid (Varibar Nectar with ≈ 300 cPs viscosity), and pudding-thick liquid (Varibar Pudding with ≈ 5000 cPs viscosity). The viscosity of boluses changed for each patient based on the speech pathologists' decision considering different factors like patients' condition at the time of examinations, patients' history including stroke history, diet, and any specific health considerations. Participants performed the swallows in a neutral head position and, in some cases, in a head-flexion ("chin-tuck") position, which has been used in dysphagia patients to prevent aspiration during swallowing [39]. Swallows with swallow maneuvers, such as the supraglottic swallow, Mendelsohn maneuver, and effortful swallow, were eliminated from the study. In this study, each swallow was rated on an accepted 8-point ordinal clinical Penetration-Aspiration (PA) scale to determine the severity of swallowing difficulty in patients, we included all swallows with different range of severity (1 < PAscore < 8). In this study, there was no specific limitation regarding the severity of dysphagia. Therefore, patients with all levels of swallowing difficulties were included in this study.

881 swallows were recorded with a tri-axial accelerometry (ADXL 327, Analog Devices, Norwood, Massachusetts) with a sensitivity of 420mV/g, resolution of 250 $\mu g/\sqrt{Hzrms}$, and an acceleration measurement of a minimum full-scale range of $\pm 2g$. Previous studies have been shown that the optimal cervical placement for digital detector to record the swallowing sounds and vibrations is on the midline of the cricoid cartilage [21], [34], [40]. Sensor was taped to the participants' anterior neck at the level of the cricoid cartilage, such that the sensitive axes of the accelerometer were aligned to the anatomical anterior-posterior (A-P), superior-inferior (S-I) and medial-lateral (M-L) axes, as shown in Fig.1. Signals from the A-P, S-I, and M-I axes were bandpass filtered from 0.1 to 3000

Hz and amplified (model P55, Grass Technologies, Warwick, Rhode Island) prior to storage on a research computer. The voltage signals for each axis of the accelerometer were fed into a National Instruments 6210 DAQ and recorded at 20 kHz by the LabView program Signal Express (National Instruments, Austin, Texas). Concurrent with the accelerometer recordings, the biomechanical activity and the bolus flow in the upper aerodigestive tract were captured by the x-ray machine (Ultimax system, Toshiba, Tustin, CA) at 30 pulses per second, and videofluoroscopic images were captured at 60 frames per second by a video card (AccuStream Express HD, Foresight Imaging, Chelmsford, MA) and recorded to a hard drive with the same Labview program.

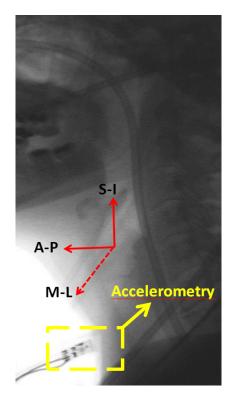


Fig. 1. Location of the sensor on a participant's neck and orientation of the three anatomical axes of the sensor.

B. Data Preprocessing

The time-marked vibration signals were collected with the tri-axial accelerometer in the S-I, A-P, and M-I directions. The time-linked onsets and offsets of each swallow were obtained via the frame-by-frame temporal analysis videofluoroscopic images by a trained experience speech-language pathologist whose inter- and intra-rater reliability in the judgment of these parameters was established a priori. The onset of the swallow segments was defined as the time at which the leading edge of the presented bolus intersected with the shadow cast on the x-ray image by the posterior border of the ramus of the mandible. The offset was the time that the hyoid bone completed motion associated with swallowing-related pharyngeal activity and returned to its resting position. After segmentation of the swallow signals, they were grouped based on the two factors including the viscosity of the liquids

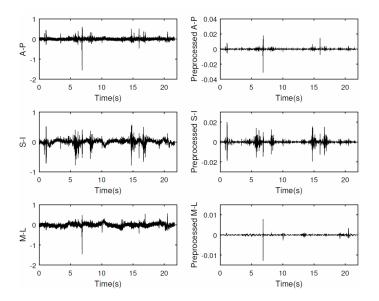


Fig. 2. Recorded swallow vibrations in three axes of accelerometer before and after applying signal processing techniques.

participants were swallowing and whether the participants have had a stroke history, since dysphagia following a stroke and the effect of viscosity on the swallowing signals have been well-documented [18], [29], [41], [42], [43], [44]. Then, each swallow was pre-processed as follow. As the first step, the mechanical and electrical thermal noise created by the accelerometer, was attenuated from the signal by applying axial-specific finite impulse response (FIR) filters [37]. We first fitted an auto-regressive (AR) model to the output of the accelerometer without any input and then use the coefficients of the AR model to generate FIR filter. In the next step, the low frequency components associated with head movement were attenuated from the signals using a least-squares spline approximation algorithm [45], [46]. We fitted a low-frequency (< 2 Hz) trend to the time-domain signal then subtracts that low-frequency trend from the time-domain recording. In the last step, to reduce the impact of both white and colored noise on a given signal, the filtered signals were denoised via a ten-level discrete wavelet transform using the Meyer wavelet with soft thresholding. The level of decomposition was determined by minimizing noise in the signal while retaining the interesting detail of the signals [47]. Figure 2 demonstrated the swallowing signals before and after applying signal processing techniques.

C. Feature Extraction

To capture key statistical differences between swallowing vibration in the A-P, S-I, and M-L axes, several features in multiple domains were extracted from each of the axes of each swallow signal. In this study, 9 features in time, information-theoretic, frequency, and time-frequency domains were evaluated. Further, the practicality of these features has been demonstrated in previous swallowing signal studies [22], [31], [32], [37], [48]. The computational details for each of these features are described in the following subsections.

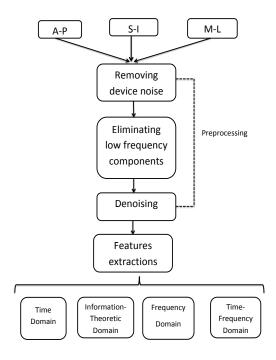


Fig. 3. Flowchart outlining the preprocessing and features extraction steps.

1) Time Domain Features: To find out whether the physics of signal behavior differs between the different anatomical axes of accelerometer, a number of time domain features are computed for the swallowing vibrations. Considering a signal $X = \{x_1, x_2, ... x_n\}$, the following features are extracted:

 The unbiased estimation of the standard deviation can be obtained as

$$\sigma_X = \sqrt{\frac{1}{n-1} \sum_{i=1}^{n} (x_i - \mu_X)^2}$$
 (1)

where μ_X denotes the mean of the signal. The standard deviation reflects how the signal fluctuates around the mean value. Here the important parameter is the ac signal power represented by the deviation from the mean. The bigger the standard deviation, the wider the distribution of data points.

 Skewness quantifies how symmetrical the amplitude distribution is. This feature can be computed as follows:

$$\xi_X = \frac{\frac{1}{n} \sum_{i=1}^{n} (x_i - \mu_X)^3}{\{\frac{1}{n} \sum_{i=1}^{n} (x_i - \mu_X)^2\}^{\frac{3}{2}}}$$
(2)

A symmetrical distribution has a skewness of zero. Whereas, an asymmetrical distribution with a long tail, higher values, to the right/left has a positive/negative skew respectively.

 Kurtosis is a measure of whether the amplitude distribution is peaked or flat relative to a normal distribution.
 A high kurtosis value tends to have a distribution with a sharp, narrow peak, declines rather rapidly, and has heavy tails, while a low kurtosis value signifies a distribution with a flattened peak and has thin tails [49]. This feature is computed as:

$$\gamma_X = \frac{\frac{1}{n} \sum_{i=1}^{n} (x_i - \mu_X)^4}{\{\frac{1}{n} \sum_{i=1}^{n} (x_i - \mu_X)^2\}^2}$$
(3)

- 2) Information-Theoretic Features: The information-theoretic features are popular metrics for characterizing biological signals. The different genres of information-theoretic features are summarized below.
 - The Lempel-Ziv complexity (LZC) is a practical tool to evaluate the randomness of finite sequences, well-suited for physiologic signals [50], [51]. To calculate the LZC, a signal has to be first transformed into a symbolic sequence, since LZC measures the complexity through an estimated number of distinct patterns obtained by parsing a symbolic sequence [51]. Quantization is a typical method used for biomedical signals to be converted into a binary sequence. The signal is converted to 100 symbols by comparing it to 99 thresholds. After the quantized signal is obtained, it can be decomposed into the *k* blocks. Then, the LZC calculated as:

$$LZC = \frac{K \log_{100} n}{n} \tag{4}$$

The logarithm of n, signal length, to the base 100 is used in the above formula as the signal is quantized into 100 symbols.

• The entropy rate is useful to compare the degree of regularity of the signal distribution [52], [53], [44] between the S-I, A-P, and M-L axes. Zero and one values of entropy rate indicate a periodic repetition of the same pattern and aperiodic dynamics, respectively. First, to normalize values of X, μ_X is subtracted from X and divided by σ_X . Then, it is quantized into 10 equally spaced levels. In the next step, sequences of consecutive points in quantized signal \hat{X} of length H, $10 \le H \le 30$, are coded as a series of integers, $\Upsilon_H = \{u_1, u_2, ...u_{n-H+1}\}$, by means of the following formula:

$$u_i = 10^{H-1}\hat{x}_i + H - 1 + 10^{H-2}\hat{x}_i + H - 2 + \dots + 10^0\hat{x}_i$$
(5)

Now the normalized entropy rate is computed as

$$\tilde{\Delta}(H) = \frac{\Delta(H) + \Delta(H-1) + \Delta(1)perc(H)}{\Delta(1)}$$
 (6)

where perc(H) is the percent of unique occurrence of coded integers in the sequence of H and $\Delta_{(H)}$ is the calculated Shannon entropy of the sequence which is obtained as follow:

$$\Delta_{(H)} = \sum_{q=0}^{10^H - 1} P \Upsilon_H(q) \ln P \Upsilon_H(q)$$
 (7)

Finally, entropy rate is presented as ρ an index of regularity

$$\rho = 1 - \min(\tilde{\Delta}(H)) \tag{8}$$

- 3) Frequency Domain Features: Three features in frequency domain were considered in this study mentioned below. First, the fast Fourier transform of the signal was computed to measure the frequency domain features.
 - Peak frequency is simply the frequency corresponding to maximum power and is computed as follows:

$$f_p = \arg_{f \in [0, f_{max}]} \max |F_X(f)|^2 \tag{9}$$

where $f_{max} = 100Hz$.

• The spectral centroid was evaluated as:

$$\hat{f} = \frac{\int_0^{f_{max}} f|F_X(f)|^2 df}{\int_0^{f_{max}} |F_X(f)|^2 df}$$
 (10)

• The bandwidth of the signal is defined as follows:

$$BW = \sqrt{\frac{\int_0^f max (f - \hat{f})^2 |F_X(f)|^2 df}{\int_0^f max |F_X(f)|^2 df}}$$
 (11)

- 4) Time-Frequency Domain Features: Time-frequency features provide simultaneous descriptions of the signal in the time and frequency domains. Recently, the wavelet transform has been considered as a powerful technique for time-scale analysis of a signal and has shown potential in biomedical signal processing [54], [55], [56], [57], [58]. It provides varying time-frequency resolution. Applying wavelet transform would have advantages since swallow are non-stationary signals.
 - The concept of entropy was used in the current study to compare the disorderly behavior in swallowing signals in three axes of accelerometer: S-I, A-P, and M-L. The wavelet entropy was defined based on the wavelet transform. To perform a wavelet transform, a 10-level Meyer wavelet decomposition is applied to a swallow signal to compute the energy contained in different time-frequency bands. Considering d_k as a vector containing wavelet approximation coefficients at kth level of decomposition, the energy at the kth level is defined as

$$E_{d_k} = \|d_k\|^2 \tag{12}$$

where $\|\cdot\|$ means the Euclidean norm. In the next step, the relative wavelet energy equal to energy at a given decomposition level to the total energy of the signal was computed as

$$\Phi_{d_k} = \frac{E_{d_k}}{E_{a_{10}} + \sum_{k=1}^{10} E_{d_k}} \times 100\%$$
 (13)

 Φ_{d_k} gives valuable information regarding the energy distribution of the signal at different frequency bands. Finally, the wave entropy rate is computed as:

$$\Theta_X = -\Phi_{a_{10}} \log_2 \Phi_{a_{10}} - \sum_{k=1}^{10} \Phi_{d_k} \log_2 \Phi_{d_k}$$
 (14)

The high/low value of Θ_X specifies disorder/order behavior for signal.

Table 1 contains the summary of all features and their definitions.

TABLE 1 SUMMARY OF ALL FEATURES

Time Domain Features				
Standard deviation (σ)	Reflects how a signal fluctuates around the mean value of the signal.			
Skewness (ξ) Describes the asymmetry of the amplitude distribution. Negative and positive skewness indicates the distribution of the signal with a long tail, higher values, to the left/right side.				
Kurtosis (γ)	Describes peaked/flat amplitude distribution relative to a normal distribution. High/low kurtosis values specify a signal distribution with a sharp/flattened peak.			
	Information-Theoretic Domain Features			
Lempel-Ziv Complexity (LZC)	Evaluate the randomness of a signal. Higher/lower values indicate more/less randomness in the signal.			
Entropy rate (ρ)	Evaluates the degree of regularity of the signal distribution. Zero and one entropy rate indicates a periodic repetition of the same pattern and aperiodic dynamics, respectively.			
	Frequency Domain Features			
Peak frequency (f)	Describes the frequency corresponding to the maximum spectral power.			
Spectral centroid (\hat{f})	The frequency that divides the spectral power into two equal parts.			
Bandwidth (BW)	The difference between the uppermost and lowermost frequencies in a signal.			
Time-Frequency Domain Features				
Wavelet entropy (Θ_X)	Evaluates the disorderly behavior in time-frequency domain. High and low value of wavelet entropy specifies disorder/order behavior for signal.			

D. Statistical Analysis

To account for multiple trials from the same participants, linear mixed models with random effects were employed as an analytic strategy in this study. First, the anatomical direction differences were computed between three axes for all extracted features in each of the trials. Then, an interceptonly model was fitted to assess statistical significance of each of the said differences in the presence of multiple trials per participant. Secondly, in order to identify the correlates of extracted features, another series of linear mixed models with each of the extracted features and their between anatomical differences (if applicable) as the dependent variable were fitted; each of the individual potential correlates age, gender, and participants' stroke history and them collectively as fixed effect(s) of interest; and a participant random effect. Statistical significance of the parameters of the fixed effects solution with interpretation analogous to regression coefficients were used as evidence of significant association. Due to the large number of extracted features examined and the opportunity for inflated type I error due to multiplicity, the false discovery rate (FDR) methodology was used to make multiplicity corrections to the p-values in all analyses. SAS version 9.3 (SAS Institute, Inc., Cary, North Carolina) was used for all statistical analyses in this study. The benchmark for statistical significance was established at p < 0.05.

III. RESULTS

The accelerometer data were presented in this study for 72 participants including 881 swallows. The total number of 162 comparisons were made between the directional couples of (A-P & S-I), (M-L & S-I), and (A-P & M-L) for all extracted features. Here, we summarize statistically significant results.

A. Demographical Analysis

Before discussing the results of features computed in this study, it should be pointed out that statistical analysis show no significant effects of age, gender, and participants' stroke history on directional comparisons of the computed features of tri-accelerometry swallowing signals.

B. Time Domain Analysis

Table 2 displays the result of time domain features calculated in this study for each of the A-P, S-I and M-L axes grouped by viscosity and participants' stroke history. Firstly, it should be mentioned that the standard deviation (σ) showed directional dissimilarity in all groups of the study (p < 0.02). As we can see in Table 2, The S-I axes demonstrated the highest σ compared to the A-P and M-L in all groups of study.

In addition, considering the asymmetry of swallowing signals, ξ_s , the A-P amplitude distributions in non-stroke participants were mostly positively skewed; whereas the S-I amplitude distributions had a tendency to be strongly negatively skewed in both thin (p < 0.05) and pudding (p < 0.05) groups. Furthermore, even though the signals in both A-P and S-I axes in non-stroke pudding group showed high kurtosis, the S-I signals were more meaningfully leptokurtic (p = 0.03), which resulted in a higher peak than the A-P signals. Other groups showed no statistical difference between the directions in swallow signals.

C. Information-Theoretic Analysis

The summary of information-theoretic features is presented in Table 3. The results for entropy rate ρ indicated the statistical difference between the anatomical axes in two cases. First, the entropy rate differed among all three directions

TABLE 2 Time-domain features (mean \pm standard deviation) extracted from the three anatomical axes of swallowing signals grouped by the participants' stroke history and the viscosity

		Stroke			Non-Stroke			
		$\sigma_s(g) * 10^{-2}$	ξ_s	γ_s*10^2	$\sigma_{ns}(g) * 10^{-2}$	ξ_{ns}	$\gamma_{ns}*10^2$	
	A-P	$0.16\pm0.13^{\S\dagger}$	0.55 ± 4.88	1.66 ± 7.92	$0.19\pm0.19^{\S\dagger}$	$0.72 \pm 2.55^{\S}$	0.37 ± 0.75	
Thin	S-I	$0.29\pm0.22^{\S\ddagger}$	-0.47 ± 1.60	0.18 ± 0.30	$0.31\pm0.21^{\S\ddagger}$	-0.57 ± 2.69 §	0.42 ± 1.50	
Г	M-L	$0.06 \pm 0.04^{\dagger \ddagger}$	0.71 ± 3.53	0.30 ± 1.75	$0.07\pm0.07^{\dagger\ddagger}$	0.31 ± 3.42	0.61 ± 4.46	
=	A-P	$0.15 \pm 0.20^{\S \dagger}$	-0.09 ± 4.57	1.48 ± 8.37	0.16 ± 0.20 §†	0.73 ± 2.24	0.49 ± 1.10	
Nectar	S-I	$0.24\pm0.19^{\S\ddagger}$	-0.30 ± 1.34	0.15 ± 0.24	$0.28\pm0.23^{\S\ddagger}$	0.14 ± 4.37	0.71 ± 2.90	
Z	M-L	$0.06 \pm 0.04^{\dagger \ddagger}$	0.13 ± 1.14	0.13 ± 0.33	$0.06\pm0.05^{\dagger\ddagger}$	0.64 ± 3.57	0.78 ± 3.79	
	A-P	$0.10 \pm 0.05^{\S\dagger}$	-1.40 ± 4.79	1.83 ± 7.96	$0.18 \pm 0.14^{\S \dagger}$	$1.12 \pm 3.21^{\S}$	0.55 ± 0.99 §	
Pudding	S-I	$0.20\pm0.11^{\S\ddagger}$	-0.13 ± 0.74	0.06 ± 0.07	$0.27\pm0.18^{\S\ddagger}$	$-0.05 \pm 1.87^{\S}$	0.24 ± 0.46 §	
	M-L	$0.05 \pm 0.03^{\dagger \ddagger}$	0.20 ± 1.07	0.08 ± 0.21	$0.08 \pm 0.10^{\dagger \ddagger}$	0.05 ± 0.45	0.41 ± 1.58	

 σ = Standard deviation, ξ = Skewness, γ = Kurtosis

Values were statistically significant when probability is less than 0.05 (*p-value* < 0.05).

of thin swallows in the non-stroke group (p < 0.005). In particular, the M-L axis for the thin non-stroke group had a greater entropy rate than the A-P (p < 0.01) and S-I (p < 0.01) signals. Secondly, the A-P entropy for nectar swallows in the non-stroke group was higher than the M-L entropy (p=0.0413).

Considering evaluation of randomness in swallow signals, the statistical analysis for LZC presented the most directional differences among the extracted information-theoretic features in this study. As demonstrated in Table 3, the S-I signals exhibited larger LZC than M-L signals in all groups of study (p < 0.005), except within the thin stroke group and the nectar the non-stroke group. In addition, the LZC for the S-I signals was higher than for A-P signals in thin and nectar swallows in the stroke group and pudding swallows in the non-stroke group (p < 0.01).

D. Frequency Domain Analysis

Table 4 represents the summary of frequency features considered in the current study. The results for most of the groups showed no significant directional difference for peak frequencies (f) as well as centroid frequencies (\hat{f}) , except for the following cases. First, considering pudding swallows in the stroke group, the A-P signals had significantly smaller peak frequencies than the S-I signals (p < 0.01) and greater peak frequencies than M-L signals (p < 0.05). Moreover, the S-I signals had lower centroid frequencies than the A-P signals for non-stroke pudding swallows (p < 0.02) and the M-L signals for stroke thin swallows (p < 0.02).

Additionally, the bandwidth of A-P signals was considerably lower than for the M-L axis in the non-stroke group for thin and nectar swallows (p < 0.03), and for the thin viscosity in

TABLE 3 Information-theoretic features (mean \pm standard deviation) extracted from the three anatomical axes of swallowing signals grouped by the participants' stroke history and the viscosity

		Stro	ke	Non-Stroke		
		LZC_s	ρ_s*10^{-1}	LZC_{ns}	$\rho_{ns}*10^{-1}$	
	A-P	$0.05\pm0.02^{\S}$	9.86 ± 0.01	0.06 ± 0.02	$9.86\pm0.01^{\S\dagger}$	
Thin	S-I	0.06 ± 0.02 §	9.87 ± 0.01	$0.07\pm0.02^{\ddagger}$	9.88 ± 0.00 §‡	
	M-L	0.06 ± 0.02	9.89 ± 0.00	$0.06\pm0.02^{\ddagger}$	$9.89 \pm 0.00^{\dagger \ddagger}$	
=	A-P	0.06 ± 0.03 §	9.89 ± 0.01	0.06 ± 0.02	$9.88 \pm 0.01^{\dagger}$	
Nectar	S-I	$0.07\pm0.02^{\S\ddagger}$	9.89 ± 0.00	0.06 ± 0.03	9.89 ± 0.00	
	M-L	$0.06 \pm 0.02^{\ddagger}$	9.89 ± 0.00	0.06 ± 0.02	$9.90\pm0.00^\dagger$	
gu	A-P	0.06 ± 0.03	9.90 ± 0.00	0.05 ± 0.02 §	9.88 ± 0.00	
Pudding	S-I	$0.07\pm0.02^{\ddagger}$	9.89 ± 0.00	$0.07\pm0.03^{\S\ddagger}$	9.89 ± 0.00	
	M-L	$0.05 \pm 0.02^{\ddagger}$	9.90 ± 0.00	$0.06 \pm 0.02^{\ddagger}$	9.89 ± 0.00	

LZC= Lempel-Ziv Complexity, ρ = Entropy rate

Values were statistically significant when probability is less than 0.05 (*p-value* < 0.05).

^{† =} Statistical difference between A-P and M-L axes

^{‡ =} Statistical difference between S-I and M-L axes

^{§ =} Statistical difference between A-P and S-I axes

^{† =} Statistical difference between A-P and M-L axes

^{‡ =} Statistical difference between S-I and M-L axes

^{§ =} Statistical difference between A-P and S-I axes

TABLE 4
FREQUENCY FEATURES (MEAN \pm STANDARD DEVIATION) EXTRACTED FROM THE THREE ANATOMICAL AXES OF SWALLOWING SIGNALS GROUPED BY
THE PARTICIPANTS' STROKE HISTORY AND THE VISCOSITY

		Stroke				Non-Stroke			
		$f_s(Hz)*10^2$	$\hat{f}_s(Hz)*10^3$	$BW_s(Hz)*10^2$	$f_{ns}(Hz)*10^2$	$\hat{f}_{ns}(Hz)*10^2$	$BW_{ns}(Hz)*10^2$		
	A-P	0.20 ± 0.68	0.41 ± 1.15	$4.22\pm5.75^\dagger$	0.15 ± 0.42	1.19 ± 2.12	$2.10 \pm 3.53^{\S\dagger}$		
Thin	S-I	0.11 ± 0.11	$0.08\pm0.07^{\ddagger}$	1.46 ± 1.46	0.12 ± 0.27	0.96 ± 3.91	1.45 ± 3.38 §		
	M-L	0.06 ± 0.11	$0.04 \pm 0.10^{\ddagger}$	$0.98\pm1.61^\dagger$	0.08 ± 0.23	0.95 ± 4.66	$1.16\pm3.30^{\dagger}$		
=	A-P	1.06 ± 8.04	0.29 ± 0.96	3.38 ± 6.12	0.08 ± 0.20	1.72 ± 3.42	$3.22 \pm 5.33^{\dagger}$		
Nectar	S-I	0.12 ± 0.28	0.04 ± 0.06	1.01 ± 1.42	0.12 ± 0.43	1.16 ± 3.98	1.90 ± 3.92		
Z	M-L	0.06 ± 0.13	0.03 ± 0.06	0.81 ± 1.02	0.06 ± 0.08	1.01 ± 4.80	$1.24\pm3.95^\dagger$		
	A-P	$0.03\pm0.02^{\S\dagger}$	0.29 ± 1.22	2.50 ± 6.25	0.12 ± 0.36	1.41 ± 2.28 §	2.67 ± 3.62		
Pudding	S-I	$0.06 \pm 0.02^{\S}$	0.03 ± 0.04	0.80 ± 1.57	0.09 ± 0.09	$0.96 \pm 2.07^{\S}$	2.06 ± 4.07		
	M-L	$0.05\pm0.03^\dagger$	0.02 ± 0.03	1.35 ± 1.93	0.08 ± 0.18	1.19 ± 4.17	2.05 ± 5.66		

 $f = \text{Peak frequency}, \ \hat{f} = \text{Spectral centroid}, \ BW = \text{Bandwidth}$

† = Statistical difference between A-P and M-L axes

‡ = Statistical difference between S-I and M-L axes

§ = Statistical difference between A-P and S-I axes

Values were statistically significant when probability is less than 0.05 (*p-value* < 0.05).

the stroke group (p < 0.05).

E. Time-Frequency Domain Analysis

In the time-frequency domain, the wavelet entropy rate (Θ_X) results for the three anatomical directions in stroke and non-stroke groups are represented in Table 5. The S-I signals had greater Θ_X than M-L signals for all cases (p < 0.01), except for pudding swallows in the stroke group.

TABLE 5 Time-frequency features (mean \pm standard deviation) extracted from the three anatomical axes of swallowing signals grouped by the participants' stroke history and the viscosity

		Stroke	Non-Stroke
		Θ_{X_s}	$\Theta_{X_{ns}}$
	A-P	1.07 ± 0.85	$1.09 \pm 0.81^{\dagger}$
Thin	S-I	$1.02\pm0.74^{\ddagger}$	$1.19 \pm 0.73^{\ddagger}$
I	M-L	$0.53 \pm 0.59^{\ddagger}$	$0.68 \pm 0.69^{\dagger\ddagger}$
	A-P	0.73 ± 0.75 §	$1.01 \pm 0.83^{\dagger}$
Nectar	S-I	$0.83 \pm 0.70^{\S}$	$1.11 \pm 0.77^{\ddagger}$
Z	M-L	0.48 ± 0.52	$0.67 \pm 0.66^{\dagger \ddagger}$
gu	A-P	0.62 ± 0.64	0.91 ± 0.75
Pudding	S-I	0.67 ± 0.56	$0.99 \pm 0.75^{\ddagger}$
Pu	M-L	0.34 ± 0.31	$0.67 \pm 0.78^{\ddagger}$

 Θ_X = Wavelet entropy

† = Statistical difference between A-P and M-L axes

‡ = Statistical difference between S-I and M-L axes

 $\S = Statistical difference between A-P and S-I axes$

Values were statistically significant when probability is less than 0.05 (p-value < 0.05).

F. Summary of All Findings

A total number of 198 comparisons were performed between the features calculated for the A-P, S-I, and M-L

directions of swallowing signals grouped by the viscosity and participants stroke history. Among 198 directional comparison cases, only 47 cases showed directional differences, which is $\approx 24\%$ of total number of comparisons. As we can see in Figure 4, most of the directional dissimilarities percentage was allocated to the standard deviation (37.5%), LCZ (16.67%) and wavelet entropy rate (14.59%).

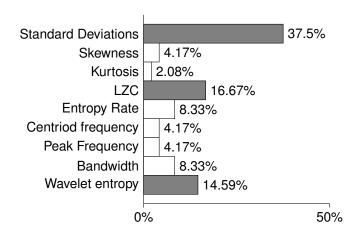


Fig. 4. Percentages of dissimilarities between axes of swallowing accelerometer for each of computed features. It is calculated by dividing the number of found dissimilarities cases to total number of comparisons for each of computed features.

Table 6 summarizes all of our findings. Each circle specifies the existing differences between the values of the computed features for the two directions. The rows and column that are highlighted in the orange and green colors indicate the features and the viscosity group, respectively, that presented the largest number of directional dissimilarities. First, we consider the features that give us the biggest numbers of directional difference among the various groups of study. Results obtained for the standard deviation (SD), Lempel-Ziv complexity (LZC), and wavelet entropy rate features showed

18, 8 and 7 directional dissimilarities cases, respectively, representing $\approx 69\%$ of the total dissimilarities. $36\%,\,34\%$ and 30% of the directional differences were found in comparisons between S-I & A-P, S-I & M-L and M-L & A-P, respectively. Generally, the S-I signals demonstrated higher values in SD, LZC, and wavelet entropy comparing with M-L and A-P signals. Secondly, while considering the viscosity and the participants' stroke history, all groups of study showed the minimum 6 directional dissimilarities in measured features. Thin swallows in the non-stroke group demonstrated the most directional dissimilarities among all of the groups of study, as highlighted in light green in the Table 6.

IV. DISCUSSION

The present study provides a quantitative characterization of the three anatomical directions of swallowing signals recorded by a tri-axial accelerometer during videofluoroscopy in patients with swallowing difficulties. This is the first study to consider the M-L axis in swallowing signals recorded during simultaneous videofluoroscopic imaging. Multiple features in time, information-theoretic, frequency, and time-frequency domains were extracted from the swallowing signals for each of the A-P, S-I, and M-L directions.

The efficiency of swallowing process is reduced with aging [59], [60] which might be reflected in swallowing vibrations recorded by accelerometer [38], [61]. However, in this study, age did not effect our result of directional differences as it was expected because of the short range of participants (72 participants with mean age of 63.94). Our data were gathered from subjects suspected of swallowing difficulties who referred for videofluoroscopic examination which includes most elderly population. In addition, we did not observe sex effects on our results. Although few studies have shown small sexrelated differences in swallowing [38], [61] due to differences in the anatomy of the neck and throat between males and females [59], [60], [62]. The absence of sex dependency in our study might be as a result of applying different methodology and most importantly recruiting unhealthy subjects who suffer from swallowing difficulties which could dissemble the effects of any other factors including sex on swallow vibrations.

Time-domain analysis showed all swallow signals in three directions demonstrated a low standard deviation, low positive or negative skewness, and high kurtosis values indicating dense symmetry distribution of swallow signals with decreasing amplitude values rapidly below and above the mean. The S-I signals had higher standard deviation values than the A-P and M-L signals, which indicates that the upward movement of the hyolaryngeal structure was actually more variable in amplitude than sideways and forward movements during swallowing. In addition, the A-P signals of pudding swallows in the non-stroke group showed higher values in skewness and kurtosis than the S-I signals. This suggests that the intensity of the forces in the forward direction were smaller for longer periods of a swallow and had a peaked distribution when compared to vertically directed forces.

Based on the results of the information-theoretic features, the swallowing signals in the three directions demonstrated predictable and regular behavior, which is representative of the presence of periodic non-random patterns in swallowing signals, in all the groups of the study as they showed high values for the entropy rate (around 0.99), and low values for the LZC (ranges from 0.05 to 0.07). However, the degree of predictability and regularity differed between the anatomical directions in some groups of study. For instance, the thin swallows in the non-stroke group, M-L and S-I directions, related to the sideways and upward movements of the hyolaryngeal structure, presented more regular behavior than the A-P direction, related to the forward movement of the hyolaryngeal structure. Furthermore, the M-L and A-P signals exhibited more predictable behavior, lower LZC, than S-I signals in most groups of the study.

In this study, the swallowing signals demonstrated few directional dissimilarities in frequency domain features, which indicates that the energy distribution over the range of the swallowing signal frequencies are approximately the same among the three axes. One of the considerable dissimilarities was the bandwidth of thin swallows in the non-stroke group. The A-P signals in this group presented a wider spread of frequency components than the S-I signals when showing a narrower spread compared to the M-L signals. Moreover, the M-L signals of the pudding swallows in the stroke group had a higher peak frequency than the A-P signals, while the A-P signals had higher frequencies than the S-I signals. A higher peak frequency indicates a higher rate of change for a signal.

Lastly, we discussed the results of wavelet entropy which provide information about the relative energy associated with the different frequency bands of the swallowing signals. In this study, the S-I signals demonstrated greater wavelet entropy than the M-L signals in all groups of the study except for the pudding swallows in the stroke group. It means that the signal related to the upward movement of the hyolaryngeal structure during swallowing behaves more disorderly in the time-frequency domain than the sideways movement.

To conclude, we investigated for the first time the existence of swallow vibrations in M-L axis in patients with swallowing difficulties. The result showed that the behavior of swallow vibrations in M-L axis, representing the sideways movement of hyolaryngeal structure during swallowing function, was pretty similar to the behavior of swallow vibrations in two other axes. In general, despite few dissimilarities obtained from the directional feature comparisons discussed before in the discussion, the swallowing signals in all three anatomical directions demonstrated similar behaviors in most computed features and among different groups of our study. Our finding seems to bring the practicability of applying dual/tri-axial accelerometer in swallowing signals analysis into question as it suggests that we can infer about the behavior of one of the axes based on the other axes by considering the viscosity of the swallows and participant's stroke history. The lack of existence of directional differences in swallowing signals also raises the question of whether the physiological sources of the three axes are dependent upon each other. Further research of swallowing physiology with tri-axial accelerometry using concurrent videofluoroscopic images must be investigated to understand the sources of swallowing-related vibrations. In

TABLE 6

Summary of directional differences among groups of study. All dots (red and black dots) indicate the existence of directional dissimilarities, which means the p-value of the comparison is statistically significant at the $p_1^*0.05$ level. Axes column demonstrates which two axes are compared to each other. Red dots indicate that first axis in the Axes column have a higher value of the computed feature in Features column than the second axis while black dots indicate the second axis have higher value than the first axis.

		stroke		Non-stroke			
		п	Nectar	Pudding	п	Nectar	Pudding
Features	Axes	Thin	Š	Puc	Thin	Š	Puc
	A-P & S-I	•	•	•	•	•	•
Standard Deviation	A-P & M-L	•	•	•	•	•	•
	M-L & S-I	•	•	•	•	•	•
	A-P & S-I				•		•
Skewness	A-P & M-L						
	M-L & S-I						
	A-P & S-I						•
Kurtosis	A-P & M-L						
	M-L & S-I						
	A-P & S-I	•	•				•
LZC	A-P & M-L						
	M-L & S-I		•	•	•		•
	A-P & S-I				•		
Entropy Rate	A-P & M-L				•	•	
	M-L & S-I				•		
	A-P & S-I						•
Centroid Frequency	A-P & M-L						
	M-L & S-I	•					
	A-P & S-I			•			
Peak Frequency	A-P & M-L			•			
	M-L & S-I						
	A-P & S-I				•		
Bandwidth	A-P & M-L	•			•	•	
	M-L & S-I						
	A-P & S-I		•				
Wavelet Entropy	A-P & M-L				•	•	
	M-L & S-I	•			•	•	•

addition, we can only interpenetrates from our results that three directions are similar in most of the computed features without finding out which one would be more successful to classify healthy and unhealthy swallows accurately. Therefore, examining the ability and validity of the three directions to classify swallows would be a valuable investigation as the final goal of swallowing vibratory studies is swallows classification.

Furthermore, the manual segmentation of swallowing signals in this study increases the rate of human error due to fatigue and decrease the consistency in segmentation criteria. Therefore, investigating a swallow segmentation algorithm that can extract each swallow from the recorded signals with high accuracy would be an important contribution. Finally, the Mendelsohn or supraglottic swallow maneuver has been used to treat patients with dysphagia for many years sometimes as a compensatory strategy to help the bolus pass more efficiently through the pharynx and sometimes as part of a rehabilitative exercise program. That would be a good research to study the differences between swallows with maneuvers and swallows

in natural situations in patients with dysphagia to evaluate the effectiveness of rehabilitation programs, and to consider the cervical method as a screening test in rehabilitation centers. Specially previous studies showed the possible effects of neutral head-neck posture and the head-neck flexion (chintuck) position on swallowing signals in the A-P and S-I axes [41], [46].

V. CONCLUSION

In this study, we have compared swallowing accelerometry signals among the anterior-posterior, superior-inferior and medial-lateral directions in patients with dysphagia. We considered 9 features from 881 swallowing accelerometry signals in various domains including: time, information-theoretic, frequency, and time-frequency domains. The results indicated that movements related to the vibrations emanating from the hyolaryngeal complex in all three directions have similar behavior in most of the extracted features in this study, but there were few directional differences as well. First, standard

deviation, LZC, and wavelet entropy values demonstrated the most directional differences among all the computed features. Secondly, S-I direction showed more complex and irregular behavior with variable amplitude than the M-L and A-P directions. Thirdly, considering viscosity and participant's pathology, thin swallows in the non-stroke group presented the most directional differences among all groups of study.

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REFERENCES

- J. A. Logemann and J. A. Logemann, "Evaluation and treatment of swallowing disorders," Nat. Student Speech Language Hearing Assoc. J., pp. 38–50, 1983.
- [2] A. J. Miller, "The neurobiology of swallowing and dysphagia," Dev. Disabil. Res. Rev., vol. 14, no. 2, pp. 77–86, 2008.
- [3] L. Sura, A. Madhavan, G. Carnaby, and M. A. Crary, "Dysphagia in the elderly: management and nutritional considerations," *Clin. Interv. Aging*, vol. 7, no. 287, p. 98, 2012.
- [4] P.-H. Chen, J. S. Golub, E. R. Hapner, and M. M. Johns III, "Prevalence of perceived dysphagia and quality-of-life impairment in a geriatric population," *Dysphagia*, vol. 24, no. 1, pp. 1–6, 2009.
- [5] J. Chai, F. Chu, T. Chow, and N. Shum, "Prevalence of malnutrition and its risk factors in stroke patients residing in an infirmary," *Singapore Med. J.*, vol. 49, no. 4, p. 290, 2008.
- [6] C. Gordon, R. L. Hewer, and D. T. Wade, "Dysphagia in acute stroke." Br. Med. J., vol. 295, no. 6595, pp. 411–414, 1987.
- [7] M. Aslam and M. F. Vaezi, "Dysphagia in the elderly," Gastroenterology & Hepatology, vol. 9, no. 12, p. 784, 2013.
- [8] D. I. Rosenthal, J. S. Lewin, and A. Eisbruch, "Prevention and treatment of dysphagia and aspiration after chemoradiation for head and neck cancer," J. Clin. Oncol., vol. 24, no. 17, pp. 2636–2643, 2006.
- [9] N. Bhattacharyya, "The prevalence of dysphagia among adults in the united states," *Otolaryngol Head Neck Surg.*, vol. 151, no. 5, pp. 765– 769, 2014.
- [10] N. Roy, J. Stemple, R. M. Merrill, and L. Thomas, "Dysphagia in the elderly: preliminary evidence of prevalence, risk factors, and socioemotional effects," *Ann. Otol. Rhinol. Laryngol.*, vol. 116, no. 11, pp. 858–865, 2007.
- [11] N. Sheth and W. C. Diner, "Swallowing problems in the elderly," *Dysphagia*, vol. 2, no. 4, pp. 209–215, 1988.
- [12] J. S. Schindler and J. H. Kelly, "Swallowing disorders in the elderly," The Laryngoscope, vol. 112, no. 4, pp. 589–602, 2002.
- [13] B. A. Mathers-Schmidt and M. Kurlinski, "Dysphagia evaluation practices: inconsistencies in clinical assessment and instrumental examination decision-making," *Dysphagia*, vol. 18, no. 2, pp. 114–125, 2003.
- [14] S. Singh and S. Hamdy, "Dysphagia in stroke patients," *Postgrad. Med. J.*, vol. 82, no. 968, pp. 383–391, 2006.
- [15] S. E. Langmore, S. M. Kenneth, and N. Olsen, "Fiberoptic endoscopic examination of swallowing safety: a new procedure," *Dysphagia*, vol. 2, no. 4, pp. 216–219, 1988.
- [16] W. Chih-Hsiu, H. Tzu-Yu, C. Jiann-Chyuan, C. Yeun-Chung, and L. Shiann-Yann, "Evaluation of swallowing safety with fiberoptic endoscope: comparison with videofluoroscopic technique," *The Laryngo-scope*, vol. 107, no. 3, pp. 396–401, 1997.
- [17] N. P. Reddy, A. Katakam, V. Gupta, R. Unnikrishnan, J. Narayanan, and E. P. Canilang, "Measurements of acceleration during videofluorographic evaluation of dysphagic patients," *Med. Eng. Phys.*, vol. 22, no. 6, pp. 405–412, 2000.
- [18] R. D. Wilson and E. C. Howe, "A cost-effectiveness analysis of screening methods for dysphagia after stroke," *Amer. J. Phys. Med. Rehabil.*, vol. 4, no. 4, pp. 273–282, 2012.
- [19] D. Zoratto, T. Chau, and C. Steele, "Hyolaryngeal excursion as the physiological source of swallowing accelerometry signals," *Physiol. Meas.*, vol. 31, no. 6, p. 843, 2010.

- [20] P. M. Zenner, D. S. Losinski, and R. H. Mills, "Using cervical auscultation in the clinical dysphagia examination in long-term care," *Dysphagia*, vol. 10, no. 1, pp. 27–31, 1995.
- [21] J. A. Cichero and B. E. Murdoch, "Detection of swallowing sounds: methodology revisited," *Dysphagia*, vol. 17, no. 1, pp. 40–49, 2002.
- [22] J. Lee, C. Steele, and T. Chau, "Time and time-frequency characterization of dual-axis swallowing accelerometry signals," *Physiol. Meas.*, vol. 29, no. 9, p. 1105, 2008.
- [23] C. M. Steele, G. L. Bailey, T. Chau, S. M. Molfenter, M. Oshalla, A. A. Waito, and D. C. Zoratto, "The relationship between hyoid and laryngeal displacement and swallowing impairment," *Clin. Otolaryngol.*, vol. 36, no. 1, pp. 30–36, 2011.
- [24] A. Perlman, P. Palmer, T. McCulloch, and D. Vandaele, "Electromyographic activity from human laryngeal, pharyngeal, and submental muscles during swallowing," *J. Appl. Physiol.*, vol. 86, no. 5, pp. 1663–1669, 1999.
- [25] Y. Kim and G. H. McCullough, "Maximum hyoid displacement in normal swallowing," *Dysphagia*, vol. 23, no. 3, pp. 274–279, 2008.
- [26] L. Lazareck and Z. Moussavi, "Swallowing sound characteristics in healthy and dysphagic individuals," in *The 26th Ann. Int. Conf. IEEE Eng. Med. Biol. Soc.*, vol. 2, San Francisco, CA, September 1-5 2004, pp. 3820–3823.
- [27] A. Yadollahi and Z. Moussavi, "Feature selection for swallowing sounds classification," in *The 29th Ann. Int. Conf. IEEE Eng. Med. Biol. Soc.*, Lyon, FR, August 22-26 2007, pp. 3172–3175.
- [28] S. Youmans and J. Stierwalt, "Normal swallowing acoustics across age, gender, bolus viscosity, and bolus volume," *Dysphagia*, vol. 26, no. 4, pp. 374–384, December 2011.
- [29] —, "An acoustic profile of normal swallowing," *Dysphagia*, vol. 20, no. 3, pp. 195–209, September 2005.
- [30] K. Takahashi, M. Groher, and K. Michi, "Symmetry and reproducibility of swallowing sounds," *Dysphagia*, vol. 9, no. 3, pp. 168–173, September 1994.
- [31] J. Lee, C. M. Steele, and T. Chau, "Classification of healthy and abnormal swallows based on accelerometry and nasal airflow signals," *Artif. Intell. Med.*, vol. 52, no. 1, pp. 17–25, 2011.
- [32] M. S. Nikjoo, C. M. Steele, E. Sejdić, and T. Chau, "Automatic discrimination between safe and unsafe swallowing using a reputationbased classifier," *Biomed. Eng. OnLine*, vol. 10, no. 1, p. 100, 2011.
- [33] J. M. Dudik, A. Kurosu, J. L. Coyle, and E. Sejdić, "A comparative analysis of dbscan, k-means, and quadratic variation algorithms for automatic identification of swallows from swallowing accelerometry signals," *Comput. Biol med.*, vol. 59, pp. 10–18, 2015.
- [34] J. M. Dudik, J. L. Coyle, and E. Sejdic, "Dysphagia screening: Contributions of cervical auscultation signals and modern signal-processing techniques," *IEEE Trans. HumanMach. Syst*, vol. 45, no. 4, pp. 465–477, 2015
- [35] J. Lee, C. M. Steele, and T. Chau, "Swallow segmentation with artificial neural networks and multi-sensor fusion," *Med. Eng. Phys.*, vol. 31, no. 9, pp. 1049–1055, 2009.
- [36] E. Sejdić, C. M. Steele, and T. Chau, "Understanding the statistical persistence of dual-axis swallowing accelerometry signals," *Comput. Biol. Med.*, vol. 40, no. 11, pp. 839–844, 2010.
- [37] E. Sejdić, V. Komisar, C. M. Steele, and T. Chau, "Baseline characteristics of dual-axis cervical accelerometry signals," *Ann. Biomed. Eng.*, vol. 38, no. 3, pp. 1048–1059, 2010.
- [38] J. M. Dudik, I. Jestrović, B. Luan, J. L. Coyle, and E. Sejdić, "A comparative analysis of swallowing accelerometry and sounds during saliva swallows," *Biomed. Eng. Online*, vol. 14, no. 3, pp. 1–15, January 2015.
- [39] T. Ayuse, S. Ishitobi, S. Kurata, E. Sakamoto, I. Okayasu, and K. Oi, "Effect of reclining and chin-tuck position on the coordination between respiration and swallowing," *J. Oral Rehabil.*, vol. 33, no. 6, pp. 402– 408, 2006.
- [40] K. Takahashi, M. E. Groher, and K.-i. Michi, "Methodology for detecting swallowing sounds," *Dysphagia*, vol. 9, no. 1, pp. 54–62, 1994.
- [41] I. Jestrovic, J. M. Dudik, B. Luan, J. L. Coyle, and E. Sejdić, "The effects of increased fluid viscosity on swallowing sounds in healthy adults," *Biomed.l Eng. OnLine*, vol. 12, no. 1, p. 90, 2013.
- [42] M. A. Holas, K. L. DePippo, and M. J. Reding, "Aspiration and relative risk of medical complications following stroke," *Arch. Neurol.*, vol. 51, no. 10, pp. 1051–1053, 1994.
- [43] D. Smithard, N. Smeeton, and C. Wolfe, "Long-term outcome after stroke: does dysphagia matter?" Age Ageing, vol. 36, no. 1, pp. 90– 94, 2007.

- [44] J. Lee, E. Sejdić, C. M. Steele, T. Chau et al., "Effects of liquid stimuli on dual-axis swallowing accelerometry signals in a healthy population," *Biomed. Eng. OnLine*, vol. 9, no. 7, 2010.
- [45] E. Sejdić, C. M. Steele, and T. Chau, "A method for removal of low frequency components associated with head movements from dual-axis swallowing accelerometry signals," *PLoS One*, vol. 7, no. 3, p. e33464, 2012.
- [46] ——, "The effects of head movement on dual-axis cervical accelerometry signals," BMC Res. Notes, vol. 3, no. 1, p. 269, 2010.
- [47] —, "A procedure for denoising dual-axis swallowing accelerometry signals," *Physiol. Meas.*, vol. 31, no. 1, p. N1, 2010.
- [48] E. Sejdić, K. A. Lowry, J. Bellanca, M. S. Redfern, and J. S. Brach, "A comprehensive assessment of gait accelerometry signals in time, frequency and time-frequency domains," *IEEE Trans. Neural Syst. Rehabil. Eng.*, vol. 22, no. 3, pp. 603–612, 2014.
- [49] L. T. DeCarlo, "On the meaning and use of kurtosis." Psychol. Methods, vol. 2, no. 3, p. 292, 1997.
- [50] M. Aboy, R. Hornero, D. Abásolo, and D. Álvarez, "Interpretation of the lempel-ziv complexity measure in the context of biomedical signal analysis," *IEEE Trans. Biomed. Eng.*, vol. 53, no. 11, pp. 2282–2288, 2006.
- [51] J. Hu, J. Gao, and J. C. Principe, "Analysis of biomedical signals by the lempel-ziv complexity: the effect of finite data size," *IEEE Trans. Biomed. Eng.*, vol. 53, no. 12, pp. 2606–2609, 2006.
- [52] A. Porta, S. Guzzetti, N. Montano, R. Furlan, M. Pagani, A. Malliani, and S. Cerutti, "Entropy, entropy rate, and pattern classification as tools to typify complexity in short heart period variability series," *IEEE Trans. Biomed. Eng.*, vol. 48, no. 11, pp. 1282–1291, 2001.
- [53] A. Porta, G. Baselli, D. Liberati, N. Montano, C. Cogliati, T. Gnecchi-Ruscone, A. Malliani, and S. Cerutti, "Measuring regularity by means of a corrected conditional entropy in sympathetic outflow," *Biol. Cybern.*, vol. 78, no. 1, pp. 71–78, 1998.
- [54] Y. Kumar, M. L. Dewal, and R. S. Anand, "Relative wavelet energy and wavelet entropy based epileptic brain signals classification," *Biomed. Eng. Lett.*, vol. 2, no. 3, pp. 147–157, 2012.
- [55] R. Q. Quiroga, O. A. Rosso, E. Başar, and M. Schürmann, "Wavelet entropy in event-related potentials: a new method shows ordering of eeg oscillations," *Biol. Cybern.*, vol. 84, no. 4, pp. 291–299, 2001.
- [56] B. Natwong, P. Sooraksa, C. Pintavirooj, S. Bunluechokchai, and W. Ussawawongaraya, "Wavelet entropy analysis of the high resolution ECG," in *Proc. of the 1st IEEE Conf. on Ind. Electron. and Applicat.*, Singapore, May 24-26, 2006, pp. 1–4.
- [57] E. Śejdić, I. Djurović, and J. Jiang, "Time-frequency feature representation using energy concentration: An overview of recent advances," *Digit. Signal Process.*, vol. 19, no. 1, pp. 153–183, 2009.
- [58] O. A. Rosso, S. Blanco, J. Yordanova, V. Kolev, A. Figliola, M. Schürmann, and E. Başar, "Wavelet entropy: a new tool for analysis of short duration brain electrical signals," *J. Neurosci. Methods*, vol. 105, no. 1, pp. 65–75, 2001.
- [59] G. McKee, B. Johnston, G. McBride, and W. Primrose, "Does age or sex affect pharyngeal swallowing?" *Clin. Otolaryngol. Allied Sci.*, vol. 23, no. 2, pp. 100–106, 1998.
- [60] O. Ekberg and L. Wahlgren, "Dysfunction of pharyngeal swallowing a cineradiographic investigation in 854 dysphagial patients," *Acta Radiol. Diagn.*, vol. 26, no. 4, pp. 389–395, 1985.
- [61] E. Sejdić, C. M. Steele, and T. Chau, "Segmentation of dual-axis swallowing accelerometry signals in healthy subjects with analysis of anthropometric effects on duration of swallowing activities," *IEEE Trans. Biomed. Eng.*, vol. 56, no. 4, pp. 1090–1097, 2009.
- [62] V. Kuhl, B. Eicke, M. Dieterich, and P. Urban, "Sonographic analysis of laryngeal elevation during swallowing," *J. Neurol.*, vol. 250, no. 3, pp. 333–337, 2003.