Wearable Healthcare Systems: A Single Channel Accelerometer Based Anomaly Detector for Studies of Gait Freezing in Parkinson's Disease

Thuy T. Pham*, Diep N. Nguyen, Eryk Dutkiewicz, Alistair L. McEwan, and Philip H.W. Leong

Abstract—The causality of gait freezing in patients with advanced Parkinson's disease is still not fully understood. Clinicians are interested in investigating the freezing of gait (FoG) histogram of patients in their daily life. To that end, one needs a real-time signal processing platform that can help record freezing information (e.g., timing and the duration of every gait freezing occurrences). Wearable wireless sensors have been proposed to monitor FoG epochs. Existing automated methods using accelerometers have been introduced with high accuracy performance only for subjectdependent settings (e.g., an individual offline training process). This is a troublesome for large scale out-of-lab deployment and time-consuming. In this work, we used spectral coherence analysis for accelerometer data to apply an anomaly detection approach. Conventional features such as energy and freezing index are introduced to help refine normal epochs while the anomaly scores from spectral coherence measures define FoG epochs. Using this new set of features, our new FoG detector for subject-independent settings achieves the mean $\pm SD$ sensitivity (specificity) of $89.2 \pm 0.3\%$ ($95.6 \pm 0.3\%$). To our best knowledge, this is the best performance for automated subject-independent approaches in literature of freezing of gait detection.

Index Terms—Anomaly detection, wearable sensor, aged care.

I. INTRODUCTION

Real-time monitoring systems using pervasive computing have been providing more effective aged care systems. One of the most common symptoms in patients with advanced Parkinson's disease (PD) is freezing of gait (FoG) that associate with falls [1], [2], [3]. FoG is described as a motor block of movement (e.g., before gait initiation, during turns, or when meeting obstacles) [1]. Self-report diaries from patients (e.g. the Unified Parkinson's Disease Rating Scale-UPDRS), that is commonly used as current clinical assessment of FoG, is subjective and has poor agreement with expert labels [4]. Freezing of Gait Questionnaire [5]) and manual video analysis of walking [6], [7] have been introduced to improve UPDRS but they are unfortunately also subjective [8]. The intra-rater reliability of existing manual video assessments was low and not robust (within or between patient sites) [8]. Hence, objective FoG detection is much desirable [9][10]. Moreover, it is difficult to invoke freezing status in participants during routine clinical exams [11], thus out-of-lab deployment is also of interest.

*correspondence: thuy.pham@uts.edu.au. T. Pham is with the School of Computing and Communications, University of Technology Sydney and the Dept. of Electrical and Information Engineering, The University of Sydney. D. Nguyen, and E. Dutkiewicz are with School of Computing and Communications, University of Technology Sydney, NSW, Australia. A. McEwan and P. Leong are with the Dept. of Electrical and Information Engineering, The University of Sydney, NSW, Australia.

Recently, small wearable devices that were designed to detect FoG events have been proposed [12], [9], [13], [14], [15], [16], [17], [18], [19], [10]. Most of these works used data from accelerometers and a few used electromyographic (EMG) data. There have been works proposed for online operation mode (e.g., [15], [16], [17]); others have been implemented in an offline methods. In terms of feature extraction, frequency analysis techniques for energy (P) [12][15] and freezing index (FI) [9] have been most used. Regarding detection algorithms, the Context Recognition Network (CRN) Toolbox [20] was utilized in [15] or machine learning based Random Forests (RF) used in [17]. Most of other works have utilized simple thresholding methods [12], [9], [13], [14], [18], [19], [10]. Nevertheless, only moderate performance was achieved in subject-independent settings (better results were obtained using subject-dependent thresholds).

When deploying as a wearable healthcare assistant in outof-lab environment, subject-independent settings are highly desirable. Avability of a specialist and training the machine for each user is not practical for a massive deployment. In one of our prelimitary works [21], we have intensively reviewed existing and our novel feature extraction methods towards the subject-independent settings. From a large pool of 244 feature candidates, we found 33 most relevant and discriminative ones by three saliency criteria (after two rounds of selection). During the third round of process, we consider FoG as a departure from normal gait therefore suggested to use anomaly detection techniques for FoG monitoring in real time. Using a simple thresholding filter to detect FoGs, the so-called anomaly score detector (ASD) [21] limited this list to seven features that have had high accuracy metrics. All of them were freezing index features. One of them is extracted from multi-channel and the rest are single channel. Though the vertical axis input at ankle sensor was often used in literature works has been also included in our final list of selection, several other single channels at other sensor locations have been showed comparable.

In this work, we aim to evaluate further the third round of feature selection in [21] in which the outcome most depends on a given detection model. Our previous model used a threshold calculated from the average values with the unity scaling factor. In this model, we investigate incorporating findings of freezing index's relevancy in a novel threshold calculation. In the previous result of feature ranking using correlation and clusterability metrics, spectral coherence of snapshots have been highly ranked. We are interested in the coherence feature as it is unitless and intiutively ranges from 0 to 1 for the

similarity metrics. Unitless features are often the best choice for subject-independent measurements as it can tolerate amplitude distortion due to artefacts during acquiring data especially for daily activities monitoring. Hence, we propose to use spectral coherence measurement for anomaly scores and incoporate abnormal freezing index for internal estimation of *normal* epochs.

The main contributions of this work are:

- A feature evaluation for combination of energy and spectral coherence features to refine the normal gait instances for FoG detector using anomaly scores is first reported.
- A novel model of FoG detection is proposed which, to the best of our knowledge, achieves the best reported performance for subject-independent settings.

II. METHODS

A. Data Set

We developed and tested our method on a recently published dataset from DAPHNet [15]. There were 10 patients with advanced Parkinson's disease (i.e., had regular freezing event). Note that the proposed method uses an unitless correlation measurement that may not suffer from amplitude-related subject variablity and the inherent scheme of dynamic threholds learned within the recording make the size of development dataset not need to be large.

Participants attached three tri-axial (x - anterior/posterior, y - medial/lateral, z - vertical) accelerometers at three locations: the shank, thigh, and lower back. Each subject implemented three walking tasks (about 10-15 minutes). Data from sensors were recorded at 64Hz and wirelessly transmitted (via a Bluetooth link). Labels/annotation were created by physiotherapists using video taping analysis to determine the timing of FoG events. According to the labels, FoG events last for $7.3\pm6.7s$ (50% less than 5.4s; 93.2% were shorter than 20s). We use this manual annotation as reference in performance evaluation for the proposed automated method.

B. Feature Extraction

1) Freezing index: Freezing gait has been found in the work [12] containing high frequency components $(6 \rightarrow 8Hz)$ compared with normal gait (2Hz). A freezing index (FI) is defined as the power in the *freeze* band $(3 \rightarrow 8Hz)$, P_H , divided by the power in the *locomotor* band $(0.5 \rightarrow 3Hz)$, P_L [9]. The power, P, and freeze index FI values are computed as in Eq.1 and Eq. 2.

$$P = P_H + P_L \tag{1}$$

$$FI = \frac{P_H}{P_L} \tag{2}$$

where

$$P_{H} = \frac{\sum_{i=H_{1}+1}^{H_{2}} [P_{XX}(i)] + \sum_{i=H_{1}}^{H_{2}-1} [P_{XX}(i)]}{2fs}$$
(3)

$$P_{L} = \frac{\sum_{i=L+1}^{H_{1}} [P_{XX}(i)] + \sum_{i=L}^{H_{1}-1} [P_{XX}(i)]}{2fs}$$
(4)

and
$$H_1 = \frac{3N_{FFT}}{fs}$$
, $H_2 = \frac{8N_{FFT}}{fs}$, $L = \frac{0.5N_{FFT}}{fs}$. $P_{XX}(\omega) = \mathfrak{F}_x(\omega).\mathfrak{F}_x(\omega)$.

2) Spectral Coherence: Let x and y be two consecutive data windows. The correlation in frequency between x and y, called spectral coherence C_{XY} , is measured using the Welch method [22] as $C_{XY}(\omega) = \frac{P_{XY}(\omega)}{\sqrt{P_{XX}(\omega).P_{YY}(\omega)}}$ where ω is frequency, $P_{XX}(\omega)$ is the power spectrum of signal x, $P_{YY}(\omega)$ is the power spectrum for signals x and y. When $P_{XX}(\omega) = 0$ or $P_{YY}(\omega) = 0$, then $P_{XY}(\omega) = 0$ and we assume that $C_{XY}(\omega)$ is zero. Note that when $C_{xy}(\omega)$ is small, x and y are weakly correlated in frequency domain.

C. Anomaly Detector

To detect freezing of gait events, we consider time periods of normal activities *normal behaviour* data and freezing epochs anomalies.

- 1) Normals and Anomalies: Normal epochs are predicted using the smoothness of activities and the dominated at lowband of the power distribution. This hypothesis has been demonstrated in our prelimitary work [21] when examining salient features from a large exploratory feature pool. Accumulating these *normal* epochs, we estimated continously the average of normal power features. Specifically in this work, the power and freezing index values P_x and FI_x have the average values μ_P and μ_{FI} respectively. If the current tested data epoch has abnormal P_x and FI_x (i.e., getting over the current thresholds) and has a failure test of the spectral coherence with the latest normal epoch, the current epoch is considered an anomaly. Otherwise, the theshold values are updated with this epoch. Note that our μ_P and μ_{FI} are only averaged continously from estimated normals thus are different from the fixed thresholds found in the literature (e.g., [9] [15] [17]) that are averaged from entire several measurements.
- 2) FoG Detection with μ_P and μ_{FI} : Let X represent a data segment recorded from a channel of an accelerometer attached to a subject. We compute a set of binary movement scores which consists of a low acceleration S_{AC} and a spectral coherence S_{speco} indicators. $S_{AC} = sgn((X (\overline{X} \sigma))_+)$ where \overline{X} and σ are the mean and standard deviation of X and sgn(x) is a sign function of x while $(x)_+$ is a function which returns x only if $x \ge 0$ otherwise it returns 0. $S_{speco} = 1$ if the number of peaks in $C_{xy}(\omega)$ that are greater than a specified correlation threshold level Θ_{speco} otherwise $S_{speco} = 0$.

From the learned information of *Normal*, dynamic thresholds μ_P and μ_{FI} are updated when a flag is asserted (Algorithm 1). Note that the flag is initialized at 10 to differentiate the beginning stage of the process from later stages of detection in which the flag changes between 1 and 0 (depending on if a FoG event is detected or not).

D. Performance Metrics

To evaluate accuracy performance of the proposed method, we note following measures. Windows which were labelled the same as annotation are True Positives (TP). Windows labelled as FoG which did not agree with the ground truth are False

Algorithm 1 FoG Detection Algorithm with μ_P and μ_{FI} .

```
Input:
X is from channel X
Output:
Out is FoG detection outcome
Parameters:
w is window size
\Theta is minimum coherence level
Main
  k=1; buff=2*w;
  flag=10;
  \Omega \Psi are empty;
  While (k < length(X) - buff)
                                        if flag = 10
    Normal =
                 X(last: last + w), if flag = 1
     [P_{Normal}, FI_{Normal}] = ComputePFI(Normal)
     \Omega \supseteq P_{Normal};
     \Psi \supseteq FI_{Normal}
     \mu_P = \text{mean}(\Omega);
     \mu_{FI} = \text{mean}(\Psi);
     [P_x, FI_x] = ComputePFI(x)
     S_{freeze} = ((P_x > \mu_P) & (FI_x > \mu_{FI}));
     S_{Ac}(k:k+buff) is computed from Eq.??
     S_{speco}=ComputeCxy(X_{buff}, Normal, w, \Theta);
     If (S_{freeze}(k:k+buff)=1)&
        ((S_{freeze}(k - buff : k - buff + w) = 0) &
        ((S_{Ac}(k-buff:k-buff+w)=0)
        ||(S_{speco}(k-buff:k-buff+w)=0)));
        Out=1; flag=1; last=k-buff;
     Else Out=0; flag=0;
     k=k+buf:
  End of while
Subfunctions:
[P,FI]=ComputePFI(x) by Eq. 1, 2
|S_{speco}| = ComputeCxy(x, Normal, w, \Theta)
     i=1; c=0;
     While (i < length(x))
        X_{win} = x(i:i+w);
        C_{XY} is computed from Normal and X_{win};
        N = \text{Count}(\text{Peak}(C_{XY}) > \Theta);
        S_{speco}(i:i+w)=(N=0);
        i = i + w;
     End of while
} end of subfunction
```

Positives (FP). Windows labelled as Normal by the proposed method but were annotated as FoG are False Negatives (FN). The window that was labelled as Normal by both are True Negative (TN). The sensitivity was calculated as $\frac{TP}{TP+FN}$ and the specificity was calculated as $\frac{TN}{TN+FP}$. We also use the harmonic mean of precision and sensitivity, F1-score [23]. F1-score= $\frac{2TP}{(2TP+FP+FN)}$ with best being 1 and worst 0.

III. RESULTS

A. Development Stage

To find the optimized parameters, we use the receiver operating characteristic (ROC) versus Θ_{speco} and w. Parameter Θ_{speco} to compute S_{speco} is varied in a wide range $(0 \rightarrow 1)$ in steps of 0.1). The window size of sliding method is also

examined from $0.2 \rightarrow 1$ s. Fig. 1 shows details of these ROCs measurement. Each illustration point in the figure, that is for a couple of (Θ_{speco}, w) , is the mean performance over three sensor locations (ankle, hip, trunk) and across subjects. When Θ_{speco} increases, the sensitivity increases dramatically from 0.28 ± 0.25 to 0.81 ± 0.29 while the specificity decreases slightly from 1.00 ± 0.05 to 0.81 ± 0.06 . This suggests a high precision configuration for any subject with a Θ_{speco} range of 0.4 - 0.6.

With the parameter settings of $(\Theta_{speco} = 0.4, w = 0.6s)$, the mean \pm SD of FoG event duration we detected was 11 ± 8 seconds with a range from 3s to 60s. These values are close to those identified from the annotations $(7.5 \pm 7s)$; range 0.5s + 40.5s. Our method and the annotations agreed that most of FoG events last less than 20s.

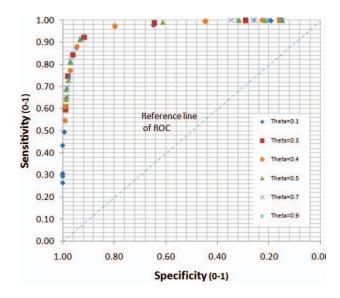


Fig. 1. a: Performance (ROC) of our method versus Θ_{speco} and window size w in a range of interest (0-1). The optimal $(\Theta_{speco}=0.4, w=0.6s)$ is suggested for a subject-independent setting.

Table I demonstrates that our dynamic threshold (settings (a): μ_P , μ_{FI}) are comparable to those personalized parameter of the literature works (settings (b): fixed subject-dependent Power-TH, Freeze-TH).

Among three locations, sensors at the knee yielded a higher sensitivity than the others. The distribution of metrics for are presented in Table. II.

B. FoG Detection Comparisons

Compared with the manual labels by specialists, we noticed the partial concordance of a moderate overall identification of freezing (e.g., fell to 86%) in the case of patient with foot drop (e.g., patient ID1). For participants who had no freezing of gait events, the proposed method also did not dectect any FoG event (e.g., patient ID10 and ID4).

According to results reported in the literature, the proposed method yields a better subject-independent performance (Table III). The compared results are reported as they were in

TABLE I

COMPARABLE VALUES OF ENERGY THRESHOLDS BETWEEN (A): MEAN DYNAMIC THRESHOLDS FOR EACH SUBJECT AUTOMATICALLY LEARNED IN OUR METHOD AND (B): RANGES OF FIXED THRESHOLDS FOR SUBJECT-DEPENDENT SETTINGS USED IN [15]. NOTE THAT OUR METHOD DOES NOT HAVE TO SPECIFY THESE VALUES BUT AUTOMATICALLY LEARNS AS VARIABLES. THIS ILLUSTRATION IS FOR EXPLANATION PURPOSES ONLY.

	(b) individual in [15]		(a) dynamic thresholds		
Subjects	PowerTH in 2 ^x	FreezeTH	μ_P in 2^x	μ_{FI}	
Patient 01	12	3	17.55	3.55	
Patient 02	16	1.8	18.78	2.11	
Patient 03	15	4	17.52	3.81	
Patient 05	11.5	1.6	17.30	1.67	
Patient 06	10	1	17.57	1.34	
Patient 07	8.5	2.5	17.04	1.36	
Patient 08	8	3	15.44	11.72	
Patient 09	7	2.5	16.7	2.19	

TABLE II
PERFORMANCE DISTRIBUTION AMONG SENSOR PLACEMENTS.

Sensor	Sensitivity	Specificity	F1-score
Ankle	$94.91 \pm 2.3\%$	$94.18 \pm 2.9\%$	$91.73 \pm 1.6\%$
Knee	$91.09 \pm 4.1\%$	$95.91 \pm 1.3\%$	$90.18 \pm 2\%$
Hip	$88.13 \pm 5\%$	$96.63 \pm 0.7\%$	$89.83 \pm 3\%$

literature works. The first compared work is the work [15] that used CNR classifier and energy features only. The second existing work in the table is also an semi-supervised method that used Random Forest classifier and statistical features apart from energy ones. The other compared methods were not online methods and only focused on freezing index features.

IV. DISCUSSION

Time-frequency domain analysis can provide an objective scheme to detect freezing of gait events while the current practices still are subjective methods such as patient's self-report questionaires (e.g., the Unified Parkinson's Disease Rating Scale-UPDRS, Freezing of Gait Questionnaire [5]) or manual specialies' video analysis of walking [6], [7].

The main difficulty of deploying objective wearable sensors for FoG detection lies on the current moderate accuracy performance in a subject-independent settings. Because of the large variability in motor performance (e.g., due to different walking styles), it is hard to accurately detect FoG in subject-dependent settings. Using anomaly score to dynamically update threshold levels during FoG detection process, our proposed method enables a high accuracy performance achievement in the subject-independent settings. Other challenges for future works of FoG detection include a better understanding of FoG pathogenesis [25] as well as the cerebral basis of FOG and real-life large scale evaluation of out-of-the-lab FoG detector device deployment.

V. CONCLUSION

In this work, a subject-independent FoG detection scheme is proposed. The main points of our system include utilising spectral coherence analysis to define anomaly scores for FoG

TABLE III

FOG DETECTION PERFORMANCE AGAINST EXISTING METHODS [15] a [17] b [18][19] c d) ACROSS CONFIGURATIONS e AND DATASETS.

PERFORMANCE IN %.

	Settings			Performance (%)	
Method	Input	Win	Tol	Sensitivity	Specificity
CNR [15]	Single	4s	2 <i>s</i>	73.1 ^a	81.6 ^a
	channel,				
	FI0y, Psum0y				
Learning	Single	4s	1s	66.25 ^b	95.38 ^b
[17]	channel,				
	Mean0y,				
	Std0y, FI0y,				
	Energy0y				
Global	Multi-	7.5s	-	84.3 ^c	78.4 ^c
[18]	channel,				
	$FI012y^d$,				
CL L L	$\overline{FI} = 3$	2		75.0.6	76.0 ^c
Global	Single	2s	-	75.0 ^c	76.0
[19]	channel, FI2x, $\overline{FI} = 1.47$				
Anomaly	FI = 1.47 Single	6s	0.4s	89.2	95.6
detection	channel,	US	0.48	09.2	93.0
based	FIOy, Std,				
(This	Spectral				
work)	coherence				

- ^a as reported in [15] using CNR classifier and LOOCV.
- ^b as reported in [17] using Random Forest classifier and LOOCV.
- ^c for event-based calculation while others were for timing-based.
- ^d the majority vote of seven sensors [18].
- ^e Input: features, sensors, and axes. 'Tol': tolerance. 'Win': window size.

events. *Normal* events are learned by using energy and freezing index features. Such a subject-independent FoG detection method is a key enabler for a more effective health care system for aged people especially for patients with advanced Parkinson's disease. We compared our FoG detector with expert annotations made by physiotherapists and other existing automated methods. We achieved an overall mean \pm SD sensitivity (specificity) of $89.2\pm0.3\%$ ($94\pm0.3\%$). For the knee, ankle, and hip sensors, the accuracies were $92\pm3\%$, $90.1\pm2\%$, and $89.8\pm3\%$, respectively. To our best knowledge, this is the best performance for subject-independent approaches.

VI. ACKNOWLEDGEMENTS

This study was supported by Endeavour Postgraduate Scholarship (Prime Ministers Australia Postgraduate Scholarship to Thuy Pham). We would like to thank Prof. Steven Moore and Prof. Simon J Lewis for their valuable advising in our larger project on Freezing of gait studies.

REFERENCES

- [1] BR Bloem, JM Hausdorff, JE Visser, and N Giladi, "Falls and freezing of gait in Parkinson's disease: a review of two interconnected, episodic phenomena," *Mov Disord*, vol. 19, no. 8, pp. 871–884, Aug 2004.
- [2] MD Latt, SR Lord, JG Morris, and VS Fung, "Clinical and physiological assessments for elucidating falls risk in Parkinson's disease," *Mov Disord*, vol. 24, no. 9, pp. 1280–1289, Jul 2009.

- [3] SS Paul, CG Canning, C Sherrington, SR Lord, JC Close, and VS Fung, "Three simple clinical tests to accurately predict falls in people with Parkinson's disease," *Mov Disord*, vol. 28, no. 5, pp. 655–662, May 2013.
- [4] J Reimer, M Grabowski, O Lindvall, and P Hagell, "Use and interpretation of on-off diaries in Parkinson's disease," J Neurol Neurosurg Psychiatr, vol. 75, no. 3, pp. 396–400, Mar 2004.
- [5] Nir Giladi, Joseph Tal, Tali Azulay, Oliver Rascol, David J Brooks, Eldad Melamed, Wolfgang Oertel, Werner H Poewe, Fabrizio Stocchi, and Eduardo Tolosa, "Validation of the freezing of gait questionnaire in patients with Parkinson's disease," *Movement Disorders*, vol. 24, no. 5, pp. 655–661, 2009.
- [6] Anke H Snijders, Vivian Weerdesteyn, Yolien J Hagen, Jacques Duysens, Nir Giladi, and Bastiaan R Bloem, "Obstacle avoidance to elicit freezing of gait during treadmill walking," *Movement Disorders*, vol. 25, no. 1, pp. 57–63, 2010.
- [7] C Moreau, L Defebvre, S Bleuse, JL Blatt, A Duhamel, BR Bloem, A Destée, and P Krystkowiak, "Externally provoked freezing of gait in open runways in advanced parkinsons disease results from motor and mental collapse," *Journal of neural transmission*, vol. 115, no. 10, pp. 1431–1436, 2008.
- [8] Tiffany R Morris, Catherine Cho, Valentina Dilda, James M Shine, Sharon L Naismith, Simon JG Lewis, and Steven T Moore, "A comparison of clinical and objective measures of freezing of gait in Parkinson's disease," *Parkinsonism & related disorders*, vol. 18, no. 5, pp. 572–577, 2012.
- [9] ST Moore, HG MacDougall, and WG Ondo, "Ambulatory monitoring of freezing of gait in Parkinson's disease," *J Neurosci Methods*, vol. 167, no. 2, pp. 340–348, Jan 2008.
- [10] E Gazit, H Bernad-Elazari, ST Moore, C Cho, K Kubota, L Vincent, S Cohen, L Reitblat, N Fixler, A Mirelman, et al., "Assessment of Parkinsonian motor symptoms using a continuously worn smartwatch: Preliminary experience," in MOVEMENT DISORDERS, 2015, vol. 30, pp. S272–S272.
- [11] JD Schaafsma, Y Balash, T Gurevich, AL Bartels, JM Hausdorff, and N Giladi, "Characterization of freezing of gait subtypes and the response of each to levodopa in Parkinson's disease," *Eur J Neurol*, vol. 10, no. 4, pp. 391–398, Jul 2003.
- [12] JH Han, WJ Lee, TB Ahn, BS Jeon, and Kwang-Suk Park, "Gait analysis for freezing detection in patients with movement disorder using three dimensional acceleration system," in *Engineering in Medicine and Biology Society, Proceedings of the 25th Annual International Conference* of the IEEE, 2003, vol. 2, pp. 1863–1865 Vol2.
- [13] A Delval, AH Snijders, V Weerdesteyn, JE Duysens, L Defebvre, N Giladi, and BR Bloem, "Objective detection of subtle freezing of gait episodes in Parkinson's disease," Mov Disord, vol. 25, no. 11, pp. 1684–1693, Aug 2010.
- [14] BT Cole, SH Roy, and SH Nawab, "Detecting freezing-of-gait during unscripted and unconstrained activity," in *Engineering in Medicine and Biology Society*, EMBC, Annual International Conference of the IEEE, 2011, pp. 5649–5652.
- [15] M Bachlin, M Plotnik, D Roggen, I Maidan, JM Hausdorff, N Giladi, and G Troster, "Wearable assistant for Parkinson's disease patients with the freezing of gait symptom," *Information Technology in Biomedicine*, *IEEE Transactions on*, vol. 14, no. 2, pp. 436–446, 2010.
- [16] E Jovanov, E Wang, L Verhagen, M Fredrickson, and R Fratangelo, "deFOG a real time system for detection and unfreezing of gait of Parkinson's patients," in *Engineering in Medicine and Biology Society*, 2009 EMBC 2009 Annual International Conference of the IEEE, 2009, pp. 5151–5154.
- [17] S Mazilu, M Hardegger, Z Zhu, D Roggen, G Troster, M Plotnik, and JM Hausdorff, "Online detection of freezing of gait with smartphones and machine learning techniques," in *Pervasive Computing Technologies* for Healthcare (PervasiveHealth), 6th International Conference on, 2012, pp. 123–130.
- [18] Steven T Moore, Don A Yungher, Tiffany R Morris, Valentina Dilda, Hamish G MacDougall, James M Shine, Sharon L Naismith, and Simon JG Lewis, "Autonomous identification of freezing of gait in Parkinson's disease from lower-body segmental accelerometry," *Journal* of neuroengineering and rehabilitation, vol. 10, no. 1, pp. 1, 2013.
- [19] Heidemarie Zach, Arno M Janssen, Anke H Snijders, Arnaud Delval, Murielle U Ferraye, Eduard Auff, Vivian Weerdesteyn, Bastiaan R Bloem, and Jorik Nonnekes, "Identifying freezing of gait in Parkinson's disease

- during freezing provoking tasks using waist-mounted accelerometry," *Parkinsonism & related disorders*, vol. 21, no. 11, pp. 1362–1366, 2015.
- [20] David Bannach, Kai Kunze, Paul Lukowicz, and Oliver Amft, "Distributed modular toolbox for multi-modal context recognition," in *Architecture of Computing Systems ARCS 2006*, Werner Grass, Bernhard Sick, and Klaus Waldschmidt, Eds., vol. 3894 of *Lecture Notes in Computer Science*, pp. 99–113. Springer Berlin Heidelberg, 2006.
- [21] Thuy T. Pham, Steven Moore, Simon Lewis, Andrew J. Fuglevand, Diep N. Nguyen, Eryk Dutkiewicz, Alistair L. McEwan, and Philip H. W. Leong, "Freezing of gait detection in Parkinson's disease: Feature review and subject-independent detection using anomaly scores," *submitted to IEEE TBME*, 2016.
- [22] RE Challis and RI Kitney, "Biomedical signal processing (part 3 of 4):the power spectrum and coherence function," *Medical and Biological Engineering and Computing*, vol. 28, no. 6, pp. 509–524, 1990.
- [23] C. J. Van Rijsbergen, Information Retrieval, Butterworth-Heinemann, Newton, MA, USA, 2nd edition, 1979.
- [24] Tiffany R Morris, Catherine Cho, Valentina Dilda, James M Shine, Sharon L Naismith, Simon JG Lewis, and Steven T Moore, "A comparison of clinical and objective measures of freezing of gait in parkinson's disease," *Parkinsonism & related disorders*, vol. 18, no. 5, pp. 572–577, 2012
- [25] John G Nutt, Bastiaan R Bloem, Nir Giladi, Mark Hallett, Fay B Horak, and Alice Nieuwboer, "Freezing of gait: moving forward on a mysterious clinical phenomenon," *The Lancet Neurology*, vol. 10, no. 8, pp. 734 – 744, 2011.
- [26] Elke Heremans, Alice Nieuwboer, and Sarah Vercruysse, "Freezing of gait in Parkinson's disease: where are we now?," Current neurology and neuroscience reports, vol. 13, no. 6, pp. 1–9, 2013.