

Detecting freezing of gait with a tri-axial accelerometer in Parkinson's disease patients

Claas Ahlrichs¹ · Albert Samà³ · Michael Lawo² · Joan Cabestany³ · Daniel Rodríguez-Martín³ · Carlos Pérez-López³ · Dean Sweeney⁴ · Leo R. Quinlan⁴ · Gearòid Ó Laighin⁴ · Timothy Counihan⁵ · Patrick Browne⁵ · Lewy Hadas⁶ · Gabriel Vainstein⁶ · Alberto Costa^{7,9} · Roberta Annicchiarico⁷ · Sheila Alcaine⁸ · Berta Mestre⁸ · Paola Quispe⁸ · Àngels Bayes⁸ · Alejandro Rodríguez-Molinero⁴

Received: 28 November 2014 / Accepted: 12 September 2015 / Published online: 1 October 2015
© International Federation for Medical and Biological Engineering 2015

Abstract Freezing of gait (FOG) is a common motor symptom of Parkinson's disease (PD), which presents itself as an inability to initiate or continue gait. This paper presents a method to monitor FOG episodes based only on acceleration measurements obtained from a waist-worn device. Three approximations of this method are tested. Initially, FOG is directly detected by a support vector machine (SVM). Then, classifier's outputs are aggregated over time to determine a confidence value, which is used for the final classification of freezing (i.e., second and third approach). All variations are trained with signals of 15 patients and evaluated with signals from another 5 patients. Using a linear SVM kernel, the third approach provides 98.7 % accuracy and a geometric mean of 96.1 %. Moreover, it is investigated whether frequency features are enough to reliably

detect FOG. Results show that these features allow the method to detect FOG with accuracies above 90 % and that frequency features enable a reliable monitoring of FOG by using simply a waist sensor.

Keywords Parkinson's disease · Freezing of Gait · Machine learning · Support vector machines

1 Introduction

PD is a chronic, progressive, neurodegenerative disorder [2, 10, 14, 16, 30], with which a great number of motor and non-motor symptoms have been associated. The disorder was first described by James Parkinson in 1817 [26]. It affects the movement, and it is typically characterized by a loss of (motor) function, increased slowness and rigidity. Presently, the cause and origin of PD remain unknown [9, 14, 17, 30] and it cannot be cured. Consequently, treatments aim at reducing severity and frequency of motor complications. The disease is generally associated with elderly people and is rarely diagnosed before the age of 40. It is estimated that the mean age of onset is about 65 years [30].

PD is a great burden as it considerably decreases the quality of life, due to a gradual loss of function and decreasing ability to take care of oneself. The World Health Organization (WHO) considers the burden of PD to be on the same disability level as an amputated arm, drug dependency, congestive heart failure, deafness and tuberculosis [20]. The cardinal symptoms are bradykinesia, rigidity, tremor and postural instability [1, 10, 14, 16, 17, 30, 32]. However, a number of non-motor-related symptoms (e.g., sleep disturbances, depression, psychosis, autonomic and gastrointestinal dysfunction as well as dementia) may occur as well [10, 14, 16, 18, 30].

✉ Claas Ahlrichs
claasahl@tzi.de

¹ neusta mobile solutions GmbH (NMS), Konsul-Smidt-Str. 24, 28217 Bremen, Germany

² Institute for Artificial Intelligence (AGKI), University of Bremen, Bremen, Germany

³ Technical Research Centre for Dependency Care and Autonomous Living (CETpD), Universitat Politècnica de Catalunya, Vilanova i la Geltrú, Spain

⁴ Electrical & Electronic Engineering Department, NUI Galway, Galway, Ireland

⁵ School of Medicine, NUI Galway, Galway, Ireland

⁶ Maccabi Healthcare Services, Tel Aviv, Israel

⁷ IRCCS Fondazione Santa Lucia, Rome, Italy

⁸ Unidad de Parkinson y trastornos del movimiento (UParkinson), Barcelona, Spain

⁹ Niccolò Cusano University of Rome, Rome, Italy

One of the motor symptoms is called FOG (also known as freezing or motor blocks). It is a form of akinesia, which presents itself as an inability to initiate or continue gait [12, 16, 24, 30, 31]. Motor blocks are a common symptom, experienced by people with Parkinson's (although it does not occur uniformly) and can affect various extremities (e.g., arms and legs) as well as the face [16]. Freezing greatly impairs the quality of life of those affected and is one of the most disabling symptoms. It is usually attributed to medium and advanced stages of PD, and it is a common cause of falls [6, 16, 30]. A single freezing episode typically lasts for several seconds. In severe cases, episodes can be apparent for as long as several minutes.

Continuous monitoring of FOG events can give neurologists information which is otherwise difficult to obtain. Clinical assessment of FOG at the doctor's office is considered to be problematic since symptoms are commonly not evident in this clinical environment [25]. Thus, a wearable device capable of ambulatory monitoring FOG could benefit patients in two ways. First, it could provide clinicians with complementary information of the disease that can be used to improve treatment [31]. Second, since patients are capable of improving gait based on specific stimulations provided as haptic, visual or auditory cues [19], real-time FOG detection would allow patients to avoid some episodes and, consequently, avoid falls, such as the system presented in [3]. Consequently, many studies have attempted to develop wearable devices for the detection of FOG.

The literature indicates that these studies typically make use of multiple sensors (i.e., accelerometers, gyroscopes) at various body locations and they usually employ some form of supervised learning approach [e.g., SVM or neural network (NN)]. Djurić-Jovičić et al. [11] achieved an error rate of up to 16 % classifying "normal" (i.e., standing and regular steps) and pathological (i.e., festination, akinesia, shuffling and small steps) walking patterns of PD patients based on a NN (using multiple inertial measurement units). The approach by Cole et al. [7] yielded to 82.9 % sensitivity and 97.3 % specificity in detecting FOG (using acceleration and electromyograph (EMG) sensors) with a multistaged algorithm that utilized a simple linear classifier and a dynamic neural network (DNN). Cole et al. employed data collected during unscripted and unconstrained activities in an apartment-like setting. However, there is no information on the activities that patients performed. The other works that can be found in the literature employed signals gathered during scripted activities, such as Niazmand et al. [23], who used an accelerometer-based smart garment [22] to extract gait-related features. They achieved 88.3 % sensitivity and 85.3 % specificity (using multiple accelerometers). The approach by Bächlin et al. [5] yielded to 73.1 % sensitivity and 81.6 % specificity for detecting

FOG events in real time (using multiple accelerometers and gyroscopes).

In this work, the authors present a multistaged approach based on an SVM and a single tri-axial acceleration sensor. Using a linear SVM kernel and the full feature set (see Table 5), an accuracy of 98.7 % and a geometric mean of 96.1 % have been achieved. The overall dataset includes signals from 20 PD patients, among who 8 of them presented FOG episodes. The overall dataset is divided into the training and the testing dataset, the latter including signals from two patients with FOG and three patients without FOG. These results have been obtained with a patient-independent methodology. Furthermore, the algorithm can be configured toward a higher sensitivity or a higher specificity. The employed movement signals were collected for the REMPARK project (Personal Health Device for the Remote and Autonomous Management of Parkinson's Disease) database [29]. This project aims to develop a closed-loop system with the purpose of monitoring PD motor and non-motor symptoms and responding to these symptoms in real time using a series of actuators. Data collection of REMPARK's database inertial signals took place in 4 different countries (Spain, Italy, Israel and Ireland). Signals were obtained at patient's home during the execution of roughly scripted tasks (e.g., walk around the apartment and show all rooms) that enabled patients to perform partially unconstrained activities.

2 Methods

Firstly, the data acquisition is described. Then, the methodology and model selection of the proposed approach are outlined.

2.1 Data acquisition and labeling

All participants (aged between 50 and 75 years) had a clinical diagnosis of idiopathic Parkinson's disease according to the UK PD Society Brain Bank [15]. Clinical fluctuations were present in all patients as well as Hoehn and Yahr stage [13] above two (moderate–severe phase of PD). Furthermore, all patients gave their signed informed consent before their participation. The experimental protocol was approved by the corresponding local ethics review committee. For this paper, signals from 20 PD patients were used, among which 8 patients presented FOG episodes and 12 did not present the symptom. The recordings are identical to those employed by Rodríguez-Martín et al. [27, 28].

As part of the experiments, participants were recorded with an HD quality camera while wearing a set of sensors (i.e., accelerometer, gyroscope and magnetometer) as they performed a set of scripted activities. However,

these activities are of a rather general nature (e.g., walking around the apartment and showing it to the researchers or carrying a full glass of water from the kitchen to another room) and they are much more variable in comparison with other typically scripted activities like hand-to-nose or similar gestures which are performed in a seated position. The recordings also include non-scripted activities that took place during the recordings, such as answering the phone or similar unexpected situations that in some cases lead to FOG episodes due to turnings or going through narrow places. The experiments took place at the participant's apartment and started in the morning. During the course of the day, two recording sessions took place: one in the "OFF" motor state and one in the "ON" motor state. For the first session, participants were asked to skip their morning dose of medication, thus recordings started while the participant was in a clinically defined OFF state [29]. After finishing the first round, participants took their normal medication and the second recording session was started once the participant had reached a clinically defined ON state. During both recording sessions, participants performed a series of short controlled activities. The activities performed by patients during their OFF state were an indoors walking test, a FOG provocation test and a gait test. During the ON state, a dyskinesia test, a dual-task test and a set of activities of daily livings (ADLs) also were performed. ADLs included brushing teeth, shaking a deodorant, erasing with an eraser, writing with a pencil, typing on a computer keyboard, cleaning a window or furniture and drying a wet glass [29].

Experienced clinicians labeled the videos based on the activities that patients performed and the symptoms shown during the video. The clinicians who performed the labeling were also physically present during the recording sessions. Each of the clinical sides (one for each country) had two clinicians with several years of experiences with PD patients (i.e., ≥ 5 years). Prior to the recording sessions, all clinicians received a training session on setting baselines for labeling of symptoms (including FOG). The group that performed the labeling is disjoint from the group that performed the analysis.

Video and inertial signals were synchronized based on the procedure described in [29]. FOG labels provided by clinicians have been treated with an automatic labeling procedure in order to consider specific peculiarities of FOG. In this sense, it is important to note that clinicians in charge of the FOG labeling had been with the patients during the experimental protocol and, in consequence, these clinicians knew whether a patient presented FOG episodes or not before examining the video recordings. However, in the labeling process of patients with FOG, clinicians may miss some FOG episodes given that, in some moments, video camera was not close enough to notice mild episodes.

Accordingly, recordings of freezers were cut to the point where only FOG labels remained. This reduced the overall amount of data for recordings of freezers but ensured that no freezing episodes (which might not have been properly labeled) were used. On the other hand, those patients without any freezing episodes were relabeled in such a way that all available data were used. Consequently, sensitivity was determined by using data from patients with freezing episodes, while specificity was determined by using data from non-freezing patients. Overall, this procedure allowed using larger portions of the recordings.

As far as the actual labeling is concerned, the presence of any type of freezing (e.g., start, turn, end) was considered to be an episode of FOG. The detection of individual types of freezing requires additional contextual information which is not contained within the database (DB). Furthermore, such a fine granularity might not provide an additional value (e.g., to a PD monitoring system). The fact that a freezing episode is happening is more relevant than the actual type of episode (e.g., for rhythmic cueing purposes). Consequently, freezing episodes are detected rather than individual types of freezing.

2.2 Methodology

The general methodology is such that acceleration signals from a waist-mounted sensor are split into equally sized windows (i.e., a sliding window is applied to the time series). Features are extracted from those windows and fed to an SVM for training or classification. The classification output of n consecutive windows s_1, \dots, s_n is then aggregated over time t (in seconds) to achieve higher accuracies. However, the volatile nature of FOG must be considered during the development of an algorithm for detecting such episodes. In contrast to resting tremor (or dyskinesia for that matter), episodes of FOG do not last for prolonged periods of time which may emphasize the importance of the chosen window size ws . In any case, the contents of the database are split into two datasets (i.e., training and testing) that are used for training an SVM as well as optimizing additional parameters and testing, respectively. Datasets stay the same for all approaches (details are listed in Table 1). The individual datasets hold 15 and 5 patients for the training and testing dataset, respectively.

Two feature sets are evaluated: a reduced feature set with only the fast Fourier transform (FFT) and a full feature set with various additional features (see Table 2). The effect of adding these additional features is quantified in Sect. 3. These features are comprised of the freezing index [4] as well as some frequency-related features for differing frequency ranges [21].

At first, varying window sizes ws were evaluated such that freezing of gait detection was optimized. The

Table 1 The number of windows (before aggregation) in each dataset that are used for signifying FOG

	Training	Test
Number of freezing windows	93	45
Number of non-freezing windows	3883	2312
Recordings with freezing	6	2
Recordings without freezing	9	3
Overall number of patients	15	5
Average duration of recordings with freezing (min)	18.38	24.32
Average duration of recordings without freezing (min)	9.59	19.48

Table 2 The full set of features used for FOG detection. In contrast, the reduced feature set is only comprised of a fast Fourier transform (i.e., index 1)

Indexes	Feature
1	FFT (raw, no filtering)
2	Mean and standard deviation of amplitude (band: 0.5–3.0 Hz)
3	Entropy of signal in time domain (band: 0.5–3.0 Hz)
4	Peak amplitude and its frequency (band: 0.5–3.0 Hz)
5	Mean and standard deviation of amplitude (band: 3.0–8.0 Hz)
6	Entropy of signal in time domain (band: 3.0–8.0 Hz)
7	Peak amplitude and its frequency (band: 3.0–8.0 Hz)
8	Freezing index

comparison of different window sizes was done on an episode level (rather than a window level). An episode of FOG was detected when at least one window within an actual FOG episode was classified as such. As far as non-freezing episodes were concerned, an aggregation of windows over a period of time that corresponds to the average length of a FOG episode plus twice the standard deviation is performed. The acceleration data are resampled to 40Hz and split into unisized chunks of data s_1, \dots, s_m with a certain length ws that overlapped to 50 %. These windows are then used to extract features which in turn were fed to an SVM for training and classification. This resembles the first and naive approach, where $freezing_j^1$ represents the j th window in the series s_1, \dots, s_m and whether FOG is present in that window.

$$freezing_j^1 = \begin{cases} 0 & \text{no freezing if } f_{SVM} \leq 0 \\ 1 & \text{freezing if } f_{SVM} > 0 \end{cases} \quad (1)$$

where $f_{svm} = \sum_{i=1}^l y_i \alpha_i K(\mathbf{x}_i, \mathbf{f}) + b$, $\mathbf{x}_1, \dots, \mathbf{x}_l$ are the support vectors (SVs), y_i, α_i are the corresponding label and Lagrange multiplier of each SV and b is the bias [8]. The number in the superscript (here: 1) indicates the

variation. The second and third variation will use 2 and 3, respectively.

The second variation aggregates the SVMs’ outputs over a time period t and calculates the degree of confidence c_j . If the confidence value exceeds a threshold th , then the aggregated time frame t is considered to be an episode of FOG, otherwise not. Here, $freezing_j^2$ covers a time frame t (starting at the j th window and covering n windows) and determines whether FOG is apparent in that time frame.

$$t = \frac{ws(n + 1)}{40 * 2} \quad (2)$$

$$c_j = \sum_{i=j}^{j+n-1} \frac{freezing_i^1}{n} \quad (3)$$

$$freezing_j^2 = \begin{cases} 0 & \text{no freezing if } c_j < th \\ 1 & \text{freezing if } c_j \geq th \end{cases} \quad (4)$$

where $c_j, th \in [0, 1]; n, j \in \mathbb{N}; n, j > 0; t \in \mathbb{R}^+$. n , as previously described, corresponds to the number of windows that are aggregated in order to span the time period of t seconds.

The third variation introduces a second threshold. The lower threshold th_l and upper threshold th_u can be used to tune sensitivity and specificity separately. The lower threshold th_l sets the maximum confidence value for “no freezing” periods, and the upper threshold th_u sets the minimum confidence value for freezing episodes. By not requiring that these thresholds need to be equal (which would essentially be variation two), the final output of the algorithm may indicate the presence of freezing as well as “undefined.” This is the case when the confidence value is between the two thresholds. Consequently, some aggregated windows may be ignored and data usage is lowered.

$$freezing_j^3 = \begin{cases} 0 & \text{no freezing if } c_j < th_l \\ 1 & \text{freezing if } c_j \geq th_u \\ -1 & \text{undefined if } th_l \leq c_j < th_u \end{cases} \quad (5)$$

where $c_j, th_l, th_u \in [0, 1]; th_l \leq th_u; j \in \mathbb{N}; j > 0$

2.3 Model Selection

The individual SVM models are trained with the features that were extracted from the training dataset. For the second and third variation, the individual parameters t, th, th_l and th_u are also optimized on the training dataset. The final results are obtained from the testing dataset.

The window size ws is determined before any of these parameters are evaluated. For each of the proposed window sizes ws (see below), the naive algorithm is applied to the training dataset. The window size that yields to the best combination of accuracy and geometric mean is chosen.

During training, varying settings for kernel, weighting, cost and gamma were considered. The weighting parameters were used to balance both classes “FOG” and “non-freezing.” The cost and gamma parameters were systematically evaluated (i.e., $10^q, q \in \{-3, -2, \dots, 2, 3\}$) depending on the chosen kernel (i.e., radial basis function (RBF) kernel or linear kernel). Additionally, a tenfold cross-validation is performed on the training dataset. However, instead of averaging the accuracy of the training set, the geometric mean of sensitivity and specificity is used (i.e., $\sqrt{\text{sensitivity} * \text{specificity}}$) to identify those parameters combinations with high sensitivity and specificity. The maximum geometric mean is used to select the optimal parameters and obtain the final SVM model to be used with the test dataset. The geometric mean was chosen as it does treat both classes equally as opposed to accuracy which implicitly weights the classes. The weighting of latter measure can be a problem if the classes have (very) different priors.

The following discrete values have been evaluated: $ws \in 2^{\{5,6,7,8\}}$; $t \in \{10, 15, 20, 25, 30, 45, 60\}$; $th, th_l \in \{0, 0.05, 0.1, \dots, 0.95, 1.0\}$; $th_u \in (th_u \geq th_l | th_u \in \{0, 0.05, 0.1, \dots, 0.95, 1\})$. The appropriate values and parameters were evaluated for each of the four conditions (two kernels and two feature sets).

3 Results

The average length of a FOG episode in our dataset was 3.48 ± 3.29 s (total: 209 freezing events). Figure 1 shows several measures for varying window sizes (i.e., sensitivity, specificity, geometric mean and accuracy). The best values for those measures were achieved with a window size of 128 samples (i.e., 2^7 samples). Accuracy and geometric mean were closest at this level. Consequently, this window size was utilized in all further analyses.

Table 3 presents the results obtained for the first variation. It was observed that, on the training dataset, both full and reduced feature sets yield a similar geometric mean regardless of the employed SVM kernel. This, however, diverges on the test dataset. The RBF kernel seems to benefit from the reduced feature set, while the linear kernel favors the full feature set. Acceptable levels of specificity are consistently achieved on the test dataset, while sensitivity was reduced by false negatives (FNs). The latter may be counteracted when windows are aggregated. Nonetheless, accuracies above 90 % were consistently reached.

The impact of window aggregation t and threshold th is highlighted in Fig. 2 for all four conditions. The subfigures indicate that a threshold close to 50 % works best in all cases. Furthermore, it is observed that the geometric mean increases with the aggregation level.

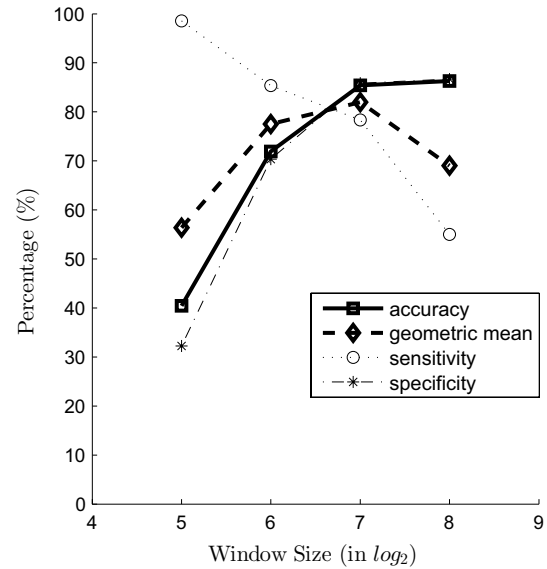


Fig. 1 Results of an evaluation for varying window sizes with respect to freezing episodes. For each window size, an SVM has been trained on the training dataset and evaluated on the test dataset

Table 3 Results in signifying FOG with the naive approach (i.e., variation 1). Various measures are listed for both datasets

Kernel	RBF	Linear	RBF	Linear
Features	Freq.	Freq.	All	All
Sensitivity (train)	0.946	0.903	0.946	0.925
Specificity (train)	0.860	0.903	0.901	0.932
Data usage (train)	1.000	1.000	1.000	1.000
Geometric mean (train)	0.902	0.903	0.924	0.928
Accuracy (train)	0.862	0.903	0.902	0.932
True positives	37	30	32	37
False positives	167	117	124	101
True negatives	2145	2195	2188	2211
False negatives	8	15	13	8
Sensitivity (test)	0.822	0.667	0.711	0.822
Specificity (test)	0.928	0.949	0.946	0.956
Data usage (test)	1.000	1.000	1.000	1.000
Geometric mean (test)	0.873	0.796	0.820	0.887
Accuracy (test)	0.926	0.944	0.942	0.954

Bold values are frequently referred to in the text

Numerical results for the second variation are shown in Table 4. All conditions yielded to a threshold close to the intuitive border of 50 %, which is consistent with the observations in Fig. 2. Moreover, the aggregation period t is the same across all four conditions. Having optimized parameters t and th on the training dataset, the results on the testing dataset show an increase by 9.4 % (on average). All conditions achieve a high specificity of 98 % or greater,

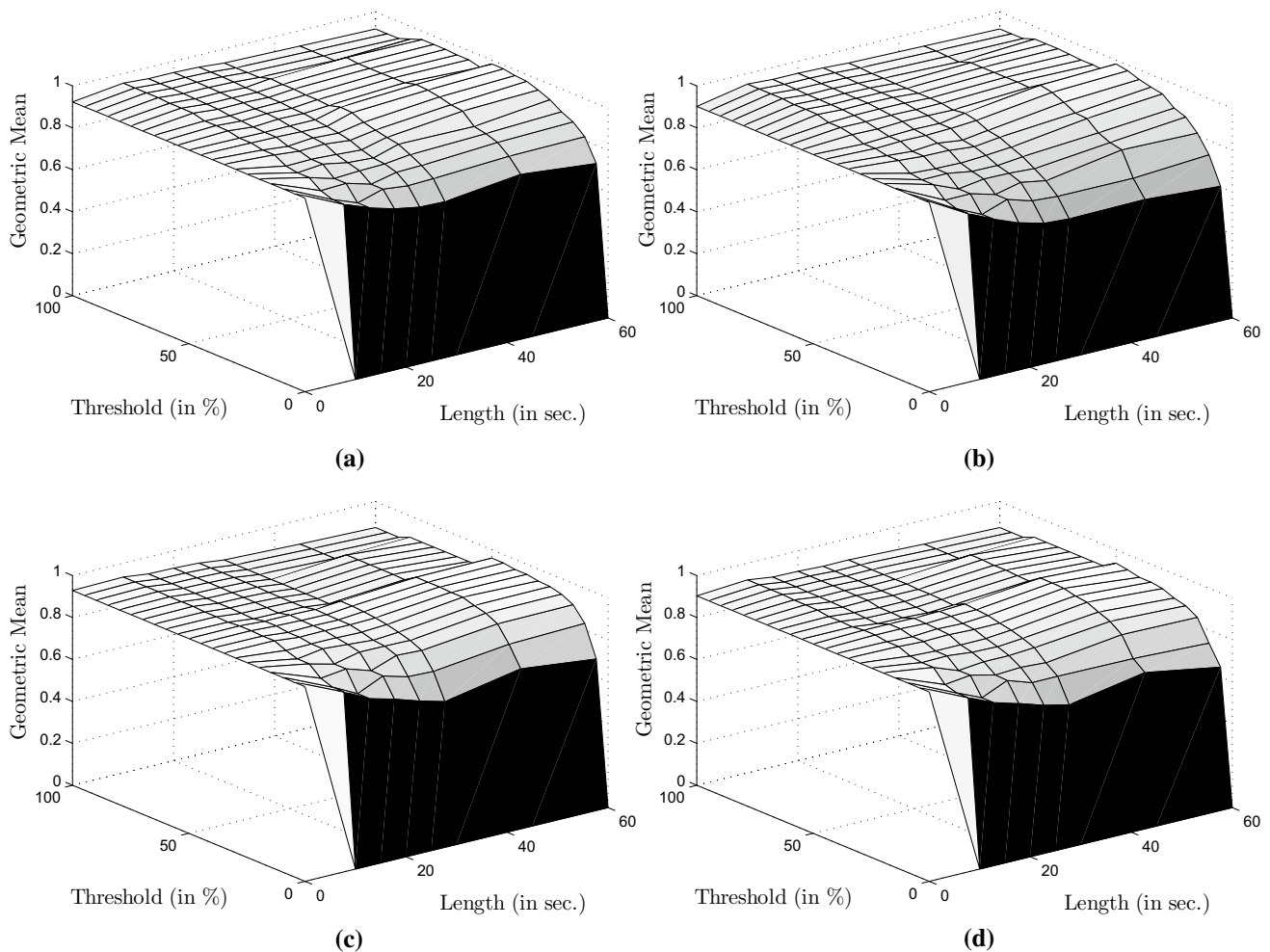


Fig. 2 Effect of window aggregation t and threshold th on geometric mean. The results are shown for all four conditions. **a** RBF with full feature set. **b** RBF with reduced feature set. **c** Linear kernel with full feature set. **d** Linear kernel with reduced feature set

and furthermore, most conditions also reached a sensitivity of 90 % or above for an aggregation period of 60 seconds.

Results in Table 5 are those of the third variation. Most conditions still favor an aggregation level of 60 s. The lower and upper thresholds (i.e., th_l and th_u) were consistently found to enclose the previously found thresholds th in the second approach (see Table 4). Allowing for two thresholds increased sensitivity and specificity values on the test dataset for the linear kernel. However, the RBF kernel did not benefit from this approach in terms of geometric mean. The average change in geometric mean from variation two to three was -1.2 and 3.7 % for the RBF and linear kernel, respectively. Nonetheless, all conditions yield to a sensitivity of roughly 90 % and a specificity well above 90 %. However, this was at the cost of a slightly reduced data usage although still above 90 % for the most parts.

4 Discussion

The presented FOG detection methods result in a geometric mean of 88.7, 96.1 and 96.1 % for each of the three proposed approaches (linear kernel with full feature set). Thus, the meta-analysis used in the second and third variation is shown to enable a better recognition of FOG episodes since it improves the overall performance (geometric mean) by 8 %. Regarding the feasibility of detecting FOG uniquely by means of frequency features, it was observed that a geometric mean of 96.1 % (one-sided approach with RBF kernel) can be achieved based on them. This way, it is concluded that frequency features enable a reliable monitoring of FOG.

The results previously obtained by Niazmand et al. [23], Cole et al. [7] and Bächlin et al. [5] were consistently lower than the obtained by the presented approach in its third variation, which has yielded to an average

Table 4 Results in signifying FOG with the one-sided approach (i.e., variation 2)

Kernel	RBF	Linear	RBF	Linear
Features	Freq.	Freq.	All	All
t (in seconds)	60	60	60	60
th (in %)	0.400	0.500	0.400	0.450
Sensitivity (train)	1.000	0.885	1.000	0.923
Specificity (train)	0.911	0.991	0.946	1.000
Data usage (train)	1.000	1.000	1.000	1.000
Geometric mean (train)	0.954	0.936	0.973	0.961
Accuracy (train)	0.928	0.971	0.957	0.986
True positives	12	10	12	12
False positives	0	1	0	0
True negatives	66	65	66	66
False negatives	1	3	1	1
Sensitivity (test)	0.923	0.769	0.923	0.923
Specificity (test)	1.000	0.985	1.000	1.000
Data usage (test)	1.000	1.000	1.000	1.000
Geometric mean (test)	0.961	0.870	0.961	0.961
Accuracy (test)	0.987	0.949	0.987	0.987

Bold values are frequently referred to in the text

Table 5 Results in detecting FOG with the two-sided approach (i.e., variation 3)

Kernel	RBF	Linear	RBF	Linear
Features	Freq.	Freq.	All	All
t (in seconds)	60	45	60	60
th_l (in %)	0.150	0.250	0.300	0.350
th_u (in %)	0.900	0.800	0.800	0.500
Sensitivity (train)	1.000	0.923	1.000	0.923
Specificity (train)	1.000	1.000	1.000	1.000
Data usage (train)	0.696	0.891	0.906	0.986
Geometric mean (train)	1.000	0.961	1.000	0.961
Accuracy (train)	1.000	0.987	1.000	0.985
True positives	9	8	9	12
False positives	0	0	0	0
True negatives	55	82	65	65
False negatives	1	1	1	1
Sensitivity (test)	0.900	0.889	0.900	0.923
Specificity (test)	1.000	1.000	1.000	1.000
Data usage (test)	0.823	0.919	0.949	0.987
Geometric mean (test)	0.949	0.943	0.949	0.961
Accuracy (test)	0.985	0.989	0.987	0.987

Bold values are frequently referred to in the text

sensitivity and specificity above 94 %. Niazmand et al. [23] achieved a sensitivity of 88.3 % and a specificity of 85.3 %. Compared to the results from Bächlin et al. [5], both sensitivity (73.1 %) and specificity (81.6 %) were much lower than the ones reported in this paper. However, Cole et al. [7] achieved a similar level of specificity

(97.3 %), but with quite lower sensitivity (82.9 %), although their signals were collected during completely unconstrained activities.

A limitation of the presented work relies in its applicability to real-time detection in order to provide rhythmic cues. In this case, a short lag between the appearance of a FOG episode and its detection is desired. The meta-analysis in the second and third variations may add a delay that could reach the aggregation time (60 s), which would not allow to be used for this particular purpose, although remaining useful in monitoring tasks. The first variation, however, could be employed, since the lag provided is roughly 3.2 s (128 samples at 40 Hz). A second limitation with respect to the work of Cole et al. [7] is that the signals employed in this work were not acquired in a completely unconstrained setting. In consequence, performances obtained may decrease with new activities in the daily life of patients. However, the REMPARK database also includes signals recorded under these specific conditions. In the future, these signals will be analyzed to determine the applicability and performance of the presented methodology on these specific conditions.

Besides the performance comparative, the proposed approach has with respect to [5] the advantage of being patient independent, given that the same classifier can be used by any patient. Moreover, we only use a single tri-axial accelerometer at the waist, while Cole et al. [7] used three tri-axial accelerometers and surface EMG, Niazmand et al. [23] five accelerometers and Bächlin et al. [5] three accelerometers and three gyroscopes. Moreover, the presented approach offers configuration capabilities since the algorithm can be tuned toward high sensitivity and high specificity by adjusting the thresholds. Finally, the optimal window size has been determined by evaluating the performance of the algorithm at episode level, as opposed to window level used in previous works, which may have increased specificity.

5 Conclusion

This work has evaluated three approaches to detecting FOG in Parkinson’s patients based on a waist-worn accelerometer. The optimal window size was determined, and it has been analyzed whether frequency features are sufficient to reliably detect FOG.

Although the linear and RBF kernel do not benefit equally from the third approach, combining the results from both variations (i.e., second and third variation) shows promising results. While the RBF kernel achieved a geometric mean greater than 95 % and an accuracy greater than 98 % with the second approach, the linear kernel reached similar levels (close to 95 % geometric mean and 98 %

accuracy) with the third approach. However, in the latter case, the data usage is slightly penalized. The findings suggest that the full feature set is not required for satisfactory results. Instead, a linear kernel that has been trained with an FFT alone can accurately detect FOG episodes. Finally, the optimal window size has been found to be 128 samples (at 40 Hz).

In comparison with the previous approaches, although the method presented in this work has obtained higher performance metrics than those previously reported, it is noted that the conditions in which each study takes place are different and, in this sense, our study suffers of a lack of completely unconstrained activities, which may decrease the method's performance during the activities of daily living of PD patients. However, the REMPARK database also includes signals recorded under these specific conditions. In the future, these signals will be analyzed to determine the applicability and performance of the presented methodology on these specific conditions. On the other hand, the present approach has the advantage of working in a patient-independent way and only requiring a single tri-axial accelerometer.

Acknowledgments This work has been performed in the framework of the FP7 project REMPARK ICT-287677, which is funded by the European Community. The author(s) would like to acknowledge the contributions of their colleagues from REMPARK Consortium (<http://www.rempark.eu>). We also like to thank all participants without whom this publication would not have been possible.

References

- Andlin-Sobocki P, Jansson B, Wittchen HU, Olesen J (2005) Cost of disorders of the brain in Europe. *Eur J Neurol* 12:1–27. doi:10.1111/j.1468-1331.2005.01202.x
- Armstrong RA (2008) Visual signs and symptoms of Parkinson's disease. *Clin Exp Optom* 91(2):129–138. doi:10.1111/j.1444-0938.2007.00211.x
- Bächlin M, Plotnik M, Roggen D, Giladi N, Hausdorff JM, Tröster G (2010) A wearable system to assist walking of Parkinson's disease patients. *Methods Inf Med* 49(1):88–95. doi:10.3414/ME09-02-0003. <http://www.schattauer.de/en/magazine/subject-areas/journals-a-z/methods/issue/special/manuscript/12447/show.html>
- Bächlin M, Plotnik M, Roggen D, Maidan I, Hausdorff JM, Giladi N, Troster G (2010) Wearable assistant for Parkinson's disease patients with the freezing of gait symptom. *IEEE Trans Inf Technol Biomed* 14(2):436–446. doi:10.1109/TITB.2009.2036165
- Bächlin M, Roggen D, Troster G, Plotnik M, Inbar N, Meidan I, Herman T, Brozgol M, Shaviv E, Giladi N, Hausdorff JM (2009) Potentials of enhanced context awareness in wearable assistants for Parkinson's disease patients with the freezing of gait syndrome. In: 2009 International Symposium on Wearable Computers (ISWC), pp 123–13. doi:10.1109/ISWC.2009.14
- Bloem BR, Hausdorff JM, Visser JE, Giladi N (2004) Falls and freezing of gait in Parkinson's disease: a review of two interconnected, episodic phenomena. *Mov Disord* 19(8):871–884. doi:10.1002/mds.20115
- Cole BT, Roy SH, Nawab SH (2011) Detecting freezing-of-gait during unscripted and unconstrained activity. In: 2011 Annual International Conference of the IEEE Engineering in Medicine and Biology Society (EMBC), pp 5649–5652. doi:10.1109/IEMBS.2011.6091367
- Cristianini N, Shawe-Taylor J (2000) An introduction to support vector machines and other kernel-based learning methods. Cambridge University Press, New York
- Dauer W, Przedborski S (2003) Parkinson's disease: mechanisms and models. *Neuron* 39(6):889–90. doi:10.1016/S0896-6273(03)00568-3. <http://www.sciencedirect.com/science/article/pii/S0896627303005683>
- Davie CA (2008) A review of Parkinson's disease. *Br Med Bull* 86(1):109–127. doi:10.1093/bmb/ldn013. <http://bmb.oxfordjournals.org/content/86/1/109.abstract>
- Djurić-Jovičić M, Jovičić NS, Milovanović I, Radovanović S, Kresojević N, Popović MB (2010) Classification of walking patterns in Parkinson's disease patients based on inertial sensor data. In: 2010 10th Symposium on Neural Network Applications in Electrical Engineering (NEUREL), pp 3–6. doi:10.1109/NEUREL.2010.5644040
- Giladi N (2006) Freezing of gait: risk factors and clinical characteristics. *Parkinsonism Relat Disord* 12(Supplement 2):S52. doi:10.1016/j.parkreldis.2006.05.015
- Hoehn MM (1967) Parkinsonism: onset, progression, and mortality. *Neurology* 17:427–442
- Hou JGG, Lai EC (2007) Non-motor symptoms of Parkinson's disease. *Int J Gerontol* 1(2):53–64. doi:10.1016/S1873-9598(08)70024-3. <http://www.sciencedirect.com/science/article/pii/S1873959808700243>
- Hughes AJ, Daniel SE, Kilford L, Lees AJ (1992) Accuracy of clinical diagnosis of idiopathic Parkinson's disease: a clinicopathological study of 100 cases. *J Neurol Neurosurg Psychiatry* 55(3):181–184. doi:10.1136/jnnp.55.3.181. <http://jnnp.bmj.com/content/55/3/181.abstract>
- Jankovic J (2008) Parkinson's disease: clinical features and diagnosis. *J Neurol Neurosurg Psychiatry* 79(4):368–376. doi:10.1136/jnnp.2007.131045. <http://jnnp.bmj.com/content/79/4/368.abstract>
- Korczyn AD (2008) Parkinson's disease. In: E. in Chief: Kris Hegggenhougen (ed) International encyclopedia of public health, Academic Press, Oxford, pp 10–17. doi:10.1016/B978-012373960-5.00028-9. <http://www.sciencedirect.com/science/article/pii/B9780123739605000289>
- Krenz A (2010) The Pathological Role of Synphilin-1 and the Therapeutic Potential of Hsp70 in Models of Parkinson's Disease Using Viral Vectors. Ph.D. thesis, Universität Tübingen, Wilhelmstr. 32, 72074 Tübingen. <http://tobias-lib.uni-tuebingen.de/volltexte/2010/4620>
- Lim I, van Wegen E, de Goede C, Deutekom M, Nieuwboer A, Willems A, Jones D, Rochester L, Kwakkel G (2005) Effects of external rhythmical cueing on gait in patients with Parkinson's disease: a systematic review. *Clin Rehabil* 19(7):695–713. doi:10.1191/0269215505cr906oa. <http://cre.sagepub.com/content/19/7/695.abstract>
- Mathers C, Fat DM, Boerma JT (2008) WHO: the Global burden of disease : 2004 update. World Health Organization, Geneva. http://www.who.int/healthinfo/global_burden_disease/GBD_report_2004update_full.pdf
- Moore ST, MacDougall HG, Ondo WG (2008) Ambulatory monitoring of freezing of gait in Parkinson's disease. *J Neurosci Methods* 167(2):340–348. doi:10.1016/j.jneumeth.2007.08.023. <http://www.sciencedirect.com/science/article/pii/S0165027007004281>
- Niazmand K, Somlai I, Louizi S, Lueth TC (2011) Proof of the accuracy of measuring pants to evaluate the activity of

the hip and legs in everyday life. In: Lin JC, Nikita KS, Akan O, Bellavista P, Cao J, Dressler F, Ferrari D, Gerla M, Kobayashi H, Palazzo S, Sahni S, Shen XS, Stan M, Xiaohua J, ZoMaya A, Coulson G (eds) *Wireless mobile communication and healthcare, lecture notes of the institute for computer sciences, social informatics and telecommunications engineering*. Springer, Berlin Heidelberg, pp 235–244. doi:[10.1007/978-3-642-20865-2_30](https://doi.org/10.1007/978-3-642-20865-2_30)

23. Niazmand K, Tonn K, Zhao Y, Fietzek UM, Schroeteler F, Ziegler K, Ceballos-Baumann AO, Lueth TC (2011) Freezing of gait detection in Parkinson's disease using accelerometer based smart clothes. In: 2011 IEEE Biomedical Circuits and Systems Conference (BioCAS), pp 201–204. doi:[10.1109/BioCAS.2011.6107762](https://doi.org/10.1109/BioCAS.2011.6107762)
24. Nieuwboer A, Giladi N (2013) Characterizing freezing of gait in Parkinson's disease: models of an episodic phenomenon. *Mov Disord* 28(11):1509–1519. doi:[10.1002/mds.25683](https://doi.org/10.1002/mds.25683)
25. Nieuwboer A, Weerdt Wd, Dom R, Lesaffre E (1998) A frequency and correlation analysis of motor deficits in Parkinson patients. *Disabil Rehabil* 20(4):142–150. doi:[10.3109/09638289809166074](https://doi.org/10.3109/09638289809166074)
26. Parkinson J (2002) An essay on the Shaking Palsy. 1817. *J Neuropsychiatry Clin Neurosci* 14(2):223–236; discussion 222. <http://www.ncbi.nlm.nih.gov/pubmed/11983801>
27. Rodríguez-Martín D, Samà A, Pérez-López C, Cabestany J, Català A, Rodríguez-Molinero A (2014) Enhancing FoG Detection By Means of Postural Context Using a Waist Accelerometer. First International Freezing of Gait Congress (IFOG 2014)
28. Rodríguez-Martín D, Samà A, Pérez-López C, Cabestany J, Català A, Rodríguez-Molinero A (2015) Posture Transition Identification on PD Patients Through a SVM-based Technique and a Single Waist-worn Accelerometer. Accepted for publication in *Neurocomputing*
29. Samà A, Pérez C, Rodríguez-Martín D, Cabestany J, Moreno Aróstegui JM, Rodríguez-Molinero A (2013) A heterogeneous database for movement knowledge extraction in Parkinson's disease. In: *European Symposium on Artificial Neural Networks, Computational Intelligence and Machine Learning*
30. Samii A, Nutt JG, Ransom BR (2004) Parkinson's disease. *Lancet* 363(9423):1783–1793. doi:[10.1016/S0140-6736\(04\)16305-8](https://doi.org/10.1016/S0140-6736(04)16305-8). <http://www.sciencedirect.com/science/article/pii/S0140673604163058>
31. Schaafsma JD, Balash Y, Gurevich T, Bartels AL, Hausdorff JM, Giladi N (2003) Characterization of freezing of gait subtypes and the response of each to levodopa in Parkinson's disease. *Eur J Neurol* 10(4):391–398. doi:[10.1046/j.1468-1331.2003.00611.x](https://doi.org/10.1046/j.1468-1331.2003.00611.x)
32. Sian J, Gerlach M, Youdim MBH, Riederer P (1999) Parkinson's disease: a major hypokinetic basal ganglia disorder. *J Neural Transm* 106:443–476. doi:[10.1007/s007020050171](https://doi.org/10.1007/s007020050171)



Claas Ahlrichs received his Ph.D. in engineering at the University of Bremen, Germany, in 2015. Since 2011, he has been involved in several national and international research projects. Currently, he is working as a software developer.



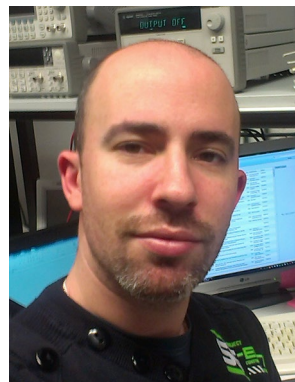
Albert Samà completed his Ph.D. at Universitat Politècnica de Catalunya (UPC) in 2013. Since 2009, he has been involved in several European projects. He has been assistant professor since 2010.



Michael Lawo professor for applied computer science at Universitaet Bremen, Managing Directors of neusta mobile solutions GmbH, involved in projects of logistics, wearable computing, eHealth, AAL and artificial intelligence.



Joan Cabestany holds currently a Professor position at the Department of Electronic Engineering of the UPC. He obtained his Ph.D. degree in Telecommunication Engineering in 1982 from UPC. Currently, he is with the staff of the CETpD, UPC Research Center, and is the responsible for applied technological research for helping people with chronic diseases. His recent activity includes the coordination of the 7FP projects REMPARK and FATE.



Daniel Rodríguez-Martín (Barcelona, Spain, 1983) is a Postdoc researcher at the Technical Research CETpD, from UPC. He received his B.Sc. degree in Industrial Automatic Control and Electronics in 2009. In 2011, he obtained the M.Sc. degree in Automatic Control and Robotics, and in 2014, he received the Ph.D. in Automatic Control, Vision and Robotics at the Automatic Control Department of the UPC. He has participated in several national and European projects carried out by CETpD since 2009.



Carlos Pérez-López (Barcelona, Spain, 1977) received his degree in Electronic Engineering in 2004. He is currently working as coordinator of research projects at the Technical Research Centre for Dependency Care and Autonomous Living of Universitat Politecnica de Catalunya. His research interests include embedded devices, online algorithms and inertial sensors.



Dr Timothy Counihan is Honorary Professor of Medicine (Neurology) at National University of Ireland Galway. He is a Fellow of the European Association of Neurology and Board Certified with the American Board of Psychiatry and Neurology. His research interests include Parkinson's disease and Huntington disease.



Dean Sweeney is a Ph.D. student in the School of Engineering and Informatics at the National University of Ireland (NUI) in Galway, Ireland. His current research interests include sensor and actuator technologies for healthcare and wellness applications. His primary focus is on NMES systems development for the use in neurological disorders.



Patrick Browne is an RANP Neurology (movement disorders) in Galway University Hospitals and is also an Hon. Clinical Fellow, School of Medicine NUIG. Research includes: assistive and therapeutic technologies, in PD and service delivery methodologies.



Leo R. Quinlan is a lecturer in Physiology in the School of Medicine at the National University of Ireland (NUI) in Galway, Ireland. His current research interests include medical device development, usability, neuromodulation, electrophysiology and ion channel function in neural and non-neural systems.



Lewy Hadas is a Director of Maccabi International Center for Research & Development. She completed her MSc in biochemistry, PhD in Human Genetics & Molecular Medicine Sackler School of Medicine, TA University, Israel. She obtained Postdoc Fellowship in Clinical Biochemistry & Molecular Medicine, Medical School Pitie-Salpetriere, Paris, France. In addition to research activities, she is responsible for multinational collaboration and EU projects, and her experience includes R&D of telemedicine-enabled medical device for cancer patients, and treatment of diabetes.



Gearóid Ó Laighin is the Professor of Electronic Engineering and Head of Electrical & Electronic Engineering at NUI Galway. He has coauthored over 50 peer-reviewed international biomedical engineering journal papers. Research interests include the design of medical devices with a particular interest in human movement sensing, the application of neuromuscular electrical stimulation, connected health and usability considerations in the design of medical devices for home health care.



Dr. Vainstein is specialized in Adult Neurology and gained his experience in general Neurology working at Meir Hospital and as a consultant specialist in Neurology at Maccabi Healthcare Services and Clalit Health Services. Dr. Vainstein is working for Maccabi and its R&D institute on new technologies (NEVET) for the aging populations. He was the first investigator in Israel of the clinical part in the EC REMPARK project and coleader in the clinical EC USEFIL project.



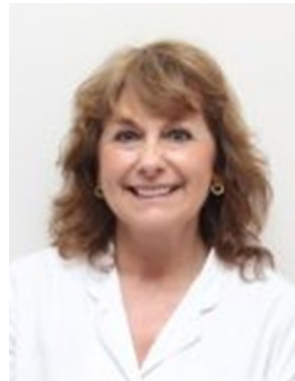
Alberto Costa was born in Rome on September 19, 1970. He holds Italian nationality. He is Associate Professor in Psychology at Niccolò Cusano University of Rome and research consultant at IRCCS Fondazione Santa Lucia, Rome.



Paola Quispe completed her MBBS at University Mayor Saint Simon, Bolivia, in 2002. Since 2008, she has been involved in the treatment of Parkinson's disease patients.



Roberta Annicchiarico MD, PhD is a geriatrician, medical assistant at Fondazione S. Lucia, Rome. She is contracted professor of Geriatrics and Int. Medicine at the University Tor Vergata, Rome. She has been involved in many EU Projects. She is chief of the Technology-assisted Neurorehabilitation Lab.

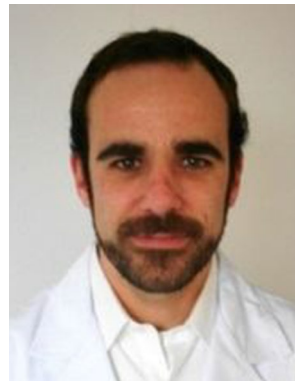


Àngels Bayes MD is a Doctor in Medicine, Neurology Specialist. She has focused her career and research on the treatment and study of Parkinson's disease, Tourette syndrome and dementias such as Alzheimer's disease. She is director of the Unit for Movement Disorders and Parkinson of Teknon Medical Center during the last 20 years. She has participated in 39 research projects and has 59 publications. She is the author of 6 books, in addition to lectures, conferences, communica-



Sheila Alcaine completed her B.Sc. in Physiotherapy at Universitat EUIF Blanquerna in 2003. Since 2008, she has been involved in the treatment of Parkinson's disease patients.

tions and teaching courses



Alejandro Rodríguez-Moliner PhD, MD graduated in medicine at the Universidad Autónoma de Madrid (UAM), in 2001. In 2006, he completed the specialization in Geriatrics at the Hospital Central de la Cruz Roja in Madrid. In 2009, he was awarded the PhD degree in Preventive Medicine and Public Health at the UAM. In 2010, he was appointed Adjunct Professor in Gerontechnology at the School of Engineering and Informatics, NUI Galway.



Berta Mestre completed her B.Sc. in Physiotherapy at Universitat Ramon Llull Blanquerna in 2010. Since 2011, she has been involved in the treatment of Parkinson's disease patients.