



STUDY PROTOCOL

REVISED Neuromuscular assessment of force development, postural, and gait performance under cognitive-motor dual-tasking in healthy older adults and early Parkinson's disease patients: Study protocol for a cross-sectional Mobile Brain/Body Imaging (MoBI) study [version 2; peer review: 1 approved, 1 approved with reservations]

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V2 First published: 17 Apr 2023, 3:58
<https://doi.org/10.12688/openreseurope.15781.1>

Latest published: 25 Aug 2023, 3:58
<https://doi.org/10.12688/openreseurope.15781.2>

Abstract

Background: Neuromuscular dysfunction is common in older adults and more pronounced in neurodegenerative diseases. In Parkinson's disease (PD), a complex set of factors often prevents the effective performance of activities of daily living that require intact and simultaneous performance of the motor and cognitive tasks.

Methods: The cross-sectional study includes a multifactorial mixed-measure design. Between-subject factor grouping the sample will be Parkinson's Disease (early PD vs. healthy). The within-subject factors will be the task complexity (single- vs. dual-task) in each motor activity,

Open Peer Review

Approval Status

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i.e., overground walking, semi-tandem stance, and isometric knee extension, and a walking condition (wide vs. narrow lane) will be implemented for the overground walking activity only. To study dual-task (DT) effects, in each motor activity participants will be given a secondary cognitive task, i.e., a visual discrimination task for the overground walking, an attention task for the semi-tandem, and mental arithmetic for the isometric extension. Analyses of DT effects and underlying neuronal correlates will focus on both gait and cognitive performance where applicable. Based on an a priori sample size calculation, a total N = 42 older adults (55-75 years) will be recruited. Disease-specific changes such as laterality in motor unit behavior and cortical control of movement will be studied with high-density surface electromyography and electroencephalography during static and dynamic motor activities, together with whole-body kinematics.

Discussion: This study will be one of the first to holistically address early PD neurophysiological and neuromuscular patterns in an ecologically valid environment under cognitive-motor DT conditions of different complexities. The outcomes of the study aim to identify the biomarker for early PD either at the electrophysiological, muscular or kinematic level or in the communication between these systems.

Clinical Trial Registration: ClinicalTrials.Gov, [NCT05477654](https://clinicaltrials.gov/ct2/show/study/NCT05477654). This study was approved by the Medical Ethical Committee (106/2021).

Keywords


Parkinson's disease (PD), Mobile Brain/Body Imaging (MoBI), dual-tasking, neuromuscular function, older adults




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Any reports and responses or comments on the article can be found at the end of the article.

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Author roles: **Marusic U:** Conceptualization, Funding Acquisition, Investigation, Methodology, Resources, Supervision, Writing – Original Draft Preparation, Writing – Review & Editing; **Peskar M:** Writing – Original Draft Preparation, Writing – Review & Editing; **Šömen MM:** Writing – Original Draft Preparation, Writing – Review & Editing; **Kalc M:** Writing – Original Draft Preparation, Writing – Review & Editing; **Holobar A:** Writing – Original Draft Preparation, Writing – Review & Editing; **Gramann K:** Conceptualization, Funding Acquisition, Writing – Review & Editing; **Wollesen B:** Writing – Original Draft Preparation, Writing – Review & Editing; **Wunderlich A:** Writing – Review & Editing; **Michel C:** Funding Acquisition, Writing – Review & Editing; **Miladinović A:** Software, Writing – Review & Editing; **Catalan M:** Investigation, Supervision, Writing – Review & Editing; **Buoite Stella A:** Writing – Review & Editing; **Ajcevic M:** Supervision, Writing – Review & Editing; **Manganotti P:** Funding Acquisition, Supervision, Writing – Review & Editing

Competing interests: No competing interests were disclosed.

Grant information: This project has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement No 952401 (TwinBrain – TWINning the BRAIN with machine learning for neuro-muscular efficiency). The authors also acknowledge financial support from the Slovenian Research Agency (research core funding no. P5-0381). AH was supported by Slovenian Research Agency (project J2-1731 and programme funding P2-0041).

The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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How to cite this article: Marusic U, Peskar M, Šömen MM *et al.* **Neuromuscular assessment of force development, postural, and gait performance under cognitive-motor dual-tasking in healthy older adults and early Parkinson's disease patients: Study protocol for a cross-sectional Mobile Brain/Body Imaging (MoBI) study [version 2; peer review: 1 approved, 1 approved with reservations]** Open Research Europe 2023, 3:58 <https://doi.org/10.12688/openreseurope.15781.2>

First published: 17 Apr 2023, 3:58 <https://doi.org/10.12688/openreseurope.15781.1>

REVISED Amendments from Version 1

According to the reviewer's request, we changed the introduction and removed part of the methods from it, moved the [Figure 1](#) below and added few additional information about the protocol.

Any further responses from the reviewers can be found at the end of the article

Introduction

Parkinson's disease (PD) is the second most common neurodegenerative disease after Alzheimer's disease, affecting 2–3% of the population over 65 years of age worldwide^{1,2}. This progressive disorder results from the depletion of dopaminergic neurons in the substantia nigra pars compacta³. In the early stages of the disease, PD patients present mainly with motor symptoms such as tremor at rest, rigidity, bradykinesia, postural instability, and gait disturbances, which become more complex in advanced stages and are accompanied by non-motor symptoms⁴. Clinically, the most striking features of patients with postural instability/gait disorder (PIGD) are postural instability with falls and freezing of gait (FOG), which occur mainly in advanced stages and are episodic and unpredictable^{5,6}. Gait impairments in PD patients encompass reduced speed and stride length, increased double limb support time, and stride-to-stride variability^{7–10}.

Although PD motor symptoms dominate clinical presentations, most patients also experience cognitive and behavioral impairments, cognitive executive deficits, and depression^{11,12}. The cognitive dysfunction observed in PD patients is primarily due to dopaminergic and cholinergic dysfunction¹³, and is typically associated with impaired frontostriatal circuitry^{11,14}, which impairs action planning and problem-solving¹⁴, attentional set-shifting¹⁵, response inhibition¹⁶, and decision making¹⁷. The three nonmotor frontostriatal circuits that connect frontal cortical regions to subcortical regions such as the basal ganglia originate in the dorsolateral prefrontal cortex (DLPFC), anterior cingulate cortex (ACC), and orbitofrontal cortex (OFC)¹⁸.

A link between global cognitive dysfunction and motor symptoms in PIGD patients has been previously suggested¹⁸. Compared to tremor-dominant patients (TD), patients with PIGD have greater impairments on measures of global cognition¹⁹, a higher frequency of mild cognitive impairment²⁰, and an increased risk of developing dementia²¹. In their study of 783 participants, Kelly and colleagues¹⁹ found an association between deficits in global cognition, executive function, memory, phonemic fluency, and more severe symptoms of postural instability/gait disturbance. While visuospatial impairments were associated only with more severe freezing and poorer memory function was associated only with greater postural instability, executive function deficits were associated with greater postural instability, more severe freezing, and gait impairments. Interestingly, gait characteristics of speed, variability, and postural control were also found to precede and

predict declines in attention and visual memory^{22,23}, as well as dementia in old age²⁴.

To understand the complexity of PIGD, locomotion control has been studied extensively. To efficiently perform a locomotor task, one should be able to (i) generate proper force for the step, (ii) maintain postural balance, and (iii) continuously adapt to environmental demands and behavioral goals²⁵. To ensure optimal walking, multiple sensorimotor processes must be functionally integrated on several levels to maintain the forward propulsion of the centre of mass while maintaining dynamic stability²⁶. At the highest level, the timing of muscle contraction between agonists, antagonists, and synergists acting on both sides of the body are precisely regulated²⁶. At lower levels, the behavior of motor units (MU) pattern is regulated following a size principle to ensure smooth muscle contractions²⁷. However, sensorimotor processes are impaired in PD patients, resulting in altered motor activity patterns during rest and locomotion²⁶. Moreover, it has been shown that the simultaneous performance of cognitive and motor tasks can increase differences in motor performance between healthy individuals and PD patients²⁶.

In daily life, balancing or walking is rarely the only task performed at a given moment (single task; ST). Most often, it is performed together with another task (dual task; DT) or sometimes even several other tasks as a multitasking activity, resulting in competition for limited resources which can degrade performance on one or all tasks²⁸. In case of performing cognitive and motor tasks simultaneously, the performance decrements are a consequence of cognitive-motor interference (CMI). By comparing the performance success of both, ST and DT conditions, we can calculate dual-task costs (DTC) using the following formula: $(DT\ performance - ST\ performance) / ST\ performance$ ^{29–32}. In PD patients, performing a cognitive serial subtraction task has been shown to impair the concurrently performed motor task as observed in decreased accuracy of force tracking³³, and promote deteriorating effects on the motor system such as inducing tremor³⁴. Studies of postural stability have shown that individuals with PD have impaired postural control under dual-task conditions^{35,36}. Impaired walking in PD patients is more pronounced under cognitive-motor DT conditions^{37–42}, with additional impairments observed in symmetry and coordination between the left and right steps^{43,44}. Walking difficulties in PD patients can additionally be exacerbated in situations where the walking path is being restricted, such as narrow pathways/corridors/doorways or when walking between objects⁴⁵. For example, Cowie and colleagues⁴⁶ have shown that PD patients slow down at doorways to an extent that is inversely proportional to doorway width and that the frequency of freeze events increases in narrower doorways.

Another important aspect is the cognitive effort involved in activities of daily living, which varies according to the complexity of the situation and the associated cognitive demands⁴⁷. Regardless of complexity, cognitive processing is highly dependent on visual information, ranging from the simplest scenario, such as recognizing and responding to a red light, to a

highly complex scenario involving navigating through an unfamiliar city on uneven surfaces while ensuring the safety of the accompanying infant. These situations illustrate the importance of visual information processing for the generation, execution, and adaptation of motor commands during walking⁴⁸⁻⁵⁰.

PD patients experience walking difficulties and gait disturbances under the ST walking condition compared to healthy individuals, which become further exacerbated under the DT condition. Despite many efforts, it remains unclear how motor and cognitive symptoms contribute to ST or DT walking difficulties²⁹. Investigating the interaction between motor and cognitive symptoms not only on behavioral and but also on neurophysiological levels in ecologically valid settings could greatly enhance our understanding of the underlying mechanism of ST and DT performance decline. Such investigations, however, have only recently been enabled by the Mobile Brain/Body Imaging (MoBI;⁵¹⁻⁵³) approach, which will be used in the present study to investigate time-synchronized electroencephalographic (EEG) and electromyographic (EMG) recordings as well as body movements in multilevel approach to understanding cognitive-motor interactions and their neuromuscular signature. The aim of the present study is to investigate the early PD disease-specific changes and laterality of neuronal rhythms during the three motor activities that demand different levels of cognitive-motor resources, namely (1) overground walking (dynamic condition), (2) semi-tandem stance (standing static condition), and (3) sustained isometric contraction (seated static condition), each as motor ST but also in combination with certain cognitive tasks in DT conditions. This will allow systematic assessment and identification of motor and cognitive factors contributing to the symptomatology of PD patients.

Compared with healthy participants, we hypothesize that early PD will have poorer ST performance on cognitive and motor tasks and increased DT costs on all motor activities. Furthermore, we expect that performance impairments with increased neural recruitment will be most pronounced in the highest motor complexity condition (walking in the narrow pathway) and even more pronounced in the corresponding DT variant (DT-narrow).

Methods

Trial design

This protocol is formulated according to the SPIRIT (Standard Protocol Items: Recommendations for Interventional Trials) statement⁵⁴. It describes a multifactorial mixed-measure design and encompasses three different motor activities. Specifically, the overground walking activity on a wide and a narrow path (ST-wide and ST-narrow) will constitute the core part of the study and will also be examined during a concurrent cognitive color discrimination task (DT-wide and DT-narrow); the cognitive task will additionally be examined in an ST condition for a comprehensive comparison (ST-visual). Second, postural control will be examined in the semi-tandem stance alone (ST-sts) and simultaneously with a cognitive task examining attention (DT-sts). The attention task will also be examined as ST in the seated position (ST-Stroop). Finally, in the seated

position, isometric force development activity over time will again be examined alone (ST-is), under dual-task cognitive conditions while performing a series-3 subtraction task (DT-is), with the series-3 subtraction also performed in the absence of the motor task (ST-sub). In each motor activity, namely the overground walking, semi-tandem stance, and isometric contractions, one between-subject factor *Group* (early PD vs. neurotypical older adults) and within-subject factor *task complexity* (ST vs. DT) will be considered. In addition, in the walking activity, the second within-subject factor *walking difficulty* (wide vs. narrow) will be implemented. This leads to a mixed between-within 2 (*Group*) x 2 (*task complexity*) study design for the isometric and stance activities and to a mixed between-within 2 (*Group*) x 2 (*task complexity*) x 2 (*walking difficulty*) design for walking activity. **Figure 1** provides an overview of the study structure and timeline, as well as motor activities with ST and DT conditions embedded. We are aware that the direct comparison between the DT conditions of the three motor activities will not be possible due to the use of different cognitive tasks, yet this decision was motivated to avoid extensive learning effects and increasing frustration during the measurement day. A detailed description of the cognitive tasks and their respective stimuli can be found below under “Cognitive task performance”.

Ethical approval and confidentiality. The study was registered at IRB of Trieste University Hospital – ASUGI, Trieste, Italy (ASUGI protocol number: 106/2021; approved 20.12.2022) and on ClinicalTrials.Gov under code: NCT05477654. The written informed consent for the patients will be obtained by a medical doctor (PM or MC), a movement disorder specialist, employed by the Neurological Clinic of Trieste University Hospital, while for the healthy participants the lead researcher will obtain it. All participant information and data will be stored securely and identified by a coded ID number only to maintain participants’ confidentiality.

Managing adverse effect. The protocol and proposed experiments were modified after the pilot trials to minimize the occurrence of factors that led to experiment termination. Specifically, the number of stimuli and the duration of conditions were kept low enough to avoid fatigue but high enough to ensure adequate data quality, while the breaks between tasks/conditions were made sufficiently long. If reasons such as fatigue will nevertheless lead to a request for an interruption, the investigation will be terminated. Similarly, if patients/participants will experience dizziness or will experience adverse effects for other reasons, the examination will be terminated. A physician will be present to ensure physical safety. The experimental day is organized in three experiments, so the recorded data will provide valuable insights despite the number of experiments completed.

Eligibility criteria

Inclusion criteria for the healthy older participants will be: (1) no diagnosis of cognitive or movement disorder, (2) Montreal Cognitive Assessment (MoCA;⁵⁵) score ≥ 24 points (3) Short Physical Performance Battery (SPPB)⁵⁶ score > 7 points

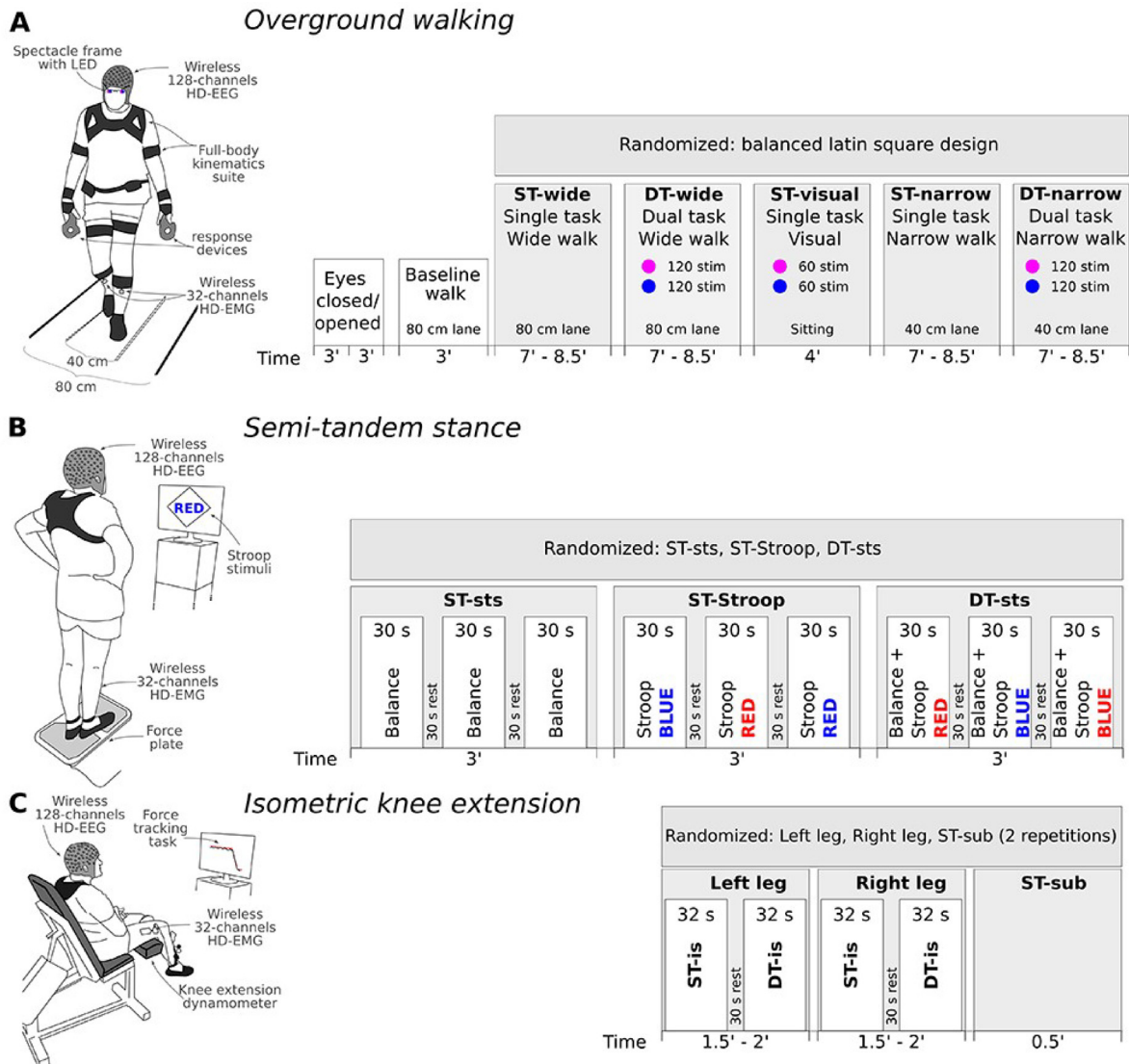


Figure 1. Graphic depiction of each motor activity, timeline of the procedure, and randomization order. (A) Overground walking will start with a resting-state EEG recording with both eyes closed and opened, followed by a baseline walk, and finally randomized order of ST and DT conditions during overground walking on either wide or narrow lanes with/or visual discrimination task (blue and purple light). (B) Next, the semi-tandem stance will be performed, again using randomization of the ST and DT conditions across participants, while the 30-second segments within each condition will be kept at fixed order. (C) Lastly, during the isometric knee contraction the order of the leg blocks and ST-sub will be randomized, while during each leg block, the ST will always be performed before the DT. Duration of each recording is denoted below the time axis. For a detailed explanation refer to the main text. Abbreviations: ST – single task, DT – dual task, sts – Semi-tandem stance, sub – subtraction or mental arithmetic task.

(4) living independently in the community, (5) age range between 50 and 75 years, (6) no color blindness.

Inclusion criteria for the early-stage PD patients: (1) diagnosed according to the last International and Movement Disorder Society (MDS) criteria (2015), (2) duration of the disease less than 5 years, (3) positive Dopamine Transporter Scan (DAT-SCAN), (4) diagnosed akinetic-rigid or mixed phenotype, (5) score of 1 or maximally 2 on the Hoehn and Yahr scale⁵⁷,

(6) MoCA score ≥ 24 points, (7) SPPB score > 7 points, (8) age range between 50 and 75 years, (9) no color blindness, (10) living independently in the community.

Exclusion criteria will be (1) advanced stages of PD (Hoehn and Yahr score > 2 points), (2) any acute or untreated chronic diseases, especially of the peripheral and central nervous system, such as heart failure, respiratory failure, severe osteoarthritis, and major psychiatric illness, such as depression,

(3) MoCA score < 24 points, (4) severe disabling tremor, (5) recurrent falls, (6) impaired vision that is not corrected with e.g., glasses, (7) SPPB score ≤ 7 points.

Outcome measures

This protocol focuses on the behavioral, cognitive, motor, and/or kinematic outcomes as well as neurophysiological markers associated with CMI during dual-tasking in healthy aged individuals and early PD patients. The accompanying information regarding the participants' demographics and general health will be treated as a secondary outcome and used to control for the confounds.

Primary outcomes. The primary outcomes suggested by this protocol will be the DT effects associated with the concurrent execution of overground walking/semi-tandem stance/isometric contraction *and* a respective secondary cognitive task. The motor task parameters, e.g., the gait parameters using full-body kinematics (step length and width, stride time variability, walking speed), the center of pressure (COP) displacement for the static stance, and deviation of the required force development over time for the isometric contractions, will be investigated in the ST and DT conditions. Additionally, each cognitive task performance will be recorded in the ST condition to allow for the investigation of the DT effect on the cognitive level too. The cognitive ST condition will be performed in a seated position for all cognitive tasks (visual discrimination task, counting of Stroop stimuli, serial-3 subtraction; for description see below). High-density surface electromyography (HD-EMG) and high-density electroencephalography (HD-EEG) will be recorded in all motor activities in both ST and DT conditions. In the overground walking experiment, the DT condition will be created using a visual discrimination task, in the semi-tandem stance experiment the cognitive secondary task will be counting the occurrences of specific visual Stroop stimuli, and finally, in the isometric contraction experiment, the secondary task will encompass mental arithmetic. All tasks include aspects of executive functions. However, to avoid learning effects the task-set up was conducted differently between all conditions. Below are detailed descriptions of the cognitive and motor tasks.

Cognitive task performance:

- **Contra- vs. Ipsi-Lateral Visual Discrimination Task performance in walking activity:** The outcome measures for a visual discrimination task will be response accuracy and response time. The implementation of such a task will allow for the recording of the event-related potentials as well as continuous brain activity. Stimuli of the cognitive task used in the ST and DT walking experiment will vary according to the presentation side (left vs. right), spectral properties (magenta vs. cyan), and presentation-response laterality (ipsi- vs. contra-lateral). DT costs and the associated neural underpinnings will be assessed in the cognitive task as well as in gait parameters.
- **Silent counting of specific Stroop stimulus occurrences in the semi-tandem stance activity:** Computerized Stroop

task words “Blue” and “Red” will be presented one at a time in either blue or red ink color. Presenting stimuli at a rate of 1/s, participants will have to count the number of specific occurrences, e.g., “Blue” in blue ink, from the four possible options across 30-second intervals. This will be repeated three times, each time focusing on and counting a different color combination of stimuli.

- **Mental arithmetic:** Sequential subtracting is typically used as a screening procedure to assess attention/working memory functions. In our study, participants will be asked to quietly perform serial-3 subtraction from a random 3-digit number and verbally indicate the resulting number after a 30-second interval. This will be performed 3 times. The resulting number will be noted and will serve as an indirect measure of success. Standardization and adult norms are available⁵⁸.

Motor performance:

- For the walking experiment, the full-body kinematics will be captured using 17 inertial Shadow Motion sensors (Motion Workshop, Seattle, WA, USA). To ensure higher precision of body position and alignment correction in space, an additional infra-red based tracker (VivePro, HTC, Taoyuan, TW) surrounded by three base stations will be used. The following gait parameters will be extracted from the feet sensors: walking speed, step length, step width, double support time, and step-to-step variability. The arm swing and hip rotation information will be extracted using the arm and torso sensors.
- Second, postural control will be measured with a force plate (AMTI HE600600-2 k, Advanced Mechanical Technology, Inc., Watertown, MA, USA). The COP displacement in antero-posterior and medio-lateral directions will be calculated in cm.
- Lastly in the sitting activity of isometric knee extension, the ST and DT performance will be investigated using the isometric dynamometer (Wise Technologies Ltd., Ljubljana, Slovenia) *Muscle activity:*

EMG activity will be recorded using two wireless 32-channel probes (MUOVI, OT Bioelettronica S.r.l., Torino, Italy). For walking and semi-tandem stance, 8 x 4 matrix electrodes (10 mm interelectrode distance, HD10MM0804, OT Bioelettronica S.r.l., Torino, Italy) will be attached to bilateral tibialis anterior muscles. For the force tracking task, the electrodes will be attached to the bilateral vastus lateralis muscles. The electrodes and probes will be taped on the skin to minimize motion artifacts.

HD-EMG signals collected during walking and semi-tandem stance will be analyzed in the time and frequency domain using HD-EMG amplitude envelopes techniques to calculate entropy, HD-EMG amplitude barycenter, and spatial differences in HD-EMG signals. HD-EMG signals collected during the isometric force tracking will be decomposed in contributions

of single MUs using blind source separation methods⁵⁹ to study individual motor unit firing patterns and early assessment of tremor.

Brain activity

EEG activity will be recorded using a mobile 128-electrode wireless system (CGX, Cognionics Inc., San Diego, USA). The electrodes will be mounted onto an elastic cap (easyCAP) and positioned according to the standard 5–10 system⁶⁰. Before recording brain activity during the three motor activities, the continuous resting-state EEG will be recorded at baseline while sitting; for 3 min with both eyes open and eyes closed. After this, the brain activity will be continuously recorded during ST and DT conditions in the overground walking, during postural semi-tandem stance, and finally for the static isometric knee extensions. The investigated event-related markers associated with visual processing in the visual discrimination task during walking activity and attentional task during the semi-tandem stance will be the visual P1, N1, and P2 components over the occipital cortex, and the P3 component over the parietal cortex assessed in both ST and DT conditions. The EEG data during all motor and motor-cognitive tasks will be analyzed by the means of time–frequency spectral analysis. For the first time, the gait-phase spectral modulation⁶¹ will be assessed during overground walking in PD, to allow for investigation of the movement-related beta band hypersynchrony which is the disease characteristic⁶². The communication between the brain and muscles (i.e., cortico-muscular connectivity) will be assessed by EEG-EMG coherence analysis.

Secondary outcomes. The secondary measures will be collected to assess the eligibility criteria and the general health status of cognitive and motor function. These measures will also be used to control for the covarying factors.

Demographic questionnaires will collect data on participants' age, sex, ethnicity, body mass and height, leg length, and socio-educational status. Also, their handedness in 10 everyday situations will be asked.

The *general health questionnaire* will include PD-specific anamnestic questions for the PD patients, while in all participants we will scan for comorbidities, history and ongoing lower limb injuries/surgeries, presence of acute pains, sleeping hours of the last night, and coffee/energy drink intake on the measurement day.

The *Short Falls-Efficacy-Scale-International* (SFES-I;^{63–65}) is a 7-item questionnaire used to assess the fear of falling associated with the execution of easy and complex physical and social activities. A translated Italian version is available and we will provide a translation for the Slovenian-speaking subsample.

The *Short Physical Performance Battery* (SPPB;⁵⁶) will assess lower extremity function in the elderly and consists of three sections. First, the ability to perform a Romberg, semi-tandem, and tandem stance. Second, the time of walking

a 4-meter track at a comfortable speed is measured. Third, participants must perform 5 sit-to-stand transfers as fast as possible, and time-to-completion is measured. Total SPPB score ranges from 0 (low mobility) to 12 (high mobility; up to 4 points per section) and takes up to 10 min to complete.

The *New Freezing of gait questionnaire* (N-FOG;⁶⁶) is a 9-item self-reported questionnaire measuring the freezing of gait and walking abilities in PD patients through various scenarios, and the impact of gait freezing on daily activities.

The *Falls History Questionnaire* (FHS) assesses the falls incidence, location, and consequence in the previous 12 months and is based on the recommendation of the Lamb and colleagues⁶⁷ and Lord and colleagues⁶⁸.

The *Movement Disorder Society Unified Parkinson's Disease Rating Scale* (MDS-UPDRS;⁶⁹) is a comprehensive assessment of various aspects of PD including motor and non-motor experience of daily living and the related complications. The evaluation characterizes the extent and burden of the disease.

The *Montreal Cognitive Assessment* (MoCA;⁵⁵) is a one-page 30-item test developed for screening for mild cognitive impairment. It involves items to assess a range of cognitive domains, including executive functions, visuospatial abilities, language, attention, working memory, abstraction, and orientation to time and place. The duration of the assessment is about 10 min. The validated Italian⁷⁰ and Slovenian versions are available⁷¹.

The *Trail Making Test* (TMT;⁷²) is a measure of psychomotor speed and attentional set-shifting. It requires the subject to connect 25 encircled numbers randomly arranged on a page in increasing order (Part A), and 25 encircled numbers and letters in alternating and increasing order (Part B) using a pen. Both part A and part B include practice exercises. About 5 to 10 minutes are needed for test administration. The outcome measure of the TMT is the time of completion.

The *Brief Mood Introspection Scale* (BMIS;⁷³) is comprised of 16 emotional/mood adjective or phrase items relating to the present moment and a participant is required to respond using a 4-point Likert scale ranging from definitely do not feel to definitely feel.

The *Beck's Depression Inventory* (BDI;⁷⁴) consists of 21 items of depressive symptoms experienced over the past week. Each symptom is described in 4 increasing intensities, and a participant must decide which one best describes their state. Zero to maximum of 3 points are given per item proportionally to the chosen intensity.

The *Dual-Task Strategy Assessment* is a modified version of the tool used by Wollesen and colleagues⁷⁵ and consists of 6 closed-type questions/statements regarding the employed strategy and allocation of one's attention during DT performance.

The *NASA Total Load Index* (NASA-TLX;⁷⁶) is a tool assessing subjective workload across 6 subscales: mental, physical, and temporal demand, as well as performance, effort, and frustration. The participant marks the degree of workload on a visual scale, ranging between 1 – 21. The overall score is calculated as a sum or average of the 6 subscales and the questionnaire is applied after each randomized condition in the overground walking activity.

Recruitment process

PD patients will be recruited at the Neurological Clinic of Trieste University Hospital, while the research will be conducted at the ASUGI Physiotherapy Gym of the Trieste University. During a regular visit to the Movement Disorders outpatient clinic, the potentially suitable candidates will be invited to participate and will have the study design, protocol as well as all procedures and treatment of clinical data fully explained, as for standard institutional practice. The control group of neurotypical older adults will be recruited via the Daily Activities Center Koper, a database of previously enrolled subjects in the experiments of the Institute of Kinesiology, Science and Research Centre Koper, and via a word of mouth. The initial telephone screening in healthy adults (demographic and general health screening) will be performed to identify the potential participants. The enrolled patients and healthy adults will undergo a comprehensive screening session including motor and neurocognitive assessment (Measurement Day 1 - M1). Having passed the inclusion/exclusion criteria confirmed on the M1, they will finally be invited to participate in Measurement Day 2 (M2). Participants will not be compensated for collaborating in the respective study in any way.

Procedure, tasks, and stimuli

The study proposed by this protocol will consist of 2 measurement days per participant. Following the recruitment and signing of the informed consent, the participants' characteristics

will be collected, and the questionnaires and tests listed under the Secondary outcomes section performed (except for the Dual-Task Strategy Assessment and NASA-TLX). The M1 will last approximately 1–1.5 hours. After the completion of the first measurement day, the fulfillment of inclusion/exclusion criteria will be carefully inspected, and the fitting participants will be invited for the M2. These participants will be shown a recording of the M2 setting embedded within the actual environment with a fully equipped volunteer performing a DT walking condition. The recording will serve to familiarize the participants with the setting and address the potential anxiety of encountering new situations in a highly controlled environment. This will also be the opportunity to respond to participants' questions and concerns.

The M2 will begin with the participant preparation; fixing EEG, HD-EMG, and full-body kinematic sensors on the participant. The order of motor activities, namely the overground walking, semi-tandem stance, and isometric force development, will remain fixed for all participants (see [Figure 1](#)). The measurement day 2 will last approximately 5 hours. Below we specify the procedure of each motor activity and the associated cognitive tasks in more detail.

Overground walking. After the resting state EEG recording (3 min with eyes closed and 3 min with eyes open), the baseline gait performance will be assessed by participants walking at self-selected speed but without carrying the response controllers in their hands along the 80 cm wide and 8m long lane for 3 minutes (excluding turn time). Then, the order of the ST and DT conditions will be randomized using a balanced Latin square formula for the 5 conditions, namely the ST-wide, DT-wide, ST-narrow, DT-narrow, and ST-visual conditions. In the visual discrimination task, the stimulus-response combinations will either share the side or not – ipsilateral or contralateral stimulus-response combinations, respectively (see [Table 1](#)). A practice session will be conducted prior

Table 1. Stimuli number of the cognitive visual discrimination task separated by motor task condition, hemifield presentation side, cognitive task modality, visual modality-specific properties, and presentation-response compatibility (in yellow).

MOTOR TASK:	Presentation side:	COGNITIVE TASK: Visual discrimination		Σ
		Magenta	Cyan	
ST-visual (Sitting)	Left	30	30	60
	Right	30	30	60
DT-wide (Walking)	Left	60	60	120
	Right	60	60	120
DT-narrow (Walking)	Left	60	60	120
	Right	60	60	120
	Σ	300	300	600
Correct response is	contra-lateral		Ipsilateral to the presentation side	

to each condition to ensure task comprehension and pacing familiarization.

The ST-visual condition will be performed in the seated position in the middle of the walking lane while wearing response devices - VIVE pro controllers. In addition, participants will carry a custom-made spectacle frame with attached LED light on each end covering the periphery of the visual field, which will serve to provide visual stimulation. A total of 240 stimuli (120 magenta and 120 cyan) will be presented on either the left or right spectacle side (50–50%; see [Table 1](#) for stimuli characteristics). The interstimulus intervals will vary between 400–800 ms, and a response window will be limited to 900 ms after the stimulus presentation and indicated by pressing the button on the left- or right-hand held controller. The trial duration will vary between 1.4 to 1.8 s. Should no response be given within the predetermined response window, the next trial will start after 900 ms.

ST-wide and ST-narrow walking multimodal data will be collected while participants walk at self-selected speed in the wide (80 cm) and narrow (40 cm;⁷⁷) lane of 8 m length, respectively. The entire walking distance excluding the turn path will amount to 384 m (approximately 420 m including the turn path). Passing the end of the lane, a turn will follow, and a new gait pass will start upon re-entering the lane. Participants will be carrying response devices (VIVE pro controllers) although they will not be used in these two conditions. The estimated time for each condition is 7–8.5 minutes with turn times included.

In both DT-wide and DT-narrow walking conditions, the timing for visual stimuli presentation and responses will be the same as in the ST-visual condition. However, at the same time, participants will be walking along either a wide or narrow lane (described above) and upon reaching the end of the lane will be required to make a turn and re-enter the lane. The walking time of 7–8.5 minutes is estimated per condition (including the turn time). Per one lane length, 5 stimuli will be presented. The condition will end as soon as a participant walked 420 m (including the turn path) and 240 stimuli were presented.

Between each condition, a 2-min rest will be assured, during which NASA-TLX will be applied and participants will be asked to refer to the past condition. At the end of the 5 conditions of the walking activity, also the dual-task strategy assessment will be asked.

Semi-tandem stance. Next, in semi-tandem postural experiment participants will perform three 30-second repetitions of semi-tandem stance for the ST-sts motor condition. This will be repeated in the DT-sts condition while simultaneously counting the occurrences of specific Stroop stimuli, and in the seated position for the ST-Stroop condition. The order of the ST and DT conditions will be counterbalanced across participants, however, the three 30-second repetitions of a certain condition will follow in succession in a fixed order.

Participants will be standing on a force plate (AMTI HE600600-2 k, Advanced Mechanical Technology, Inc., Watertown, MA, USA) in a semi-tandem position with their preferred (but always the same) foot in front and with their hands fixed on their hips. Over the 30-second runs in both ST-sts and DT-sts conditions, an average COP sway length (in cm) will be taken as an outcome measure. In DT-sts and ST-Stroop conditions the outcome measure will (also) be the reported number of the required observed Stroop-stimuli. In each of the three 30 sec repetitions, different combinations of Stroop stimuli will have to be attended, (e.g. “Blue” in blue ink, “Red” in red ink, and “Blue” in red ink). We choose silent counting of the Stroop-stimuli occurrences over loud to reduce the talking-related EEG artifacts. To motivate the participants into performing the cognitive task, they are required to report the result at the end of each 30-second trial. During the ST-sts condition, participants will be asked to direct their gaze to a fixation cross presented on the screen at the level of their eyes.

Isometric knee extension. Finally, for the trapezoidal isometric force tracking participants will undergo two 32-second force tracking sessions of knee extension (6 s rising phase, 20 s sustained phase, 6 s decline phase) per both right and left lower limbs in the ST-is motor condition. The participants will be sitting in a knee extension dynamometer (Wise technologies, Ljubljana, Slovenia) equipped with a force sensor and they will be instructed to actively contract their knee extensors to produce force up to 30% of their maximum voluntary contraction (MVC). The force tracking task will be repeated while concurrently performing the secondary serial-3 subtraction task quietly during the DT-is condition for both right and left lower limbs. The serial 3-subtraction task will also be quietly performed in an ST-sub condition for 32 seconds. Each time, a random 3-digit number between 300 and 500 will be selected as a starting point and at the end of the 32 second period, participants will be asked to report the number they had reached. The order of the right and left limbs and the ST-sub will be counterbalanced across the participants, while they will always perform the ST-is before the DT-is with a respective lower limb. The task will consist of 2 rounds of described conditions.

Randomization sequence. The randomization sequence will be created via the online [balanced Latin square tool](#) separately for each experiment.

Data streaming and management

All data collected through paper-pencil tests and questionnaires will be digitized for further processing. Functional testing will be controlled through Unity3D (Unity Technologies, San Francisco, USA), which will also assign basic demographic data to each participant’s measurement data. All functional measurements will be streamed and synchronized with the Lab Streaming Layer (Swartz Center for Computational Neuroscience, UCSD, USA). The synchronized data streams will comprise gait analysis (Motion Workshop, Seattle,

WA, USA and VivePro, HTC, Taoyuan, TW), EEG (CGX, Cognionics Inc., San Diego, USA), and HD-EMG (OT Bioelettronica S.r.l., Torino, Italy) recordings, and secondary task performance (VivePro, HTC, Taoyuan, TW & Raspberry Pi, Raspberry Pi Foundation, Cambridge, UK). Day-to-day management of the study, including data management, will be coordinated by the principal investigator of the study. All data collected will be fully anonymized. Each participant will be assigned a unique study identification number. Access to the data will be restricted to TwinBrain researchers and staff, in possession of authorization rights. Directly identifiable data (e.g., name, address, phone number) are kept secure and separate from other study data. Data procedures will be in accordance with the General Data Protection Regulation (GDPR). In accordance with the Ethics Committee commitment, we will obtain informed consent from participants to share their anonymous data with other researchers. Data management will be regularly reviewed by an internal data monitoring committee composed of data experts, engineers, psychologists, and movement disorder physicians.

Statistical analysis and power calculation

Descriptive data will be presented as means (M) and standard deviations (SD). The homogeneity of variances and normality of distribution of parameters will be tested with Levene's test and Shapiro-Wilk test, respectively. Using univariate analyses of variances (ANOVA), the between-subject factor grouping the sample will be *PD* (early PD vs. no PD), while the within-subject factors will be the *task complexity* (ST vs. DT) in each motor activity, i.e., the overground walking, semi-tandem stance, and isometric force knee extension, as well as the *walking condition* (wide vs. narrow) for the overground walking only. Significance will be set at $\alpha = 0.05$. The effect size will be presented as the unbiased adjusted partial eta squared⁷⁸. All post-hoc comparisons will control for multiple comparisons (e.g., Bonferroni). Statistical analyses will be done using SPSS (IBM, SPSS Inc, Chicago, IL, United States) or R (R Core Team, 2013).

Sample size estimation calculated using *g*Power* software (a priori: F-tests; ANOVA: repeated measures, within-between interaction; effect size $f = 0.2$; alpha error probability = 0.05; power (1-beta error probability) = 0.8; number of groups = 2; number of measurements = 3) revealed a total of $N = 42$ subjects, half of who will be healthy older adults, and the other half will belong to early PD patient population.

Discussion

Recent advances in neuroimaging, wearables and wireless technologies offer a range of opportunities to better understand human behavior and link it to cortical and muscular activation patterns. MoBI enables the investigation of brain and muscle dynamics in conjunction with biomechanical parameters and secondary sensorimotor processes. The lack of scientific evidence in clinical populations has prompted us to conduct the present study with early-diagnosed PD patients. Therefore, the aim of this study is to investigate the differences in functional static and dynamic performance between healthy

and age-matched early-PD populations. The effect of interest will be investigated through DT effects and associated neuromuscular underpinnings. Specifically, using the MoBI approach, EEG, HD-EMG, and whole-body kinematics will be recorded under various ST and DT conditions to uncover the underlying mechanisms of force production, balance, and gait control in early-diagnosed PD patients. The underlying processes of interaction between motor and cognitive tasks will be identified at behavioral and neurophysiological levels under conditions of varying complexity. The general hypothesis is that there are performance differences and corresponding neuromuscular correlates between subgroups, with the highest neuromuscular compensations occurring in the PD group. We hypothesize that the performance differences are related to the different motor-cognitive processes such as stimulus input, resource allocation, and movement execution. Our goal is to find a neuromuscular biomarker that distinguishes the two populations and thus reflects an early onset of neuromuscular deficit in PD.

We anticipate several challenges in conducting this study. First, recruitment of early-stage PD will be a relatively lengthy process because of the limited number of such patients in the Trieste region (Italy). Second, the complexity of the test blocks is planned in such a way that subjects will have enough breaks and will be able to rest between trials. However, the long protocol (about 5 hours) will require careful preparation and handling with constant medical monitoring. We will try to maintain motivation throughout the measurement and avoid fatigue by taking multiple breaks. Finally, in our sample of early PD, we do not expect the occurrence of FOG during our measurements. A hypothetical explanation for the occurrence of FOG is that the narrowness of the corridor causes an increase in optic flow, as the visual image of near objects moves faster on the retina than that of more distant objects⁷⁹. Thus, visual and vestibular signals provide conflicting information to the brain in such situations. In our case, participants will be walking in a wide and narrow lane that gives the impression of a narrower path but does not enforce the narrow space/corridor.

In summary, the current study employs a state-of-the-art methodology to assess neuromuscular outcomes that aim to elucidate important underlying mechanisms of neural control of movement during dual motor-cognitive tasks in healthy older adults and in patients with early-diagnosed PD. Thus, we aim to uncover an early biomarker of movement (in)efficiency.

Dissemination

The communication about the running, upcoming or past activities for the general and specific public are regularly uploaded to the webpage of the TwinBrain project (www.twinbrain.si). In addition, the webpage and social media contain links to all media press, podcast series, and newsletters which are intended to reach a wide audience.

The scientific output of the study will primarily target researchers and medical doctors. The output of the proposed

protocol will produce at least three scientific papers, which will be submitted to peer-review scientific journals. In addition, the results of the clinical trial will be disseminated at content-related conferences, seminars, and congresses, such as the International Congress of Parkinson's Disease and Movement Disorders and the Mobile Brain/Body Imaging Conference.

The website will also be used for the long-term sustainability of the dissemination which will continue to communicate research findings and ongoing updates, and partnering with relevant organizations to continue research and dissemination efforts.

Study status

The protocol has been tested on 5 pilot healthy young subjects, intended to detect potential issues with the software and recorded data. Upon resolving all the issues, three healthy older adults underwent the protocol to assess subjective fatiguing effects in the target age group and to identify the strategies to ameliorate the fatigue. Currently, 6 subjects had been tested, of which 4 belonged to the health group and 2 were PD patients.

Ethics and consent

The study will be conducted in accordance with the ethical standards of the 1964 Declaration of Helsinki and the guidelines of Good Clinical Practice. IRB of Trieste University Hospital – ASUGI, Trieste, Italy approved this study protocol (ASUGI protocol number: 106/2021). The study was registered on ClinicalTrials.Gov under code: NCT05477654.

The informed consent for each study participant will be obtained by a medical doctor (MC or PM), a movement disorder specialist, employed by the Neurological Clinic Trieste. All participant information and data will be stored securely and identified by a coded ID number only to maintain participants' confidentiality.

Data availability

Underlying data

No underlying data are associated with this article.

Extended data

Zenodo: Demographic and General Health Questionnaire. <https://doi.org/10.5281/zenodo.7768228>⁸⁰

Data are available under the terms of the [Creative Commons Attribution 4.0 International license \(CC-BY 4.0\)](#).

Authors' contributions

UM, KG, CM, and PM conceived of the study and are grant holders. UM with help of MP, MMŠ, and MK drafted the manuscript. BW supervised and validated the DT settings. All authors contributed to the refinement of the study protocol and approved the final manuscript.

Acknowledgements

We thank Benjamin Paulisch for programming the experimental paradigm and building the customized hardware as well as being a great support in the background.

References

- Poewe W, Seppi K, Tanner CM, et al.: **Parkinson disease**. *Nat Rev Dis Primers*. 2017; **3**(1): 17013. [PubMed Abstract](#) | [Publisher Full Text](#)
- Tysnes OB, Storstein A: **Epidemiology of Parkinson's disease**. *J Neural Transm (Vienna)*. 2017; **124**(8): 901–905. [PubMed Abstract](#) | [Publisher Full Text](#)
- Gibb WR, Lees AJ: **The relevance of the Lewy body to the pathogenesis of idiopathic Parkinson's disease**. *J Neurol Neurosurg Psychiatry*. 1988; **51**(6): 745–52. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Jankovic J: **Parkinson's disease: clinical features and diagnosis**. *J Neurol Neurosurg Psychiatry*. 2008; **79**(4): 368–76. [PubMed Abstract](#) | [Publisher Full Text](#)
- Bloem BR, Hausdorff JM, Visser JE, et al.: **Falls and freezing of gait in Parkinson's disease: a review of two interconnected, episodic phenomena**. *Mov Disord*. 2004; **19**(8): 871–84. [PubMed Abstract](#) | [Publisher Full Text](#)
- Müller MLTM, Marusic U, van Emde Boas M, et al.: **Treatment options for postural instability and gait difficulties in Parkinson's disease**. *Expert Rev Neurother*. 2019; **19**(12): 1229–1251. [PubMed Abstract](#) | [Publisher Full Text](#)
- Blin O, Ferrandez AM, Serratrice G: **Quantitative analysis of gait in Parkinson patients: increased variability of stride length**. *J Neurol Sci*. 1990; **98**(1): 91–7. [PubMed Abstract](#) | [Publisher Full Text](#)
- Hausdorff JM, Cudkowicz ME, Firtion R, et al.: **Gait variability and basal ganglia disorders: stride-to-stride variations of gait cycle timing in Parkinson's disease and Huntington's disease**. *Mov Disord*. 1998; **13**(3): 428–37. [PubMed Abstract](#) | [Publisher Full Text](#)
- Morris ME, Iansek R, Matyas TA, et al.: **The pathogenesis of gait hypokinesia in Parkinson's disease**. *Brain*. 1994; **117**(Pt 5): 1169–81. [PubMed Abstract](#) | [Publisher Full Text](#)
- O'Sullivan JD, Said CM, Dillon LC, et al.: **Gait analysis in patients with Parkinson's disease and motor fluctuations: influence of levodopa and comparison with other measures of motor function**. *Mov Disord*. 1998; **13**(6): 900–6. [PubMed Abstract](#) | [Publisher Full Text](#)
- Zgaljardic DJ, Borod JC, Foldi NS, et al.: **An Examination of Executive Dysfunction Associated with Frontostriatal Circuitry in Parkinson's Disease**. *J Clin Exp Neuropsychol*. 2006; **28**(7): 1127–44. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Chaudhuri KR, Odin P, Antonini A, et al.: **Parkinson's disease: the non-motor issues**. *Parkinsonism Relat Disord*. 2011; **17**(10): 717–23. [PubMed Abstract](#) | [Publisher Full Text](#)
- Mollenhauer B, Rochester L, Chen-Plotkin A, et al.: **What Can Biomarkers Tell Us About Cognition in Parkinson's Disease?** *Mov Disord*. 2014; **29**(5): 622–33. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Owen AM, James M, Leigh PN, et al.: **Fronto-striatal cognitive deficits at different stages of Parkinson's disease**. *Brain*. 1992; **115**(Pt 6): 1727–51. [PubMed Abstract](#) | [Publisher Full Text](#)
- Cools R, Barker RA, Sahakian BJ, et al.: **Enhanced or impaired cognitive function in Parkinson's disease as a function of dopaminergic medication and task demands**. *Cereb Cortex*. 2001; **11**(12): 1136–43. [PubMed Abstract](#) | [Publisher Full Text](#)
- Bokura H, Yamaguchi S, Kobayashi S: **Event-related potentials for response inhibition in Parkinson's disease**. *Neuropsychologia*. 2005; **43**(6): 967–75. [PubMed Abstract](#) | [Publisher Full Text](#)
- Cools R, Barker RA, Sahakian BJ, et al.: **L-Dopa medication remediates**

- cognitive inflexibility, but increases impulsivity in patients with Parkinson's disease.** *Neuropsychologia*. 2003; **41**(11): 1431–41.
[PubMed Abstract](#) | [Publisher Full Text](#)
18. Middleton FA, Strick PL: **A revised Neuroanatomy of Frontal-Subcortical Circuits.** In: Lichten DG Cummings JL editors. *Frontal-Subcortical Circuits in Psychiatric and Neurological Disorders*. New York: The Guildford Press; 2000; 44–58.
[Reference Source](#)
 19. Kelly VE, Johnson CO, McGough EL, et al.: **Association of cognitive domains with postural instability/gait disturbance in Parkinson's disease.** *Parkinsonism Relat Disord*. 2015; **21**(7): 692–7.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 20. Verbaan D, Marinus J, Visser M, et al.: **Cognitive impairment in Parkinson's disease.** *J Neurol Neurosurg Psychiatry*. 2007; **78**(11): 1182–7.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 21. Poletti M, Frosini D, Pagni C, et al.: **Mild cognitive impairment and cognitive-motor relationships in newly diagnosed drug-naive patients with Parkinson's disease.** *J Neurol Neurosurg Psychiatry*. 2012; **83**(6): 601–6.
[PubMed Abstract](#) | [Publisher Full Text](#)
 22. Williams-Gray CH, Foltynie T, Brayne CEG, et al.: **Evolution of cognitive dysfunction in an incident Parkinson's disease cohort.** *Brain*. 2007; **130** (Pt 7): 1787–98.
[PubMed Abstract](#) | [Publisher Full Text](#)
 23. Morris R, Lord S, Lawson RA, et al.: **Gait Rather Than Cognition Predicts Decline in Specific Cognitive Domains in Early Parkinson's Disease.** *J Gerontol A Biol Sci Med Sci*. 2017; **72**(12): 1656–1662.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 24. Beauchet O, Anweiler C, Callisaya ML, et al.: **Poor Gait Performance and Prediction of Dementia: Results From a Meta-Analysis.** *J Am Med Dir Assoc*. 2016; **17**(6): 482–90.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 25. Balasubramanian CK, Clark DJ, Fox EJ: **Walking Adaptability after a Stroke and Its Assessment in Clinical Settings.** *Stroke Res Treat*. 2014; **2014**: 591013.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 26. Islam A, Alcock L, Nazarpour K, et al.: **Effect of Parkinson's disease and two therapeutic interventions on muscle activity during walking: a systematic review.** *NPJ Parkinsons Dis*. 2020; **6**(1): 22.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 27. Henneman E, Somjen G, Carpenter DO: **Functional significance of cell size in spinal motoneurons.** *J Neurophysiol*. 1965; **28**: 560–80.
[PubMed Abstract](#) | [Publisher Full Text](#)
 28. Al-Yahya E, Dawes H, Smith L, et al.: **Cognitive motor interference while walking: A systematic review and meta-analysis.** *Neurosci Biobehav Rev*. 2011; **35**(3): 715–728.
[PubMed Abstract](#) | [Publisher Full Text](#)
 29. Kelly VE, Eusterbrock AJ, Shumway-Cook A: **A review of dual-task walking deficits in people with Parkinson's disease: Motor and cognitive contributions, mechanisms, and clinical implications.** *Parkinsons Dis*. 2012; **2012**: 918719.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 30. Li KZH, Lindenberger U: **Relations between aging sensory/sensorimotor and cognitive functions.** *Neurosci Biobehav Rev*. 2002; **26**(7): 777–83.
[PubMed Abstract](#) | [Publisher Full Text](#)
 31. Marusic U, Kavcic V, Giordani B, et al.: **Computerized spatial navigation training during 14 days of bed rest in healthy older adult men: Effect on gait performance.** *Psychol Aging*. 2015; **30**(2): 334–40.
[PubMed Abstract](#) | [Publisher Full Text](#)
 32. Wollesen B, Scrivener K, Soles K, et al.: **Dual-task walking performance in older persons with hearing impairment: Implications for interventions from a preliminary observational study.** *Ear Hear*. 2018; **39**(2): 337–43.
[PubMed Abstract](#) | [Publisher Full Text](#)
 33. Pradhan SD, Brewer BR, Carvell GE, et al.: **Assessment of fine motor control in individuals with Parkinson's disease using force tracking with a secondary cognitive task.** *J Neurol Phys Ther*. 2010; **34**(1): 32–40.
[PubMed Abstract](#) | [Publisher Full Text](#)
 34. Dideriksen JL, Gallego JA, Holobar A, et al.: **One central oscillatory drive is compatible with experimental motor unit behaviour in essential and Parkinsonian tremor.** *J Neural Eng*. 2015; **12**(4): 046019.
[PubMed Abstract](#) | [Publisher Full Text](#)
 35. Nocera JR, Roemmich R, Elrod J, et al.: **Effects of cognitive task on gait initiation in Parkinson disease: evidence of motor prioritization?** *J Rehabil Res Dev*. 2013; **50**(5): 699–708.
[PubMed Abstract](#) | [Publisher Full Text](#)
 36. Yogeve-Seligmann G, Rotem-Gallili Y, Mirelman A, et al.: **How does explicit prioritization alter walking during dual-task performance? Effects of age and sex on gait speed and variability.** *Phys Ther*. 2010; **90**(2): 177–86.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 37. Brown LA, de Bruin N, Doan JB, et al.: **Novel Challenges to Gait in Parkinson's Disease: The Effect of Concurrent Music in Single- and Dual-Task Contexts.** *Arch Phys Med Rehabil*. 2009; **90**(9): 1578–83.
[PubMed Abstract](#) | [Publisher Full Text](#)
 38. Galletly R, Brauer SG: **Does the type of concurrent task affect preferred and cued gait in people with Parkinson's disease?** *Aust J Physiother*. 2005; **51**(3): 175–80.
[PubMed Abstract](#) | [Publisher Full Text](#)
 39. Lord S, Rochester L, Hetherington V, et al.: **Executive dysfunction and attention contribute to gait interference in 'off' state Parkinson's Disease.** *Gait Posture*. 2010; **31**(2): 169–74.
[PubMed Abstract](#) | [Publisher Full Text](#)
 40. O'Shea S, Morris ME, Ianssek R: **Dual Task Interference During Gait in People With Parkinson Disease: Effects of Motor Versus Cognitive Secondary Tasks.** *Phys Ther*. 2002; **82**(9): 888–97.
[PubMed Abstract](#) | [Publisher Full Text](#)
 41. Plotnik M, Dagan Y, Gurevich T, et al.: **Effects of cognitive function on gait and dual tasking abilities in patients with Parkinson's disease suffering from motor response fluctuations.** *Exp Brain Res*. 2011; **208**(2): 169–79.
[PubMed Abstract](#) | [Publisher Full Text](#)
 42. Rochester L, Hetherington V, Jones D, et al.: **Attending to the task: interference effects of functional tasks on walking in Parkinson's disease and the roles of cognition, depression, fatigue, and balance.** *Arch Phys Med Rehabil*. 2004; **85**(10): 1578–85.
[PubMed Abstract](#) | [Publisher Full Text](#)
 43. Plotnik M, Giladi N, Hausdorff JM: **Bilateral coordination of gait and Parkinson's disease: the effects of dual tasking.** *J Neurol Neurosurg Psychiatry*. 2009; **80**(3): 347–50.
[PubMed Abstract](#) | [Publisher Full Text](#)
 44. Yogeve G, Plotnik M, Peretz C, et al.: **Gait asymmetry in patients with Parkinson's disease and elderly fallers: when does the bilateral coordination of gait require attention?** *Exp Brain Res*. 2006; **177**(3): 336–46.
[PubMed Abstract](#) | [Publisher Full Text](#)
 45. Rahman S, Griffin HJ, Quinn NP, et al.: **The factors that induce or overcome freezing of gait in Parkinson's disease.** *Behav Neurol*. 2008; **19**(3): 127–36.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 46. Cowie D, Limousin P, Peters A, et al.: **Doorway-provoked freezing of gait in Parkinson's disease.** *Mov Disord*. 2012; **27**(4): 492–9.
[PubMed Abstract](#) | [Publisher Full Text](#)
 47. Colcombe S, Kramer AF: **Fitness effects on the cognitive function of older adults: A meta-analytic study.** *Psychol Sci*. 2003; **14**(2): 125–30.
[PubMed Abstract](#) | [Publisher Full Text](#)
 48. Logan D, Kiemel T, Dominici N, et al.: **The many roles of vision during walking.** *Exp Brain Res*. 2010; **206**(3): 337–50.
[PubMed Abstract](#) | [Publisher Full Text](#)
 49. Mahoney JR, Verghese J: **Visual-Somatosensory Integration and Quantitative Gait Performance in Aging.** *Front Aging Neurosci*. 2018; **10**: 377.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 50. Rosano C, Studenski SA, Aizenstein HJ, et al.: **Slower gait, slower information processing and smaller prefrontal area in older adults.** *Age Ageing*. 2012; **41**(1): 58–64.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 51. Gramann K, Gwin JT, Ferris DP, et al.: **Cognition in action: Imaging brain/body dynamics in mobile humans.** *Rev Neurosci*. 2011; **22**(6): 593–608.
[PubMed Abstract](#) | [Publisher Full Text](#)
 52. Gramann K, Ferris DP, Gwin J, et al.: **Imaging natural cognition in action.** *Int J Psychophysiol*. 2014; **91**(1): 22–9.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 53. Makeig S, Gramann K, Jung TP, et al.: **Linking brain, mind and behavior.** *Int J Psychophysiol*. 2009; **73**(2): 95–100.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 54. Chan AW, Tetzlaff JM, Gatzsche PC, et al.: **SPiRiT 2013 explanation and elaboration: guidance for protocols of clinical trials.** *BMJ*. 2013; **346**: e7586.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
 55. Nasreddine ZS, Phillips NA, Bédirian V, et al.: **The Montreal Cognitive Assessment, MoCA: A Brief Screening Tool For Mild Cognitive Impairment.** *J Am Geriatr Soc*. 2005; **53**(4): 695–9.
[PubMed Abstract](#) | [Publisher Full Text](#)
 56. Guralnik JM, Simonsick EM, Ferrucci L, et al.: **A short physical performance battery assessing lower extremity function: association with self-reported disability and prediction of mortality and nursing home admission.** *J Gerontol*. 1994; **49**(2): M85–94.
[PubMed Abstract](#) | [Publisher Full Text](#)
 57. Hoehn MM, Yahr MD: **Parkinsonism: onset, progression and mortality.** *Neurology*. 1967; **17**(5): 427–42.
[PubMed Abstract](#) | [Publisher Full Text](#)
 58. Bristow T, Jih CS, Slabich A, et al.: **Standardization and adult norms for the sequential subtracting tasks of serial 3's and 7's.** *Appl Neuropsychol Adult*. 2016; **23**(5): 372–8.
[PubMed Abstract](#) | [Publisher Full Text](#)
 59. Holobar A, Farina D: **Blind source identification from the multichannel surface electromyogram.** *Physiol Meas*. 2014; **35**(7): R143–65.
[PubMed Abstract](#) | [Publisher Full Text](#)
 60. Oostenveld R, Praamstra P: **The five percent electrode system for high-resolution EEG and ERP measurements.** *Clin Neurophysiol*. 2001; **112**(4): 713–9.
[PubMed Abstract](#) | [Publisher Full Text](#)

61. Seeber M, Scherer R, Wagner J, et al.: **EEG beta suppression and low gamma modulation are different elements of human upright walking.** *Front Hum Neurosci.* 2014; **8**: 485.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
62. Jenkinson N, Brown P: **New insights into the relationship between dopamine, beta oscillations and motor function.** *Trends Neurosci.* 2011; **34**(12): 611–8.
[PubMed Abstract](#) | [Publisher Full Text](#)
63. Kempen GJM, Yardley L, van Haastregt JC, et al.: **The Short FES-I: a shortened version of the falls efficacy scale-international to assess fear of falling.** *Age Ageing.* 2008; **37**(1): 45–50.
[PubMed Abstract](#) | [Publisher Full Text](#)
64. Kempen GI, Todd CJ, van Haastregt JC, et al.: **Cross-cultural validation of the Falls Efficacy Scale International (FES-I) in older people: Results from Germany, the Netherlands and the UK were satisfactory.** *Disabil Rehabil.* 2007; **29**(2): 155–62.
[PubMed Abstract](#) | [Publisher Full Text](#)
65. Yardley L, Beyer N, Hauer K, et al.: **Development and initial validation of the Falls Efficacy Scale-International (FES-I).** *Age Ageing.* 2005; **34**(6): 614–9.
[PubMed Abstract](#) | [Publisher Full Text](#)
66. Nieuwboer A, Rochester L, Herman T, et al.: **Reliability of the new freezing of gait questionnaire: Agreement between patients with Parkinson's disease and their carers.** *Gait Posture.* 2009; **30**(4): 459–63.
[PubMed Abstract](#) | [Publisher Full Text](#)
67. Lamb SE, Jørstad-Stein EC, Hauer K, et al.: **Development of a common outcome data set for fall injury prevention trials: the Prevention of Falls Network Europe consensus.** *J Am Geriatr Soc.* 2005; **53**(9): 1618–22.
[PubMed Abstract](#) | [Publisher Full Text](#)
68. Lord SR, Sherrington C, Menz HB, et al.: **Falls in Older People: Risk Factors and Strategies for Prevention.** 2nd edition. New York, USA: Cambridge University Press, 2011.
69. Goetz CG, Fahn S, Martinez-Martin P, et al.: **Movement Disorder Society-sponsored revision of the Unified Parkinson's Disease Rating Scale (MDS-UPDRS): Process, format, and clinimetric testing plan.** *Mov Disord.* 2007; **22**(1): 41–7.
[PubMed Abstract](#) | [Publisher Full Text](#)
70. Santangelo G, Siciliano M, Pedone R, et al.: **Normative data for the Montreal Cognitive Assessment in an Italian population sample.** *Neurol Sci.* 2015; **36**(4): 585–91.
[PubMed Abstract](#) | [Publisher Full Text](#)
71. Potocnik J, Ovcar Stante K, Rakusa M: **The validity of the Montreal cognitive assessment (MoCA) for the screening of vascular cognitive impairment after ischemic stroke.** *Acta Neurol Belg.* 2020; **120**(3): 681–5.
[PubMed Abstract](#) | [Publisher Full Text](#)
72. Reitan RM: **Validity of the Trail Making Test as an Indicator of Organic Brain Damage.** *Percept Mot Skills.* 1958; **8**(3): 271–6.
[Publisher Full Text](#)
73. Mayer JD, Gaschke YN: **The Experience and Meta-Experience of Mood.** *J Pers Soc Psychol.* 1988; **55**(1): 102–11.
[PubMed Abstract](#) | [Publisher Full Text](#)
74. Beck AT, Ward CH, Mendelson M, et al.: **An inventory for measuring depression.** *Arch Gen Psychiatry.* 1961; **4**: 561–71.
[PubMed Abstract](#) | [Publisher Full Text](#)
75. Wollesen B, Mattes K, Schulz S, et al.: **Effects of Dual-Task Management and Resistance Training on Gait Performance in Older Individuals: A Randomized Controlled Trial.** *Front Aging Neurosci.* 2017; **9**: 415.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
76. Hart SG, Staveland LE: **Development of NASA-TLX (Task Load Index): Results of Empirical and Theoretical Research.** *Adv Psychol.* 1988; **52**(C): 139–83.
[Publisher Full Text](#)
77. Hennah C, Ellis G, Doumas M: **Dual task walking in healthy aging: Effects of narrow and wide walking paths.** *PLoS One.* 2021; **16**(12): e0261647.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
78. Mordkoff JT: **A Simple Method for Removing Bias From a Popular Measure of Standardized Effect Size: Adjusted Partial Eta Squared.** *Adv Methods Pract Psychol Sci.* 2019; **2**(3): 228–32.
[Publisher Full Text](#)
79. Shaikh AG, Zee DS, Taube J, et al.: **Visual-vestibular interactions.** *Multisensory Percept From Lab to Clin.* 2020; 201–19.
[Publisher Full Text](#)
80. Peskar M: **Demographic and General Health Questionnaire.** *Zenodo.* 2023. <http://www.doi.org/10.5281/zenodo.7768228>

Open Peer Review

Current Peer Review Status:  

Version 2

Reviewer Report 08 September 2023

<https://doi.org/10.21956/openreseurope.17851.r34667>

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 **Nicola J Ray** 

Manchester Metropolitan University, Manchester, England, UK

The authors have much improved the organisation of their proposal. I still have some comments:

Minor: Primary and secondary outcomes has a special meaning that in clinical research, and the term is used incorrectly here. Primary/secondary study outcomes are those that arise from any primary/secondary study aim. Rather, in the current proposal, the authors have listed their full testing battery under Primary and Secondary outcomes.

Minor: Reviewer 2 suggested not to use 'Patients' to describe your 'Participants'. I agree with this and suggest this wording is changed throughout.

Minor: In my previous report I identified an issue with the screening procedure. There remains an issue with this:

Inclusion criteria are PD diagnosis, PD duration, Dopamine Transporter Scan (DATSCAN), diagnosed akinetic-rigid or mixed phenotype, Hoehn and Yahr scale, MoCA , SPPB score, age, no color blindness, independent living. Additional exclusions are (3) MoCA score < 24 points, (4) severe disabling tremor, (5) recurrent falls, impaired vision.

The first testing session lasts 1.5 hours. Does this mean more tests are planned than are included in the exclusion criteria? This would mean that ineligible participants are taking part in research elements of the study. Will the authors discard this data? It is ethically dubious to ask a person with Parkinson's to take part in a 1.5 hour testing session for a study that their data cannot contribute to. Many of the tests included in their inclusion criteria are likely done routinely in the clinic? Additionally, the questionnaires are all self-report, and could be sent to people's homes before asking them to come in for testing?

I understand the testing needs to take place over two visits since the battery is extensive, but screening is usually done first and no tests that do not contribute to screening should be completed in participants who are not eligible. If I have misunderstood the proposal, and only

screening tests will be done on M1, I suggest the proposal lists explicitly 'M1 testing battery' and 'M2 testing battery'.

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Parkinson's disease

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Version 1

Reviewer Report 10 July 2023

<https://doi.org/10.21956/openreseurope.17047.r32804>

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Nicola J Ray 

Manchester Metropolitan University, Manchester, England, UK

The authors propose to use a MoBI approach to investigate cognitive and motor interactions in People with Parkinson's compared with controls. Participants will perform motor tasks under ST and DT conditions.

I believe the presentation of this study requires closer attention. There are methods presented in the intro, for example. The description of the primary outcomes are very vague, and the hypotheses presented would only confirm findings already known in the literature. What does the MoBI approach add, and how will the authors analyse the data in order to add new knowledge to the field.

The study design also requires closer attention. The different cognitive tasks will present a larger problem than acknowledged in comparing across tasks. I suggest to use a well practised task for all.

Additionally, some of these tasks will interfere with ability to perform the motor tasks for reasons not related to CMI. For example, requiring people perform a visual task during walking will change their ability to use visual information for the walking task itself. This will deteriorate performance during DT conditions, but not because of cognitive interference.

There also seems to be two levels of eligibility - one for M1 and the other for M2. However no information is provided to understand why this is.

Is the rationale for, and objectives of, the study clearly described?

Partly

Is the study design appropriate for the research question?

Partly

Are sufficient details of the methods provided to allow replication by others?

No

Are the datasets clearly presented in a useable and accessible format?

Not applicable

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Parkinson's disease

I confirm that I have read this submission and believe that I have an appropriate level of expertise to state that I do not consider it to be of an acceptable scientific standard, for reasons outlined above.

Author Response 18 Aug 2023

Uros Marusic

\\\\\\\\\\\\\\\\The authors propose to use a MoBI approach to investigate cognitive and motor interactions in People with Parkinson's compared with controls. Participants will perform motor tasks under ST and DT conditions.

I believe the presentation of this study requires closer attention. There are methods presented in the intro, for example. The description of the primary outcomes are very vague, and the hypotheses presented would only confirm findings already known in the literature. What does the MoBI approach add, and how will the authors analyse the data in order to add new knowledge to the field. /////

Thank you for the remark. As suggested, we reorganized the introduction section and moved the methods-oriented parts to the methods. As stated under the *Data streaming and management* chapter, our recordings of the hdEEG, hdEMG, and full-body kinematics will be time-synchronized while a participant will perform an overground walking task. Time synchronization will allow us to investigate the brain-muscle cross-talk for the first time in such a population in addition to unraveling the physiological dynamics in each modality separately. Because the technology and the whole MoBI field are relatively new (Gramann et al., 2010, doi: 10.3389/fnhum.2010.00202) such cross-modal investigations hold promise to better understanding the fundamental surface physiology and their potential compensatory mechanisms in PD patients during tasks resembling everyday functioning. It has been shown that PD patients demonstrate a hypersynchrony in the beta oscillatory range which is related to bradykinesia (Jenkinson & Brown, 2011; doi:10.1016/j.tins.2011.09.00). We made modifications to the *Brain Activity* chapter, where

we highlight the novel approach of performing a gait-phase modulations analysis (Seeber et al. 2014, doi: 10.3389/fnhum.2014.00485) that has not yet been performed in PD patients and would allow investigating how and when is beta modulated relative to the gait cycle. Due to its association with symptom manifestation, the outcomes of such analysis carry great functional significance.

\\\\\\\\\\\\The study design also requires closer attention. The different cognitive tasks will present a larger problem than acknowledged in comparing across tasks. I suggest to use a well practised task for all. //////////////

We appreciate you pointing out this aspect, as we spent quite some time discussing this issue ourselves. We wrote in the manuscript: *»We are aware that the comparison between the DT conditions of the three motor activities is complicated by the use of different cognitive tasks, yet this decision was motivated to avoid extensive learning effects and increasing frustration during the measurement day.«* We are fully aware that a direct comparison across tasks will not be possible, however after a thorough consideration we decided upon taking this tradeoff instead of having participants perform thousands of trial repetitions of a single task which would promote a massive learning effect and automaticity. Such experimental (non)manipulation would in addition mean a step away from the ecological validity, which is one of the primary intents in the MoBI field. Being aware of this, we never intended on comparing across motor activities, that is overground walking vs balancing vs isometric knee extensions, but rather within each motor activity, meaning single-task walking vs dual-task walking, for instance. The main comparison will always be performed within a certain motor activity experiment. Examining the performance across all motor activities will, however, allow for making general claims, such as cognitive-motor interference was/wasn't observed in the respective motor activity/ies. For the balance and isometric experiments, the cognitive tasks we decided to use were derived from our previous work (for instance: Marušič et al., 2019, DOI: [10.1186/s11556-018-0209-7](https://doi.org/10.1186/s11556-018-0209-7) ; Peskar et al. 2023, doi: [10.1016/j.biopsycho.2023.108543](https://doi.org/10.1016/j.biopsycho.2023.108543)) and our research interests to better understand how the executive function as a complex and advanced cognitive system support our voluntary actions. For the walking activity, the visual secondary task was used because visuo-motor coordination holds great ecological validity. However, because exploring visual processing while overground walking is rather challenging due to the movement itself and controlling for the constant stimulus distance, evidence in this cross-domain remains rather scarce. For this reason, we decided to use a relatively simple visual working memory task (visual discrimination) not to overload the visual system. We believe our findings will contribute significantly to this as-of-yet vastly unexplored field. [Please consider also the answer to your next comment below.] Additionally, one of our goals (which exceeds the present protocol) is to compare the walking data in PD patients and in patients with hearing impairment (see protocol Wunderlich et al 2021, doi: 10.3389/fnagi.2021.773287). Lastly, to avoid recording adaptation to the task, all tasks will be well-trained during the practice run prior to recording.

\\\\\\\\\\\\Additionally, some of these tasks will interfere with ability to perform the motor tasks for reasons not related to CMI. For example, requiring people perform a visual task during walking will change their ability to use visual information for the walking task itself. This will deteriorate performance during DT conditions, but not because of cognitive

interference. //////////////

»Cognitive-motor interference (CMI) is evident when simultaneous performance of a cognitive task and a motor task results in deterioration in performance in one or both of the tasks, relative to performance of each task separately (Plummer et al., 2013 (doi: [10.1016/j.apmr.2013.08.002](https://doi.org/10.1016/j.apmr.2013.08.002))).« According to this definition, CMI as a consequence of the limited attentional resources is precisely *the reason* why the visual task during walking would result in deteriorated gait performance. Not only it offers a theoretical model for understanding the mechanism behind performance decrements, but, in addition, tackles the functional capacities of PD patients in real-world situations. For instance, in reality, on a busy pavement surrounded by heavy traffic, the motor actions have to be continuously adapted with respect to the ever-changing environment. Our perception and appropriate reactions to the environment will ensure our safety (such as reacting to or withdrawing action in response to passing cars or traffic lights). Here, vision arguably demonstrates the paramount importance to guide the translations of perception to motor actions. The ability to retain independence in PD patients can be legitimately assessed using a visual cognitive-motor dual-task walking experiment. In practical terms, however, the spectacle frame with LED lights attached to each end was custom-made with the awareness to avoid the coverage of a significant portion of the visual field, from which important information about the environment (and safety) could be detected. Also, because the frame (together with the lights) is attached to a participant, they are free to move the head around in all directions and use all visual angles to uncover potential occlusion to the walking lane, if needed. When using MoBI, we are translating the overly-controlled laboratory settings to more ecologically valid conditions, and through these lenses, the proposed paradigm is at the frontier of what technology currently allows. The outcomes of the present study hold great translational promise.

\\\\\\\\\\\\There also seems to be two levels of eligibility - one for M1 and the other for M2. However no information is provided to understand why this is. //////////////

Under the *Recruitment Process* we wrote »*The enrolled patients and healthy adults will undergo a comprehensive screening session including motor and neurocognitive assessment (Measurement Day 1 - M1). Having passed the inclusion/exclusion criteria confirmed on the M1, they will finally be invited to participate in Measurement Day 2 (M2).*« M1 constitutes the first measurement day during which all the secondary outcomes are collected; these outcomes at the same time confirm either the inclusion or exclusion criteria. Measurement day M2 then constitutes the 3 motor activity experiments in combination with hdEEG, hdEG, and full-body kinematics. If you suggest to add additional information to make the procedure clearer, we are happy to do so.

Competing Interests: No competing interests were disclosed.

Reviewer Report 13 June 2023

<https://doi.org/10.21956/openreseurope.17047.r31835>

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Breiffni Leavy 

Department of Neurobiology, Karolinska Institutet, Huddinge, Sweden

This protocol describes the planned test conditions for a cross-sectional investigation of the early PD disease-specific changes in motor-cognitive performance, assessed during three test conditions, compared to healthy age-matched controls.

Background and rationale are clearly outlined, relevant and in line with the current evidence in the field.

Methods and test protocols are clearly outlined and described in detail with an ambitious and widely encompassing list of outcome measures. I cannot see anything lacking.

In summary, this is a clearly defined and scientifically robust plan to investigate motor-cognitive performance in mild PD.

Suggestion: One minor semantic alteration I suggest to bring the test in line with use of a person-centered language is not to refer to 'PD patients' (occurs frequently in the text) but people with PD. People are most often not patients, and not in the context of cross-sectional research.

Is the rationale for, and objectives of, the study clearly described?

Yes

Is the study design appropriate for the research question?

Yes

Are sufficient details of the methods provided to allow replication by others?

Yes

Are the datasets clearly presented in a useable and accessible format?

Not applicable

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Rehabilitation for Parkinson's disease

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
