

BMJ Open Video game therapy on mobility and dual tasking in multiple sclerosis: study protocol for a randomised controlled trial

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To cite: Baroni A, Fregna G, Milani G, *et al.* Video game therapy on mobility and dual tasking in multiple sclerosis: study protocol for a randomised controlled trial. *BMJ Open* 2021;**11**:e052005. doi:10.1136/bmjopen-2021-052005

► Prepublication history and additional supplemental material for this paper are available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2021-052005>).

Received 02 April 2021

Accepted 23 September 2021



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ABSTRACT

Introduction Multiple sclerosis (MS) is one of the major causes of disability in young adults and affects mobility, compromising daily living activities and participation in social life. Cognitive domain is also frequently impaired in people with MS (PwMS), particularly the capacity to perform dual-task activities. Impaired cognitive processing abilities need to be treated, and motor and cognitive aspects need to be considered together. Recently, video game therapy (VGT) has been used in rehabilitation to improve motor outcomes and cognitive processing speed. The aim of this study is to test the efficacy of commercially available VGT on mobility and dual tasking in PwMS compared with standardised balance platform training (BPT).

Methods and analysis This will be a parallel-assignment, double-blinded, randomised control trial. Forty-eight (24 per arm) PwMS with Expanded Disability Status Scale 4–5.5 will be randomly assigned to receive 1 hour training session over 4 weeks (three sessions/week) of either: (1) VGT on commercial video game console to train balance and mobility-related activities or (2) BPT to perform balance, postural stability and weight-shifting exercises with and without visual feedback. The same assessor will evaluate outcome measures at points: before and after the 12 training sessions and at 3 months of follow-up. The primary outcome will be functional mobility, assessed by the Timed Up and Go test. We will also evaluate gait, risk of fall, fatigue and health-related quality of life as well as cognitive and psychological aspects (depression, anxiety and attentional performance) and stability through posturographic evaluation. Dual-tasking assessment will be performed combining posturographic and neuropsychological tests. Data analysis will be performed to compare the efficacy of the two treatments.

Ethics and dissemination Ethical approval have been granted from the local Ethics Committee. Study results will be communicated through high-quality journals and national and international conferences.

Trial registration number NCT03353974

INTRODUCTION

Multiple sclerosis (MS) is characterised by multifocal inflammatory demyelinated plaques distributed over time and space within

Strengths and limitations of this study

- This trial will use objective evaluation as a posturographic assessment to test balance and dual tasking.
- Motor and cognitive outcomes will be assessed separately and in association.
- Results from this trial will need a larger sample to be confirmed.
- This study protocol includes only young subjects (aged under 60) with good comprehensive functioning.
- This study does not analyse long-term effects (over 3 months).

the central nervous system (CNS). It affects approximately 1.3 million people worldwide and is a major cause of chronic neurological disability in young adults aged 18–50 years.^{1,2} Deficits in balance control and cognition are prevalent impairments in people with MS (PwMS),³ even at an early stage and without clinical disability.⁴ Previous studies have reported that 30%–63% of PwMS experience a fall event between 1 and 12 months since the onset of the disease.⁵ Balance maintenance is a complex task that depends on the continuous flow of proprioceptive information from the muscles, tendons, joints, skin, vestibular and visual systems toward the CNS.³ In PwMS, the extended damage caused in the CNS leads to a decreased ability in integrating the afferent proprioceptive information, thus negatively influencing postural response and the capability to maintain balance safely.^{5,6} Balance impairment can consistently limit the activities of daily living and the active participation in social life.

Cognitive impairment is various among PwMS, with current prevalence rates ranging from 43% to 70%.⁷ The following cognitive domains are frequently impaired: memory



processing speed, attention/concentration and executive functioning.⁸ Cognitive impairments are associated with reduced functional status in MS. They have a deleterious impact on the individual's personal, occupational and social functioning and the comprehensive quality of life.⁹

The role of cognitive functioning on motor performance and balance control is widely known.¹⁰ However, the effects of impaired cognitive processes on balance efficacy have not been extensively investigated in PwMS.¹¹ Besides, the comprehensive link between attention and motor-action has been supported by several studies.¹² Selective attention allows the execution of a correct motor response by selecting relevant information between the task and the distractors, and it is essential for action planning. Concerning the cognitive-motor interference, this relation is directly measurable by the dual-task cost,^{13 14} investigated in PwMS through the Stroop Test.¹⁵ Dual-task performance is the capacity to do two tasks simultaneously, particularly motor and cognitive tasks. The subject's attention is drawn to an external source of attention while the primary task is ongoing, resulting in cognitive-motor interactions.¹⁶ Concerning the constrained action hypothesis, this attentional change may lead motor systems to react automatically, thus increasing the performance effectiveness.¹⁷ The processing capacity required for doing dual-task activities may be affected by cognitive impairments.¹⁸ In subjects with neurological disorders, such as PwMS, the ability of doing a motor task simultaneously to a cognitive one is frequently affected.¹⁷

It is plausible that motor deficits enhance the cognitive demands necessary to execute functional movements, and the concurrent performance of tasks may exceed cognitive processing abilities.¹⁹ Considering the strong impact of dual tasking on activities of daily living (ADL),⁷ rehabilitative treatment does not have to consider the motor and cognitive aspects separately.

In recent years, active video games technologies have begun to be used as a treatment tool in rehabilitation given their low cost, high portability, off-the-shelf nature and ability to deliver engaging, high-repetitive, task-oriented, standardised, active learning therapies.²⁰ Moreover, PwMS defined gaming experience as fun, challenging and self-motivating, critical elements for successful motor learning.²¹ Active video games' multisensory feedback provided to patients may potentiate the use-dependent plasticity processes in the sensorimotor cortex, promoting functional recovery.²² Furthermore, combined training of cognitive and motor abilities in constantly changing virtual environments is particularly suited to address dual-tasking as required for the constantly changing situations of everyday life.²³

The evidence of interactive video game therapeutic exercises for improving balance and motor functions in PwMS were inconclusive, even if few studies showed a possible positive effect on balance and cognitive functions such as processing speed.²⁴⁻²⁷ Furthermore, patients' motivation seems to be capable of being increased during

active video game rehabilitation, allowing patients to exercise more consistently.²²

All previous studies did not investigate the effects on motor and cognitive functions simultaneously, particularly when compared with conventional instrumental balance training.

Aims

This study aims to test the efficacy of a commercially available video game therapy (VGT) (Xbox Kinect) on mobility and dual-tasking in ambulatory PwMS in comparison to a standardised balance platform training (BPT).

We hypothesise that augmented feedback during VGT, in terms of intensity and type, would activate the cognitive components of motor learning more effectively than BPT. Moreover, a challenging and engaging approach contributes to enhance treatment adherence and patient's satisfaction.

METHODS AND ANALYSIS

Study design and setting

This is a parallel-assignment, double-blinded, randomised control trial. The outcome assessor and the data analyst will be blinded to the group allocation of participants. PwMS who meet the inclusion criteria and provide written informed consent will be assigned to two treatments, the VGT or the BPT. The trial protocol has been registered on ClinicalTrials.gov.

The protocol of this clinical trial is reported following the Standard Protocol Items: Recommendations for Interventional Trials (SPIRIT) guidelines.²⁸ A SPIRIT checklist is available as an additional file (online supplemental additional file 1). Subjects will be recruited from the patients afferent to Outpatient Rehabilitation Clinic at University Hospital of Ferrara. Patients' recruitment started on 1 December 2017, and it is going to finish on 1 October 2022.

Selection criteria and recruitment of participants

People affected by MS will be included if they meet the following inclusion criteria:

- ▶ Men and women, aged 18–60 years;
- ▶ Diagnosis of MS (primary or secondary progressive, relapsing–remitting), without relapses in the preceding 3 months;
- ▶ Disability rate defined by Expanded Disability Status Scale (EDSS) score from 4 to 5.5²⁹;
- ▶ Balance impairments with increased fall risk, defined as Timed Up and Go (TUG) >8.4 s³⁰;
- ▶ Mini-Mental Status Examination score $\geq 24/30$.³¹

Exclusion criteria will be:

- ▶ Other (neurological) conditions that may affect motor function;
- ▶ Visual impairments (daltonism and visual acuity deficit);
- ▶ Medical conditions might interfere with the ability to complete the study protocol safely.

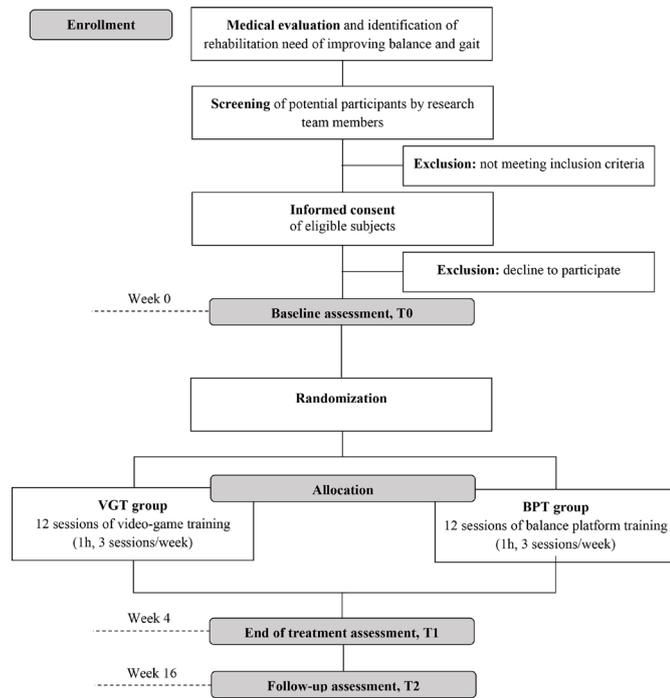


Figure 1 Consolidated Standards of Reporting Trials flow diagram of the study. BPT, balance platform training; VGT, video game therapy.

During the first appointment, potential participants will be informed about all the study procedures and screened following the inclusion criteria, suppose they meet inclusion criteria and are interested in taking part in the study. In that case, the physician will give them a letter explaining study purpose and procedures, time commitments, risks, potential benefits, treatment alternatives, study staff contact information and the Consent Form. A copy of the Consent Form is available as an additional file (online supplemental additional file 2). In the following 3 days, the potential participant will be contacted and asked about its decision; if the subject decides to take part in the study, the research staff will give him/her an appointment for the consign to the signed Consent Form and for the baseline outcome measures, conducted by a physiotherapist. If the subject rejects participation in the study, the research staff remains available for further information. The total number of subjects screened for participation and the number of subjects who decline to participate will be recorded, according to the Consolidated Standards of Reporting Trials (CONSORT) guidelines (figure 1).³²

Randomisation and blinding

People meeting inclusion criteria who decided to participate will be assigned to one of the two treatment groups through a block randomisation approach (1:1 ratio). The randomisation scheme will be set up in permuted blocks of four to ensure a similar number of participants between groups. The randomisation scheme will be generated using the website <http://www.randomization.com> and managed by an external administrator to the research group to prevent selection bias. The outcome

assessor will be blinded about the subject's group allocation. All outcome data and assignment groups will be organised in different data sets to maintain the blinding during data analysis. The privacy of the participants and their medical records will be guaranteed by treating the data according to the Italian Law n. 196/2003, to the 'Safe Harbor Act' (2000/520/CE) and to the 'European Union Data Protection Directive (95/46/EC 24 October 1995)'.

Intervention

All participants will receive twelve 1-hour training sessions over 4 weeks, resulting in a three sessions/week scheme. To manage possible absence lasting one or more treatment sessions, a potential window of 5 weeks will be set to ensure the achievement of all 12 sessions. Subjects who miss a training session will be contacted by phone to determine the absence's reasons and maintain adherence to following treatment sessions. Subjects who miss more than three consecutive treatment sessions will be excluded from the study.

Every training session consists of about 50 min of exercise and about 10 min of mobility and flexibility activity to prevent muscle soreness due to movement. All interventions will be delivered at the Rehabilitation Clinic of the University Hospital of Ferrara.

VGT

VGT will be delivered with a commercial video game console, Kinect for Xbox 360 game system (Microsoft Corporation, Redmond, Washington, USA). Pre-selected games were chosen from 'Kinect Adventures!' and 'Kinect Sports' (Microsoft Game Studios) that encompassed a wide range of motor activities in a standing position. Specifically, balance and mobility-related motor tasks were trained, such as sidestepping, lateral weight shifting, jumping, walking (lateral, forward and backward), and arm goal reaching.

Kinect is a commercial device therefore calibration of the instrument on the individual patient is not provided. The physiotherapy treatment with this device will be set by the physiotherapist basing on clinical observations of the patient's characteristics, preferences and level of functioning.

A list of games will be tested according to the patient's abilities and preferences. In the following sessions, games will be proposed with a randomised practice approach. Progression proceeds over time according to the patients' motor and balance improvements. Each task will consist of 2–5 min of training and a rest period will be given if necessary. During sessions, the patients will be carefully supervised by a physiotherapist who monitored the patient's safety (eg, risk of falls, impulsive reactions). The physiotherapist will also give performance feedback and those provided by VGT: visual and augmented (knowledge of both results and performance).³³ Despite variability among treatment protocols available in the literature, our treatment dosage in terms of number of

sessions and intervention duration is in line with other studies on the efficacy of exergame in people with neurological disorders.^{25 34–36}

BPT

Balance/rebalancing, postural stability and weight-shifting exercises with and without visual feedback will be administered using a Balance Platform (Biodex Medical Systems, Shirley, New York, USA) that had been previously tested in patients with MS.³⁷ Each task will be trained for about 2–3 min, followed by a rest period when necessary. During the first session, the tasks will be set to an ‘entry-level’. The exercise progression will be adjusted over time according to the patient’s capabilities (intermediate and difficult level). BPT offers visual feedback about reaching goals (augmented feedback). The physiotherapist will carefully supervise the patient and monitor his safety, providing additional external feedback during the activities.

Concomitant care and recommendations

All the subjects will be advised to not undertake other physical treatments until the end of the assessment period. Subjects will also be encouraged to not use the video game console at home for leisure to prevent confounding effects. It will be asked to patients to wear the same shoes and orthosis during all the outcome assessment and training sessions.

Intervention fidelity and monitoring of adverse events

All the interventions will be delivered by a physiotherapist with at least 5 years of experience in the treatment of PwMS, properly formed about VGT or BPT. Training sessions features and comments will be tracked in a precompiled form. Any unpredictable adverse events will

be recorded in the patient’s registry and the electronic study database. Their management will agree with the related hospital policies, with a referral for appropriate medical follow-up.

Outcome assessment and data collection

All the clinical evaluations will be performed at the Ferrara University Hospital by the same blinded assessor at the three time-point evaluations: (T0) baseline, before the first intervention; (T1) end of treatment, after the 12 therapy sessions; (T2) follow-up, 3 months after the end of treatment. Clinical and posturographic assessment will be delivered by a physiotherapist trained adequately about evaluation procedures. A neuropsychologist with years of experience in the assessment and treatment of PwMS will provide cognitive tests. A physician member of the research team will define the patient’s EDSS score. A team member will record the general demographic information (age, gender), comorbidities and medical history of every participant. A summary of the study plan is reported in [table 1](#).

Primary outcome: functional mobility

This function will be assessed by the TUG test, a reliable and valid performance-based measure of functional mobility.³⁰ The patient will be instructed to stand up from a chair, walk for 3 m, cross a line marked on the floor, turn around, walk back and sit down. The time used to complete the task is recorded using a chronometer. During the assessment, the subject can use any necessary gait aid (not physical assistance). This test will be the first performed during the assessment session to reduce variability due to the subject’s fatigue. The TUG test

Table 1 Schedule of enrolment, interventions and assessments

	Study period				
	Enrolment	Allocation	Post allocation	Close-out	
Time point	T-1		T0	T1	T2
Enrolment					
Eligibility screen	X				
Informed consent	X				
Allocation		X			
Interventions					
Video game therapy				↔	
Balance platform therapy				↔	
Assessments					
Primary outcome					
Timed Up and Go test			X	X	X
Secondary outcome					
Clinical measures and questionnaires			X	X	X
Cognitive and psychological assessment			X	X	X
Posturographic assessment			X	X	X

T-1 enrolment, T0 before treatment, T1 after treatment, T2 3-month follow-up.

will be repeated three times, and the mean value will be recorded.

Secondary outcome measures

Secondary outcomes will include: (1) clinical measures and questionnaires, (2) cognitive and psychological assessment and (3) posturographic assessment. All secondary outcome measures will be carried out in random order.

Clinical measures and questionnaires

- ▶ Dynamic Gait Index (DGI): gait, balance and risk of fall are measured using DGI. DGI will evaluate not only usual steady-state walking but also during a more challenging task (ie, cross obstacles, slalom). The subject will perform eight functional walking tests and score out of three (maximum total score is 24).³⁸
- ▶ Four Square Step Test (FSST): this timed test is intended to challenge the rapid change in direction while stepping forward, backward and sideways over a low obstacle. The faster the time measured to perform the task signifies a superior level of dynamic balance abilities. The minimal detectable change estimate for the FSST in MS is 4.6 s, and it was found to be a valid assessment tool in MS.³⁹
- ▶ Functional Reach Test (FRT): this test assesses the subject's stability by measuring the maximum distance an individual can reach forward while standing in a fixed position. A longer reaching distance indicates better postural control.⁴⁰
- ▶ Multiple Sclerosis Impact Scale—29 (MSIS-29): this health-rated quality of life questionnaire assesses the impact of MS on physical and psychological functions. It is formed by 29 items on ADL I and II: 20 about physical activity and 9 about psychological status. Each item can be scored with a value from 0 to 5; the total score is given by the sum of all the items and then is transformed in a range from 0 to 100.⁴¹
- ▶ Multiple Sclerosis Walking Scale—12 (MSWS-12): this questionnaire assesses the impact of MS on walking ability. It is formed by 12 items, asking the patient about his perception on gait speed, running, confidence ascending/descending stairs, balance and fatigue. The total score is obtained by the sum of the score of each item (0–5) and then transformed into a value from 0 to 100.⁴²
- ▶ Modified Fatigue Impact Scale (MFIS): this 21-item questionnaire assesses the perceived impact of fatigue on the subscale physical, cognitive and psychological functioning during the past 4 weeks. MFIS has been recommended for use by the Multiple Sclerosis Council for Clinical Practice Guidelines.^{43 44}

Cognitive and psychological assessment

- ▶ Beck Depression Inventory—Second Edition (BDI-II): this is a 21-item self-report measure that quantifies the severity of depression and over behavioural characteristics of depression.⁴⁵

- ▶ State Trait Anxiety Inventory (STAI-Y): this self-report questionnaire measures the presence and severity of current symptoms of anxiety and a generalised propensity to be anxious. There are two subscales: 20 items allocated to each of the State Anxiety (S-Anxiety) and Trait Anxiety (T-Anxiety).⁴⁶
- ▶ Test of Attentional Performance (T.A.P. V.2.3): attentional performance will be evaluated using a neuropsychological computer test. Errors, omissions and reaction times will be recorded as outcomes of performance. Three modules of the T.A.P. will be administered: Go-No Go subtest as it allows assessment of the specific ability of subjects to suppress undesired responses; alertness subtest that measures the simple reaction time in response to a visual stimulus and, divided attention subtest, that explored with 'dual-task' test the ability to attend simultaneously two stimuli (visual and acoustic) processed in parallel.⁴⁷
- ▶ Stroop Color-Word Test (SCWT): this neuropsychological test is used to assess the ability to inhibit cognitive interference when processing a specific stimulus feature impedes the simultaneous processing of a second stimulus attribute. Subjects are required to read three different tables as fast as possible. Two of them represent the 'congruous condition', in which participants are required to read names of colours printed in black ink and name different colour patches. Conversely, the third table represents the incongruent condition, in which participants are required to name the colour of the ink instead of reading the word.⁴⁸
- ▶ Symbol Digit Modalities Test (SDMT): this neuropsychological test quantifies cognitive processing speed. It consists of orally report the correct number corresponding to a symbol in a pseudorandom sequence of nine symbols as quickly as possible.⁴⁹

Posturographic assessment

Instrumented basic balance evaluation: Force platform measurements are routinely used as objective markers of subjects' balance ability.^{50 51} Several parameters can be extracted from the force platforms that correlate with balance ability and risk of falls in PwMS. During the instrumented tasks, subjects will be asked to stand on a force plate (BERTEC Model 4080–10, Bertec, Columbus, Ohio, USA) with arms parallel to their body. Each subject will undergo five repetitions (each lasting 60 s) of two tasks that will consist in standing with the eyes open and standing with the eyes closed. The movements of the centre of pressure (CoP) of the subjects will be recorded. A series of features will be extracted from the CoP traces to inform on different balance characteristics of each patient and the specific effect of each of the proposed therapies. Features that will be analysed include different measures of CoP displacement in the anteroposterior (AP) and mediolateral (ML) directions, the CoP path length (total, AP and ML) and the CoP average and maximum speed.³³

Dual-tasking assessment will be performed combining posturographic and neuropsychological tests: subjects will be asked to complete SCWT standing on the force platform, ensuring a similar condition to single-task SCWT. Head position will be fixed using a large panel fixed on the wall facing the force plate in order to prevent subjects lowering their head and distort the data. The ability to inhibit cognitive interferences during quiet and standing conditions will be compared.

Data management

Data analysis will be performed according to the research hypothesis mentioned. Stata Statistical Software (Release 13.1.: StataCorp LP) will be used for data analysis.

Sample size and power

The primary outcome of this study is to highlight differences in the time used to perform the TUG test between PwMS who underwent VGT and BPT. Our preliminary results from an unpublished pilot study (n=6) shown a VGT effect size of 0.93 in people with MS and EDSS <6. Given equal allocation between treatment and control arms, and using 80% power and alpha of 5%, we would need 40 subjects to complete the study. Conservatively, we expect a 10% rate of drop-out. Thus the sample size will be increased by 10% to 48 subjects (24 per arm).

Statistical analyses

Descriptive statistics (mean and 95% CI) will be reported before treatment, after treatment and at 3 months follow-up for all the selected variables (clinical, instrumental and questionnaires). Specifically, TUG changes after treatments will be considered as our primary endpoint, whereas changes of all the other outcome measures (DGI, FSST, FRT, MSIS-29, MSWS-12, MFIS, BDI-II, STAI-Y, SCWT, SDMT, T.A.P., posturographic variables) will be treated as secondary endpoints. Between-group differences will be explored with the Wilcoxon rank-sum test. Moreover, a repeated-measures analysis of variance (within-group factor: TIME; between-group factor: TREATMENT) will be conducted to detect the main effects for treatment and time for all the available outcomes. To calculate the effect size of both treatments, we will use Cohen's d.⁵² Results will be reported as mean and 95% CI. Significance will be recognised for $p < 0.05$.

Intention-to-treat

Every attempt will be made to avoid missing data through a careful check of self-reported measures, as self-administered questionnaires. An intention-to-treat analysis was carried out on all outcome measures. Missing data will be treated using the last observation carried forward approach.

Data monitoring and interim analysis

The trial does not include Data Monitoring Committee. An update on trial progress will be shared with Ferrara University Hospital Research office every 6 months. The research coordinator will be responsible for interim

analysis to determine if the trial should stop, modify or carry out. The research group will discuss any subsequent modifications and communicate to the funding agency and Ethics Committee.

Patient and public involvement

The research question in this study starts from years of experience in rehabilitation of PwMS and previous research study on the use of VGT in balance impairment. The drafting of the study was submitted to the Multiple Sclerosis Italian Society (FISM) and reviewed to better meet patients' needs. The final version of the study reflects the collaboration between the research group and the patients' association.

Ethics and dissemination

The Ethics Committee of Ferrara approved this study with approval number 170 691 on 19 October 2017. Communication of results and conclusions will be assigned to high-quality journals and national and international conferences. Results will also be disseminated through FISM annual conference. People with MS will be informed about the possible efficacy of the proposed treatment through MS support groups.

DISCUSSION

This trial may highlight the role of gaming in the rehabilitation of PwMS, enforcing the utilisation of new technologies in daily clinical practice among subjects with mild-to-moderate disability.⁵³

Our expectation from the proposed research is to observe a more significant effect on mobility, balance and dual-tasking through virtual reality training compared with a conventional approach. A meta-analysis by Casuso-Holgado *et al* found that active video game training could be considered as effective as conventional training in improving balance and gait abilities in PwMS, but treatment modalities' variability among the included studies may give rise to various interpretations.⁵⁴ Furthermore, we expect different results due to the dosage of our treatment compared with those of included studies in the review and treatment administration modalities that include home-based interventions and one intervention based on telerehabilitation. Instead, our expected results are in line with what was found by Nascimento *et al* in their systematic review regarding fatigue, quality of life and balance.⁵⁵ Making the rehabilitative session more engaging for patients may increase involvement and adherence in the rehabilitation process. Increased participation and motivation were already being observed in PwMS treated with gamified training.^{53 56} Traditional rehabilitation approaches is often repetitive and boring, decreasing patient's interest and exercise participation. However, patients' satisfaction represents one of the key features for treatment adherence and rehabilitation success, particularly in PwMS.⁵⁷ Virtual reality-based therapy has been proposed to overcome the drawbacks

of conventional rehabilitation, providing augmented feedback during training that contribute to a more effective motor learning.⁵⁸ Active video games may offer an enriched environment useful for subjects with neuropsychological disorders, like attentional deficit or impaired alertness. Exergaming has recently been studied as an effective strategy to improve dual-task performance in people with neurological disabilities.⁵⁹ Indeed, active video games not only engage patients in motor activities, but it simultaneously require subjects to use cognitive abilities for managing inputs into an enriched environment.⁵ Therefore, we expect to observe greater modifications on cognitive and psychological evaluation of people who received rehabilitation through video game platforms. Recently, gaming consoles introduced in clinical and research settings may represent a low-cost opportunity of delivering virtual-reality training. For this reason, the use of VGT may be delivered at home, promoting self-management strategies to improve mobility function and long-term outcomes. Our study may have several limitations. First, the absence of evaluation over 3 months could not give any information about the long-term effects of active video game training. Second, we will not use any instrument to assess patients' satisfaction with the experimental treatment. Third, any neuroimaging technique will be used to show the possible neuroplastic changes in the brain due to VGT. Finally, we will not study the effects of combined treatment of VGT and other rehabilitative techniques for balance and mobility, despite combining treatments seems to augment training efficacy and boost effects of a single approach.²² Further studies should consider these possible limitations and confirm the results related to physical and neuropsychological outcomes.

Contributors SS, GZ, AB, NB and GS conceptualised the protocol. AB, GM and GF drafted the manuscript. AB, GF, GM, NB, SS and GS read and corrected the manuscript. All authors have read and approved the final manuscript.

Funding This research was supported by the Multiple Sclerosis Italian Society (Grant 2018/R/19).

Competing interests None declared.

Patient consent for publication Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

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Università
degli Studi
di Ferrara

ARCISPEDALE S. ANNA
DIPARTIMENTO DI NEUROSCIENZE/RIABILITAZIONE

Foglio informativo per il paziente

Gentile Signora/e,

Lei è invitata/o a partecipare ad uno studio che viene effettuato presso questo ospedale cui lei si è rivolto per motivi di diagnosi o cura. Si prenda tutto il tempo necessario per leggere questo foglio. Se qualcosa non le è chiaro, non esiti a porre tutte le domande che vorrà.

Il titolo dello studio è:

Il ruolo della video game terapia sull'equilibrio e le funzioni cognitive in pazienti con sclerosi multipla e disabilità lieve o moderata. Uno studio pilota randomizzato controllato.

Perché questo studio clinico viene proposto?

Spesso la Sclerosi Multipla si caratterizza per la presenza sia di disordini dell'equilibrio, sia di disordini cognitivi, cioè quelle funzioni del cervello quali l'attenzione e la memoria. Gli aspetti cognitivi non devono essere considerati come una dimensione separata dagli aspetti motori, ad esempio l'attenzione agisce sul controllo dell'equilibrio. Inoltre, nella vita quotidiana è possibile osservare la compresenza dei due aspetti: basti pensare a quando si cammina e contemporaneamente si parla. Il trattamento riabilitativo dovrebbe quindi tenere conto di entrambi questi aspetti, proponendo attività che potenzino la performance sia cognitiva che motoria. La Video-Game Terapia potrebbe essere uno strumento utile in tal senso, in quanto può favorire il trattamento sia dell'equilibrio sia degli aspetti cognitivi contemporaneamente. Inoltre, rispetto a trattamenti più "classici", la Video-game Terapia aumenta la motivazione, favorendo l'acquisizione di nuove competenze. E' inoltre meno dispendiosa e più "versatile" in quanto è possibile proseguire il trattamento riabilitativo anche al domicilio. Con questo studio esplorativo vogliamo confrontarne gli effetti rispetto ai trattamenti "classici" per l'equilibrio, sia per quanto riguarda il miglioramento dell'equilibrio, sia delle funzioni cognitive in generale.

Lo scopo principale di questo studio è quello di verificare gli effetti di un terapia mediante un dispositivo commercialmente disponibile sull'equilibrio e sulle funzioni cognitive nei pazienti SM deambulanti rispetto a un trattamento riabilitativo "standard". In secondo luogo, esploreremo gli effetti in altri settori che sono generalmente compromessi nella popolazione MS e che possono essere migliorati mediante riabilitazione motoria, come il benessere psicologico, la stanchezza e la qualità di vita.

Quali sono le caratteristiche di questo studio?

Lo studio prevede di raccogliere dati di sicurezza ed efficacia di una procedura confrontata con quella attualmente in uso secondo le norme di buona pratica clinica.

Verrà reclutato un campione di pazienti con Sclerosi Multipla presso l'Azienda Ospedaliero - Universitaria di Ferrara. I soggetti reclutati saranno attribuiti in maniera casuale in gruppi che riceveranno la Video Game Terapia (VGT) o la Terapia mediante Piattaforma dell'Equilibrio (BPT).

Il gruppo sperimentale riceverà 12 sessioni di allenamento VGT (3 volte alla settimana) per 4 settimane. Il gruppo di controllo riceverà la stessa quantità di sessioni BPT. Selezioneremo le misure di equilibrio che esplorano un'ampia gamma di compiti motori. Saranno inoltre somministrati test per valutare l'affaticamento, le funzioni cognitive e il benessere fisico e psicologico. L'equilibrio verrà valutato mediante una serie di test

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clinici funzionali che indagano un'ampio spettro di compiti motori. Inoltre verrà eseguito un esame della postura attraverso una pedana in grado di misurare le oscillazioni corporee in posizione eretta. Verranno somministrati questionari per valutare la fatica, il benessere psicofisico e la qualità di vita. Le misure verranno raccolte prima del trattamento (T0), alla fine (T1) e a distanza di 3 mesi (T2) per valutare il mantenimento dei risultati.

Chi propone lo studio?

Lo Studio è proposto dal Settore di Medicina Riabilitativa "S. Giorgio" – Dipartimento di Neuroscienze/Riabilitazione dell'Azienda Ospedaliero-Universitaria di Ferrara.

Perché sono invitato a partecipare allo studio?

Le stiamo proponendo di partecipare a questo studio perché è affetto da Sclerosi Multipla. È previsto che partecipino a questo studio 48 pazienti.

Cosa comporta la partecipazione allo studio, rispetto al normale percorso diagnostico-terapeutico per la mia malattia?

Lo studio prevede la creazione di due gruppi di pazienti: il primo gruppo riceverà la Video-Game Terapia, mentre il secondo gruppo riceverà un trattamento mediante Piattaforma dell'Equilibrio. L'attribuzione ad un gruppo o ad un altro è casuale. Prima dell'inizio del trattamento, alla fine del trattamento e a distanza di tre mesi, verranno effettuate misurazioni attraverso specifici test, così da permettere un confronto tra i due gruppi.

Quali rischi o inconvenienti potrei avere dalla partecipazione a questo studio?

Non vi sono rischi legati allo studio.

Eventuali nuove informazioni che potrebbero influenzare la sua volontà di partecipazione le verranno comunicate il più presto possibile. Lo stesso vale per una eventuale interruzione o sospensione dello studio.

Quali vantaggi potrei avere nel partecipare a questo studio?

Lei potrebbe non avere beneficio diretto dalla partecipazione. Questo studio potrà contribuire a migliorare la diagnosi e la comprensione della sua malattia, e portare allo sviluppo di nuovi trattamenti riabilitativi per i disturbi dell'equilibrio e cognitivi. I risultati dello studio saranno poi pubblicati su riviste scientifiche nazionali o internazionali. Per la partecipazione allo studio non è previsto alcun compenso.

Sono obbligato a partecipare allo studio?

No. La decisione di partecipare è assolutamente libera. Se lei acconsente ha la possibilità di contribuire alla ricerca medica attraverso questo studio. Se però non vuole partecipare, non deve fornire alcuna spiegazione. Il suo rifiuto non influenzerà in alcun modo il trattamento che le verrà proposto, e riceverà comunque tutte le terapie previste dalla buona pratica clinica per la sua patologia.

Potrò cambiare idea dopo aver accettato di partecipare?

Sì. La decisione di partecipare allo studio è volontaria e libera, e lei ha il diritto di revocare il suo consenso in qualunque momento lo desidera, senza fornire spiegazioni e senza che questo influenzi in alcun modo il trattamento che le verrà proposto, che sarà comunque il migliore disponibile.

Se partecipo allo studio, miei dati personali e clinici saranno noti a tutti?

No. I suoi dati clinici saranno resi anonimi. Il suo nome e cognome saranno sostituiti da un codice che solo il responsabile dello studio conoscerà.

Trattamento dei dati

Se Lei deciderà di partecipare allo studio, tutti i dati raccolti (età, sesso, origine etnica e i dati clinici), saranno archiviati elettronicamente in maniera rigorosamente anonima, ai sensi del Decreto Legislativo n. 196/03 sulla tutela delle persone rispetto al trattamento dei dati personali e saranno trattati in modo assolutamente riservato. I dati verranno conservati presso l'unità Operativa di Medicina Riabilitativa S.

Giorgio dell'Azienda Ospedaliero-Universitaria di Ferrara per un periodo di 2 anni dalla chiusura dello studio. La persona responsabile della gestione dei suoi dati per questo studio e reparto è la dr.ssa Sofia Straudi. L'accesso diretto alla sua documentazione sarà consentito a tutti coloro che sono coinvolti nell'effettuazione dello studio (personale sanitario, personale che elabora i dati, personale ispettivo e quant'altri abilitati dal protocollo di studio e/o dalle normative vigenti) e alle autorità regolatorie nella misura permessa dalle leggi senza violare la sua riservatezza.

Il medico della ricerca le consegnerà una lettera rivolta al suo medico di base, per informarlo della sua partecipazione allo studio, per la migliore conduzione clinica dello stesso.

Chi ha approvato lo studio?

Il protocollo dello studio è stato redatto in accordo con la dichiarazione di Helsinki sull'etica della ricerca in medicina ed è stato approvato dal Comitato Etico di questo ospedale.

Se Lei accetterà di partecipare a questo studio Le verrà chiesto di **firmare e datare** di suo pugno questo foglio informativo di cui Le sarà consegnata copia assieme alla copia del foglio di consenso.

A chi posso rivolgermi se ho dei problemi durante lo studio?

Il medico referente per questo studio è la Dr.ssa Sofia Straudi Tel. 0532-236185 (s.straudi@ospfe.it)

Lo staff operativo coinvolto nello studio è costituito dai seguenti operatori:

- 1) Andrea Baroni (a.baroni@ospfe.it)
- 2) Nino Basaglia
- 3) Anna Scotti
- 4) Giada Milani
- 5) Giulia Fregna

Nome in stampatello del partecipante allo studio

Data e Firma

Nome in stampatello del Medico ricercatore

Data e Firma

Il ruolo della video game terapia sull'equilibrio e le funzioni cognitive in pazienti con sclerosi multipla e disabilità lieve o moderata. Uno studio pilota randomizzato controllato.

Modulo di consenso allo studio e al trattamento dei dati

Questo modulo deve essere firmato da Lei solo nel caso decida di partecipare allo studio. E' importante che Lei abbia discusso approfonditamente con il Medico prima di firmare questo consenso, anche sulla base del foglio informativo a cui esso si riferisce. Partecipano allo studio solo i Pazienti che accettano. Il Paziente può ritirare il suo consenso in ogni momento.

Dichiaro di ricevere copia firmata del presente modulo di consenso unitamente a copia datata e firmata del foglio informativo. Dichiaro inoltre di:

- aver ricevuto dal medico esaurienti spiegazioni in merito alla richiesta di partecipazione allo studio, in particolare sulle finalità e sulle procedure;
- aver letto e compreso il foglio informativo che mi è stato consegnato con sufficiente anticipo e che conferma quanto mi è stato spiegato a voce;
- aver avuto la possibilità di porre domande ed aver avuto risposte soddisfacenti;
- farmi carico della consegna della lettera relativa allo studio per il mio medico di famiglia;
- essere consapevole che la partecipazione è volontaria, con l'assicurazione che il rifiuto a partecipare non influirà sulla scelta della terapia migliore per me;
- essere consapevole che, se ritirerò il mio consenso, i dati raccolti prima del ritiro del consenso saranno utilizzati dal ricercatore;
- autorizzare il trattamento dei miei dati personali ai sensi del Decreto legislativo n. 196/2003 (codice privacy) con le finalità indicate nello studio
- Acconsento a partecipare allo studio clinico suddetto

Nome in stampatello del partecipante allo studio

Luogo e Data nascita _____

Residenza:

Data e Firma

Nome in stampatello del Medico ricercatore

Data e Firma

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