

Efficacy of Wii Balance Board-Based Exergame Training Among Individuals with Cerebellar Ataxia: A Feasibility Study

Sayan Pratihari^{1*}, Karthiga Rajasekaran¹, Shanmuga Priya Raji Reddy Parasuraman¹

¹SRM College of Physiotherapy, Faculty of Medicine and Health Sciences, SRM Institute of Science and Technology, Chennai, India.

* Corresponding Author: Sayan Pratihari, SRM College of Physiotherapy, Faculty of Medicine and Health Sciences, SRM Institute of Science and Technology, Chennai, India. Email: pratiharsayan0@gmail.com

ORCID ID: <https://orcid.org/0009-0008-7378-0366>

Received 2024 December 08; Accepted 2025 February 16.

Abstract

Background: Conventional rehabilitation methods have shown limited and transient improvements, necessitating personalized approaches in the diverse population of cerebellar ataxia (CA). Wii balance board exergame training, integrating physical exercise with interactive video games, presents a novel and engaging neuro-rehabilitation strategy.

Objectives: The primary objective of this study was to assess the clinical feasibility of implementing Wii Balance Board-based exergame training among individuals with various forms of CA. The secondary objective was to investigate the preliminary efficacy and assess the enjoyment of the intervention.

Methods: The study incorporates a pilot randomized control trial and feasibility study design. We recruited 10 patients using a block randomization method. The Wii balance board training was administered for 18 sessions, 3 sessions per week, over 6 weeks. The primary outcomes of feasibility testing were evaluated through clinical research log documentation, while secondary outcomes of balance, ataxia severity rate, functional independence, and enjoyment were assessed with the mini-BESTest, Scale for Assessment and Rating of Ataxia (SARA), Functional Independence Measure Scale (FIMs), and Exergame Enjoyment Questionnaire (EEQ). Data were analyzed using descriptive statistics and non-parametric tests to evaluate changes in outcomes.

Results: The study enrollment rate was 77% (n = 10). The Wii intervention group showed a 100% (n = 5) retention rate compared to 80% (n = 4) in the control group (CG). The Wii intervention group demonstrated a tendency towards better outcomes at follow-up in SARA (P = 0.063, effect size/R_M = 0.84) and Mini-BESTest (P = 0.071, effect size/R_M = 0.79) but not in the case of FIM (P = 0.794, effect size/R_M = 0.14), along with reporting a moderate level of enjoyment.

Conclusions: Wii Balance Board-based exergame training is considered feasible for implementation in clinical settings among individuals with various forms of CA, suggesting the conduction of a larger definitive study to further explore the intervention's efficacy.

Keywords: Balance; Cerebellar Ataxia; Exergames; Virtual Reality Rehabilitation; Wii Balance Board

1. Background

Cerebellar ataxia (CA) includes a range of disorders that cause problems with coordination, balance, and motor control, which can severely affect the quality of life for individuals experiencing these issues (1). The CA is categorized into three main types: Acquired, degenerative nonhereditary, and inherited. The inherited forms include autosomal recessive (ARCA), autosomal dominant (ADCA), and X-linked ataxias (2). Hereditary ataxia affects about 2.7 out of every 100,000 people for autosomal dominant hereditary cerebellar ataxia (AD-HCA) and around 3.3 out of every 100,000 for autosomal recessive hereditary cerebellar ataxia (AR-HCA) (3). Spino-cerebellar ataxia (SCA) type 12 and type 2 are more frequently found in the northern regions of India, whereas SCA-1 is predominantly seen in

the southern part of the country (4).

An ataxic patient often struggles with functional disabilities that further challenge their socio-economic life (5). They experience a progressive loss of functional independence due to frequent falls, requiring long-term rehabilitation and caregiving. This imposes a considerable impact on their families, increasing both emotional and financial stress. Additionally, the costs associated with medical care, assistive devices, and therapy can be substantial, while the disease's disabling nature often leads to reduced workforce participation and productivity. Studies across different countries have reported substantial annual costs per ataxia patient, ranging from €18,776 in Spain to HKD 146,832 (for 6 months) in Hong



Copyright © 2025 Tehran University of Medical Sciences.

This work is licensed under a Creative Commons Attribution-NonCommercial 4.0 International license (<https://creativecommons.org/licenses/by-nc/4.0/>). Noncommercial uses of the work are permitted, provided the original work is properly cited.

Kong (6-8).

Several current treatment approaches for CA, including standard care physiotherapy methods, vestibular rehabilitation, intensive physiotherapy, and other advanced techniques like repetitive transcranial magnetic stimulation (r-TMS), have demonstrated varying degrees of effectiveness. However, these methods often face challenges such as limited accessibility, high resource demand, and inconsistent adherence (9-11). Moreover, conventional rehabilitation approaches for CA frequently lack individualized interventions tailored to patient-specific needs, which are critical for optimizing rehabilitation outcomes (12).

The combination of physical exercise and interactive video games in exergame training offers a novel, distinctive, and engaging approach to neuro-rehabilitation (13). With advancements in technology, the cost-effective Wii Balance Board-based exergame training provides real-time feedback on balance and posture, making it an effective rehabilitation tool for individuals with CA (14). Despite its potential, prior studies emphasized the need for extended monitoring to fully understand the long-term benefits of exergame training programs (15, 16). Furthermore, recent studies frequently lack patient-specific adaptations and personalized feedback mechanisms or enjoyment by involving Wii Balance Board-based exergame training, which could significantly enhance therapeutic outcomes (17).

It is also necessary to include outcome measures that encompass not only motor enhancements but also take into account the functional status experienced after engaging in exergame training (18). On the other hand, the feasibility and long-term efficacy of implementing Wii Balance Board-based exergame training in clinical settings remain underexplored across the wide range of CA types induced by various pathophysiological factors (19).

A major challenge in researching therapeutic interventions for CA is the heterogeneity of the patient population, which includes various inherited and non-inherited forms with differing etiologies, progression rates, and severity levels. This diversity can confound study outcomes, making it difficult to isolate the effects of specific interventions (20). Furthermore, conducting large-scale trials on such varied groups can be ethically and logistically challenging, especially for innovative interventions like Wii Balance Board-based exergame training, particularly when different pathological mechanisms may respond differently to the same treatment (21).

Given these complexities, there is a critical need to first evaluate the feasibility and safety of using Wii balance board exergame training in a more controlled yet diverse group of CA patients. Conducting a pilot feasibility study would allow us to address these challenges on a smaller scale, ensuring that the intervention is practical and acceptable before committing resources to a larger, more definitive trial.

2. Objectives

The primary objective of this study was to assess the clinical feasibility of implementing Wii Balance Board-based exergame training in clinical settings for individuals with different types of CA. The secondary objective was to investigate the preliminary efficacy of the Wii balance board training intervention in terms of improving balance, ataxia severity rate, and functional independence. Furthermore, the study also sought to assess the degrees of enjoyment experienced by individuals from Wii Balance Board-based exergame training.

3. Methods

3.1. Study Design and Randomization

This study utilized a pilot randomized control trial (parallel arm) and feasibility study design with limited statistical power (as per the assumption of a 10 - 15% rate of actual power calculated sample of 86 using a two-tailed Laplace distribution at $\alpha = 0.05$, power $(1 - \beta) = 80\%$, effect size $d = 0.5$) involving the experimental group (EG) or Wii balance board training group and a control group (CG). An independent researcher implemented a 1:1 allocation ratio for the group allocation. The recruitment procedure employed a block randomization approach with a block size of 4 and a numerical sequence. A computer-based random allocation software was used to execute the randomization, which was produced by an independent researcher who was not affiliated with the trial and had no involvement in its conduct. The computerized database concealment was adhered to until the intervention started. The original group allocation was concealed from the outcome assessor, who was a field expert involved in this study. Due to the limited sample size, stratification could not be performed. However, block randomization ensured balanced allocation of key demographics across groups. Baseline characteristics were compared to assess any residual imbalances. The study follows the CONSORT 2010 guidelines for the reporting of a pilot and feasibility trial (22).

The inclusion criteria of the study were specified as follows: (1) Both males and females aged 30 - 60 years; (2) the condition of CA was diagnosed by a neurologist; (3) participants who could stand and ambulate independently or with the use of mobility aids such as a cane or walker; (4) participants who could visually observe a display screen; (5) participants who could comprehend the therapist's instructions and conversation; (6) the Mini-Mental State Examination (MMSE) score considered as 24 or higher; (7) no previous VR-based rehabilitation training experience.

The study's exclusion criteria encompassed the following: (1) Individuals with amputations wearing prosthetic devices on their lower limbs; (2) any congenital anomalies impacting the lower extremities and spine; (3) experience of any recent injuries, back pain, or serious joint disorders in the lower limbs that would make it difficult for weight-bearing and standing upright; (4) history of pre-existing vestibular disorders, receptive aphasia, and global apha-

sia; (5) sensory ataxia; (6) history of psychological disorders; (7) history of peripheral neuropathy; (8) previous occurrence of epilepsy or seizure; (9) pregnant women; (10) previous medical history includes severe cardiovascular conditions and respiratory ailments.

3.2. Participants, Screening, and Enrollment

We concentrated on both the individuals who sought medical care as out-patients at the Neurology OPD and those who were hospitalized as in-patients in the Neurology ward at SRM Medical College Hospital and Research Centre (SRM MCH&RC) in Chennai, India. Between November 2023 and January 2024, 13 CA patients were screened for

eligibility in the Neurology department. Two participants did not meet the inclusion criteria, and one was excluded based on the exclusion criteria. Finally, 10 patients successfully enrolled in the study (Figure 1). Of these, six had ataxia originating from cerebellar stroke, two had ataxia induced by metabolic causes (hypothyroidism, Wilson disease), and two had inherited ataxia (Friedreich's ataxia, SCA-1). The EG included three cerebellar stroke cases, one metabolic ataxia case (hypothyroidism-induced ataxia), and one hereditary ataxia case (SCA-1), while the CG had a similar enrollment of three cerebellar strokes, one metabolic ataxia case (Wilson disease-induced ataxia), and one hereditary ataxia case (Friedreich's ataxia).

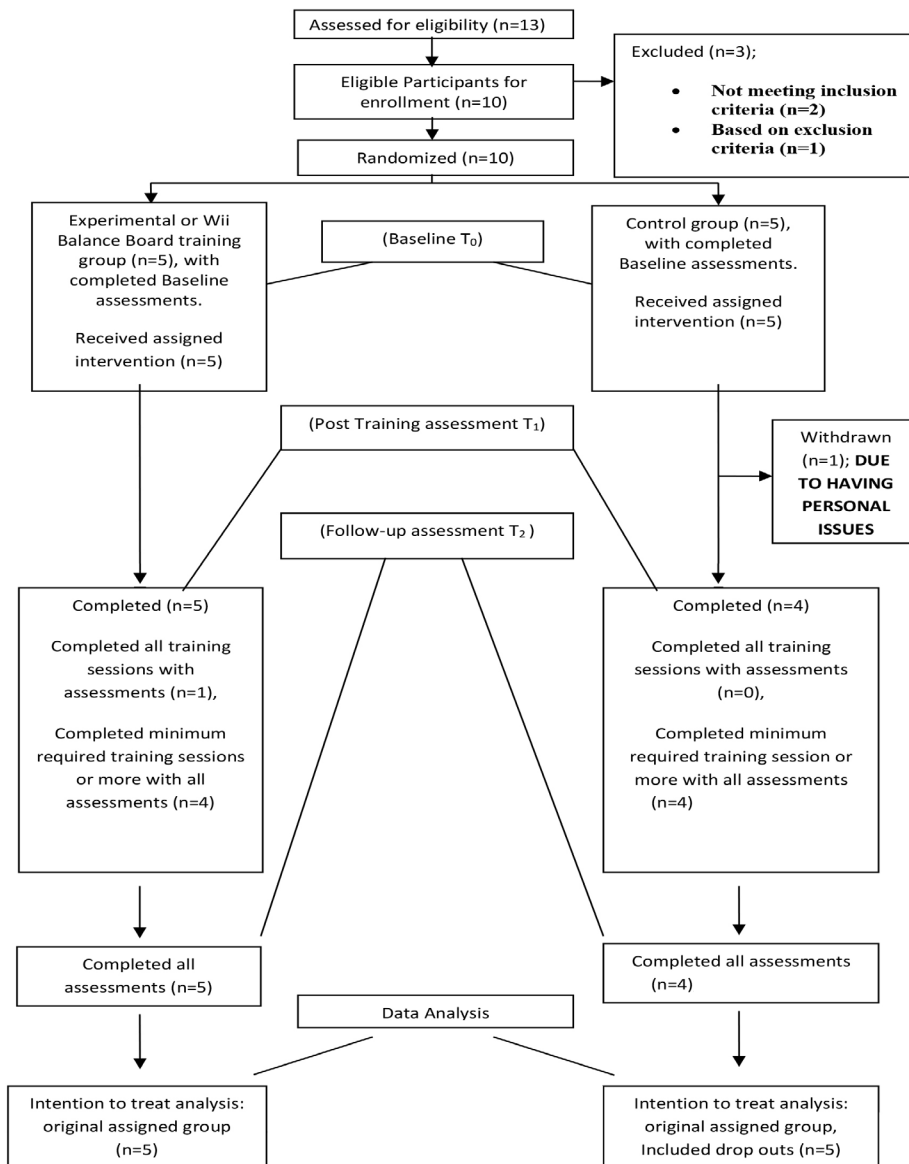


Figure 1. Participants' flow chart

As shown in Table 1, the groups were generally matched across key influential variables, including age, sex, cogni-

tion status (MMSE scores), types of ataxia, ataxia severity [Scale for Assessment and Rating of Ataxia (SARA) scores],

balance impairment (mini-BESTest scores), and functional independence [Functional Independence Measure Scale (FIM) scores]. Minor differences were observed in age and balance impairment baseline values, although these differences fall within the overlapping range of both groups. Baseline statistical analysis (Table 1) suggests no significant imbalance between the groups. Fur-

thermore, other key variables, such as sex distribution, cognition status (MMSE scores), and types of ataxia, were evenly distributed across groups. Given the pilot nature of this feasibility study, these differences are unlikely to bias the outcomes or affect the study's primary aim of evaluating the feasibility and acceptability of the intervention (23).

Table 1. Participants' Demographic Characteristics a

Baseline Variables	EG (N = 5)	CG (N = 5)
Age (y)	48.40 ± 10.99	55.40 ± 6.30
Min-max	35 - 60	45 - 60
Sex		
Male	4 (80)	4 (80)
Female	1 (20)	1 (20)
Cognition status		
MMSE	28.40 ± 2.07	28.40 ± 1.81
Types of ataxia		
Acquired brain injury	3 (60)	3 (60)
Metabolic ataxia	1 (20)	1 (20)
Inherited ataxia	1 (20)	1 (20)
Ataxia severity rate		
SARA score	15.30 ± 6.07	15.70 ± 2.38
Balance impairment rate		
Mini-BESTest score	17 ± 4.84	15.20 ± 1.30
Functional independence status		
FIM score	101.20 ± 12.21	102.60 ± 9.28

^a Abbreviations: EG, experimental group; CG, control group; MMSE, mini-mental state examination score; SARA, Scale for Assessment and Rating of Ataxia; FIM, Functional Independence Measure Scale.

^a Values are expressed as mean ± SD or No. (%).

3.3. Ethical Consideration

This study was approved by the SRM Institutional Ethical Committee (approval No: SRMIEC-ST0523-657) and conducted in accordance with the ethical standards of the Declaration of Helsinki. Prior to enrollment, all eligible participants provided written informed consent after receiving detailed explanations about the study's purpose, procedures, potential risks, and benefits. They were informed of their right to withdraw from the study at any point without consequences. To ensure participant confidentiality, all personal and clinical data were anonymized and securely stored, accessible only to authorized researchers. Precautions were taken to protect the well-being of the participants, including careful monitoring during interventions and immediate access to medical care if required. Additionally, this study was prospectively registered on the Clinical Trial Registry of India (ctri. nic.in) with the registration number CTRI/2023/11/059589.

3.4. Intervention

The exergame training took place in a 150 sq. ft. room at

the Department of Physical Medicine and Rehabilitation (SRM MCH&RC), using the Nintendo® Wii and Wii balance board connected to a projector (Figure 2). The EG underwent 18 sessions over 6 weeks, with 3 sessions per week, each session lasting 20 minutes. Participants played four Wii Fit Plus standing balance games (table tilt, ski slalom, tightrope walk, soccer heading) at different difficulty levels (beginner, advanced, professional), with a 1-minute rest period between games. Hemodynamic status (such as blood pressure, heart rate, respiratory rate, SPO₂) was monitored before and after every training session to safeguard against the occurrence of any unwanted harmful cardiovascular events. Each training session included a 2-minute warm-up and a 2-minute cool-down period. A 30-minute practice session was scheduled for all the EG participants to understand the concepts of game control before starting their training session. Assistive devices (like a cane and walker) were available during the training sessions for safety holding, though participants were encouraged to minimize the use of those aids as much as possible. Additional support from two therapists was provided to prevent falls throughout the training ses-

sions. Alongside the exergame training, the participants in the EG also received routine physiotherapy treatments such as upper and lower limb strengthening exercises using a Thera-Band, equilibrium and non-equilibrium coordination exercises, and parallel bar gait training inside the same intervention setting for the duration of 20 minutes in each session. The CG participants received standard standing balance training on a wobble board, Thera-Band strengthening exercises for upper and lower

limbs, coordination exercises, and parallel bar gait training for 40 minutes at the frequency of 3 sessions per week for 6 weeks in a usual care setting at the Department of Physiotherapy (SRM MCH&RC). During the study period, participants were not constrained from other treatment programs related to their health conditions either outside the study settings or inside the study hospital due to ethical concerns.



Figure 2. Wii balance board training

3.5. Outcome Measures

The primary outcome of feasibility testing was measured with the recruitment capability and retention rate, treatment-specific compliance rate, adherence rate, and adverse events through clinical research log-book documentation. We used the Pragmatic Explanatory Continuum Indicator Summary (PRECIS-2) scores of the trial domain or PRECIS-2 tool to evaluate the pragmatic versus explanatory nature and the applicability of our pilot feasibility study design (24, 25). The secondary outcome measures of balance, ataxia severity rate, and functional independence were measured by the mini version of the Balance Evaluation System Test Scale or mini-BESTest

Scale (26), the SARA (27-29), and the FIMs (30).

Initially, the outcomes were measured at Baseline (T0) during the recruitment of samples, and the post-test (T1) assessments were conducted following the completion of either the minimum required training session or more as per prescribed sessions (adhering to a 70% benchmark adherence rate guideline for total training sessions participation to ensure feasibility generalization) (31). A follow-up assessment (T2) was also conducted 4 weeks after the post-test (T1) assessments. The additional secondary outcome regarding patient enjoyment, experience, and satisfaction from Wii balance board training was evaluated by a patient self-reported measure — the Exergame Enjoyment Questionnaire (EEQ) — after the completion of the trial.

3.6. Data Analysis

We performed the statistical data analysis using IBM® SPSS® version 27 software. The non-parametric Friedman test was employed to evaluate changes in outcomes over time within each group across three time points. Post-hoc pairwise comparisons were conducted using the Wilcoxon signed-rank test, with a Bonferroni correction applied to adjust for multiple comparisons. The Mann-Whitney U test was used to compare differences in outcomes between the experimental and CGs at each time point. Effect sizes (R_w , R_m) were calculated to determine the magnitude of the difference between groups. Data regarding usefulness were analyzed using descriptive methods.

4. Results

4.1. Feasibility Outcomes

The number of CA patients screened month-wise was as follows: Seven patients in November 2023, 3 patients in December 2023, and 3 patients in January 2024. Recruitment for this study started on November 20, 2023, and the final baseline assessments were completed on January 19, 2024. The outcome assessments were conducted from February 12, 2024, to March 30, 2024. The rate of enrollment for qualifying screens was 77% ($n = 10$ qualified for successful enrollment into groups among 13 screened patients). The percentage of participants who both enrolled and attended at least one session was 100%.

The mean attendance of our participants over the six-week period was 14 sessions ($n = 5$, range = 13 - 18) out of a total of 18 training sessions in the EG. Four participants successfully completed the minimum required 13 or more consecutive Wii balance board training sessions along with the required assessments, meeting the benchmark adherence rate of 70% of the total number of sessions in a pilot study (31). In contrast, only one participant in the EG successfully finished all of the designated sessions and assessments. In the CG, four participants completed the minimum required training sessions, although nobody could complete all the training sessions. One individual in the CG withdrew from the study during the ongoing trial and did not attend the post-trial assessments or follow-up visits due to personal issues. The EG did not have any dropouts.

The treatment-specific retention rate in the EG was 100% ($n = 5$), while the CG had a retention rate of 80% ($n = 4$). The proportion of planned assessments that were completed in this study remained 100% for the EG, while it was 90% for the CG. The average duration to complete total assessment visits per participant was around 45 minutes in both groups. The compliance rate for the Wii balance board training varied depending on the difficulty levels of the specific exergame. At the beginner level, the compliance rates were as follows: GAME I (table tilt) = 91%, GAME II (ski slalom) = 90%, GAME III (rope walk) = 77%, GAME IV (soccer heading) = 78%. At the advanced level, the compliance rates for GAME II, GAME III, and GAME IV were 82%, 52%, and 82%, respectively. At the professional level, the compliance rate in Game III was 48% (Table 2).

Table 2. Gameplay Events a

Events	Values (N = 5)
Level attained in games	
Table tilt	100% beginner level, 0% advanced level, and 0% pro level
Ski slalom	20% beginner level only, 80% beginner + advanced level, and 0% pro level
Rope walk	60% beginner level only, 40% beginner + advanced + pro level
Soccer heading	20% beginner level only, 80% beginner + advanced level, and 0% pro level
Number of playing attempts	
Beginner level	
Table tilt	64 ± 15.23
Ski slalom	22.60 ± 18.74
Rope walk	38.60 ± 18.66
Soccer heading	15.60 ± 21.01
Advanced level	
Ski slalom	36.80 ± 22.68
Rope walk	7.80 ± 15.30
Soccer heading	41.20 ± 24.50
Pro level	
Rope walk	2.4 ± 3.57
Number of sessions	
Beginner level	
Table tilt	14 ± 2.23

Ski slalom	5 ± 4.60
Rope walk	10 ± 4.38
Soccer heading	4 ± 5.27
Advanced level	
Ski slalom	9 ± 5.16
Rope walk	3 ± 3.89
Soccer heading	10 ± 6.42
Pro level	
Rope walk	1 ± 1.41

^a Values are expressed as percentage or mean ± SD.

Formula for calculation of treatment-specific compliance rate: (Estimated number of actual playing attempts at various levels of specified games in Wii balance board Training sessions)/(Total prescribed playing attempts for the individual games in Wii balance board training sessions) × 100%.

The PRECIS-2 scores of trial domains for our study indicate a predominantly pragmatic (reflecting real-world practice) but also explanatory approach to evaluate the practical implementation of the study design, especially in the following domains. Eligibility criteria, sample recruitment process, study setting, intervention organization, experimental intervention-delivery flexibility, and experimental interven-

tion adherence-flexibility incorporated both pragmatic and explanatory aspects in this feasibility study. As the outcome assignment focused primarily on a certain component of the ICF domains, it was less directive towards the pragmatic aspect. The follow-up assessments were anticipated to be very pragmatic in terms of the operational convenience of the interventions and the regular monthly hospital visits for health evaluations. The data analysis conducted with the intention-to-treat analysis according to the original assigned group showed great practicability, particularly in the CG, as the retention rate was limited (Table 3).

Table 3. Pragmatic Explanatory Continuum Indicator Summary Scores of Trial Domains ^a

Domain	Score	Rationale
Eligibility criteria	3	P: Inclusion of both male and female. E: Inclusion of age group 30 - 60, inclusion of all types of CA excluding degenerative CA
Recruitment path	3	P: Recruitment of participant from neurology OPD units and admitted patients in neurology Ward of the study hospitals. E: Checklist to assess recruitment eligibility
Setting	3	P: Catchment area is three neurology OPD units and two neurology wards (including male and female ward). E: Single training center trial (feasibility)
Organization of intervention	3	P: Resource, expertise, and delivery of care in both the arm are similar. E: Requirement of technical staffs/ground staffs for the maintenance of the research equipments and ensuring treatment surveillance in the experimental arm. Appointment of a neurologist who is having expertise in dealing with CA to assess any reported disease specific adverse events for need of special care
Flexibility of experimental intervention-delivery	3	P: Warm up exercises, vitals monitoring before and after of the training at both the arm. Clinical log documentations of the individuals tasks attempts and completions of tasks. E: Delivering practice sessions for understanding the concept of virtual reality, patient handling during standing on the Wii balance board, and expert supervision while playing exergames in the experimental arm
Flexibility of experimental intervention-adherence	3	P: Usual encouragement to adhere to routine PT. E: Providing incentives to encourage participants' adherence
Follow-up	5	P: Follow-up assessments during usual monthly visit to the hospital by our participants for general health checkups (4 weeks after post-test assessment)
Outcome	4	P: Outcomes are measured at the impairment and activity domain of ICF
Analysis	5	P: Intention to treat analysis including drop out participants' data in the CG for preliminary efficacy

^z Abbreviations: P, pragmatic; E, explanatory; CA, cerebellar ataxia; CG, control group.

^a Scores: 1 = very explanatory; 2 = rather explanatory; 3 = equally pragmatic/explanatory; 4 = rather pragmatic; 5 = very pragmatic.

4.2. Exergame Training Safety and Participant Experience/Satisfaction

Three individuals reported experiencing adverse events, including dizziness, headaches, and eye pain, after playing Wii Fit games at the end of a few training sessions. No accidental falls or other adverse consequences were documented. The participants reported a moderate level of enjoyment (54.40 ± 15.69) as documented by the EEQ from playing Wii Fit Plus standing balance games.

4.3. Preliminary Efficacy

Over time, both the experimental and CGs demonstrated notable enhancements in balance control (Mini-

BESTest), ataxia rate (SARA), and functional independence (FIMs) within their respective groups. In comparison to the CG (Kendall's $W = 0.80$, $P = 0.005$), the EG exhibited greater effect sizes across all measures (Kendall's $W = 1.00$, $P < 0.001$) (Table 4, Figure 3). Although significant gains were observed in pairwise comparisons within the EG, these improvements did not remain significant following Bonferroni correction ($P < 0.016$) (Table 5). No statistically significant differences were seen between the groups at baseline, post-test, or follow-up. However, the EG demonstrated a tendency towards better outcomes at follow-up in SARA ($P = 0.063$, $R_M = 0.84$) and Mini-BESTest ($P = 0.071$, $R_M = 0.79$) but not in the case of FIM ($P = 0.794$, $R_M = 0.14$) (Table 6).

Table 4. Results of Balance Control, Ataxia Rate and Functional Independence Measures Comparison Within Groups a, b, c

Outcome Measures	EG (N = 5)						CG (N = 5)					
	Mean Rank			Chi-Square d	P-Value e	Kendall's Wf	Mean Rank			Chi-Square d	P-Value e	Kendall's Wf
	T0	T1	T2				T0	T1	T2			
Mini-BESTest	1.00	2.00	3.00	10.00	< 0.001	1.000	1.00	2.00	3.00	8.00	0.005	0.800
SARA	3.00	2.00	1.00	10.00	< 0.001	1.000	3.00	2.00	1.00	8.00	0.005	0.800
FIMs	1.00	2.00	3.00	10.00	< 0.001	1.000	1.00	2.00	3.00	8.00	0.005	0.800

^a Abbreviations: EG, experimental group; CG, control group; SARA, Scale for Assessment and Rating of Ataxia; FIM, Functional Independence Measure Scale.

^a Analysis made including drop out participants' data assuming the same value measured at baseline.

^b T0 = baseline, T1 = post-test, and T2 = follow-up.

^c Rank 3 = highest enhancement, rank 2 = modest enhancement, and rank 1 = lowest enhancement.

^d K^2 .

^e P-values refer to 2-tailed Exact Sig. (< 0.05) of the non-parametric Friedman test.

^f Effect size.

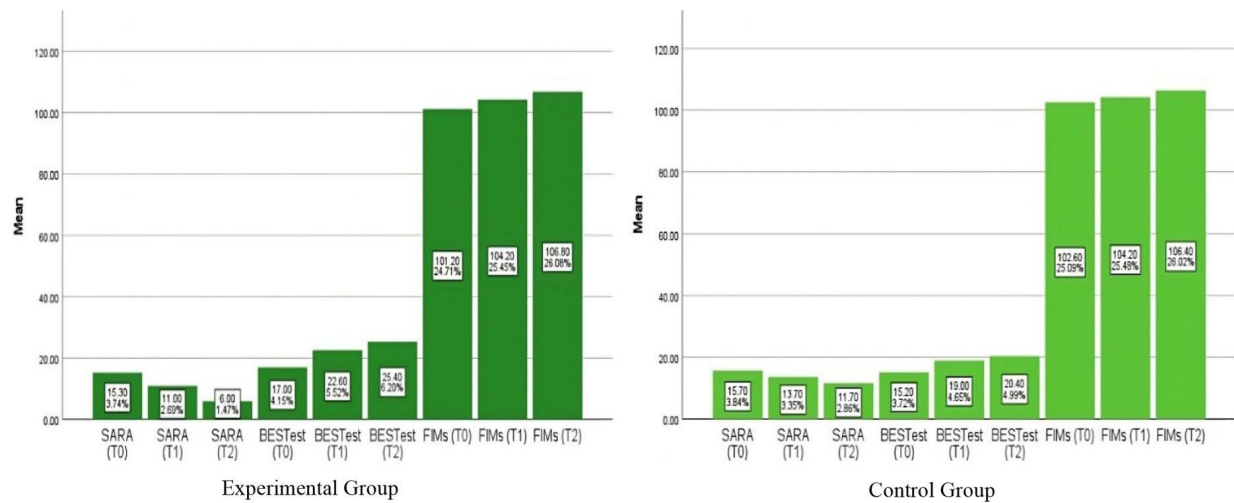


Figure 3. Observed changes in outcome variables over time in experimental group (EG) and control group (CG)

Table 5. Results of Post-hoc Paired Wilcoxon Signed-Rank Test with Bonferroni Correction Within Groups a, b

Outcome Measures; Comparison	EG (N = 5)				CG (N = 5)			
	Z-Value c	Original P-Value d	Bonferroni Adjusted P-Value	R _w e	Z-Value c	Original P-Value d	Bonferroni Adjusted P-Value	R _w e
Mini-BESTest								
T0 vs. T1	-2.041	0.041	0.123	0.91	-1.826	0.068	0.204	0.81
T1 vs. T2	-2.032	0.042	0.126	0.90	-1.841	0.066	0.198	0.82
T0 vs. T2	-2.032	0.042	0.126	0.90	-1.826	0.068	0.204	0.81
SARA								
T0 vs. T1	-2.023	0.043	0.129	0.90	-1.826	0.068	0.204	0.81
T1 vs. T2	-2.060	0.039	0.117	0.92	-1.841	0.066	0.198	0.82
T0 vs. T2	-2.032	0.042	0.126	0.90	-1.826	0.068	0.204	0.81
FIMs								
T0 vs. T1	-2.023	0.043	0.129	0.90	-1.857	0.063	0.189	0.83
T1 vs. T2	-2.060	0.039	0.117	0.92	-1.841	0.066	0.198	0.82
T0 vs. T2	-2.023	0.043	0.129	0.90	-1.826	0.068	0.204	0.81

z Abbreviations: EG, experimental group; CG, control group; SARA, Scale for Assessment and Rating of Ataxia; FIM, Functional Independence Measure Scale.

a Analysis made including drop out participants' data assuming the same value measured at baseline.

b T0 = baseline, T1 = post-test, and T2 = follow-up.

c Direction of pair wise difference.

d P-value refers to Asymp. Sig.~ 2-tailed (< 0.05) of post-hoc paired Wilcoxon signed rank test and adjusted significance level at 0.016 using Bonferroni correction.

e Effect size or magnitude of observed changes within each pair wise comparison of Wilcoxon Signed-Rank Test (small effect ≈ 0.1 , medium effect ≈ 0.3 , large effect ≈ 0.5).

Table 6. Results of Balance Control, Ataxia Rate, and Functional Independence Measures Comparison Between the Experimental Group (N = 5) and Control Group (N = 5) a, b

Time Point	Mean Rank	Sum of Rank	U-Value c	Z-Value d	P-Value e	R _M f
SARA (T0)			10	-0.522	0.690	0.23
Experimental	6	30				
Control	5	25				
Mini BESTest (T0)			5.50	-1.471	0.183	0.65
Experimental	6.90	34.50				
Control	4.10	20.50				
FIMs (T0)			11	-0.313	0.841	0.13
Experimental	5.20	26				
Control	5.80	29				
SARA (T1)			8	-0.949	0.397	0.42
Experimental	4.60	23				
Control	6.40	32				
Mini BESTest (T1)			5	-1.571	0.135	0.70
Experimental	7	35				
Control	4	20				
FIMs (T1)			12.50	0.000	1.000	0.00
Experimental	5.50	27.50				
Control	5.50	27.50				
SARA (T2)			3.50	-1.886	0.063	0.84

Experimental	3.70	18.50				
Control	7.30	36.50				
Mini BESTest (T2)			4	-1.786	0.071	0.79
Experimental	7.20	36				
Control	3.80	19				
FIMs (T2)			11	-0.319	0.794	0.14
Experimental	5.80	29				
Control	5.20	26				

^z Abbreviations: SARA, Scale for Assessment and Rating of Ataxia;; FIM, Functional Independence Measure Scale.

^a Analysis made including drop out participants 'data assuming the same value measured at baseline.

^b T0 = baseline, T1 = post-test, and T2 = follow-up.

^c Rank comparison between the groups.

^d Direction of differences between groups.

^e P-values refer to 2-tailed Exact Sig. (< 0.05) of the non-parametric Mann-Whitney U test.

^f Effect size between group at a measure point of Mann-Whitney U test (small effect ≈ 0.1 , medium effect ≈ 0.3 , large effect ≈ 0.5).

5. Discussion

5.1. Feasibility of Wii Intervention

The results obtained from our investigation have yielded valuable insights regarding the practicality and possible advantages of the Wii exergame intervention. The high enrollment rate and the mandatory attendance of all participants in at least one session underscore the significant initial interest in the Wii balance board as a training tool for this specific population. Technology-based solutions, such as the Wii Balance Board, have the potential to be embraced by neurological patients because of their captivating and hands-on qualities (32).

The impressive commitment to the training program among those who completed the minimum requisite training sessions was encouraging, particularly given the challenges individuals with CA might face in consistently participating in physical training programs (33). The high retention rate in the EG compared to the CG further indicates the acceptability and engagement potential of Wii exergames. Such high retention and adherence rates are critical for the success of rehabilitation programs, as sustained active participation is often correlated with better outcomes (34). The intervention's compliance rate, which varied by exergame difficulty levels, suggests that certain Wii Fit Plus balance games may be better suited for skill progression among different CA patients, potentially due to differing motor skills demands (35).

The incidence of adverse events did not create any severe harm to the participants and, therefore, did not require special medical attention under the circumstances. These findings complement prior research indicating that motion sickness and similar symptoms are very common adverse events resulting from virtual reality and exergame-based training (36).

5.2. Participants' Experience Related to the Intervention

Enjoyment is a crucial factor in the success of gamified rehabilitation programs, as it influences adherence and motivation (37). The reported moderate enjoyment level suggests that while Wii exergames were engaging, there is still room to make the sessions more enjoyable for participants. Customization of exergame content to better suit individual preferences could enhance the user experience and potentially improve adherence further (38).

5.3. Preliminary Efficacy

In all outcome measures, including balance control (Mini-BESTest), ataxia rate (SARA), and functional independence (FIMs), participants in both the experimental and CGs significantly improved over time. However, the EG demonstrated greater relative improvements compared to the CG. This indicates that Wii exergame training may offer additional benefits over conventional balance training (39).

Standard-care or conventional balance training involves structured and therapist-driven exercises, which, while effective, have drawbacks due to limited patient engagement and adherence (40). In contrast, the interactive and gamified nature of Wii balance board training introduces an enjoyable and motivating element, which likely contributed to the better improvements observed in the EG. The high effectiveness of interactive rehabilitation methods in enhancing both motor skills and cognitive functions among individuals with neurological conditions has been documented in previous CA studies (41).

The notable enhancement observed in the EG can be attributed to the dynamic and individualized exercises offered by the Wii Fit Plus balance games, which potentially provide more engaging and comprehensive training for the vestibular and proprioceptive systems. These systems are crucial for ataxia patients when designing a

rehabilitation plan, whereas conventional balance training often does not adequately address them (42). Additionally, the real-time visual and auditory feedback provided by the Wii games can enhance motor learning and promote better postural adjustments during training, a vital component that is often lacking in many conventional therapies (43, 44).

Beyond conventional therapy, vestibular rehabilitation is another commonly used method for managing balance disorders, including those associated with CA. Vestibular rehabilitation typically targets gaze stabilization, vestibulo-ocular reflex (VOR) adaptation, and habituation exercises. While effective in improving vestibular compensation, its reliance on static exercises may not fully address dynamic balance challenges or engage proprioceptive systems comprehensively (45). In contrast, Wii Balance Board-based exergame training involves dynamic and multidirectional movements, which likely contribute to training the vestibular and proprioceptive systems in a more integrated and functional manner.

Intensive physiotherapy programs, often involving task-specific training or over-ground walking interventions, are also well-documented for improving postural stability. However, such approaches demand high resource utilization and therapist involvement, making them less feasible for long-term use in outpatient or home-based settings (46, 47). The cost-effectiveness and accessibility of Wii exergame training make it a more practical alternative for regular rehabilitation, where resources are limited.

Recent advancements in ataxia rehabilitation alongside Wii balance board training include r-TMS, which has shown promise in modulating cortical excitability and promoting neuroplasticity. While r-TMS can facilitate specific neural pathways, it does not directly provide functional balance training or engage the motor systems through physical practice (48). Wii balance board exergames, on the other hand, offer a dual benefit by combining physical training with interactive engagement, addressing both motor and cognitive domains simultaneously (49). However, the integration of r-TMS with physical rehabilitation, including exergame training, warrants further exploration to maximize therapeutic potential in future studies.

Though the EG showed significant improvements in pairwise comparisons, it remains unable to reach statistical significance in the Bonferroni-corrected analysis ($P < 0.016$), likely due to limited statistical power. Improvements in ataxia severity rate and balance were observed at follow-up, but not in functional independence. This finding aligns with previous research suggesting that while exergame training effectively improves specific motor skills like balance, it might not directly involve the transfer of acquired skills to the performance of everyday activities (50). Functional independence encompasses a wide range of tasks that may require more than just improved balance, including muscle strength, en-

durance, and coordination (51). Standard physiotherapy programs often integrate such components, which could explain why improvements in functional independence require additional or complementary training (52). Thus, it is suggested that rehabilitation programs for ataxia patients should also incorporate other forms of training that address a broader range of functional abilities.

5.4. Clinical Implementations

5.4.1. Patient Selection and Screening

The current feasibility study ensures that Wii balance board exergame training can be safely administered among patients affected by inherited, acquired, and metabolic forms of ataxia, within the 30 - 60 years age limit. Cognitive (in cases of pre-existing cognitive deficit) and postural control assessments are necessary and should be included in the initial screening of ataxia patients prior to referring them to Wii balance board training to facilitate efficient and safe administration.

5.4.2. Exergame Design and Customization

A six-week intervention with three weekly sessions (minimum 13 sessions) seems to be feasible and well-tolerated in our study. The selection of the Wii fit plus exergames needs to be more patient-centric, and a gradual progression in difficulty level could be more ideal to attain sustained adherence in the rehabilitation program. Patients with severe ataxia should be the targeted community for beginner-level games. However, they could move on to the advanced level as they grasp the present level of difficulty. The professional level of gameplay could not be executed well due to excessive challenges experienced by the participants. Thus, this stage is suggested to be useful only for ataxia patients with a low severity rate and more postural control. Nevertheless, a patient may require additional support and adaptation to play at this level.

5.4.3. Session Structure and Safety Considerations

The Wii balance board training should include practice sessions of Wii Fit Plus games for the patients, along with warm-up exercises, vital sign monitoring, safe patient handling while stepping on the Wii board, and supervised gameplay to ensure safety. To reduce adverse effects (such as dizziness and headache), a gradual adaptation period and longer rest intervals between exergames are recommended. Encouragement strategies, including verbal reinforcement, progress feedback, and motivational incentives, should be implemented to enhance participant engagement and adherence to the training. Dedicated technical staff are required to maintain research equipment properly and oversee treatment implementation in the experimental arm. Additionally, a neurologist with expertise in CA was appointed in our study to monitor and assess any disease-specific adverse events requir-

ing special care.

5.4.4. Integration with Existing Rehabilitation Programs

Wii Balance Board-based exergame training should be incorporated as a supplementary intervention alongside conventional or standard-care physiotherapy for balance training. Given the findings that functional independence did not improve significantly, future rehabilitation programs should combine Wii training with progressive resistance exercises and cardiovascular endurance training to target broader functional domains.

5.5. Conclusions

Despite various limitations, the study provides evidence of the feasibility of implementing Wii balance board training in clinical settings, as well as a modest level of enjoyment experienced by CA patients. This supports the necessity for a more extensive definitive study to further assess the effectiveness of Wii Balance Board-based exergame training. With balanced pragmatic and explanatory elements, the findings are relevant to clinical practice while maintaining scientific rigor. Preliminary findings of efficacy indicate enhancements in balance control and improvement in ataxia symptoms. However, additional study is required to refine the current Wii intervention and maximize its clinical applicability, along with optimizing functional outcomes in the ataxia population.

5.6. Future Research Directions

5.6.1. Larger and Multicenter Trials

Expanding the recruitment of samples beyond a single study center could enhance the generalizability and external validity of the current findings. A more diverse ataxia population with stratified group allocation should be considered to assess variations in response in future studies.

5.6.2. Optimizing Training Duration and Follow-up Assessments

Future trials should extend intervention periods (beyond 6 weeks) to determine the long-term benefits on functional independence. Additional second follow-up assessments should also be conducted to evaluate sustained motor and functional improvements.

5.6.3. Personalization Strategies in Wii Fit Plus Balance Games

It is crucial to investigate adapting difficulty algorithms to tailor exergames to patients' individual progress and motor abilities in future studies.

5.7. Study Limitations

This study has several limitations. First, the small sample size limits statistical power, making it difficult

to detect significant differences, especially after Bonferroni correction. Second, the short intervention period (6 weeks) restricts conclusions about long-term effects on functional independence. The age restriction (30- 60 years) excluded individuals with late-onset degenerative ataxias, limiting the applicability of findings to this population. Complete blinding of both the participants and investigator was not feasible due to the nature of the intervention, posing a risk of subtle performance bias. Additionally, participants might have engaged in other treatment activities outside the study setting or uncertain lifestyle activities at home. Such potential confounding variables might also influence the study's outcomes, although these confounders could not be controlled due to ethical constraints. While adverse events were minimal, individual tolerance to exergame training may vary, necessitating further safety evaluations. Lastly, our study did not include a second follow-up beyond the first follow-up assessment, limiting insights into sustained benefits. Future studies should address these limitations through the inclusion of a more diverse population of ataxia, powered sample sizes, and multicenter trials with extended follow-up, double-blind assessments, and better tracking of additional rehabilitation activities.

Acknowledgments

We express our gratitude to all the participants for their voluntary participation in the study and their cooperation in the intervention and also to the SRM Medical College Hospital and Research Centre (SRM MCH&RC) for providing valuable infrastructure support, which has facilitated the timely execution of our study. We also would like to extend our sincere gratitude to Mr. Balamurugan Janakiraman (Research associate, SRM College of Physiotherapy) for his continuous guidance and support in this research.

Authors' Contribution: S. P.: Conceptualization, methodology, formal analysis, investigation, data curation, writing-original draft, visualization, and funding acquisition; K. R.: Validation, resources, writing- review, and editing; Sh. P. R. P.: Supervision and project administration. All authors reviewed and approved the final version of the manuscript. We comply with the ICMJE criteria for defining authorship.

Clinical Trial Registration: This study was prospectively registered on the Clinical Trial Registry of India (ctri.nic.in) with the registration number CTRI/2023/11/059589.

Conflict of Interests: The authors declare no conflict of interest.

Data Reproducibility: The data presented in this study are available on request to the corresponding author. The data are not publicly available due to privacy reasons of the study participants.

Ethical Approval: This study was approved by the SRM Institutional Ethical Committee (approval No: SRMIEC-ST0523-657) and conducted in accordance with the ethical standards of the Declaration of Helsinki.

Funding/Support: The present study received no funding/support.

Informed Consent: All eligible participants provided written informed consent after receiving detailed explanations about the study's purpose, procedures, potential risks, and benefits.

References

- Manto M, Gandini J, Feil K, Strupp M. Cerebellar ataxias: an update. *Curr Opin Neurol*. 2020;33(1):150-60. [PubMed ID:31789706]. <https://doi.org/10.1097/WCO.0000000000000774>.
- Manto M, Marmolino D. Cerebellar ataxias. *Curr Opin Neurol*. 2009;22(4):419-29. [PubMed ID:19421057]. <https://doi.org/10.1097/WCO.0b013e32832b9897>.
- Rudaks LI, Yeow D, Ng K, Deveson IW, Kennerson ML, Kumar KR. An Update on the Adult-Onset Hereditary Cerebellar Ataxias: Novel Genetic Causes and New Diagnostic Approaches. *Cerebellum*. 2024;23(5):2152-68. [PubMed ID:38760634]. [PubMed Central ID:PMC11489183]. <https://doi.org/10.1007/s12311-024-01703-z>.
- Bhuin S, Biswas S, Roy A, Mukherjee A, Pandit A, Gangopadhyay G. Etiology and Course of Cerebellar Ataxia: A Study from Eastern India. *Med J Dr DY Patil Vidyapeeth*. 2023;16(4):591-8. https://doi.org/10.4103/mjdrdypu.mjdrdypu_314_21.
- Gorcenco S, Karremo C, Puschmann A. Patients' Perspective in Hereditary Ataxia. *Cerebellum*. 2024;23(1):82-91. [PubMed ID:36525215]. [PubMed Central ID:PMC10864479]. <https://doi.org/10.1007/s12311-022-01505-1>.
- Stanley WJ, Kelly CKL, Tung CC, Lok TW, Ringo TMK, Ho YK, et al. Cost of Cerebellar Ataxia in Hong Kong: A Retrospective Cost-of-Illness Analysis. *Front Neurol*. 2020;11:711. [PubMed ID:32765413]. [PubMed Central ID:PMC7380245]. <https://doi.org/10.3389/fneur.2020.00711>.
- Lowit A, Greenfield J, Cutting E, Wallis R, Hadjivassiliou M. Symptom burden of people with progressive ataxia, and its wider impact on their friends and relatives: a cross-sectional study. *AMRC Open Res*. 2021;3:28. [PubMed ID:38708068]. [PubMed Central ID:PMC11064976]. <https://doi.org/10.12688/amrcopenres.13036.1>.
- Lopez-Bastida J, Perestelo-Perez L, Monton-Alvarez F, Serrano-Aguilar P. Social economic costs and health-related quality of life in patients with degenerative cerebellar ataxia in Spain. *Mov Disord*. 2008;23(2):212-7. [PubMed ID:17999424]. <https://doi.org/10.1002/mds.21798>.
- Milne SC, Roberts M, Williams S, Chua J, Grootendorst AC, Agostinelli G, et al. Goal-Directed Rehabilitation Versus Standard Care for Individuals with Hereditary Cerebellar Ataxia: A Multicenter, Single-Blind, Randomized Controlled Superiority Trial. *Ann Neurol*. 2025;97(3):409-24. [PubMed ID:39520242]. <https://doi.org/10.1002/ana.27130>.
- Grobe-Einsler M, Bork F, Faikus A, Hurlmann R, Kaut O. Effects of cerebellar repetitive transcranial magnetic stimulation plus physiotherapy in spinocerebellar ataxias - A randomized clinical trial. *CNS Neurosci Ther*. 2024;30(6):e14797. [PubMed ID:38887169]. [PubMed Central ID:PMC11183922]. <https://doi.org/10.1111/cns.14797>.
- Mitoma H, Manto M, Gandini J. Recent Advances in the Treatment of Cerebellar Disorders. *Brain Sci*. 2019;10(1). [PubMed ID:31878024]. [PubMed Central ID:PMC7017280]. <https://doi.org/10.3390/brainsci10010011>.
- Gupta A, Prakash NB, Rahman H. Rehabilitation in Ataxia. *Indian J Physical Med Rehabil*. 2023;33(1):21-9. https://doi.org/10.4103/ijpmr.ijpmr_42_22.
- Ringgenberg N, Mildner S, Hapig M, Hermann S, Kruszewski K, Martin-Niedeken AL, et al. ExerG: adapting an exergame training solution to the needs of older adults using focus group and expert interviews. *J Neuroeng Rehabil*. 2022;19(1):89. [PubMed ID:35974409]. [PubMed Central ID:PMC9382774]. <https://doi.org/10.1186/s12984-022-01063-x>.
- Rohof B, Betsch M, Rath B, Tingart M, Quack V. The Nintendo((R)) Wii Fit Balance Board can be used as a portable and low-cost posturography system with good agreement compared to established systems. *Eur J Med Res*. 2020;25(1):44. [PubMed ID:32972447]. [PubMed Central ID:PMC7517684]. <https://doi.org/10.1186/s40001-020-00445-y>.
- Ayvat E, Onursal Kilinc O, Ayvat F, Savcun Demirci C, Aksu Yildirim S, Kursun O, et al. The Effects of Exergame on Postural Control in Individuals with Ataxia: a Rater-Blinded, Randomized Controlled, Cross-over Study. *Cerebellum*. 2022;21(1):64-72. [PubMed ID:33973141]. [PubMed Central ID:PMC8110432]. <https://doi.org/10.1007/s12311-021-01277-0>.
- Barbuto S, Martelli D, Isirame O, Lee N, Bishop L, Kuo SH, et al. Phase I Single-Blinded Randomized Controlled Trial Comparing Balance and Aerobic Training in Degenerative Cerebellar Disease. *PM R*. 2021;13(4):364-71. [PubMed ID:32383352]. [PubMed Central ID:PMC7647960]. <https://doi.org/10.1002/pmrj.12401>.
- Franzo M, Pica A, Pascucci S, Serrao M, Marinozzi F, Bini F. A Proof of Concept Combined Using Mixed Reality for Personalized Neurorehabilitation of Cerebellar Ataxic Patients. *Sensors (Basel)*. 2023;23(3). [PubMed ID:36772721]. [PubMed Central ID:PMC9920853]. <https://doi.org/10.3390/s23031680>.
- Winser S, Chan HK, Chen WK, Hau CY, Leung SH, Leung YH, et al. Effects of therapeutic exercise on disease severity, balance, and functional Independence among individuals with cerebellar ataxia: A systematic review with meta-analysis. *Physiother Theory Pract*. 2023;39(7):1355-75. [PubMed ID:35212247]. <https://doi.org/10.1080/09593985.2022.2037115>.
- Pacheco TBF, de Medeiros CSP, de Oliveira VHB, Vieira ER, de Cavalcanti FAC. Effectiveness of exergames for improving mobility and balance in older adults: a systematic review and meta-analysis. *Syst Rev*. 2020;9(1):163. [PubMed ID:32682439]. [PubMed Central ID:PMC7368979]. <https://doi.org/10.1186/s13643-020-01421-7>.
- Pilotto F, Del Bondio A, Puccio H. Hereditary Ataxias: From Bench to Clinic, Where Do We Stand? *Cells*. 2024;13(4). [PubMed ID:38391932]. [PubMed Central ID:PMC10886822]. <https://doi.org/10.3390/cells13040319>.
- Paul NW, Mahdiani H. Ethical challenges in clinical studies with adaptive design in oncology. *Clinical Ethics*. 2022;18(2):148-54. <https://doi.org/10.1177/1477509221133974>.
- Eldridge SM, Chan CL, Campbell MJ, Bond CM, Hopewell S, Thabane L, et al. CONSORT 2010 statement: extension to randomised pilot and feasibility trials. *BMJ*. 2016;355:i5239. [PubMed ID:27777223]. [PubMed Central ID:PMC5076380]. <https://doi.org/10.1136/bmj.i5239>.
- Moore CG, Carter RE, Nietert PJ, Stewart PW. Recommendations for planning pilot studies in clinical and translational research. *Clin Transl Sci*. 2011;4(5):332-7. [PubMed ID:22029804]. [PubMed Central ID:PMC3203750]. <https://doi.org/10.1111/j.1752-8062.2011.00347.x>.
- Loudon K, Treweek S, Sullivan F, Donnan P, Thorpe KE, Zwarenstein M. The PRECIS-2 tool: designing trials that are fit for purpose. *BMJ*. 2015;350:h2147. [PubMed ID:25956159]. <https://doi.org/10.1136/bmj.h2147>.
- Darker C, Loudon K, O'Connell N, Castello S, Burke E, Vance J, et al. An application of PRECIS-2 to evaluate trial design in a pilot cluster randomised controlled trial of a community-based smoking cessation intervention for women living in disadvantaged areas of Ireland. *Pilot Feasibility Stud*. 2022;8(1):19. [PubMed ID:35078530]. [PubMed Central ID:PMC8787878]. <https://doi.org/10.1186/s40814-022-00969-6>.
- Miyata K, Kondo Y, Bando K, Hara T, Takahashi Y. Structural Validity of the Mini-Balance Evaluation Systems Test in Individuals With Spinocerebellar Ataxia: A Rasch Analysis Study. *Arch Phys Med Rehabil*. 2024;105(4):742-9. [PubMed ID:38218308]. <https://doi.org/10.1016/j.apmr.2023.12.015>.
- Lawerman TF, Brandsma R, Verbeek RJ, van der Hoeven JH, Lunsing RJ, Kremer HPH, et al. Construct Validity and Reliability of the SARA Gait and Posture Sub-scale in Early Onset Ataxia. *Front Hum Neurosci*. 2017;11:605. [PubMed ID:29326569]. [PubMed Central ID:PMC5733344]. <https://doi.org/10.3389/fnhum.2017.00605>.
- Kim BR, Lim JH, Lee SA, Park S, Koh SE, Lee IS, et al. Usefulness of the Scale for the Assessment and Rating of Ataxia (SARA) in Ataxic Stroke Patients. *Ann Rehabil Med*. 2011;35(6):772-80. [PubMed ID:22506205]. [PubMed Central ID:PMC3309386]. <https://doi.org/10.1186/s40001-020-00445-y>.

- org/10.5535/arm.2011.35.6.772.
29. Weyer A, Abele M, Schmitz-Hubsch T, Schoch B, Frings M, Timmann D, et al. Reliability and validity of the scale for the assessment and rating of ataxia: a study in 64 ataxia patients. *Mov Disord.* 2007;22(11):1633-7. [PubMed ID:17516493]. <https://doi.org/10.1002/mds.21544>.
 30. Matsugi A. Physical Therapy for Cerebellar Ataxia. In: Suzuki T, editor. *Neurological Physical Therapy*. Rijeka, Croatia: IntechOpen; 2017.
 31. National Center for Complementary and Integrative Health. Pilot Studies: Common Uses and Misuses. National Center for Complementary and Integrative Health; 2024. [Cited:2024]. Available from: <https://www.nccih.nih.gov/grants/pilot-studies-common-uses-and-misuses>.
 32. Moulaei K, Sharifi H, Bahaadinbeigy K, Dinari F. Efficacy of virtual reality-based training programs and games on the improvement of cognitive disorders in patients: a systematic review and meta-analysis. *BMC Psychiatry.* 2024;24(1):116. [PubMed ID:38342912]. [PubMed Central ID:PMC10860230]. <https://doi.org/10.1186/s12888-024-05563-z>.
 33. Nagar A, Vij J S. Optimizing rehabilitation strategies for cerebellar ataxia in indian population : A systematic review of physical therapy interventions. *Int J Sci Res.* 2023;12.
 34. Pajaro-Blazquez M, Pons JL. Research highlights in neurorehabilitation. *J Neuroeng Rehabil.* 2014;11:21. [PubMed ID:24594120]. [PubMed Central ID:PMC3996103]. <https://doi.org/10.1186/1743-0003-11-21>.
 35. Lin CR, Kuo SH, Opal P. Cognitive, Emotional, and Other Non-motor Symptoms of Spinocerebellar Ataxias. *Curr Neurol Neurosci Rep.* 2024;24(3):47-54. [PubMed ID:38270820]. [PubMed Central ID:PMC10922758]. <https://doi.org/10.1007/s11910-024-01331-4>.
 36. Conner NO, Freeman HR, Jones JA, Luczak T, Carruth D, Knight AC, et al. Virtual Reality Induced Symptoms and Effects: Concerns, Causes, Assessment & Mitigation. *Virtual Worlds.* 2022;1(2):130-46. <https://doi.org/10.3390/virtualworlds1020008>.
 37. Dan B. Gamification of therapy: the fun factor in rehabilitation. *Dev Med Child Neurol.* 2022;64(3):276. [PubMed ID:35120264]. <https://doi.org/10.1111/dmcn.15126>.
 38. Alexiou A, Schippers MC. Digital game elements, user experience and learning: A conceptual framework. *Educ Inform Technol.* 2018;23(6):2545-67. <https://doi.org/10.1007/s10639-018-9730-6>.
 39. Bonanno M, De Pasquale P, De Marchis C, Lombardo Facciale A, Paladina G, Fonti B, et al. Might patients with cerebellar ataxia benefit from the Computer Assisted Rehabilitation ENvironment (CAREN)? A pilot study focusing on gait and balance. *Front Bioeng Biotechnol.* 2024;12:1385280. [PubMed ID:39011156]. [PubMed Central ID:PMC11247328]. <https://doi.org/10.3389/fbioe.2024.1385280>.
 40. Loomis KJ, Roll SC, Hardison ME. The role of therapist-patient relationships in facilitating engagement and adherence in upper extremity rehabilitation. *Work.* 2023;76(3):1083-98. [PubMed ID:37248936]. <https://doi.org/10.3233/WOR-220384>.
 41. Marin-Medina DS, Arenas-Vargas PA, Arias-Botero JC, Gomez Vasquez M, Jaramillo-Lopez MF, Gaspar-Toro JM. New approaches to recovery after stroke. *Neurol Sci.* 2024;45(1):55-63. [PubMed ID:37697027]. [PubMed Central ID:PMC10761524]. <https://doi.org/10.1007/s10072-023-07012-3>.
 42. Heusel-Gillig LL, Hall CD. Effectiveness of Vestibular Rehabilitation for Patients with Degenerative Cerebellar Ataxia: A Retrospective Cohort Study. *Brain Sci.* 2023;13(11). [PubMed ID:38002480]. [PubMed Central ID:PMC10669586]. <https://doi.org/10.3390/brainsci13111520>.
 43. Ghazavi Dozin SM, Mohammad Rahimi N, Aminzadeh R. Wii Fit-Based Biofeedback Rehabilitation Among Post-Stroke Patients: A Systematic Review and Meta-Analysis of Randomized Controlled Trial. *Biol Res Nurs.* 2024;26(1):5-20. [PubMed ID:37247514]. <https://doi.org/10.1177/10998004231180316>.
 44. Alves de Souza G. The Benefits of Game Therapy in Neurological Physiotherapeutic Treatment. *Health Soc.* 2023;3(03):149-75. <https://doi.org/10.51249/hs.v3i03.1365>.
 45. Andreev A, Mindova S. Advancements in Vestibular Physiotherapy: A Comprehensive Review. *J IMAB - Ann Proceed (Sci Papers).* 2024;30(4):5829-33. <https://doi.org/10.5272/jimab.2024304.5829>.
 46. Paillard T. The optimal method for improving postural balance in healthy young and older people: specific training for postural tasks encountered in personal physical practice. *Front Physiol.* 2023;14:1188496. [PubMed ID:37449015]. [PubMed Central ID:PMC10338096]. <https://doi.org/10.3389/fphys.2023.1188496>.
 47. Mohammed AA, Ghalib Abdulah A, Abdulkarim Sulaiman A. Investigate the Principles and Applications of Task-Specific Training in Neuro-Physiotherapy Rehabilitation. *J Adv Scholarly Res Allied Educ.* 2024;21(3):153-8. <https://doi.org/10.29070/ms9xcn64>.
 48. Afonso M, Sanchez-Cuesta F, Gonzalez-Zamorano Y, Pablo Romero J, Vourvopoulos A. Investigating the synergistic neuromodulation effect of bilateral rTMS and VR brain-computer interfaces training in chronic stroke patients. *J Neural Eng.* 2024;21(5). [PubMed ID:39419104]. <https://doi.org/10.1088/1741-2552/ad8836>.
 49. Graciani Z, Moraes IAP, Alberissi CAO, Prado-Rico JM, Silva TDD, Martinez JP, et al. The effect of different interfaces during virtual game practice on motor performance of individuals with genetic ataxia: A cross-sectional study. *PLoS One.* 2024;19(11):e0312705. [PubMed ID:39485822]. [PubMed Central ID:PMC11530066]. <https://doi.org/10.1371/journal.pone.0312705>.
 50. Bonney E, Jelsma LD, Ferguson GD, Smits-Engelsman BC. Learning better by repetition or variation? Is transfer at odds with task specific training? *PLoS One.* 2017;12(3):e0174214. [PubMed ID:28333997]. [PubMed Central ID:PMC5363924]. <https://doi.org/10.1371/journal.pone.0174214>.
 51. Mlinac ME, Feng MC. Assessment of Activities of Daily Living, Self-Care, and Independence. *Arch Clin Neuropsychol.* 2016;31(6):506-16. [PubMed ID:27475282]. <https://doi.org/10.1093/arclin/acw049>.
 52. Newell A, Cherry S, Fraser M. Principles of Rehabilitation: Occupational and Physical Therapy. In: Nowicki PD, editor. *Orthopedic Care of Patients with Cerebral Palsy: A Clinical Guide to Evaluation and Management across the Lifespan*. Cham, Germany: Springer International Publishing; 2020. p. 221-50.