Robot-assisted arm training in patients with Parkinson’s disease: a pilot study

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Abstract

Background: Despite the growing diffusion of robotic devices in neurorehabilitation, no previous study investigated the effects of robotic training on arm impairment due to Parkinson’s disease. The aim of this pilot study was to evaluate whether robot-assisted arm training might improve upper limb function in patients with Parkinson’s disease.

Findings: Ten patients with Parkinson’s disease (Hoehn and Yahr stage 2.5-3) received ten, 45-minute, treatment sessions, five days a week, for two consecutive weeks. Robot-assisted arm training was performed with the Bi-ManuTrack (Reha-Stim, Berlin, Germany) that provides a computer-controlled, repetitive, bilateral, mirror-like practice of forearm pronation/supination and wrist extension/flexion. Patients were trained according to the following modalities: passive-passive (both arms moved by the machine) and active-active (both arms actively moving against resistance). The dominant upper limb was evaluated before and immediately after treatment as well as at two weeks of follow-up. Outcomes were the nine-hole peg test, the Fugl-Meyer assessment (upper limb section) and the Unified Parkinson’s Disease Rating Scale. After treatment, a significant improvement was found in the nine-hole peg test ($P = 0.007$) as well as in the upper limb section of the Fugl-Meyer assessment ($P = 0.012$). Findings were confirmed at the 2-week follow-up evaluation only for the nine-hole peg test ($P = 0.007$). No significant improvement was found in the Unified Parkinson’s Disease Rating Scale at both post-treatment and follow-up evaluations.

Conclusions: Our findings support the hypothesis that robot-assisted arm training might be a promising tool in order to improve upper limb function in patients with Parkinson’s disease.

Keywords: Rehabilitation, Physical therapy, Basal ganglia

Introduction

Parkinson’s disease (PD) is an idiopathic neurodegenerative disorder due to a progressive loss of dopaminergic neurons in the substantia nigra pars compacta [1]. Typical features of PD are bradykinesia (slowed movement), hypokinesia (poverty of movement), rigidity and resting tremor [1,2]. Even if impaired manual dexterity with progressive limitations in reaching, grasping and fine motor tasks has been described in PD, to date evidence base for upper limb intervention strategies in parkinsonian patients is lacking [3].

Robotic arm training (RAT) was found to effectively improve upper limb function in patients with neurological disorders, such as stroke [4]. As to PD, forced use, task-specific, intensive, training programs based on robotic devices were found to effectively improve lower limb function [5-10]. Despite the growing diffusion of robotic devices in neurorehabilitation, to date no previous study investigated the effects of robotic training on arm impairment due to PD. The aim of this pilot study was to evaluate whether RAT might improve upper limb function in patients with PD.

Methods

This study was performed in the Neurorehabilitation Unit of the Azienda Ospedaliera-Universitàrìa Integrata of Verona, Italy. Inclusion criteria: confirmed diagnosis of idiopathic PD according to the UK Brain Bank Criteria [11]; Hoehn and Yahr (H&Y) stage of 2.5 or 3 determined in the “on” phase [12]; Mini Mental State Examination score $>24$ [13]. Exclusion criteria: severe dyskinesias or “on-off” fluctuations; change of PD medication during the
study; deficits of somatic sensation involving the upper limbs; other neurological or orthopedic conditions involving the upper limbs.

All participants were outpatients and gave their informed written consent for participation in the study, which was carried out according to the Declaration of Helsinki and was approved by the Ethics Committee of the Department of Neurological and Movement Sciences of Verona University.

During the study, participants were instructed to take their normal PD medications: they were tested and trained during the on phase (1 to 2.5 hours after taking morning dose). Participants did not perform any type of rehabilitation in the three months before the study, nor undergo any form of training other than that scheduled in the study protocol.

**Treatment procedures**

Each patient underwent a training program consisting of ten, 45-minute sessions (including rest periods), five days a week (from Monday to Friday) for two consecutive weeks. Robot-assisted arm training was performed with the Bi-Manu-Track (Reha-Stim, Berlin, Germany) that provides a computer-controlled, repetitive, bilateral, mirror-like practice of forearm pronation/supination and wrist extension/flexion according to three modalities: passive-passive (both arms moved by the machine), active-passive (one arm driving the other), and active-active (both arms actively moving against resistance) [14]. As shown in Figure 1, patients sat at a height-adjustable table with their elbow bent at 90°, putting their forearms into an arm trough and grasping a handle (written informed consent for the publication of this image was obtained). Each training session consisted of two parts with a 5-minute rest between them. First we trained forearm pronation/supination for 20 minutes: 10 minutes of passive-passive mode (100 repetitions) followed by 10 minutes of active-active mode (100 repetitions). Then we trained wrist extension/flexion for 20 minutes: 10 minutes of passive-passive mode (100 repetitions) followed by 10 minutes of active-active mode (100 repetitions). Amplitude and resistance were set individually.

**Testing procedures**

Patients were evaluated before (T0), immediately after treatment (T1) (primary endpoint) and at two weeks of follow-up (T2). The same rater evaluated all patients.

**Outcome measures**

The nine-hole peg test (NHPT) was used to assess dominant hand dexterity. We required patients to take 9 pegs from a container placing them into 9 holes on a board and vice versa as quickly as possible. Score was the time taken to complete the test activity [15].

The Fugl-Meyer assessment (FM) was used to evaluate dominant upper limb motor ability to perform selective movements. The FM upper limb section allows a maximum score of 66, with sub-scores of 36 for the upper arm, 10 for the wrist, 14 for the hand, and 6 for coordination and speed of movement [16,17].

The Unified Parkinson’s Disease Rating Scale (UPDRS) was used to measure disease severity in PD. It has three subscales: I—mentation, behavior, and mood (range 0–16); II—activities of daily living (range 0–52); III—motor examination (range 0–108). Total score is the sum of these subscales (range 0–176) [18,19].

**Statistical analysis**

Statistical analysis was carried out with SPSS 21.0 (SPSS Inc, Chicago, IL). The Friedman test was used to analyze overall changes in performance between the different evaluation sessions. In the presence of significant main effects, the Wilcoxon signed ranks test was used on the T1 vs. T0, T2 vs. T0 and T2 vs. T1 comparisons to determine any significant difference. Descriptive analysis was used to evaluate the effect size measures (Cohen’s d calculation) and the 95% confidence intervals [20]. The level for significance was $P < 0.05$. The Bonferroni correction was used when investigating multiple comparisons ($P < 0.016$) [21].

**Results**

Ten right-handed subjects (7 males and 3 females; mean age 70.7 years) presenting with idiopathic PD (mean duration 7.1 years) were recruited from among 18 outpatients consecutively admitted to our Neurorehabilitation Unit.
from April to October 2012. No dropout was observed. No adverse event occurred during the study. Row data (medians and interquartile ranges) of patients’ performance at T0, T1 and T2 evaluations are reported in Table 1.

As to the NHPT, a significant overall change was found between the three time points ($P = 0.007$; $\chi^2 = 9.800$). A significant difference was found at both T1 vs. T0 ($P = 0.007$) and T2 vs. T0 ($P = 0.007$) comparisons. As to the FM, a significant overall change was found between the three time points ($P = 0.003$; $\chi^2 = 11.400$). A significant difference was found only at T1 vs. T0 ($P = 0.012$) comparison. As to the UPDRS, no significant overall change was found between the three time points ($P = 0.062$; $\chi^2 = 5.568$) at the Friedman test. Treatment effects in all outcome measures are reported in Table 2.

**Discussion**

Our results show that ten, 45-minute sessions of RAT may improve fine and gross motor function of the dominant upper limb in patients with PD (H&Y 2.5–3). As measured by the NHPT, improvements of fine motor function were maintained also at the follow-up examination. Conversely, no significant change was found in the UPDRS.

In people with PD, altered upper limb function generally manifests as impaired timing and force modulation, progressively affecting the quality of hand movement [3]. Even if intensive, task-specific, practice has been proposed to reduce arm impairment due to PD, the most effective rehabilitative approach in order to best facilitate upper limb skill learning has not yet been defined [3]. Our preliminary findings about the role of RAT in PD are in keeping with those evidences about the effective use of robotic devices provide task-specific, intensive, training programs in patients with progressive lower limb functional impairment due to PD [5–10]. Furthermore, our findings are in line with those of Lee and colleagues, which examined the effects of constraint-induced movement therapy in twenty patients with PD (H&Y 2–3), observing significant improvements of fine and gross motor performance of the upper limb after twenty, 3-hour, treatment sessions [22].

In order to understand why RAT showed to improve upper limb function in PD, we hypothesize that several repetitions of rhythmic arm movements could act as an external proprioceptive cue, by reinforcing the neuronal circuits that contribute to the upper limb movements. In particular, RAT provides an external rhythm that could improve motor output bypassing the deficient internal motor generation system (including the supplementary motor area and basal ganglia) that would support the generation on actions based on intention and internal reference frame [22]. In addition, it is plausible that the strengthening effect of RAT would play a role. In line with this issue, a previous case series study by Combs and colleagues, described a significantly reduction of disability as well as an improvement of quality of life after a training program based on 24 to 36 boxing sessions in 6 patients with PD (H&Y 1–4) [23]. Unfortunately, the Authors did not evaluate upper limbs function before.

### Table 1 Row data of patients’ performance in all outcome measures

| Outcomes                      | Before treatment Median (IQR) | After treatment Median (IQR) | Follow-up Median (IQR) |
|-------------------------------|-------------------------------|------------------------------|------------------------|
| Nine-hole peg test (s)        | 16.20 (15.80 to 18.18)        | 13.80 (13.06 to 14.92)       | 13.80 (13.08 to 15.13) |
| Fugl-Meyer assessment (0–66)  | 60.00 (57.25 to 62.00)        | 65.00 (62.50 to 65.75)       | 63.50 (62.00 to 65.75) |
| UPDRS motor examination (0–108)| 16.00 (12.25 to 18.50)        | 14.50 (13.25 to 15.75)       | 15.00 (13.25 to 16.00) |
| UPDRS total (0–176)           | 34.50 (25.27 to 37.75)        | 30.00 (24.00 to 34.00)       | 27.50 (23.00 to 34.00) |

Abbreviations: IQR Interquartile range, s seconds, UPDRS Unified Parkinson’s Disease Rating Scale.

### Table 2 Treatment effects in all outcome measures

| Outcomes                      | Comparisons | 95% Confidence interval (Effect size) |
|-------------------------------|-------------|--------------------------------------|
|                               | Wilcoxon signed ranks test | T1 vs. T0 (P value (Z)) | T2 vs. T0 (P value (Z)) | T2 vs. T1 (P value (Z)) | T1 vs. T0 (r) | T2 vs. T0 (r) | T2 vs. T1 (r) |
| Nine-hole peg test (s)        | 0.007 (−2.701)* | 0.007 (−2.701)* | 0.359 (−0.918) | 1.90 to 4.78 (−0.60) | 1.73 to 4.34 (−0.53) | −1.07 to 0.47 (0.08) |
| Fugl-Meyer assessment (0–66)  | 0.012 (−2.527)* | 0.018 (−2.371) | 0.606 (−0.516) | −6.31 to −1.28 (0.45) | −5.53 to −1.26 (0.41) | −1.62 to 2.42 (−0.07) |
| UPDRS motor examination (0–108)| 0.097 (−1.658) | 0.174 (−1.358) | 0.334 (−0.966) | −0.34 to 2.54 (−0.14) | −0.40 to 2.00 (−0.11) | −1.05 to 0.45 (0.04) |
| UPDRS total (0–176)           | 0.046 (−1.995) | 0.037 (−2.082) | 0.813 (−0.214) | 0.26 to 6.73 (−0.20) | 0.54 to 6.65 (−0.20) | −1.30 to 1.50 (−0.01) |

Abbreviations: UPDRS Unified Parkinson’s Disease Rating Scale. *= statistically significant after Bonferroni correction ($P < 0.016$).
and after treatment [23]. Thus, we cannot directly compare their data with ours.

This pilot study was limited by the lack of a control group and the small sample size. Furthermore, there is no long-term follow-up evaluation and no assessment of activities of daily living.

Conclusions

Our preliminary findings support the hypothesis that RAT might be a promising tool in order to improve upper limb function in patients with PD. However, there is the possibility that changes observed in this pilot study might be due to a placebo effect. Furthermore, it would be useful to evaluate RAT not only in terms of effectiveness but also in terms of costs and time taken to prepare the treatment setting. On these bases, future, properly sized, randomized controlled trials dealing with RAT compared to conventional/non-robotic rehabilitation are needed in order to further validate our results.

Abbreviations

PD: Parkinson's disease; RAT: Robotic arm training; H&Y: Hoehn and Yahr; NHPT: Nine-hole peg test; FM: Fugl-Meyer assessment; UPDRS: Unified Parkinson's Disease Rating Scale.

Competing interests

The authors declare that they have no competing interest.

Authors’ contributions

AP conceived the study, made substantial contribution to its design, performed the statistical analysis and drafted the manuscript. ST participated in the design of the study and helped to draft the manuscript. MP carried out the acquisition of data. AW revised critically the manuscript for important intellectual content. NS coordinated the study and revised critically the manuscript for important intellectual content. All authors read and approved the final manuscript.

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