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Cognitive Rehabilitation in Parkinson's Disease: A systematic review

Hamad Alzahrani [a], Annalena Venneri [b]

a. National Neuroscience Institute, King Fahad Medical City, Riyadh, Saudi Arabia
b. Department of Neuroscience, Faculty of Medicine, University of Sheffield, UK

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Address for correspondence:
Dr Hamad Alzahrani
National Neuroscience Institute
King Fahad Medical City
P.O. Box 59046, Riyadh 11525, Saudi Arabia
E-mail: hsalzahrani@kfmc.med.sa
Tel. +966 11 2889999 ext. 18781

Abstract

Introduction: Cognitive impairments are the most common non-motor symptoms in Parkinson's Disease (PD). These symptoms have a negative impact on patients' quality of life and daily living activities. This review will focus on published articles that investigated the efficacy of cognitive rehabilitation in PD.

Objectives: To review the existing literature on the efficacy of cognitive rehabilitation in PD and highlight the most effective form of intervention to prevent cognitive decline. This review will also point out any limitations and provide directions for future research.

Methods: Published articles available in the Web of Science and PubMed databases up to November 2017 were reviewed for possible inclusion. We identified 15 articles that examined the effects of cognitive rehabilitation in PD and met inclusion criteria.

Results: The main outcomes of this review indicated that, although previous studies used different cognitive rehabilitation methodologies, all studies reported cognitive improvements on at least one cognitive domain. Additionally, the most frequent cognitive domains showing improvements are executive functions and attention.

Conclusion: This review reports the outcomes of studies that examined the effectiveness of cognitive rehabilitation in PD. It also points out the limitations of the studies indicating the limited availability of follow up data on the long-term effects of cognitive interventions. The review also highlights the fact that some of the studies did not include a PD group who did not undergo training. There remains, therefore, a need for longitudinal studies to investigate the potential long term benefits of cognitive training. In addition, future investigations should examine whether any disease characteristics such as disease stage, degree of cognitive impairment and/or the dominant side (right/left) or specific motor symptoms (rigidity/tremor) influence treatment efficacy.

Keywords: cognitive rehabilitation; cognitive training; executive function; attention;

Cognitive impairments are the most common non-motor symptoms in Parkinson's Disease (PD) [1, 2]. These symptoms have a negative impact on patients' quality of life and interfere with daily living activities [3]. It has been reported that the prevalence of mild cognitive impairment (MCI) among PD patients ranges from 17% to 53% [1]. Patients who have MCI could be at higher risk of developing dementia. The prevalence of dementia associated with the progression of PD ranges from 48% to 78% [4]. The most frequent cognitive domains affected in patients with PD are executive functions such as planning and shifting abilities, working memory [1, 5, 6], episodic memory [6, 7], attention and visuo-spatial skills [1, 6]. There is a growing interest in research in the development of strategies to prevent more severe cognitive decline in those patients with PD who show mild cognitive impairment or in having ways to stabilise or improve cognitive dysfunctions when they occur.

Since there is no approved pharmacological treatment for cognitive decline in PD, the possibility of using non-pharmacological interventions for improving cognitive functions in this patient population was recently introduced [8]. These non-pharmacological interventions include cognitive training, physical exercise and the combination of both. So far, little is known about the efficacy of these varieties of interventions on cognitive deficits in PD, and several researchers have emphasised the need for effective techniques particularly for long-term efficacy [9].

To our knowledge there are only five review articles on this topic. One is a general review of non-pharmacological interventions in PD [10], and another included only four studies that have investigated the effect of cognitive training on a single cognitive domain (executive function) in this patient population [11]. The most recent reviews all had a specific focus. The Leung et al [12] included only randomized controlled trials (RCT) studies, the Walton et al [13] reviewed articles that focused only on interventions implemented late in the disease course and computer-based cognitive training techniques, while the van der Weijer et al [14] reported on their theoretical perspective of cognitive training in PD more than performing a systematic

review. In summary, most of the articles reporting the findings of studies of cognitive rehabilitation in PD have been published after the first two reviews were published [8, 9, 15-19], while the most recent ones [12-14] had a more specific focus rather than including a thorough review of most available studies. None of the previous review articles focused on cognitive rehabilitation of PD in all disease stages, different cognitive training techniques as well as including randomized controlled trials and other methodological designs. There is, therefore, a need for an up to date systematic review of studies that have focused on cognitive rehabilitation in patients with PD and that overcomes all limitations mentioned above.

Although the testing of the efficacy of cognitive rehabilitation in patients with PD is still in the early stages, all of the previous studies have revealed a positive impact of cognitive interventions in PD patients [8, 9, 15-27]. However, most published studies have used different methodologies including differences in patients' characteristics and experimental design. There is a growing number of studies that has investigated the effect of various non-pharmacological interventions in PD. This review will focus exclusively on the effect of cognitive rehabilitation in this group of patients. Therefore, the aim of this systematic review was to search for all the existing literature that studied the efficacy of cognitive rehabilitation in patients with PD and to highlight which method seems to be the most effective on preventing cognitive decline. Furthermore, this review will attempt to determine which cognitive skills are more susceptible to the benefits of cognitive rehabilitation, will point out any limitations of existing approaches and will provide directions for future research.

Method

A systematic review of published research articles that have focused on cognitive rehabilitation interventions in patients with PD was carried out. An on-line literature search of the PubMed and Web of Science databases was carried out using the term Parkinson disease with each of the following: cognitive stimulation, cognitive rehabilitation, cognitive enhancement and cognitive training. We also used the term Parkinson disease in combination with each of the above terms

and with each of the following: language, executive function, memory, attention, working memory, learning and problem solving. All published papers up to November 2017 were searched. The initial search identified 791 titles and abstracts. Then 478 duplicate publications were excluded. The abstracts and complete reports were reviewed to eliminate articles according to the following exclusion criteria: (1) review articles, (2) not cognitive intervention, (3) papers that included participants with other neurological condition, (4) studies of healthy participants, (5) reports published only in abstract format, (6) non-peer reviewed articles, (7) case reports, and (8) articles written not in the English language (see Figure 1 and Table 1). A total of 15 articles met our inclusion criteria, those articles had to: have cognitive rehabilitation/training as their main focus, be studies of cognitive rehabilitation/training even if they were not RCT, be studies that included all PD stages and used different cognitive training techniques (not only computer-based). This methodological decision was taken to give a full picture of the kind of cognitive interventional studies currently available for this patient population. These articles were assessed for scientific suitability for inclusion in the present review, by using a set of 12 criteria adapted from Welton et al. [28] (these criteria are listed in Table 2). Each article was rated from 0 to 12.

- Insert Figure 1 and Table 1 about here -

Results

The process of literature search is illustrated in Figure 1. In total, 791 studies were reviewed including duplicate publications from the two databases. After we excluded duplicate papers, 313 full copies were retrieved and evaluated for eligibility. Initially, we identified 17 articles that reported intervention/rehabilitation studies in PD. However, a closer inspection of the full papers identified two articles that did not match the main inclusion criterion as they reported the findings of non-cognitive interventions and were excluded on that basis. There were 15 studies included in this review and the time span of search was from 2004 to 2017. Cognitive tests, cognitive domains targeted, techniques/design, outcome measures, duration and

frequency of training and results are summarized in Table 1. Table 2 includes a quality

assessment of the reviewed articles.

- Insert Table 2 about here -

An in depth review of the findings of the included studies is given below. We will first focus on

intervention parameters and design, domains that were studied, and then will review the

studies according to outcome measures looking, in turn, at cognitive, imaging and mood, fatigue

and quality of life as outcome measures.

Intervention Parameters and Design

The length of all cognitive rehabilitation/training interventions ranged from 3 weeks to 6

months, with the number of sessions varying from 4 to 180. The frequency of these training

sessions was from once a week to every day over a period of 6 months. Only two studies tested

the long-term effects of cognitive intervention: one study had a follow up after 6 months,

whereas the other had a follow up after one year. Furthermore, there was heterogeneity and

variations of the intervention strategies between studies. Most of the studies used computerized

based training programs, whereas other forms of treatment included different strategies such as

the use of paper-and-pencil methods or multimodal cognitive rehabilitation. Moreover, in four

studies cognitive training was compared to other active treatments (e.g. specific versus non-

specific or structured versus unstructured). For instance, Zimmermann et al [19] compared two

different computerized cognitive rehabilitation programs, specific cognitive training (CogniPlus,

19 patients) versus non-specific computer sport games with motion-capturing controllers

(Nintendo Wii, 20 patients). This study aimed at finding a possible positive effect in five

cognitive domains (attention, working memory, executive functions, visuo-construction and

episodic memory) as measured with neuropsychological testing before and after training. All

patients received a 40-minute training session three times per week over four weeks, either

with the specific or non-specific training programs. The CogniPlus rehabilitation program

included four modules: FOCUS, that trained focused attention; NBACK, that trained working

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memory; PLAND, that trained planning and action skills, and HIBIT, that trained response inhibition (last two modules trained executive functions), whereas the Nintendo Wii training was a game console with movement-capturing controllers. This training included four sports games: Table Tennis, Swordplay, Archery, and Air Sports. The results showed that greater improvement in attention skills was triggered by the nonspecific training (Nintendo Wii) rather than by the specific training (CogniPlus). No positive effects were found on tests assessing other cognitive skills. Furthermore, another study by Petrelli et al [18] examined the effect of different computerized cognitive group trainings: a structured training program (NEUROvitalis) with sessions targeting specific cognitive functions (attention, memory and executive functions) was contrasted with an unstructured training program (Mentally fit) similar to brain jogging. All treatment groups (22 patients in the structured training groups and 22 patients in the unstructured training group) had a 90-minute session twice a week over 6 weeks whereas a third PD group had no training at all (N=21). The results revealed that compared to the no training control group, patients in the structured training groups improved in short-term memory and working memory. In addition, the "NEUROvitalis" group improved significantly in working memory compared to the "Mentally fit" group. Petrelli and colleagues [8] conducted a one-year follow up with these groups of patients. However 18 patients from the original samples could not be re-assessed due to difficulties in contacting them. The one year follow up, therefore, was available only for 16 patients in the structured training group, for 17 patients in the unstructured training group and for 14 in the no training control group. The findings showed that, compared with the no training control group, both training groups maintained better overall cognitive functions as assessed by the DemTect. However, only the structured training group appeared to maintain their cognitive level when assessed with the MMSE. This study also concluded that cognitive training might prevent cognitive decline or onset of MCI in PD.

Cognitive Domains Targeted by Rehabilitation

Most of the studies targeted one or more specific cognitive domains (N=13), whilst two studies used a non-specific method of cognitive rehabilitation. The majority of the studies focused on improvements of executive function and attention. For instance, although studies that focused on executive functions used different duration and frequency of sessions, all results showed improvements in this specific cognitive domain. The evidence from these studies, therefore, suggests that all cognitive intervention programs targeting executive functions were effective in treating some component of executive abilities in PD. Furthermore, another cognitive domain that seemed to benefit from non-specific cognitive intervention was attention. Zimmermann et al. [19] demonstrated that a nonspecific training program (Nintendo Wii) improved attentions skills more than a specific one (CogniPlus). In addition, Petrelli et al. [18] found that a structured cognitive training program improved short-term memory and working memory skills in patients with PD.

Measures of Cognition as Outcome Measures

All investigations focused on cognitive outcomes and reported improvements, although there was diversity in the cognitive domain/s targeted and the duration of each intervention. Only one study, that of Petrelli et al. [8], intended to follow up the training groups after one year of intervention and they found that the training groups maintained their overall cognitive functions better than the control group. Most of the studies were carried out with non-demented patients (N=10), three studies included patients with MCI, whereas, two studies did not state the overall cognitive status of their samples. In terms of severity of disease, nine out of 15 studies included patients who were in the mild to moderate disease stages; two more studies included patients who were in the mild to severe disease stages, whereas the remaining four studies did not specify the disease severity stage of their samples. As for the type of intervention, there was a considerable diversity among studies and all appeared to focus on different cognitive domains. For instance, four studies focused their intervention on executive functions; one study focused on non-specific multiple skills, one study focused on cognitive

processing speed, another study focused on shifting ability, while the other studies targeted

different cognitive domains such as attention (in four studies), working memory (in four

studies) and verbal fluency (in two studies). Other domains were mentioned at least once in

different studies including: abstract reasoning, visuospatial, sustained, selective, alternating and

divided attention, mental flexibility, episodic memory, mental speed, verbal and visual memory

and short/long term memory.

Imaging parameters as outcome measures

Only one study has investigated the role of neuroimaging techniques in the assessment of the

effects of cognitive rehabilitation in PD. Nombela and others [23] examined whether cognitive

training improved cognitive dysfunction and they also assessed whether any cognitive changes

were correlated with changes in any measure of brain function. One easy level of Sudoku was

used (4-by-4 grid with 2-by-2 blocks). These authors measured cortical activation tested in

response to a modified version of the Stroop test performed while participants were in a

functional magnetic resonance imaging (fMRI) scanner. There were 10 patients with PD and 10

healthy controls; half of the PD patients had 6 months of cognitive daily training based on

Sudoku exercises that mainly focused on working memory and attention skills. The results

revealed that the training program improved cognitive performance in the Stroop task of the

trained PD group during fMRI (in terms of reaction time, and of correct and missed answers).

Furthermore, in the untrained PD group, there were reduced cortical activation patterns similar

to the patterns of activation that were observed in the controls. Therefore, from this study it

appears that neuroimaging techniques might provide evidence of the positive impact of

cognitive rehabilitation in PD. Further investigations with larger samples are needed, however,

to confirm this finding.

Mood, Fatigue and Quality of Life assessments as outcome measures

There is evidence that the cognitive deficits experienced by patients with PD have a negative

impact on their quality of life [3]. The majority of the cognitive rehabilitation studies also

evaluated other important non-cognitive factors such as depression (N=7), fatigue (N=2), anxiety (N=2), and quality of life (N=3). However, some studies evaluated these aspects only at the baseline stage (depression N=2, anxiety N= 2, fatigue N=1). The outcomes of these studies were inconsistent. For instance, out of five studies, two found improvement in depressive symptoms after cognitive rehabilitation. In addition, out of three studies, two found improvement in quality of life. Due to the small number of studies that evaluated these factors, it is very difficult to draw any firm conclusion as to the impact of cognitive intervention on quality of life and on neuropsychiatric symptoms in PD. Furthermore, the difference among the results of different studies might be explained by the use of different methodological approaches. For example, Adamski et al. [20] concluded that depressive symptoms assessed by a general depression scale were improved in PD patient after cognitive intervention, whereas fatigue that was assessed by the Fatigue Scale for Motor and Cognitive Functions did not improve. Edwards and colleagues [17] used the Center for Epidemiological Studies Depression Scale to assess depression and found no improvement in the PD group following cognitive training. Furthermore, Paris and others [24] found no improvement in depressive symptoms in PD after cognitive rehabilitation. This latter study used the Geriatric Depression Scale to assess depression. From the review above, it appears that the various studies that assessed the effect of cognitive training on depression used different tools to measure depression that may explain the inconsistency in outcomes. In addition, based on prior investigations, it seems that type of cognitive intervention, duration and frequency of training have no influence on improving depression in PD. Further studies to examine the effect of cognitive rehabilitation on quality of life and other neuropsychiatric symptoms are needed.

Conclusions and future directions

The previous review by Hindle et al [10] was a more general review of the effect of a variety of forms of non-pharmacological interventions in PD. The search strategy for this previous review included terms that were not exclusive to cognitive interventions such as physical activity,

exercise and motor. Although the review by Hindle and others [10] found that many studies reported positive outcomes for executive function in particular, the findings of our review indicate that several studies of cognitive training reported a positive impact on executive functions and attention. For the second review by Calleo and colleagues [11], their work focused on only one cognitive domain that was executive function. Unlike our review they did not include terms such as attention, language, memory, working memory, learning and problem solving in their search strategy, potentially missing effects on other cognitive domains. Overall, converging evidence of the benefit of cognitive rehabilitation on executive functions is reported both by the previously published reviews and our own. Our review, however, given the broader search including other cognitive domains, also highlights the benefits on attention, a finding that was missed both by the Hindle et al and the Calleo et al reviews. The newly published reviews showed different findings based on their research focus. For instance, Leung et al. [12] looked at only RCT studies (7 papers) and they found that cognitive training resulted in improvements on measures of working memory, processing speed and executive functions. Whereas Van de Weijer et al [14] review was focused on the late stage of PD and they provided limited evidence of effectiveness of cognitive training. The results indicated improvements in non-demented patients in HY-stage 3-4, but not in HY-stage 5 and they did not state which cognitive domains were improved. Walton et al [13] discussed the theoretical perspective of cognitive training in PD; they illustrated cognitive training as a potential therapeutic technique and the efficacy of cognitive training in general not only in PD. They also reported some results from the previous studies to provide evidence of the efficacy of cognitive training in PD without reaching any conclusion on which cognitive domains appeared to benefit more from cognitive training.

This review aimed to examine the effects of cognitive rehabilitation in PD, describe the present situation in this field and provide directions for future research. From this review it is evident that most of the articles were published in the last decade. Thus, this research area can be considered in its early stages of investigations. The main outcomes of this review indicate that, although previous studies used different cognitive rehabilitation methodologies, all studies

revealed cognitive improvements on at least one cognitive domain. Additionally, the most frequent cognitive domains showing positive effects were executive functions and attention. Therefore, it seems that cognitive interventions have a positive impact in patients with PD irrespective of methods, duration and frequency of training.

Due to the use of various cognitive rehabilitation techniques, we cannot draw any firm conclusion on which method might trigger the largest improvement in cognition, mood, fatigue and quality of life. However, cognitive improvements were observed even in the studies that included a small sample size. This finding might be explained by the inclusion of patients in the early stages of the disease, as well as by the inclusion of non-demented patients.

This review has also pointed out the limitations of previous studies. The main shortcoming appears to be the absences of long term follow up data, the absence of a control group of patients with PD who did not undergo training. Some cognitive training programs included other elements such as psycho-education, physical exercise, medications or unspecific conversations that make it difficult to determine which specific elements had more influence on the training results. Thus, more studies are needed to have a clearer picture of which techniques are more effective to improve cognitive abilities in PD.

Only two studies included follow up of their samples after cognitive interventions to see the long-term effects on cognitive functions. Sinfiorini et al. [27] found that after a 6 month follow up PD patients maintained their improved performance and no changes were observed on the neuropsychological tests scores. A second study by Petrelli et al [8] reported that after a one year follow up the trained groups maintained their overall cognitive function levels. However, this study did not do an extensive cognitive assessment at the follow up stage (after one year) as they did immediately after cognitive training, but applied only some limited testing, a strategy that limits our knowledge about the long term effects of the training programs on specific cognitive domains. Therefore, longitudinal studies are needed to investigate the potential benefits of cognitive training long term. In addition, future investigations should also examine

whether any disease characteristics such as disease stage, degree of cognitive impairment and/or the dominant side (right/left) or specific motor symptoms (rigidity/tremor) influence treatment efficacy.

Furthermore, only one neuroimaging study by Nombela et al [23] examined whether the potential cognitive improvement might be correlated with neural alterations in PD. The results showed that training improved cognitive performance on the Stroop test during fMRI. Specifically, trained PD patients showed alterations in activation patterns that involved the left precentral gyrus, left medial frontal gyrus, right precuneus, and left inferior parietal gyrus when compared with the control group. Further imaging studies could provide a better understanding of these neural changes and more clear evidence of the efficacy of cognitive rehabilitation in PD.

While some studies concluded that neuropsychiatric symptoms such as depression were not responsible for the different treatment outcomes, other studies reported that depression had an effect on the cognitive rehabilitation findings. Also in this case more investigations with improved design and more comprehensive outcome measures are needed to provide clearer evidence of the role of depression and other neuropsychiatric symptoms on the effects of cognitive rehabilitations in PD.

Despite some reports finding an association between improvement of cognitive functions and quality of life, other investigations found no significant benefit of cognitive interventions on quality of life. Of course, cognitive training should result in improved daily live activities to have a significant impact on improvement of quality of life. Therefore, future research may investigate whether the improvement in cognitive skills can be transferred to improve daily life activities and consequently quality of life in this patient population.

Overall, cognitive rehabilitation in PD is still in its infancy stage. No firm conclusion can, therefore, be drawn at this stage. However, future research should take into account the shortcoming and unanswered questions highlighted in this review to design better trials. Future studies should also investigate whether PD patients may benefit more from a cognitive

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rehabilitation strategy that involves intervention on multiple cognitive domains or from one focused on only one cognitive skill.

Conflict of interest: The authors declare that they have no conflict of interest.

References

[1]   Aarsland D, Bronnick K, Fladby T (2011) Mild cognitive impairment in Parkinson's disease. Current neurology and neuroscience reports **11**, 371-378.

[2]   Litvan I, Aarsland D, Adler CH, Goldman JG, Kulisevsky J, Mollenhauer B, Rodriguez-Oroz MC, Troster AI, Weintraub D (2011) MDS Task Force on mild cognitive impairment in Parkinson's disease: critical review of PD-MCI. Movement disorders : official journal of the Movement Disorder Society **26**, 1814-1824.

[3]   Leroi I, McDonald K, Pantula H, Harbishettar V (2012) Cognitive impairment in Parkinson disease: impact on quality of life, disability, and caregiver burden. J Geriatr Psychiatry Neurol **25**, 208-214.

[4]   Emre M, Aarsland D, Brown R, Burn DJ, Duyckaerts C, Mizuno Y, Broe GA, Cummings J, Dickson DW, Gauthier S, Goldman J, Goetz C, Korczyn A, Lees A, Levy R, Litvan I, McKeith I, Olanow W, Poewe W, Quinn N, Sampaio C, Tolosa E, Dubois B (2007) Clinical diagnostic criteria for dementia associated with Parkinson's disease. Mov Disord **22**, 1689-1707; quiz 1837.

[5]   Cools R (2006) Dopaminergic modulation of cognitive function-implications for L-DOPA treatment in Parkinson's disease. Neuroscience and Biobehavioral Reviews **30**, 1-23.

[6]   Pfeiffer HC, Lokkegaard A, Zoetmulder M, Friberg L, Werdelin L (2014) Cognitive impairment in early-stage non-demented Parkinson's disease patients. Acta Neurol Scand **129**, 307-318.

[7]   Costa A, Monaco M, Zabberoni S, Peppe A, Perri R, Fadda L, Iannarelli F, Caltagirone C, Carlesimo GA (2014) Free and cued recall memory in Parkinson's disease associated with amnestic mild cognitive impairment. PLoS One **9**, e86233.

[8]   Petrelli A, Kaesberg S, Barbe MT, Timmermann L, Rosen JB, Fink GR, Kessler J, Kalbe E (2015) Cognitive training in Parkinson's disease reduces cognitive decline in the long term. Eur J Neurol **22**, 640-647.

[9]   Naismith SL, Mowszowski L, Diamond K, Lewis SJ (2013) Improving memory in Parkinson's disease: a healthy brain ageing cognitive training program. Mov Disord **28**, 1097-1103.

[10]   Hindle JV, Petrelli A, Clare L, Kalbe E (2013) Nonpharmacological enhancement of cognitive function in Parkinson's disease: a systematic review. Mov Disord **28**, 1034-1049.

[11]   Calleo J, Burrows C, Levin H, Marsh L, Lai E, York MK (2012) Cognitive rehabilitation for executive dysfunction in Parkinson's disease: application and current directions. Parkinsons Dis **2012**, 512892.

[12]   Leung IH, Walton CC, Hallock H, Lewis SJ, Valenzuela M, Lampit A (2015) Cognitive training in Parkinson disease: A systematic review and meta-analysis. Neurology **85**, 1843-1851.

[13]   Walton CC, Naismith SL, Lampit A, Mowszowski L, Lewis SJ (2017) Cognitive Training in Parkinson's Disease. Neurorehabil Neural Repair **31**, 207-216.

[14]   Van de Weijer SCF, Hommel A, Bloem BR, Nonnekes J, De Vries NM (2018) Promising non-pharmacological therapies in PD: Targeting late stage disease and the role of computer based cognitive training. Parkinsonism Relat Disord **46 Suppl 1**, S42-S46.

[15]   Angelucci F, Peppe A, Carlesimo GA, Serafini F, Zabberoni S, Barban F, Shofany J, Caltagirone C, Costa A (2015) A pilot study on the effect of cognitive training on BDNF serum levels in individuals with Parkinson's disease. Front Hum Neurosci **9**, 130.

[16]   Costa A, Peppe A, Serafini F, Zabberoni S, Barban F, Caltagirone C, Carlesimo GA (2014) Prospective memory performance of patients with Parkinson's disease depends on shifting aptitude: evidence from cognitive rehabilitation. J Int Neuropsychol Soc **20**, 717-726.

[17]   Edwards JD, Hauser RA, O'Connor ML, Valdes EG, Zesiewicz TA, Uc EY (2013) Randomized trial of cognitive speed of processing training in Parkinson disease. Neurology **81**, 1284-1290.

[18]   Petrelli A, Kaesberg S, Barbe MT, Timmermann L, Fink GR, Kessler J, Kalbe E (2014) Effects of cognitive training in Parkinson's disease: a randomized controlled trial. Parkinsonism Relat Disord **20**, 1196-1202.

[19]   Zimmermann R, Gschwandtner U, Benz N, Hatz F, Schindler C, Taub E, Fuhr P (2014) Cognitive training in Parkinson disease: cognition-specific vs nonspecific computer training. Neurology **82**, 1219-1226.

[20]   Adamski N, Adler M, Opwis K, Penner IK (2016) A pilot study on the benefit of cognitive rehabilitation in Parkinson's disease. Ther Adv Neurol Disord **9**, 153-164.

[21]   Disbrow EA, Russo KA, Higginson CI, Yund EW, Ventura MI, Zhang L, Malhado-Chang N, Woods DL, Sigvardt KA (2012) Efficacy of tailored computer-based neurorehabilitation for improvement of movement initiation in Parkinson's disease. Brain Res **1452**, 151-164.

[22]   Mohlman J, Chazin D, Georgescu B (2011) Feasibility and acceptance of a nonpharmacological cognitive remediation intervention for patients with Parkinson disease. J Geriatr Psychiatry Neurol **24**, 91-97.

[23]   Nombela C, Bustillo PJ, Castell PF, Sanchez L, Medina V, Herrero MT (2011) Cognitive rehabilitation in Parkinson's disease: evidence from neuroimaging. Front Neurol **2**, 82.

[24]   Paris AP, Saleta HG, de la Cruz Crespo Maraver M, Silvestre E, Freixa MG, Torrellas CP, Pont SA, Nadal MF, Garcia SA, Bartolome MV, Fernandez VL, Bayes AR (2011) Blind randomized controlled study of the efficacy of cognitive training in Parkinson's disease. Mov Disord **26**, 1251-1258.

[25]   Reuter I, Mehnert S, Sammer G, Oechsner M, Engelhardt M (2012) Efficacy of a multimodal cognitive rehabilitation including psychomotor and endurance training in Parkinson's disease. J Aging Res **2012**, 235765.

[26]   Sammer G, Reuter I, Hullmann K, Kaps M, Vaitl D (2006) Training of executive functions in Parkinson's disease. J Neurol Sci **248**, 115-119.

[27]   Sinforiani E, Banchieri L, Zucchella C, Pacchetti C, Sandrini G (2004) Cognitive rehabilitation in Parkinson's disease. Arch Gerontol Geriatr Suppl, 387-391.

[28]   Welton T, Kent D, Constantinescu CS, Auer DP, Dineen RA (2015) Functionally relevant white matter degradation in multiple sclerosis: a tract-based spatial meta-analysis. Radiology **275**, 89-96.

Table 1 Summary of reviewed articles

Authors	N	PD stages	Cognitive status	Cognitive domains targeted	Technique/ design	Outcome measures	Duration and frequency	Results
Sinfiorini et al. (2004)	20	Mild to moderate	Not demented	Attention, abstract reasoning, visuospatial	Computerised software for cognitive training (Pre-post- follow up)	Digit span, FAS, Corsi-test, Babcock's story, Raven's matrices, WCST and Stroop test	1 hour, 12 sessions over 6-week. Follow up after 6 months	Improvement on FAS, Babcock's story and Raven's matrices, maintained after 6 months. No changes on MMSE, digit span, Corsi-test, WCST, and Stroop after training
Sammer et al. (2006)	26	Mild to moderate	Average MMSE= 27.15; SD= 1.49	Executive functions	Working memory tasks for cognitive training group N=12, no cognitive training for the control group N=14 (Pre-post- no further follow up)	BADS, cognitive estimation test, trail making test ZVT, GNL, AKT and MWT	30-minute, 10 sessions during a 3-4 week hospital stay	Cognitive training group improved on the BADS, both groups improved on the 6-element task but training group tended to gain more improvement, no effect on other tests
Mohlman et al. (2011)	14	Not stated	MMSE >23	Sustained, selective, alternating and divided attention	Attention process training, participants also rate their level of fatigue, effort, progress and enjoyment after the completion of each training block (Pre-post-no further follow up)	Six-point Likert scale to rate fatigue, effort, progress, enjoyment; MMSE, BNT, digit span, Stroop test, TMT, COWAT	90-minute, 4 sessions over one month	Average Likert responses across participants was some to much on progress, enjoyment, effort and a little to some on fatigue; patients improved on digits backward, Stroop, TMT-B and COWAT
Paris et al. (2011)	26	Mild to moderate	MMSE >22	Non-specific/ multiple skills	Multimedia software and paper-and-pencil cognitive exercises for experimental group N=16 while control group N=12 had speech therapy (Pre-post-no further follow up)	MMSE, digit span, CVLT, SDMT, TMT, Stroop test, Logical memory test, ROCFT, verbal fluency, RBANS-line orientation, TOL, mood and quality of life	45-minute, 3 sessions per week over 4 weeks	Improvement on digit span forward (attention), Stroop test (information processing speed), ROCFT (visual memory), RBANS-line orientation (visuospatial), semantic verbal fluency, TMT-B and TOL (executive functions), no changes in mood or quality of life
Nombela et al. (2011)	20	Mild to moderate	MMSE >25	Attention	One easy level of Sudoku (4-by-4 grid with 2-by-2 blocks; cortical activation	Modified Stroop test in fMRI scanner and brain activation	90-120 minutes, everyday over 6 months	Trained group improved in Stoop test during MRI, in reaction time, correct and missing responses, and they had

Study	N	Severity	Inclusion	Domain	Intervention	Outcome measures	Dose	Results
					tested by fMRI, 10 PD patients (5 trained, 5 untrained) and 10 healthy controls (Pre-post)			stronger activation in right temporal gyri than untrained group, and stronger activation in some brain areas including right anterior cingulate gyrus, left frontal and parietal cortex than controls
Disbrow et al. (2012)	51	Not stated	Impaired and unimpaired groups based on motor test and TMT	Executive functions	Computer-based training of externally and internally generated sequences of number, there were 3 groups, 14 impaired PD, 16 unimpaired PD and 21 healthy controls (Pre-post)	TMT, D-KEFS, TUG, TIADL, sequence initiation, completion time and error rate	40-minute, 5 days per week over 12-14 days	Improvement in impaired PD group in sequence initiation, completion time and error rate, all groups improved on TMT-B-A
Reuter et al. (2012)	222	Mild to severe	MCI	Executive functions	Multimodal cognitive rehabilitation, there were 3 groups, group A n=71 received cognitive training only, group B n=75 received cognitive training and transfer training, group C n=76 conducted cognitive training, transfer and psychomotor training, (Pre-post)	PANDA, MMSE, primary outcome: ADAS-COG, secondary outcome: SCOPA-COG, PASAT, BADS, HADS and PDQ-39	60-minute, 14 sessions over 4 weeks, hospital stay, follow up after 6 months at home	All groups improved on the ADAS-COG and SCOPA-COG. Group C improved most indicated by a significant interaction between groups and assessments. After 6 months Group B and especially group C maintained better than group A. Group C improved on BADS test compared with group A and B. group B and C improved on the PASAT and group C had higher rate on the PDQ-39 than the other groups
Naismith et al. (2013)	50	Mild to severe	Average MMSE= 27.5; SD= 3.3	Memory and general cognition e.g. psychomotor speed, mental flexibility and verbal fluency	Computer-based cognitive training and psychoeducation for training group n=35, no intervention for control group n=15, (Pre-post)	Logical memory test immediate and delay recall subtest of the Wechsler memory scale, TMT A and B-A, COWAT, BDI and	1-hour cognitive training and 1-hour psychoeducation, twice a week over 7 weeks	Improvement in training group on logical memory test both in learning and memory retention skills, no effect of cognitive training on psychomotor speed, mental flexibility, verbal fluency,

						knowledge test		knowledge or depressive symptoms
Edwards et al. (2013)	74	Mild to moderate	MMSE >23	Cognitive speed of processing	Self-administered version (InSight software) of cognitive speed of processing training was completed by training group n=32, no intervention for control group n=41, (Pre-post)	UFOV, this test evaluate speed of processing, cognitive self-report questionnaire for cognitive self-perceptions, Depression scale (CES-D)	1-hour cognitive training, about 3 sessions per week over 3-month, participants were encouraged to complete at least 20 hours of training	Improvement in training group on UFOV compared to control group, no effect of training on the other assessments
Zimmermann et al. (2014)	39	Mild to moderate	MMSE >27	Attention, working memory, executive functions, visuo-construction and episodic memory	Computer-based training for specific (CogniPlus software) n=19 and nonspecific (Nintendo Wii) n=20 cognitive training, (Pre-post)	Tests of attentional performance (alertness and working memory), TMT, Block-Design Test and CVLT	40-minute, 3 sessions per week over 4 weeks	Improvement in nonspecific training (Nintendo Wii) than specific training (CogniPlus) in attention skills, no differences on other tests
Costa et al. (2014)	17	Mild to moderate	MCI	Shifting ability	Paper and pencil exercises of shifting training for experimental group n=9 and language training for control group n=8, (Pre-post)	Prospective memory procedure, verbal fluency (phonemic, semantic and alternating tasks) and TMT	45-minute, 3 sessions pre week over 4 weeks	Improvement in the experimental group on the alternate task and the accuracy indices of the prospective memory procedure, no differences on the control group performance was observed
Petrelli et al. (2014)	65	Mild to moderate	MMSE >24	Attention, verbal and visual memory and executive functions (working memory and verbal fluency)	Computer-based cognitive training, first group n=22 had structured training, second group n=22 had unstructured training and control group n=21 had no training (Pre-post)	Brief test of attention word list learning test, phonemic and semantic fluency test, digit span reverse ROCFT, BDI and PDQ-39	90-minute, 12 sessions over 6 weeks	Improvement in the structured training group on the tasks of short-term memory and working memory whereas depression scores reduced in unstructured training group, no improvement in quality of life in all groups
Petrelli et al. (2015)	47	Mild to moderate	MMSE >24	Overall cognitive functions	A one year follow up of the previous study (Petrelli et al. 2014) to see the long-term effects on cognitive function, however number of	MMSE and DemTect (Cognitive screening tool to detect MCI and dementia)	No training just a follow up study	Compared to control group both training groups maintained the overall cognitive functions assessed by DemTect, however only structured training group

				participates was smaller in this follow up study, structured training group n=16, unstructured training group n=17 and control group n=14			maintained the cognitive level assessed by the MMSE compared to the other two groups. Cognitive training also may prevent cognitive decline or onset of MCI in PD
Angelucci et al. (2015)	15	Not stated	MCI	Executive functions	Paper and pencil exercises focused on shifting abilities that involved various stimuli (e.g. numbers, letters and shapes) for training group n=7 and simple cognitive tests of attention and language for control group n=8 (Pre-post)	Zoo map test, part of the BADS	45-minute, 12 sessions over one month	Experimental group improved in the first trial (no instructions was given) of the zoo map test but not in the second trail (Instructions was given), whereas no improvement was observed in the control group in both trails
Adamski et al. (2016)	6	Not stated	Baseline assessment showed worse performance in PD group in information processing speed, short term memory, verbal long term memory and working memory	Working memory, short term memory, mental speed and long term memory	Computerized cognitive training tool (BrainStim), this training focused on working memory. There were 3 groups, 6 PD patients, 19 healthy controls (12 of them were trained and 7 were without training (pre-post after 6 weeks and ager 3 months)	Self-report questionnaires for depression, The Centre for Epidemiologic Studies Depression Scale, Fatigue Scale for Motor and Cognitive Functions, BRB-N, TAP, WMS-R and SDMT	45-minute, 4 sessions per week over a period of 4 weeks	Both training groups showed improvement in verbal and visuospatial short term, and long-term memory. After 3 months both training groups revealed stable results in all short-term visuospatial tasks and PD group had low depression scores.

N: number of participants, FAS: phonological word fluency test, WCST: Wisconsin Card Sorting Test, MMSE: Mini Mental State Examination, SD: Standard Deviation, BADS: Battery of Behavioural Assessment of the Dysexecutive Syndrome, ZVT: Zahlenverbindungstest, GNL: face-name-learning test, AKT: Alters-Konzentrations Test, MWT: test of verbal intelligence, BNT: Boston Naming Test, TMT: Trail Making Test, COWAT: Controlled Oral Word Association Test, ROCFT: Rey Osterrieth Complex Figure Test, RBANS: Repeatable Battery for the Assessment of Neuropsychological Status, TOL: Tower of London, fMRI: functional Magnetic Resonance Imaging, CVLT: California Verbal Learning Test, SDMT: Symbol-Digit Modalities Test, D-KEFS: Delis-Kaplan Executive Function Scale, TUG: Timed-Up-and–Go test, TIADL: Timed Instrumental Activities of Daily Living Tasks, MCI: Mild Cognitive Impairment, PANDA: Parkinson neuropsychometric dementia assessment, ADAS-COG: Alzheimer Assessment Scale-Cognition, SCOPA-COG: Scales for outcome of Parkinson's Disease-Cognition, PASAT: Paced Auditory Serial Addition Test, BADS: Behavioural Assessment of the Dysexecutive Syndrome, HADS: Hospital Anxiety and Depression Scale, PDQ-39: Parkinson's Disease Quality of life, BDI: Beck Depression Inventory, UFOV: Useful Field of View Test, CES-D: The Centre for Epidemiological Studies Depression Scale, BADS: Behavioural Assessment of the Dysexecutive Syndrome, BRB-N: Brief Repeatable Battery of Neuropsychological Tests, TAP: Test Battery for Attention Performance, WMS-R: Wechsler Memory Scale- revised,

Table 2. Quality assessment of the cognitive rehabilitation studies included.

Study	1	2	3	4	5	6	7	8	9	10	11	12	Total
Sinfiorini et al. (2004)	0	1	0	1	0	0	1	0	1	1	1	0	6
Sammer et al. (2006)	1	0	0	1	1	1	1	0	1	0	1	1	8
Mohlman et al. (2011)	1	1	0	0	0	0	1	1	1	0	1	1	7
Paris et al. (2011)	1	1	0	1	1	1	1	1	1	0	1	1	10
Nombela et al. (2011)	1	1	1	1	0	1	1	1	1	0	1	1	10
Disbrow et al. (2012)	0	0	0	0	1	1	1	1	1	0	1	0	6
Reuter et al. (2012)	1	1	0	1	1	1	1	1	1	1	1	1	11
Naismith et al. (2013)	1	1	1	1	1	1	1	0	1	0	1	1	10
Edwards et al. (2013)	1	1	1	1	1	1	1	1	1	0	1	1	11
Zimmermann et al. (2014)	1	1	1	1	1	0	1	0	1	0	1	1	9
Costa et al. (2014)	1	1	1	1	0	1	1	1	1	0	1	1	10
Petrelli et al. (2014)	0	1	1	1	1	1	1	0	0	0	1	1	9
Petrelli et al. (2015)	1	1	1	1	1	0	0	0	0	1	0	1	7
Angelucci et al. (2015)	1	1	1	0	0	1	1	1	1	0	1	1	9
Adamski et al. (2016)	1	1	0	0	0	0	0	1	1	1	1	1	7

Appendix 1. Quality assessment criteria used to assess reviewed studies on cognitive rehabilitation in PD

Questions	Values
1. Were aims related to study clearly stated?	no = 0; yes = 1
2. Were demographic clearly provided for the PD patients?	no = 0; yes = 1
3. Were clinical features of PD clearly stated?	no = 0; yes = 1
4. Were PD stages indicated?	no = 0; yes = 1
5. How large was the sample size?	< 25 = 0; ≥ 25= 1
6. Was there a PD control group who did not undergo training?	no = 0; yes = 1
7. Was cognitive status clearly defined prior to intervention?	no = 0; yes = 1
8. Was the study focus only on cognitive training?	no = 0; yes = 1
9. Was the cognitive training technique well described?	no = 0; yes = 1
10. Did patients have pre, post and follow up assessment?	no = 0; yes = 1
11. Were duration and frequency of training clearly stated?	no = 0; yes = 1
12 Were limitations of the studies clearly stated?	no = 0; yes = 1

Figure 1 Flow chart of the study selection process

Titles and abstracts identified and screened N= 791

Excluded N= 478 (Duplicate

Full copies retrieved and assessed for eligibility N= 313

Excluded N= 296
Reason for exclusion:
(1) Review articles
(2) Not cognitive intervention
(3) Papers that included participants with other neurological conditions
(4) Studies of healthy participants
(5) Reports published only in abstract format
(6) Non-peer reviewed articles
(7) Case reports

Publications meeting inclusion criteria N=

Excluded N= 2
Not cognitive intervention

Number of studies included in the review N= 15