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# Evidence-based cognitive rehabilitation after acquired brain injury: A systematic review of content of treatment

# Caroline van Heugten<sup>1,2</sup>, Gisela Wolters Gregório<sup>1</sup>, and Derick Wade<sup>3,4</sup>

<sup>1</sup>School for Mental Health and Neuroscience, University of Maastricht, Maastricht, The Netherlands

<sup>2</sup>Department of Neuropsychology and Psychopharmacology, University of Maastricht, Maastricht, The Netherlands

<sup>3</sup>Care and Public Health Research Institute, University of Maastricht, Maastricht, The Netherlands

We reviewed all randomised trials on cognitive rehabilitation in order to determine the effective elements in terms of patients' and treatment characteristics, treatment goals and outcome. A total of 95 random controlled trials were included from January 1980 until August 2010 studying 4068 patients in total. Most studies had been conducted on language (n = 25), visuospatial functioning (n = 24), and memory (n = 14). Stroke patients were the commonest subjects (57%; overall mean age = 52.2, SD = 15.0 years). Of the interventions 39% were offered more than 12 months after onset and 23% were offered within two months of onset. The mean (SD) number of hours of treatment actually delivered was 4.1 (3.6) per week; treatment was mostly offered individually. No papers gave specific information on the expertise or competences of the staff involved. With 95 RCTs there is a large body of evidence to support the efficacy of cognitive rehabilitation, and the current study can serve as a database for clinicians and researchers. But most studies have given little information about the actual content of the treatment which makes it difficult to use the studies when making treatment decisions in daily

<sup>&</sup>lt;sup>4</sup>Oxford Centre for Enablement, Oxford, United Kingdom

Correspondence should be addressed to Caroline van Heugten, Department of Psychiatry and Neuropsychology, Maastricht University, PO Box 616, 6200 MD Maastricht, The Netherlands. E-mail:c.vanheugten@maastrichtuniversity.nl

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clinical practice. We suggest developing an international checklist to make standardised description of non-pharmacological complex interventions possible.

*Keywords*: Brain injuries; Stroke; Cognition disorders; Rehabilitation; Randomised controlled trial.

# INTRODUCTION

Impairments of cognitive function are frequent after acquired brain injury and cause problems in activities of daily life and participation in society. Cognitive rehabilitation is the treatment of choice for these deficits and can be defined as "a systematic, functionally oriented service of therapeutic activities that is based on assessment and understanding the patients' brain—behavioral deficits" (Cicerone et al., 2000, pp. 1596–1597). Rehabilitation of cognitive deficits is recognised as a standard component of rehabilitation programmes for patients with acquired brain injury as can be seen in many national clinical guidelines (i.e., American Heart Association, UK Royal College of Physicians).

The Brain Injury Interdisciplinary Special Interest Group (BI-ISIG) of the American Congress of Rehabilitation Medicine (ACRM) previously conducted a systematic review of the literature up to 2008 on cognitive rehabilitation for people with traumatic brain injury (TBI) or stroke (Cicerone et al., 2000, 2005, 2011). The review found that there was a differential benefit in favour of cognitive rehabilitation in almost 80% of all treatment comparisons. Recommendations for clinical practice were formulated accordingly. In addition, Rohling, Faust, Beverly, and Demarkis (2009) performed a metanalytic re-examination of Cicerone et al.'s (2000, 2005) reviews to provide treatment effect sizes and showed that there is sufficient evidence for the effectiveness of attention training after traumatic brain injury, and the effectiveness of language and visuo-spatial training for aphasia and neglect syndromes after stroke.

However, when a healthcare professional wants to apply evidence from a study in everyday clinical practice, or when a researcher wishes to replicate the study, the details listed below are needed to allow practical use of the information generated by research studies:

- What were the clinical characteristics of the patients in the study?
- What was the treatment studied (content, process, resources, etc.)?
- What was the treatment setting?
- What are the costs/benefits for this patient (on the basis of the prognosis and condition of the patient)?

In practice this information is not easily found in the published literature, either in reviews or in the original papers. We therefore set out to review systematically what detailed information is available about the cognitive rehabilitation interventions that have been evaluated. As a necessary part of this, we also had to establish how many randomised controlled trials evaluating interventions designed to ameliorate cognitive impairments are now published.

#### **METHODS**

A systematic literature search was performed to identify randomised controlled studies evaluating interventions targeted at people with cognitive deficits associated with acquired brain injuries, using PubMed and PsycINFO which are the primary electronic databases covering this area of research. We searched the period from January 1980 until August 2010. Details of the search strategy are presented in Table 1 and are available from the corresponding author. The starting point for the search strategy was the one by the BI-ISIG to make sure the same 46 class I studies identified by Cicerone et al. (2000, 2005) would be identified. In addition, we hand-searched the reference lists of the reviewed articles, and of relevant systematic reviews from the Cochrane database (i.e., reviews on memory, neglect, aphasia, apraxia, and attention). The second author performed the literature search as well as the primary selection of articles based on the title and abstracts.

# Study selection

Figure 1 shows the selection of papers. Studies were included only when they addressed the effect of cognitive rehabilitation in an RCT related to non-

#### TABLE 1 Search terms

Cognitive domains	Attention/concentration, information processing/ slowness, awareness/ insight, cognition, communication, executive/ planning/organisation/ regulation, language/aphasia, memory, perception/perceptual/agnosia/ neglect/visual, problem solving, reasoning, apraxia/dyspraxia
Rehabilitation	Rehabilitation, remediation, education, training, retraining, paging system, treatment, treatments, therapy
Acquired brain injury	Stroke, brain injuries/brain-injuries/brain injury/brain-injury/brain injured/brain-injured, head injuries/head-injuries/head injury/head-injury/head injured/head-injured, brain damage/brain damaged, tbi/head trauma/traumatic, cerebrovascular disorders/cerebrovascular accident/cerebrovascular accidents/cva/stroke/poststroke/post-stroke/post stroke, chronic aphasia

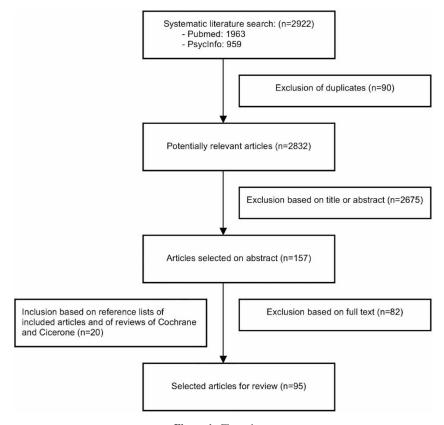


Figure 1. Flow-chart.

progressive acquired brain injury, defined operationally in the search terms. Quasi or semi-randomised trials, controlled comparative studies or uncontrolled cohort studies were excluded. This process yielded 959 published articles in PsycINFO and 1963 in PubMed.

The abstracts were reviewed to eliminate reports according to seven exclusion criteria: (1) no intervention; (2) design other than RCT; (3) reports that did not explicitly describe randomisation of patients (i.e., randomised was used in the title but the procedure was not described in the paper itself); (4) subjects other than persons with acquired brain injury; (5) pharmacological interventions; (6) interventions not aimed at cognitive deficits; and (7) non-English language papers. Through this screening process, 157 articles were selected for inclusion in the study. After reviewing the complete reports enabling us to check all criteria, 95 articles were finally included in the review.

These articles were then assigned to categories according to their primary area of intervention as was done in the previous reviews (Cicerone et al., 2000; 2005): awareness, visuospatial functioning, memory, attention, executive functioning, apraxia, language and communication, and multi-domain studies (i.e., targeting more than one cognitive domain, such as memory and attention). Studies investigating comprehensive-holistic cognitive rehabilitation programmes aimed at community reintegration were not included, because a recent systematic review of these programmes described patient and treatment characteristics extensively (Geurtsen, van Heugten, Martina, & Geurts, 2010).

# Description of treatment given

Each paper included was specifically reviewed to discover details on the treatment given. Three sources were used. First, the original paper itself; any description given was reviewed and data were extracted. Second, if the report gave a reference to a paper or publically available manual or document, the document was reviewed and data on treatment were extracted. Third, a brief search of the same databases was made using the name of the first author only (but accepting the name in any position in identified papers), restricting the search to four years before and after the publication of the main study. The titles and abstracts were read to determine whether the paper described the treatment investigated, and if so data on treatment were extracted.

# Data extraction

The patient and treatment characteristics that we felt should be available were formulated by the authors on the basis of clinical experience, available guidelines for description of treatments (for instance, Netherlands Association for Medical Rehabilitation), and consensus. These variables were then discussed with the national committee on cognitive rehabilitation in The Netherlands (consisting of rehabilitation physicians and psychologists) and the final set of variables was determined on the basis of consensus. The first and the second author reviewed the articles separately after selection and independently extracted the following data: (1) patient characteristics (number of patients, age, type of injury, time since injury); (2) treatment characteristics (treatment setting; duration and intensity of treatment; type of treatment, group and/or individual versus direct therapy or applied in daily life; discipline(s) of the therapists involved; involvement of the caregiver in treatment); and (3) treatment goals and outcome. Any disagreements were resolved by discussion until consensus was reached. Disagreements were mainly caused by incomplete description of characteristics in the literature.

### **RESULTS**

Ninety-five RCTs were selected (described in 96 papers; references Dirette and Hinojosa [1999] and Dirette, Hinojosa, and Carnevale [1999] describe the same study), published up to August 2010. The frequency of studies per domain (including all references) is shown in Table 2. Most studies were conducted in the area of language and communication (n = 25), visuospatial functioning (n = 24) and memory (n = 14).

### Patient characteristics

In total 4068 patients were studied in the 95 studies. The mean (SD) number of patients per study was 42.8 (47.0) varying from 4 to 360. The largest study by Vanderploeg et al. (2008), for instance, compared a protocol-specific cognitive didactic approach with a functional experimental approach in a group of 360 adult veterans or active duty military service members with moderate to severe traumatic brain injury. Most studies (n=37) were small with only 4 to 20 patients. Only seven studies included more than 100 patients.

The mean age of the total sample (n = 94) was 52.2 (SD 15.0; range 17–77) years. The older age groups were mostly stroke patients as in the study of Kalra, Perez, Gupta, and Wittink (1997) in which visual neglect was treated in a group with a mean age of 77.0 (8.6) years. Most studies (n = 56; 59%) selected patients aged between 20 and 60 years.

The time since brain injury was described in 87 studies and was 24.0 months on average (SD 34.0; range 5 days-82.5 months). The most acute patients were studied by Carter, Howard, and O'Neil (1983); inclusion of patients in the programme was at a mean (SD) of 4.8 (1.6) days. In most of the studies (n = 37; 43%), however, the time since injury was more than 1 year, while in 23% the patients were included less than 2 months after onset (n = 22).

Stroke was the most studied diagnosis (n = 54; 57%); in 22 studies (23%) patients with traumatic and closed head injuries were included. The remaining studies dealt with heterogeneous populations (n = 16) or other forms of brain injuries (n = 3) such as solvent-induced chronic toxic encephalopathy (van Hout, Wekking, Berg, & Deelman, 2008).

Further detailed information on patient characteristics can be obtained from the authors.

# Intervention characteristics

We applied the three sources for retrieval of treatment information as described in the methods section. In 40 papers the only available information was found in the paper itself. For instance, in the paper by Cheng and Man (2006) the Awareness Intervention Program (AIP) is described in the methods section, but no further information to other sources is given. In

TABLE 2 Study domains

Domain	N	%	<i>Author/year</i>
Awareness	2	2	Cheng & Man, 2006; Goverover, Johnston, Toglia, & Deluca, 2007
Visuospatial functioning	24	25	Antonucci et al., 1995; Cherney, Halper, & Papachronis, 2003; Dirette & Hinojosa, 1999; Dirette, et al., 1999; Edmans, Webster, & Lincoln, 2000; Fanthome, Lincoln, Drummond, & Walker, 1995; Jacquin-Courtois et al., 2010; Kalra et al., 1997; Kasten et al., 1998; Katz et al., 2005; Lincoln, Whiting, Cockburn, & Bhavnani, 1985; Luukkainen-Markkula, Tarkka, Pitkänen, Sivenius, & Hämäläinen, 2009; Neistadt, 1992; Poggel, Kasten, & Sabel, 2004; Robertson, McMillan, MacLeod, Edgeworth, & Brock, 2002; Rossi, Kheyfets, & Reding, 1990; Rusconi, Meinecke, Sbrissa, & Bernardini, 2002; Si Hyun et al., 2009; Taylor, Schaeffer, Blumenthal, & Grisell, 1971; Tsang, Sze, & Fong, 2009; Weinberg et al., 1977; 1979; Weinberg, Piasetsky, Diller, & Gordon, 1982; Wiart et al., 1997; Zeloni, Farnè, & Baccini, 2002
Memory	14	15	Berg, Koning-Haanstra, & Deelman, 1991; Doornhein & De Haan, 1998; Dou, Man, Ou, Zheng, & Tam, 2006; Freeman, Mittenberg, Dicowden, & Bat-Ami, 1992; Hildebrandt, Bussmann-Mork, & Schwendemann, 2006; Kaschel et al., 2002; Kerner & Acker, 1985; Ownsworth & McFarland, 1999; Rose et al., 1999; Ryan & Ruff, 1988; Schmitter-Edgecombe, Fahy, Whelan, & Long, 1995; Thickpenny-Davis & Barker-Collo, 2007; Twum & Parente, 1994; Westerberg et al., 2007
Attention	10	11	Barker-Collo et al., 2009; Couillet et al., 2010; Fasotti, Kovacs, Eling, & Brouwer, 2000; Gray, Robertson, Pentland, & Anderson, 1992; Malec, Jones, Rao, & Stubbs, 1984; Mazer et al., 2003; Niemann, Ruff, & Baser, 1990; Novack, Caldwell, Duke, Bergquist, & Gage, 1996; Sohlberg et al., 2000; Winkens, Van Heugten, Wade, Habets, & Fasotti, 2009
Executive functioning	6	6	Hewitt et al., 2006; Levine et al., 2000; Man et al., 2006; McPherson, Kayes, & Weatherall, 2009; Soong, Tam, Man, & Hui-Chan, 2005; Spikman et al., 2010
Apraxia	3	3	Donkervoort, Dekker, Stehmann-Saris, & Deelman, 2001; Smania et al., 2006; Smania, Girardi, Domenicali, Lora, & Aglioti, 2000
Language and communication	25	26	Bakheit et al., 2007; Dahlberg et al., 2007; David, Enderby, & Bainton, 1982; Denes, Perazzolo, Piani, & Piccione, 1996; Di Carlo, 1980; Doesborgh et al., 2004a; 2004b; Drummond & Rentschler, 1981; Elman & Bernstein-Ellis, 1999; Hinckley, Patterson, & Carr, 2001; Katz & Wertz, 1997; Lincoln et al., 1984; Lincoln & Pickersgill, 1984; Lincoln, Pickersgill, Hankey, & Hilton, 1982; Lyon et al., 1997; MacKay,

TABLE 2. Continued.

Domain	N	%	Author/year
Multi-domain	11	12	Holmes, & Gersumky, 1988; Meikle et al., 1979; Meinzer, Streiftau, & Rockstroh, 2007; Prins et al., 1989; Pulvermuller et al., 2001; Rochon, Laird, Bose, & Scofield, 2005; Shewan & Kertesz, 1984; Wertz et al., 1981; Wertz, Weiss, Aten, & Brookshire, 1986; Worrall & Yiu, 2000  Carter et al., 1983; Ruff et al., 1994; Salazar et al., 2000; Sarkamo et al., 2008; Soderback, 1988; Sturm & Willmes, 1991; Thomas-Stonell, Johnson, Schuller, & Jutai, 1994; Tiersky et al., 2005; van Hout et al., 2008; Vanderploeg et al.,
Total	95	100	2008; Watanabe, Black, Zafonte, Millis, & Mann, 1998

Dirette and Hinojosa (1999) and Dirette, Hinojosa, and Carnevale (1999) is the same study.

studies using medical or other devices (prisms, software programs or virtual reality programs; n = 7), the information should include the type of device used; in those cases further information is not necessary, for instance, type of prisms (Jacquin-Courtois et al., 2010) or a software program (Dirette, Hinojosa, & Carnevale, 1999). In 45 papers a reference is given to another study where a specific task which is practised during treatment or elements from the treatment are described, or the complete treatment is described as it had been evaluated before. This is the case in the study by Antonucci et al. (1995) who state that the training is specified in detail elsewhere. Most of the papers however state that "the treatment was derived from" or "the treatment was based on" and going to the reference given does not lead to specific information. The same holds for references to more general methods such as "role playing" where the reference does not give sufficient information making replication or application in one's own setting possible (i.e., a handbook). In only two papers a reference to a treatment manual is given which is retrievable through open access. This concerns the Attention Process Training, evaluated in the paper by Sohlberg, McLaughlin, Pavese, Heidrich, and Posner (2000) which can be obtained. We could find only one paper where the authors mention that further information about the treatment can be requested through one of the authors (Spikman, Boelen, Lamberts, Brouwer, & Fasotti, 2010). A summary table of the intervention characteristics can be obtained from the authors.

# Frequency, intensity and duration

Information on frequency, intensity and duration of treatment is summarised in Table 3. It is obvious that this information was not always given and that

		Total		Planned	Actual		
	n	M (SD)	n	M (SD)	n	M (SD)	
Hours	79	37.5 (56.2)	72	35.9 (54.7)	21	30.4 (40.0)	
Weeks	88	9.1 (9.1)	80	8.7 (9.0)	22	10.5 (9.4)	
Hours/week	78	4.0 (4.0)	70	3.9 (3.8)	21	4.1 (3.6)	

TABLE 3 Frequency, duration and intensity of treatment

the treatments described varied greatly. For instance, Hewitt, Evans, and Dritschel (2006) offered only 30 minutes of planning training, while in the study by Wertz et al. (1981) a group of veterans received 48 weeks of aphasia treatment with 8 hours of treatment each week (352 hours in total).

The total number of treatment hours was known for 79 studies (83%); the number of weeks the treatment was offered was known for 88 studies (93%), and the number of hours per week for 78 studies (82%). Planned treatment was rarely the same as actual treatment delivered and both are shown in Table 3. The mean (SD) actual number of hours treatment given was only reported in 21 studies where it was 30.4 (40.0) hours in total, equivalent to 4.0 (3.6) hours a week.

In 46 studies (48%) treatment was offered individually, while in 8 studies (8%) it was offered in groups. In some cases individual and group treatment was combined in one treatment programme (n = 7; 7%) and occasionally individual treatment and group treatment were compared in an experimental setting (n = 2; 2%) as in the study by Vanderploeg et al. (2008). Information on treatment form was not available for one third of the studies (n = 32).

In 18 studies the treatment was not restricted to therapy sessions, but homework was also arranged (19%). Practising only at home without therapy sessions was studied in 5 cases (5%). For instance, in the study by Kasten, Wust, Behrens-Baumann, and Sabel (1998) a computer was placed at the patient's home enabling daily practice. This kind of information was lacking in 75% of the studies (n = 71).

In 73 studies (77%) there was no information available about the involvement of caregivers in the study or in treatment. In only four studies was the caregiver actually involved in therapy (4%), while in 12 studies (13%) the caregiver was involved in outcome measurements but not therapy. Occasionally the caregiver was asked to give informed consent, or was only involved in the control treatment (n = 5; 5%).

In Table 4 the setting in which treatment was offered is presented. This information was missing in 16 studies (17%). In most cases patients were treated in the experimental setting of the study after being referred from a

Setting	N	%
Hospital/stroke unit	20	21
Laboratory/research centre/experimental setting	19	20
No information available	16	17
Multiple settings	11	12
Community/home	7	7
Rehabilitation centre (inpatient and outpatient combined)	9	10
Rehabilitation centre (inpatient)	6	6
Rehabilitation centre (outpatient)	5	5
Other	2	2
Total	95	100

TABLE 4
Treatment setting

hospital or outpatient clinic (n = 19; 20%). Patients in hospitals (n = 20; 21%) and rehabilitation centres (inpatient as well as outpatient programmes) were also studied frequently (n = 20 in total; 21%).

In Table 5 the disciplines involved in treatment are presented. In 19 studies (20%) the therapist was not specified and in 22 studies (23%) information about the clinicians involved was not available at all. We could not find information about the necessary experience, knowledge or skills of the staff in any studies. Speech and language pathologists were most often specified (n = 12; 13%) and in 13% of the cases a computer with or without a therapist involved was used for treatment (n = 12).

Detailed information on stated treatment goals and outcome measures, and type of intervention can also be obtained from the authors. Treatment goals were not specified in 16 studies (18%), while in many cases the goals for treatment were only specified very generally such as "to improve memory function". In some studies the treatment goals can be derived from the outcome measures that were used to evaluate the treatment effect. For instance, Antonucci et al. (1995) did not describe treatment goals explicitly, but the treatment effect was measured in terms of the presence of neglect disorders in daily life situations; one could derive from this information that the training was supposed to influence the level of activities. In many studies outcome measures on the level of impairments (i.e., neuropsychological tests) were combined with outcome measures at the level of activities. In only a few studies treatment was directed at participation and well-being (for instance, in the study of Lyon et al., 1997).

Intervention studies were designed differently in terms of the treatment to which the experimental treatment was compared. In most cases a control treatment was offered (n = 28; 30%) or the experimental treatment was compared to care as usual (n = 29; 31%). In 14 studies the control groups

Discipline	N	%
Clinician/therapist	22	23
No information available	19	20
Computer, in combination with therapist	12	13
Speech and language pathologist	12	13
Combinations of disciplines	12	13
Experimenter/researcher/examiner	6	6
Occupational therapist	5	5
Psychologist	5	5
Volunteer/student	2	2
Total	95	100

TABLE 5
Disciplines offering treatment

received no treatment (15%) or two experimental treatments were compared (n = 11; 12%). In the remaining studies combinations of control treatments were offered (n = 13; 14%).

Where the content of the experimental treatment was described, the information was structured in many different ways, with no common headings or format. In 19 studies only the type of treatment was given without any detail about the content of treatment, for instance, "speech therapy" (Lincoln et al., 1984), while in other studies the content of treatment was presented in appendices in which the content per session was described, for instance, in the study of Man, Soong, Tam, and Hui-Chan (2006). In some studies a reference to a specific training is given and thus to be found in detail elsewhere; for instance, Dahlberg et al. (2007) refer to a workbook on group treatment for social skills.

Details about the control intervention were even less. In 34 studies only "control treatment" or "conventional treatment" was mentioned. In a few studies also a reference is given for a description of conventional therapy, for instance, in the study of Prins, Schoonen, and Vermeulen (1989). Additionally, in 19 studies there was no treatment given in the control group. A remarkable observation is that in many studies the outcome measures are all described in high detail while the intervention is only described in very general terms. A detailed description of the differences between control and experimental treatments was not available in any report, but where two experimental treatments were given, differences were sometimes tabulated.

#### Effectiveness of treatment

In more than half of the studies, the experimental treatment was more effective than the control treatment (n = 53), while in one quarter of the studies there was no difference between experimental and control treatment (n = 22).

In the remaining studies (n = 17; 20%) the effects showed more variation (i.e., treatment altered function in some domains more than in others).

# DISCUSSION

The evidence relating to the effects of cognitive rehabilitation has evolved rapidly since 2005: the Cicerone reviews included only 46 class I studies up to 2002 (review of 2000 included 29 class I studies, and in 2005 another 17 class I studies were included) (Cicerone et al., 2000; 2005) while we could include 95 class I studies up to 2010 representing 4068 patients in total. Just recently, a new update of the Cicerone review (Cicerone et al., 2011) was published in which another 19 class I studies were added. Our review has thus shown that many more randomised studies have been carried out, albeit the number of larger studies (100 patients or more) remains low at seven. Cicerone et al. concluded that 37 out of the 47 studies (79%) showed a beneficial effect of cognitive rehabilitation. Cicerone et al. (2011) conclude in their latest update that there is now sufficient information to support evidence-based clinical protocols, and to design and implement empirically-supported treatments for cognitive disabilities after stroke and TBI, but is that really the case?

In our review in 22 out of 95 studies (23%) there was no difference between the experimental treatment and control treatment and in another 20% the effects were partial (i.e., not on the primary or all outcome measures); making this distinction in treatment effectiveness, our conclusions are less promising. In this systematic review, the 95 RCTs on the efficacy of rehabilitation for cognitive deficits have been analysed thoroughly and give the opportunity to build a detailed database on evidence-based cognitive rehabilitation which can be used by clinicians in selecting the current best evidence for treatment decisions and by researchers in formulating new research objectives. However, the review also shows that the treatments are only occasionally described in sufficient detail to allow replication of the study or use of the treatment clinically after reading the paper, and the control treatment is similarly hardly ever described. However in 54% of studies, a statistically significant effect of the experimental treatment was shown, but even this positive conclusion is qualified by the small size of most studies – only seven studies included 100 or more patients and 39% had less that 20 patients – and by the heterogeneity of the cognitive domains studied.

The details given concerning the treatment investigated should be set in the general context of the description of other healthcare interventions. The usual model used is that of a drug intervention, where the physical, pharmacological and other characteristics of the treatment are always carefully described and controlled. However in practice almost all other healthcare interventions,

including surgery, behavioural therapy and rehabilitation therapies are poorly described, if described at all. In a semi-systematic review of non-pharmacological treatment trials, Glasziou, Meats, Heneghan, and Shepperd (2008) concluded that the clinician trying to replicate non-pharmacological treatments in practice hardly ever has sufficient detail about the "how to". Thus the findings in this review should not be surprising. Boutron et al. (2005) acknowledged the specific methodology of non-pharmacological trials and suggested an alternative checklist to report such trials.

There are several reasons for this state of affairs. First, there is no agreed framework for describing rehabilitation treatments (Wade, 2005) or indeed complex interventions. In the absence of a standardised way to describe an intervention many people simply do not attempt it. Second, researchers are likely to face great difficulties in publishing the intervention itself. Moreover, it will be especially difficult to publish a description of an intervention proven not to work, but this is of equal or greater importance. Thirdly, although the importance of complex interventions has been recognised, the standards applied to randomised studies do not require full description of the intervention. Even the CONSORT statement structured for complex interventions requires only one item saying "sufficient details to allow replication, including how and when they were actually administered" should be given in which the operationalisation of "sufficient details" is up to the author, and only one item in the registration of RCTs on clinicaltrial gov concerns the treatment, and no guidance is given. Perera, Heneghan, and Yudkin (2007) proposed a graphical method for depicting RCTs of complex interventions which should support the comparison of interventions in the trial. This is a first step to a clearer description of non-drug treatments.

The difficulty of knowing what rehabilitation treatment is has been likened to a Russian doll (Whyte & Hart, 2003), However, at least one suggestion has been made on how to improve the description of therapies (Wade, 2005) and some recently evaluated treatments have been published in detail (Winkens, Van Heugten, Wade, & Fasotti, 2009). In addition, the American Congress of Rehabilitation Medicine just recently published the beta version of a "Cognitive rehabilitation manual" of which the authors state it provides step-by-step instructions for the interventions recommended in the Cicerone reviews (www.acrm.org/cognitive-rehabilitation-manual). The first edition of this manual, which is expected in the Spring of 2012, should show whether the lack of knowledge on the content of treatment from the journal articles reviewed here is indeed sufficiently covered.

Drawing general conclusions about "cognitive rehabilitation" is further complicated by the wide range of cognitive problems studied. We categorised outcomes into eight domains used by others, but the severity and complexity of problems faced by patients included varied considerably. Moreover the outcome measures used in cognitive rehabilitation studies are also often

limited to the level of impairment, making it difficult to judge the clinical relevance and importance of any effect. The underlying disease also varies and, with this, it is likely that the associated impairments (cognitive and non-cognitive) vary greatly. Lastly the patients studied are usually young and quite selected, but most patients with cognitively based problems are old and have multiple problems. In contrast there are many more studies of physiotherapy; they tend to study only two domains, mobility and arm function, and they usually include more patients.

TABLE 6
Checklist for description of rehabilitation interventions

Item	Aim
Patient characteristics (input)	Is my patient comparable to the patients in the study?  • Age  • Gender  • Type of injury  • Time since injury  • Indications for treatment  • Contra-indications
Intervention characteristics (process, structure, context)	What is the treatment?  • Activity: - frequency, intensity, duration - Individual or group - involvement of caregivers/others - direct or contextual treatment  Can I offer this intervention in my own setting?  • Context: - organisation (healthcare setting) - theoretical basis  • Necessary resources: - physical (equipment) - knowledge (of staff) - skills (of staff)
Treatment goals and outcome (anticipated outcome)	<ul> <li>What are the costs/benefits for my patient and his family?</li> <li>Treatment goal (proximal, immediate)</li> <li>Treatment goal (distal, general)</li> <li>Expected effect size <ul> <li>for patient</li> <li>for caregivers</li> </ul> </li> <li>Differential effects for subgroups</li> <li>Adverse effects</li> <li></li> </ul>

This review has its own limitations. It is quite possible that we failed to identify some RCTs, although in fact our strategy identified those found in other systematic reviews. It is possible that detailed descriptions of some interventions were published somewhere, but were not referred to in the original study. It is also possible that some studies had their own written protocols.

We decided that we would limit our attempts to find details to those (a) that most clinicians would make, and (b) that were most likely to succeed (because about half of all studies were undertaken over 10 years ago, making it unlikely that we would find the researcher or that they would still have the treatment protocol). We checked whether a reference to a treatment description was given but could not search systematically for descriptions, particularly if published after the main study. In searching we put ourselves in the position of a working neuropsychologist or other professional in following up a reference to an accessible journal.

Third, our data extraction obviously may have been weak in terms of the items extracted. However, none of the studies gave any additional details not included in the categories we used, and we feel that the information we searched for was a reasonable minimum.

The main implication of this review is that the evidence on cognitive rehabilitation is growing rapidly, but a clinician cannot pursue true "evidence-based" cognitive rehabilitation at present because there is insufficient published detail about the nature of any treatment that has been evaluated, and the evidence for effectiveness for any single treatment is not sufficient.

In future, studies need to give a clear description of the treatment offered both in the experimental group and in the control group. We would suggest that the characteristics shown in Table 6 would be a reasonable minimum. Such a checklist should ideally be considered by an international group, such as the CONSORT group or the Special Interest Group on Neuropsychological Rehabilitation (SIG-NR) of the World Federation for Neurorehabilitation (WFNR), making standardised description of interventions a fixed habit of transparent reporting of trials.

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