

**A Systematic Review of physical activity and exercise interventions to improve health,
fitness and wellbeing of children and young people who use wheelchairs**

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ABSTRACT

Aim: To perform a systematic review establishing the current evidence base for physical activity and exercise interventions that promote health, fitness and wellbeing, rather than specific functional improvements, for children who use wheelchairs. **Design:** A systematic review using a mixed methods design. **Data sources:** A wide range of databases including Web of Science, PubMed, BMJ Best Practice, NHS EED, CINAHL, AMED, NICE, PsychINFO were searched for quantitative, qualitative and health economics evidence. **Eligibility:** participants: children/young people aged >25 years who use a wheelchair, or parents and therapists/carers. Intervention: home or community-based physical activity to improve health, fitness and wellbeing. **Results:** Thirty quantitative studies that measured indicators of health, fitness and wellbeing, and one qualitative study were included. Studies were very heterogeneous preventing a meta-analysis, and the risk of bias was generally high. Most studies focused on children with cerebral palsy and utilised an outcome measure of walking or standing, indicating that they were generally designed for children with already good motor function and mobility. Improvements in health, fitness and wellbeing were found across the range of outcome types. There were no reports of negative changes. No economics evidence was found. **Conclusions:** It was found that children who use wheelchairs can participate in physical activity interventions safely. The paucity of robust studies evaluating interventions to improve health and fitness is concerning. This hinders adequate policy making and guidance for practitioners, and requires urgent attention. However, the evidence that does exist suggests that children who use wheelchairs are able to experience the positive benefits associated with appropriately designed exercise

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WHAT IS ALREADY KNOWN?

Physical activity is essential for health and wellbeing, but for children who use wheelchairs participation rates are low.

Health policy and guidance on physical activity and exercise for disabled children is lacking.

WHAT ARE THE NEW FINDINGS?

No reports of adverse outcomes were found, indicating that children who use wheelchairs can participate in exercise safely.

Models of delivery utilising an initial period of individual or group supervision followed by home-based programmes appear effective and could be sustainable within health and wheelchair service practice.

Evidence on the effectiveness of health and fitness focused exercise interventions for children who use wheelchairs is weak, yet suggests that these children can experience the usual benefits associated with well-designed interventions.

INTRODUCTION

Participation in exercise and physical activity is vital to maintain health and fitness, and for children it is essential for emotional and physical development [1]. However, children with mobility impairments face multiple barriers to participation and their uptake rates are lower than those of typically developing children [2,3]. Children with mobility impairments also tend to be more frequently overweight or obese, physically weaker and less fit than their typically developing peers [4,5]. This predisposes them to long-term health risks including type-2 diabetes and cardiovascular disease, and results in a “debilitative cycle” of poor fitness, increased disability, and further reduced health and wellbeing [6]. The consequence is impaired quality of life and future health prospects for the children, and spiralling demands on families and health and social care services. Given that there are 770,000 children and young people under the age of 16 years living in the UK with a disability (1 in 20 children [7]), this is an important issue that must be tackled.

Rationale

Government policy and guidance on physical activity is well established for adults and typically developing children [8]. This is also reflected in professional body position statements [9], but there is a lack of guidance on physical activity for children with mobility impairments. In the UK, in response to the NICE guidance which omitted disabled children, a specific briefing paper was published and promoted access to play and physical activity for disabled children and young people [10], citing physiological benefits as well as enhanced confidence, self-esteem, resilience and social inclusion. Whilst the benefits of increased physical activity levels are clear, it is not apparent what activities actually promote the health and fitness benefits desired for this group of children.

A number of systematic reviews have been undertaken to evaluate a range of interventions to improve physical function in children with mobility impairments [e.g., 11-16]. However, the studies reviewed primarily focused on occupational- or physiotherapy-type therapeutic interventions and outcomes, by which they treated a specific functional impairment such as a joint range of motion or walking function, with a targeted intervention or exercise. Typically such interventions do not provide insights in the wider benefits of exercise and physical activity for health (which we define as the composite of physical, mental and social aspects required for an individual to function, adapt and self-manage) and fitness (which we define as the set

attributes required for physical function, that can each be related to specific tests, e.g., muscle strength or maximal cardiovascular capacity). By their definitions, health and fitness are interdependent. Consequently, there is a gap in our knowledge relating to (i) intervention effectiveness for health and fitness outcomes, (ii) what are the most suitable primary outcomes to assess health and fitness, (iii) engagement and long-term adherence, (iv) psychological preparation/intervention to promote readiness and motivation to exercise in this group of children, (v) a lack of focus on the most physically disabled children (which for the purposes of this review, we define as those who use wheelchairs all or some of the time), and (vi) evaluation of health economics (a further discussion of these points is presented in the protocol of this review [17]). We are not aware of any published systematic reviews of physical activity interventions with the aim of improving health and fitness of children and young people who use a wheelchair.

Objectives

The purpose of this systematic review was to establish the current evidence base for exercise or physical activity interventions to improve the health and fitness of children and young people who use wheelchairs. We specified the following objectives:

1. To determine the effectiveness and cost-effectiveness of physical activity interventions for children and young people who use a wheelchair within a home and community setting.
2. To describe programme theories/concepts or models and identify appropriate outcome measures in current interventions.
3. To explore the views and experiences of children, young people and their parents, and key stakeholders about physical activity interventions for wheelchair users.

METHODS

Design

We set out to undertake a mixed methods systematic review of quantitative, qualitative and health economics evidence to identify and evaluate effectiveness of physical activity interventions. However, we did not find any economic evidence that met our inclusion criteria. The search strategies and type of evidence to be included in this stream is described in the

protocol [17] (PROSPERO registration: CRD42013003939), but it will not be described further in this review. The review broadly followed design, methods and processes of the Cochrane Effective Practice and Organisation of Care Group (EPOC) [18], for the synthesis of effect evidence, and Reporting guidance set out in Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) [19].

Inclusion criteria

Population

Our primary objective was to review studies with children and young people who use wheelchairs. Wheelchair use was broadly defined as requiring the use of a wheelchair all or some of the time, even if only for long distances. However, initial pilot searches found that many studies include physically disabled children who do not use wheelchairs along with those who do, or that wheelchair use is not reported for groups that would be expected to use wheelchairs, based on functional classification or medical condition. Consequently, in addition to the studies that reported their samples included “wheelchair users”, studies with probable but “undefined wheelchair use”, as judged by our multi-disciplinary team, were also included. These separate groups of studies are handled independently in reporting and synthesis.

Children and young people aged <25 years (this upper age limit was chosen to ensure that data from young people under 18 years were not lost, if they were part of a sample including young adults).

Types of intervention

Any intervention that included physical activity or exercise at home or in the community. Physical activity and exercise were defined as being general or structured movement of the body that would increase energy expenditure.

To allow a more detailed examination of effectiveness of specific types of exercise or physical activity, interventions were grouped according into activity mode (Table 1). Interventions aiming to improve specific disabilities or impairments with targeted programmes (such as physiotherapy or occupational therapy) were excluded.

Table 1. Physical activity modes and definitions used for grouping and analysis in this review.

Exercise mode	Further definition for this review
Walking	Including treadmill
Cycling	Either outdoor or stationary, and peddle or hand/crank
Progressive resistance training (PRT)	Lifting of an external resistance, performed across multiple sets (3-5) of a low number of repetitions (<20)
Functional / closed kinetic chain exercises	Lifting, controlling and moving one's own body weight, during movement's replicating real-life movements, e.g., stepping forwards or up on a step, excluding walking only.
PRT using functional / closed kinetic chain exercises	Functional exercises, with the addition of external weight; either held in hands or as weighted vest
Sport/games-based	Including group or individual game or play based activities
Spirometry exercise	Externally resisted inspiratory muscle training
Nintendo Wii	Utilising "off-the-shelf" games
Motivation	Any study providing psychological or emotional feedback, or music to enhance motivation during a physical activity
Education	Education to enhance participation levels
Complex programme	An intervention the broadly reflects a combination of any above interventions, that cannot be defined as primarily one or another for any reason. This includes counselling to determine individualised activities. The combination is defined for each intervention in included study tables.

Types of outcome measures

Any evidence related to the effectiveness or cost-effectiveness of an intervention was considered. To evaluate an intervention's potential to improve specific components of health, fitness and wellbeing, quantitative data were categorised according to outcome type. All outcome measures reported in the included studies could be defined as being in one of the following categories. These categories align with those described for assessment in our main project study protocol (20), and were used to organise the data only and not as inclusion criteria.

- 1) Exercise capacity / fitness
- 2) Metabolism
- 3) Body composition / weight
- 4) Quality of Life
- 5) Respiratory function
- 6) Physical function
 - a. Gross motor function
 - b. Mobility
 - c. Strength
- 7) Physical activity levels, including self-reported, and attitudes to physical activity

Several of the included studies evaluated walking capacity in tests ranging from 6-10 min. These studies were inconsistent in their classification of this outcome as either exercise capacity (outcome type 1) or tests of ambulation and mobility (outcome type 6b). For clarity and consistency in the present systematic review, we classified these to be exercise capacity. Walking tests of less than one minute or over short distances were classified as mobility.

In most cases, a single study provided data of several outcome types.

Types of studies

Preliminary pilot searches of existing literature, found few randomised controlled trials (RCTs) of exercise or physical activity interventions for children and young people who use wheelchairs. Therefore, we decided to include all relevant study designs (e.g., RCT, non-RCT, controlled before-after, survey, qualitative) to draw from as wide an evidence base as appropriate. Consequently, in some studies there was no comparison or control group included, and in others the control group consisted of usual care. In cases where the comparison was between two different targeted interventions, these were treated as separate treatments in the same study.

Only publications in English, Welsh or German language were included, reflecting translation capacity within the study team. Grey literature, including contemporary local government/agency, charity reports and intervention programme evaluations which have not been subject to peer review, were also included to limit publication bias and ensure that all relevant literature may be located. Grey literature was screened against criteria of Completeness, Accuracy, Relevance and Timeliness (CART) to determine inclusion or exclusion.

Search methods

In addition to using the population, interventions, comparators, outcomes and study design approach (PICOS), the Setting, Perspective, Phenomena of Interest, Comparison and Evaluation (SPICE) Framework [21] was used to refine the design the search methods for qualitative and grey literature exploring views and experiences of exercise and interventions as the phenomena of interest. Medical Subject Heading (MeSH) keywords were combined with free-text keywords using Boolean operators (and/or/not) to search multiple databases chosen

to reflect the multidisciplinary nature of this review, including but not limited to Web of Science, PubMed, BMJ Best Practice, NHS EED, CINAHL, AMED, NICE, PsychINFO (see supplementary material of [17] for SPICE framework, complete list of databases and specific search terms). Articles included in the review were also searched for ancestral references. Grey literature was searched for by hand searching of Opengrey.eu, key journals, website searches (including Google and Google scholar) and approaching personal contacts. An example search log is reported in supplementary material of this publication.

Selection of studies

All identified studies from the searches were assessed by two reviewers independently for inclusion. Results were initially screened according to title and abstract, the full text of those considered suitable were then further assessed for relevance against the inclusion/exclusion criteria. Disagreement between reviewers was resolved by discussion or arbitration by a third person when necessary.

Quality assessment

The risk of bias of quantitative evidence from each included study was assessed using the Cochrane risk of bias tool in line with GRADE (Grading of Recommendations, Assessment, Development and Evaluation) approach [22]. Risk of bias was rated as low, unclear or high depending on whether the randomisation sequence is adequate, randomisation is concealed, baseline characteristics and outcomes are similar in treatment and control groups, missing data was handled appropriately, experimenters were blinded, all outcome measures were reported, and the study was protected from possible contamination and any other risks of bias. The GRADE approach specifically assesses:

- Methodological flaws within the component studies
- Consistency of results across different studies
- Generalisability of research results to the wider patient base
- How effective the treatments have been shown to be.

When using the full GRADE approach, the evidence is not rated study by study, but a composite rating is made across studies for specific outcomes. In this review, almost all included studies were single stand-alone studies with heterogeneous interventions and a very high number of different outcome measures (these can be seen in the data extraction tables).

Therefore, it was not possible to apply the full GRADE assessment for every heterogeneous outcome.

Qualitative studies were assessed with the Critical Appraisal Skills Programme (CASP) tool for qualitative research. As only one study was located it was not possible to apply GRADE CERQual to assess the confidence in the synthesised finding [23].

Data abstraction

Data were extracted by the two reviewers, including study design, participants, intervention type, outcome measures and results. Data extracted by each reviewer were combined and discrepancies were discussed and where necessary resolved by arbitration by a third person.

The intervention studies included in this review were too heterogeneous in terms of participant characteristics, intervention design and outcome measures to allow a meta-analysis. Therefore, extracted data were summarised and presented in harvest plots, which offer the possibility to present complex and diverse data within easily interpreted graphical formats, analogue to forest plots [24]. These plots were constructed to indicate whether the specific intervention type (listed in Table 1) used by each study was able to improve at least one measure of health or fitness (listed above). Full data extraction tables are included in the supplementary materials.

The studies reporting interventions including wheelchair users were managed separately from those with “undefined wheelchair use”; they were differentiated in harvest plots and considered separately in data analysis and synthesis.

Validity, reliability and rigour

The review has been designed to follow Cochrane methods and processes, including the use of double independent data processing and assessment of risk of bias using validated tools. None of the authors have any conflicts of interest that would affect their interpretation of evidence.

RESULTS

Included studies

Searches returned 9,806 hits (excluding duplicates), of which 31 papers from 28 studies [25-55] were deemed to meet the criteria for inclusion in this review. Only one qualitative study [55] and no health economics studies that met the inclusion criteria were found. The flow chart of article screening and inclusion is in Figure 1. Tables 2, 3 and 4 present characteristics of all included studies, their article number (#) and risk of bias. Of the 31 included articles, 11 (#1-10 and #31) reported their sample to include wheelchair users. Only seven quantitative papers were considered to be low risk of bias (articles #3,4,13,15,19,20,23), seven were considered to have an unclear risk of bias (#2,11,12,14,17,21,22), meaning 16 articles were rated as high risk of bias (#1,5-10,16,18,24-30).

A wide range of interventions were utilised by the included studies (Table 1). The most common categorisation of intervention type was “complex” (n=8), which entailed combinations of exercise modes (#5,7,10,28) or life-style interventions including counselling and individualised physical activity (#3,4,6,21). Interestingly, the majority (n=6) of these complex interventions were with “wheelchair users”. The variability in intervention was even apparent amongst those categorised similarly, which often had disparate training protocols, e.g. exercise intensity, duration or number of repetitions performed, and location of training. This led to variability in intervention effectiveness (reported below).

Outcome measures all fell within the categories listed in Methods under *types of outcome measures*. However, there was little consistency in the tools or protocols used to assess each outcome type (see data extraction tables in supplementary material). For instance, assessments of cardiovascular fitness/exercise capacity included a 6-min cycle, peak aerobic power, 9-min run, energy expenditure index and self-assessment. Mobility assessments included walking speed and distance in 30 s-2 min, time to raise from the floor or a chair, step-up repetitions and balance stability. Standard tools were used to assess quality of life and wellbeing (CAPE and PEDS-QL were common), and domains D and E were frequently selected from the Gross Motor Function Measure. Similar variability existed amongst all other outcome types, making comparison between interventions difficult.

Sample size in the treatment group of included studies ranged from 2-52, with a total of 725 children and young people reported in the included studies. Two studies included only males (#1,29), it was not possible to determine sex distribution in four, and in the rest the sample

typically consisted of ~60 % males. Age of participants ranged from 3-25 years, although ~7-14 years was common. It was notable that cerebral palsy was the most frequent cause of disability and that most participants were ambulant to some extent. Gross Motor Function Classification System, where reported, was typically level 1-3. Only four studies reported including GMFCS level 4 or 5. There were no major concerns about methodological limitations in the qualitative study (31)

Table 2. Characteristics and risk of bias of included quantitative studies including “wheelchair users”. Organised by study design.

Article #	Author (Year)	Intervention type (exercise mode)	Setting; nationality	Participants: cause of impairment; intervention and control <i>n</i> , % male; age; GMFCS [when available]	Duration; follow-up	Risk of bias
<i>RCTs including wheelchair users</i>						
1	Jansen, et al. (2013)	Cycling: arm and peddle	Home; Netherlands	Duchenne muscular dystrophy; I: 17 C: 13 ; 100%; range 7-13 yrs; late ambulatory phase or wheelchair dependent	24 weeks; 24 weeks	High
2a	Van Den Berg-Emons, et al. (1998)	Sport/games-based aerobic. (4 times per week)	Community, group; Netherlands	Cerebral palsy (spastic); I:10 C:10; 55%; range 7-13 yrs; “half of the children was ambulant, the other half wheelchair-bound”	9 months; 3 months	Unclear
2b	Van Den Berg-Emons, R. J., et al 1998	Sport/games-based aerobic. (2 times per week)	Community, group; Netherlands	Cerebral palsy (spastic); I:9 C:9; unknown%; range 7-13 yrs; “half of the children was ambulant, the other half wheelchair-bound”	9 months; -	Unclear
3	Van Wely, et al. (2014a) [part of same study as #4]	Complex: Life-style counselling and fitness training	Combined home and supervised group; Netherlands	Cerebral palsy; I: 23 C: 23; 57%; range 7–13 yrs; GMFCS 1-3	6 months; 6 months	Low
4	Van Wely, et al. (2014b) [part of same study as #3]	Complex: Life-style counselling and fitness training	Combined home and supervised group; Netherlands	Cerebral palsy; I: 23 C: 23; 57%; range 7–13 yrs; GMFCS 1-3	6 months; 6 months	Low
<i>Non-RCTs including wheelchair users</i>						
5	Andrade, et al. (1991)	Complex programme: PRT and cardiovascular exercise, and psychosocial component	Community, group; Canada	Spina bifida; I : 8 C: 5; 50%; range 7-13 yrs; 6 “wheelchair users” and 7 “community ambulator”	10 weeks; -	High
6	Buffart, et al. (2010)	Complex: Life-style counselling and personalised activities	Mixture of home and supervised; Netherlands	A 17-year-old male with myelomeningocele and hydrocephalus, non-functional ambulator and a 23-year-old female with unilateral cerebral palsy, GMFCS 1	10 weeks; -	High
7	Fragala-Pinkham, et al. (2005)	Complex programme: PRT and aerobic exercise.	Group supervised phase followed by home phase; USA	Physical or other developmental disabilities; group phase: 9 home phase: 7; 78%; range 5-9 yrs; GMFCS 1-3	14 weeks group; 12 weeks home	High
8	Gordon, et al. (2013)	Nintendo Wii	Supervised; Jamaica	Cerebral palsy; I: 7 C: -; 57%; range 9-12 yrs; 4 wheelchair users.	6 weeks; -	High
9	Odman & Oberg (2005)	Functional exercises (combined Lemo + Move&Walk interventions)	Group, location unclear; Sweden	Cerebral palsy; I:52 C: -; 59%; unknown%; range 3-16 yrs; GMFCS 1-5	4 weeks intensive; 1 year voluntary participation	High
10	Unnithan, 2007	Complex programme: PRT and walking	Group; Greece	Cerebral palsy; I: 7; C: 6; 31%; range 14–18 yrs; GMFM (D&E) ~30%	12 weeks; -	High

Table 3. Characteristics and risk of bias of included quantitative studies with “undefined wheelchair use”. Organised by study design.

Article #	Author (Year)	Intervention type (exercise mode)	Setting; nationality	Participants: cause of impairment; intervention and control <i>n</i> , % male; age; GMFCS [when available]	Duration; follow-up	Risk of bias
<i>RCT with population with undefined wheelchair use</i>						
11	Choi, et al. (2016)	Spirometer exercise with vs without additional therapy	Mixture of individual and supervised in clinic; Korea	Cerebral palsy; I:25 C:23; 52%; range: 8-15 yrs; GMFCS 1-5	4 weeks; -	Unclear
12	De Groot, et al. (2011)	Walking	Home, supervised; Netherlands	Spina Bifida; I: 18 C: 14, 56%; mean ~10.7 SD ~2.8 yrs; “community ambulatory”	12 weeks; -	Unclear
13	Demuth, et al. (2012) [part of same study as #15]	Cycling: using a complex structure of strengthening and cardiorespiratory phases in each session	Clinic; USA	Cerebral palsy (spastic diplegic); I: 28 C: 29, 48%; range 7–18 yrs; GMFCS 1-3	12 weeks; -	Low
14	Dodd, 2003	PRT with functional exercises	Home; Australia	Cerebral palsy (spastic diplegic); I: 11 C: 10; 48%; range 8 to 18 yrs; GMFCS 1-3	6 weeks; 12 weeks	Unclear
15	Fowler, et al. (2010) [part of same study as #13]	Cycling: using a complex structure of strengthening and cardiorespiratory phases in each session	Clinic; USA	Cerebral palsy (spastic diplegic); I: 26 C: 26, 48%; range 7-18 yrs; GMFCS 1-3	12 weeks; -	Low
16a	Gates, et al. (2012)	Walking	Home; USA	Cerebral palsy (spastic bilateral); I: 14 C: -; 54%; range 6-13 yrs; GMFCS 2-4	12 weeks; 4 weeks	High
16b	Gates, et al. (2012)	PRT (including some functional exercises)	Home; USA	Cerebral palsy (spastic bilateral); I: 12 C: -; 54%; range 7-13 yrs; GMFCS 2-4	12 weeks; 4 weeks	High
17	Katz-Leurer, et al. (2009)	Functional exercises	Home with weekly phone support; Israel	Traumatic brain injury or cerebral palsy; I:8, C:10; 70%; range 7-13 yrs; GMFCS 1-2	6 weeks; 6 weeks for intervention group only	Unclear
18	Maher, et al. (2010)	Education (internet-based)	Home; Australia	Cerebral palsy; I:20 C:21; 63%; range 11-17 yrs; GMFCS 1-3	10 weeks; 10 weeks	High
19	Scholtes, et al. (2010) [part of same study as #20]	PRT with functional exercises	Community, small groups; Netherlands	Cerebral palsy (spastic); I: 24 C: 25; 59%; 6-13 yrs; GMFCS 1-3	12 weeks; 6 weeks	Low
20	Scholtes, et al. (2012) [part of same study as #19]	PRT with functional exercises	Community, small groups; Netherlands	Cerebral palsy (spastic); I: 24 C: 25; 59%; 6-13 yrs; GMFCS 1-3	12 weeks; 6 weeks	Low
21	Slaman, et al. (2014)	Complex: Life-style; counselling and cardio-pulmonary fitness.	Combined supervised in	Cerebral palsy; I:17 C:19; 90%; range 16–25 yrs; GMFCS 1-5	6 months; 6 months	Unclear

			centre and home; Netherlands			
22	Verschuren, et al. (2007)	Functional exercises	Community, group; Netherlands	Cerebral palsy (spastic); I: 32 C: 33; 65%; range 7-18 yrs; GMFCS 1-2	8 months; -	Unclear
23	Wang, et al. (2013)	Motivation: Music during PRT	Home; Taiwan	Cerebral palsy; I:18 C:18; 75 %; 5-13 yrs; GMFCS 1-3	6 weeks; 12 weeks	Low
<i>Non-RCT with population with undefined wheelchair use</i>						
24	Blundell, et al. (2003)	Functional exercises	Group; Australia	Cerebral palsy; I: 7 C: -; 88% range 4 -8 yrs; Motor-Assessment Scale, sit-to-stand item 1.5	4 weeks; 8 weeks	High
25	Damiano, et al. (1995)	PRT	Setting unclear; USA	Cerebral palsy (spastic diplegia); I: 14 C: -; 71%; range 6-14 yrs; 11 independent ambulators, 3 required hand-held assistive devices	6 weeks; -	High
26	Chen, et al. (2012)	Cycling (in Virtual Reality)	Home; Taiwan	Cerebral palsy; I: 13 C: 14; 68%; range 6–12 yrs; GMFCS 1-2	12 weeks	High
27	Crompton, et al. (2007)	PRT with functional exercises	Clinic, group; Australia	Cerebral palsy; I: 7 C: 6; 40%; range 6–14 yrs; GMFCS 1-3	6 weeks; 6 weeks	High
28	Kelly & Legg (2009)	Complex programme: PRT, aerobic exercise and sports	Community, group; Canada	Cerebral palsy; I: 5 C: -; unknown%; range 8–12 yrs; GMFCS 1-2	10 weeks	High
29	Lancioni, et al. (2004)	Motivation: feedback	Home centre; Europe	Profound disability range; 2 boys, 15.6 and 22.1 yrs; not reported	~7 months	High
30a	Shinohara, 2002	Cycling: peddle	Unclear; Japan	Cerebral palsy; I: 6, C: -; range 13-15 yrs; not reported, but could peddle a bike.	16.7 ± 4.7 weeks; -	High
30b	Shinohara, 2002	Cycling: arm	Unclear; Japan	Cerebral palsy; I: 5, C: -; unknown%; range 11-16 yrs; not reported, but could not peddle a bike.	12.7 ± 6.3 weeks; -	High

Table 4. Characteristics and quality assessment of included qualitative study.

Article #	Author (Year)	Methods	Intervention / Context	Participants	Findings	Study quality
31	Carter et al. (2014)	Participant observation, focus groups and interviews. Thematic analysis	Children's wheelchair sports club England	Total = sixty-three 37 children including wheelchair users (age not specified), 14 stakeholders, 10 parents, two older siblings	One unifying theme (realising potential) and four main themes: invisibility of disability; ambivalence and attraction of the chair; fun and fellowship; and thrills and skills. The Sports club created opportunities for meaningful participation in wheelchair sports for children with and without disabilities.	No major concerns about methodological limitations

Outcomes

Positive changes in health, fitness and wellbeing were reported across all outcome types. Wide variability existed in effectiveness of each intervention type. Many interventions were reported to be effective in some studies, but other studies found a similar intervention type to have no effect on the specific outcome. It was noted that there were no reports of changes favouring control, or a negative change in uncontrolled trials. Additionally, none of the included articles reported adverse outcomes during or following exercise intervention.

Fitness and exercise capacity was improved by walking (n=1/1 studies), cycling (n=1/3), functional exercises with body weight (n=1/2) and added weight (functional PRT) (n=1/1), sports and games (n=1/4) and complex programmes (n=2/2) (Figure 2). Quality of life was improved by walking (n=1/2), cycling (n=1/2), PRT (n=1/1) and functional exercises (n=2/2) but not functional PRT (n=0/1) or complex life-style programmes (n=0/2), or the addition of music for motivation (n=0/1) (Figure 3).

Gross motor function and mobility (Figures 4 and 5) were improved by programmes containing functional exercises (n=2/2), including walking (n=1/2) and Nintendo Wii (n=1/1), when using only the participant's own body weight. When additional weight was added to create a functional resistance training, no effect was seen (n=0/4 studies). The only exception to this was functional resistance training at 20-50% of 1-repetition maximum accompanied with rhythmically-matched music, which temporarily improved gross motor function but not mobility. Cycling preserved gross motor function in boys with duchenne muscular dystrophy but was not effective in other groups. Resistance training was not effective to improve gross motor function or mobility.

Most intervention types (n=8/15 studies), including functional resistance training, increased strength in some studies but not others (Figure 6). Walking did not to increase strength (n=1/1). The addition of music during PRT did not increase strength any more than PRT alone (n=1/1).

Changes in physical activity levels were variable in response to different interventions (Figure 7). What was interesting was that articles #4 and #21, which implemented the same complex intervention of counselling and individualised physical activities, found increased self-reported physical activity levels, but detected no change in objectively measured physical activity levels.

Three RCTs quantifying changes in body composition were identified, all were considered to present an unclear risk of bias. Article #2 delivered a sports and games based intervention to wheelchair users four-times per week for three months (#2a) and two-times per week for nine months (#2b); in both cases fat mass increased in the control group ($p < 0.05$) but did not change in the exercise group (p-values not reported). However, only in the two-times per week intervention was a significant interaction detected ($p < 0.05$). A follow-up was performed in the four-time per week arm and fat mass increased similarly in the experimental and control groups. Articles #12 and #22 used walking and functional open kinetic-chain exercises, respectively, with undefined wheelchair users and found no improvements post-intervention.

One study (#10) measured resting respiratory exchange ratio as an indicator of metabolic function. Following a complex programme of resistance training and walking, no change was found (p-value not reported). This study was considered to be at high risk of bias.

In an RCT with a sample of undefined wheelchair use, article #11 found spirometer training significantly improved multiple components of respiratory function.

When considering only the few studies that explicitly reported including wheelchair users (#1-10), a similar picture of high variability of intervention effectiveness generally emerges. Of note, all “wheelchair user” studies that measured gross motor function found improvements following cycling, Wii or complex programmes (Figure 4), but the only article to report improved quality of life was an uncontrolled before-after study using functional exercises (Figure 3).

Fifteen (~50%) of the quantitative articles (from 13 studies) included a follow-up period of some form, either complete intervention withdrawal or recommended self-directed activity. In all cases when quality of life was improved following intervention (#9,16a,b) the improvements were retained at follow-up (Figure 3). However, for other outcome measures any improvements were almost all lost by follow-up.

Children’s experiences

Only one study reported children’s views and experiences of the benefits of attending a UK-based sports club with typically developing children. This experience contrasts to those children recruited to quantitative intervention studies where all the samples consisted of only

disabled children. In addition to participating in sports, keeping fit and experiencing fun and thrills and wider social benefits, children learned to be more independent and became more aware of what they were capable of achieving. Some children also developed a more positive relationship with their wheelchair.

DISCUSSION

This systematic review included 31 articles from 28 studies measuring a wide range of quantitative health-, fitness- and quality of life-related outcomes, and qualitative experiences before and after exercise and physical activity interventions. These studies demonstrate that children who use wheelchairs are able to participate in various physical activity and exercise, and do so safely. Although the evidence is weak, it appears that these children can experience the same improvements in fitness, strength, mobility and quality of life as usually associated with appropriate exercise.

All intervention types were able to elicit some improvements in health, fitness and/or wellbeing. Conversely, most intervention types also failed to improve health, fitness and/or wellbeing in other studies. This reflects the highly inconsistent way that interventions are designed and delivered, and outcomes are assessed. Undoubtedly, delivering physical activity interventions to children with severe mobility impairments who use wheelchairs is challenging, but investigators must make efforts to follow established exercise prescription guidance to maximise the effectiveness of interventions. Failure to do so is particularly evident in the resistance training interventions, where several studies do not meet standard recommendations [56]. Indeed, in their introduction, Scholtes et al. (#20) also identified that many previous studies provided insufficient training loads or appropriate progression. Amongst the studies reviewed here, some used low weights (#10,23), a high number of lifting repetitions more akin to circuit training (#27), and achieved progression by increasing number of repetitions rather than load (#10,17).

The evidence included in this review indicate that work is required to define appropriate primary outcome measures that are specific and applicable to the intervention and population studied. This is complex due to the multi-faceted constructs of health and fitness and the heterogeneity of the samples tested, but failure to do so prevents robust evidence synthesis and identification of the most effective interventions. Best systematic review practise is to pre-

specify the outcomes of interest and only include evidence concerning these. GRADE can then be used to assess the overall quality of evidence within each outcome type. We were unable to apply GRADE in full due to the many different outcome measures used across studies. This, along with inconsistent and sometimes sub-optimal intervention design, reflects the relative infancy of this field.

It is clear that efforts to develop appropriately designed interventions and a standardised framework for evaluating interventions consistently across different subgroups of children are required. The great heterogeneity within the current evidence makes it challenging to make detailed recommendations for policy makers and practitioners at present. Nonetheless, the evidence gathered in the present review does allow us to reach some more specific conclusions and make recommendations beyond general conclusion that physical activity is safe and appears effective.

First, it appears possible to improve the gross motor function of children who use wheelchairs by several different types of physical activity. This may reflect the great capacity this subgroup of children have for improvement, and shows the potential they may be able to achieve. However, these effects did not appear to improve self-reported quality of life. It is hard to determine whether this is because the improvement in function is not large enough or the nature of the improvements are not transferable to improve overall quality of life, or whether the quality of life tools are not sensitive to the functional improvements.

Second, if the desired outcome is to improve mobility or gross motor function, functional closed-kinetic chain exercises against the child's own body weight are required. The lack of effectiveness of resistance training suggests that strength was not the limiting factor for most children who participated in the studies reviewed. This finding is in line with a previous systematic review on physical therapy interventions for children with cerebral palsy [14]. Performing functional exercise against normal body weight appears important for specificity of postural control strategies developed and transferability from training into actual functional tasks. However, the participants within the current evidence were mostly diagnosed with cerebral palsy and generally had good gross motor function and few severe cognitive impairments, therefore they do not reflect the whole range of children who use wheelchairs. Future work should evaluate the benefits of strengthening exercises for children with more severer or different causes of mobility impairment. In addition, improving strength may be

important for other activities required for independent living, e.g., self-transfer in and out of wheelchair

Third, the use of spirometer training, alongside other physical activity interventions or as part of regular therapy, may provide an effective and simple way to improve respiratory function. Given that supporting the child's posture so as not to restrict respiratory capacity is an important consideration in wheelchair fitting, this training mode may provide an effective way to increase seating options, as well as possibly health and fitness. Further work in this area is required to explore this possibility.

Forth, children experience and benefit from wider developmental and social impacts that are not currently assessed in most quantitative studies. More qualitative studies are needed to better understand how and why these impacts motivate some children to engage in exercise whereas others do not.

Most of the included articles (n=21, from 19 studies) consisted of group or supervised activities, and most of these were delivered or supervised by a therapist. There were few interventions based at the participants' home and fewer that integrated physical activity into daily routine and self-caring activities. This makes it difficult to evaluate programme theory (objective 2) for construct and delivery, although it is apparent that such interventions requiring intensive supervision are costly for health care services and likely not sustainable.

The study by Katz-Leurer, et al. (#17) developed a possibly relevant model of a home visit to deliver simple and acceptable exercises, with weekly telephone support to monitor participation, provide encouragement and ensure progression. This intervention was successful in improving a range of balance and mobility outcomes, which were consistent with other studies delivering similar exercises. Alternately, a model of short-term group-based intervention to initiate the programme followed by home-based self-directed participation may be possible, similar to that implemented by Fragala-Pinkham, et al. (#7). This study found that after 14 weeks of supervised group exercise measures of fitness, mobility and strength were improved, and many participants retained some of those improvements following 12-weeks of unsupervised, home-based participation. The Fragala-Pinkham, et al. study was considered to pose a high risk of bias, while the risk of bias in the Katz-Leurer, et al. study was unclear but with a sample with undefined wheelchair use.

A group of studies from the Netherlands (#3,4,6) with both wheelchair users and undefined wheelchair use (#21) evaluated a complex life-style intervention, combining counselling/motivational interviewing, physiotherapy and individualised physical fitness interventions. This model is instinctively appealing as it address individual barriers and facilitators to participation, as well as providing opportunities to participate in a variety of suitable physical activities. However, although some improvements were seen in gross motor function and the children's attitude to physical activity, they failed to improve objectively assessed physical activity levels, mobility, strength and most aspects of self-perceived quality of life. Moreover, the benefits that were gained were lost by follow up. This model of intervention delivery may still warrant further development, as it appears to encompass many important elements. However, much like the intensively supervised interventions, this may be resource intensive.

Given that the National Health Service, public health, primary care, local authorities, social services, schools and wheelchair services should deliver integrated care and support to enhance health, wellbeing and quality of life, there are opportunities for greater integration and more cost-effective delivery of effective intervention models. A previous review by our group [57] reported that wheelchair services generally met their key performance indicators by providing an appropriate wheelchair but did not commonly include lifestyle and keep-fit advice or mentoring to encourage healthy lifestyles and weight management. Utilising social care practitioners to deliver appropriate physical activities (such as exercise referrals dispensed by General Practitioners) that are integrated into the child's social activities, underpinned by an individualised aspirational programme theory and supported by schools to provide individually tailored keep fit activities should engender motivation and discipline from an early age.

Most studies did not report a programme theory as to how the interventions were intended to change behaviour by instilling motivation to keep fit in the short, medium or long term. Even the lifestyle interventions (#3,4,6,21) failed to elicit long-term changes in physical activity. Given that the benefits of physical activity interventions were lost once participation ceases, long-term solutions are required.

No health economics studies that met our inclusion criteria were found, this is an important omission as these children place high demands on services across the life course. Thus, we were not able to achieve our objective of evaluating cost-effectiveness (part of objective 1).

This evidence is a vital component in the development of effective and sustainable interventions and to provide evidence-based health promotion policy. In designing trials, exercise scientists need to reflect on the complex nature of delivering interventions in health services and incorporate economic and process evaluations, as well as measures of effectiveness.

The conclusions of this systematic review are limited primarily by the methodological quality of evidence, which was very low with concerns about high risk of bias. In many cases small sample size contributed to the high/unclear risk of bias, with few *a priori* or *post-hoc* power calculations. It is of course challenging to recruit disabled children (and families) for a long-term intervention, but future efforts must ensure larger and better justified sample sizes are achieved to ensure robustness. Quality of reporting was low and did not consistently follow CONSORT reporting guidelines [58] either because the trial predated CONSORT or if post-CONSORT authors did not adhere to the requirements. Overall, few estimates of precision such as confidence intervals were reported, thereby not meeting a key CONSORT reporting requirement. The quality of design and reporting was higher in more recent studies. Overall many studies lacked rigorous methodology and awareness of best-practice frameworks for designing research, as is available in Medical Research Council Guidance for designing complex public health interventions [59,60]. As already discussed, many studies did not pre-specify a single primary outcome and measured multiple outcomes. To allow comparison and to perform a meta-analysis, questions need to be consistently refined using a PICOS structure and outcomes need to be specified and measured as outlined in trial design principles.

Children who use wheelchairs constitute a complex heterogeneous group with multiple and varied diagnoses and wide ranges of fluctuating impaired mobility. Current evidence mostly includes relatively physically able children with cerebral palsy rather than more dependent wheelchair users, as indicated by how few studies included GMFCS 4 and 5, and the number of studies using walking capacity as an outcome. Few studies included children with learning impairments and no studies included children with severe learning impairments. Therefore, the applicability of the conclusions and recommendations of the present systematic review to the full heterogeneous population with a high level of children with significant learning impairments is limited.

Summary

Current evidence indicates that children with use wheelchairs can improve their health, fitness and wellbeing by participating in physical activity interventions, without adverse outcomes. However, the evidence base on the most effective intervention types is weak and lacks the necessary range of theoretical underpinnings. The samples included were typically small in number and with relatively less severe mobility impairments. Consequently, it is not surprising that government policy and practice guidance is lacking and, as a consequence, insufficient attention has been paid to the fitness of children who use wheelchairs who are known to be less fit than typically developing children. Future investment and effort is required to conduct high quality trials of promising intervention types and models of delivery to establish a robust evidence base to support government and health policy to improve the fitness and wellbeing of children who use wheelchairs.

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Competing Interests statement

The authors have no competing interests to disclose

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Contributorship

The review was conceived by all authors and the protocol developed by JN, TOB with LHS; searches were undertaken by LHS; article screening by LHS, TOB, HK and RH, with mediation by JN; Risk of bias assessment was undertaken by TOB and JN; Data were interpreted by all authors; the manuscript was drafted by TOB and critically reviewed by all authors.

REFERENCES

1. World Health Organisation, Global recommendations on physical activity for health; 5–17 years old. Accessed from <http://www.who.int/dietphysicalactivity/pa/en/> 2011; accessed August 2015
2. Tsan-Hon L, F-Xavier P, Blandine L. Physical disability and obesity. *Nutrition Reviews* 2005; 63: 321–360.
3. Maher CA, Williams MT, Olds T, Lane AE. Physical and sedentary activity in adolescents with cerebral palsy. *Developmental Medicine and Child Neurology* 2007; 49: 450–457.
4. Buffart LM, Roebroek ME, Rol M, Stam HJ, van den Berg-Emons RJG. Triad of physical activity, aerobic fitness and obesity in adolescents and young adults with myelomeningocele. *Journal of Rehabilitation Medicine* 2008; 40: 70–75.
5. Winnick J, Short F. The physical fitness of youngsters with spinal neuromuscular conditions. *Adapted Physical Activity Quarterly* 1984;1: 37–51.
6. Hoffman MD. Cardiorespiratory fitness and training in quadriplegics and paraplegics. *Sports Medicine* 1986; 3: 312–330.
7. Papworth Trust. Disability in the United Kingdom 2011: facts and figures. Papworth Trust, Cambridge.
http://www.papworth.org.uk/downloads/factsandfigures_disabilityintheuk_july2011_110_110721132605.pdf 2012; accessed September 2012.
8. National Institute for Health and Clinical Excellence (NICE). Public health guidance 17. Promoting physical activity for children and young people.
<http://guidance.nice.org.uk/PH17/Guidance/pdf/English> 2009; accessed October 2012.
9. American Academy of Pediatrics, Committee on Nutrition. Policy statement on prevention of pediatric overweight and obesity. *Pediatrics* 2003; 112: 424.
10. Scott R. The health benefits of play and physical activity for disabled children and young people: Kids briefing paper.
http://www.sensoryintegration.org.uk/Websites/sinetwork/images/Other_images/Health_PlayPhysicalActivitysmall2.pdf 2010; accessed December 2012.
11. Novak I, Cusick A. Home programmes in paediatric occupational therapy for children with cerebral palsy: where to start? *Australian Occupational Therapy Journal* 2006; 53: 251–264.
12. Hoare B, Imms C, Carey L, Wasiak J. Constraint induced movement therapy in the treatment of the upper limb in children with hemiplegic cerebral palsy: a Cochrane systematic review. *Clinical Rehabilitation* 2007; 21: 675–685.
13. Snider L, Korner-Bitensky N, Kammann C, *et al.* Horseback riding as therapy for children with cerebral palsy: is there evidence of its effectiveness? *Physical and Occupational Therapy in Pediatrics* 2007; 27: 5–23.

14. Anttila H, Autti-Ramo I, Suoranta J, *et al.* Effectiveness of physical therapy interventions for children with cerebral palsy: a systematic review. *BMC Paediatrics* 2008; 8: 14.
15. Mockford M, Caulton JM. Systematic review of progressive, strength training in children and adolescents with cerebral palsy who are ambulatory. *Pediatric Physical Therapy* 2008; 20: 318–333.
16. Scianni A, Butler JM, Ada L, Teixeira-Salmela LF. Muscle strengthening is not effective in children and adolescents with cerebral palsy: a systematic review. *Australian Journal of Physiotherapy* 2009; 55: 81–87.
17. O'Brien TD, Noyes J, Spencer LH, *et al.* 'Keep fit' interventions to improve health and fitness of children and young people who use wheelchairs: mixed-method systematic review protocol. *Journal of Advanced Nursing* 2014; 70: 2942–2951.
18. Cochrane Effective Practice and Organisation of Care (EPOC) Review Group. The Data Collection Checklist. <http://epoc.cochrane.org/sites/epoc.cochrane.org/files/uploads/datacollectionchecklist.pdf> 2002, accessed January 2014.
19. Moher D, Liberati A, Tetzlaff J, *et al.* Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Medicine* 2009; 6: e1000097
20. O'Brien TD, Noyes J, Spencer LH, Kubis HP, Edwards RT, Bray N & Whitaker R. Wellbeing, health and fitness of children who use wheelchairs: Feasibility study protocol to develop child-centred 'keep-fit' exercise interventions. *J Adv Nurs* 2015; 71: 430-40.
21. Booth A, Brice A. Evidence based practice for information professionals: a handbook. London, Facet Publishing 2003.
22. Effective Practice and Organisation of Care (EPOC). Suggested risk of bias criteria for EPOC reviews. EPOC Resources for review authors; Available at: <http://epoc.cochrane.org/epoc-specific-resources-review-authors> 2012; accessed 2014.
23. Lewin S, Glenton C, Munthe-Kaas H, *et al.* Using Qualitative Evidence in Decision Making for Health and Social Interventions: An Approach to Assess Confidence in Findings from Qualitative Evidence Syntheses (GRADE-CERQual). *PLoS Med* 2015; 12: e1001895.
24. Ogilvie D, Fayter D, Petticrew M, *et al.* The harvest plot: A method for synthesising evidence about the differential effects of interventions. *BMC Medical Research Methodology* 2008; 8: 8

25. Jansen M, van Alfen N, Geurts ACH, & de Groot IJM. Assisted bicycle training delays functional deterioration in boys with duchenne muscular dystrophy: The randomized controlled trial “No use is disuse”. *Neurorehabilitation and Neural Repair* 2013; 27: 816-827.
26. van den Berg-Emons RJ, van Baak MA, Speth L, Saris WH. Physical training of school children with spastic cerebral palsy: Effects on daily activity, fat mass and fitness. *International Journal of Rehabilitation Research* 1998; 21: 179-194.
27. van Wely L, Balemans ACJ, Becher JG, & Dallmeijer AJ. The effectiveness of a physical activity stimulation programme for children with cerebral palsy on social participation, self-perception and quality of life: A randomized controlled trial. *Clinical Rehabilitation* 2014a; 28: 972-982.
28. van Wely L, Balemans ACJ, Becher JG, & Dallmeijer AJ. Physical activity stimulation program for children with cerebral palsy did not improve physical activity: A randomised trial. *Journal of Physiotherapy* 2014b; 60: 40-49.
29. Andrade C, Kramer J, Garber M, Longmuir P. Changes in self-concept, cardiovascular endurance and muscular strength of children with spina bifida aged 8 to 13 years in response to a 10 week physical-activity programme: A pilot study. *Child Care Health and Development* 1991; 17: 183-196.
30. Buffart LM, van den Berg-Emons RJG, van Mechelen W, et al. Promoting physical activity in an adolescent and a young adult with physical disabilities. *Disability and Health Journal* 2010; 3: 86-92.
31. Fragala-Pinkham M, Haley SM, Rabin J, Kharasch VS. A fitness program for children with disabilities. *Physical Therapy* 2005; 85: 1182-1200.
32. Gordon C, Roopchand-Martin S, & Gregg A. Potential of the Nintendo Wii™ as a rehabilitation tool for children with cerebral palsy in a developing country: A pilot study. *Physiotherapy* 2012; 98: 238-242.
33. Odman P, Oberg B. Effectiveness of intensive training for children with cerebral palsy - a comparison between child and youth rehabilitation and conductive education. *Journal of Rehabilitation Medicine* 2005; 37: 265-270.
34. Unnithan VB, Katsimanis G, Evangelinou C, et al. Effect of strength and aerobic training in children with cerebral palsy. *Medicine & Science in Sport & Exercise* 2007; 39: 1902-09.

35. Choi JY, Rha D, & Park ES. Change in pulmonary function after incentive spirometer exercise in children with spastic cerebral palsy: A randomized controlled study. *Yonsei Medical Journal* 2016; 57: 769-775.
36. de Groot JF, Takken T, van Brussel M, et al. Randomized controlled study of home-based treadmill training for ambulatory children with spina bifida. *Neurorehabilitation and Neural Repair* 2011; 25: 597-606.
37. Demuth SK, Knutson LM, Fowler EG. The PEDALS stationary cycling intervention and health-related quality of life in children with cerebral palsy: A randomized controlled trial. *Developmental Medicine and Child Neurology* 2012; 54: 654-661.
38. Dodd KJ, Taylor NF, Graham HK. A randomized clinical trial of strength training in young people with cerebral palsy. *Developmental Medicine & Child Neurology* 2003; 45: 652-657.
39. Fowler EG, Knutson LM, de Muth SK, et al. Pediatric endurance and limb strengthening (PEDALS) for children with cerebral palsy using stationary cycling: A randomized controlled trial. *Physical Therapy* 2010; 90: 367-381.
40. Gates PE, Banks D, Johnston TE, et al. Randomized controlled trial assessing participation and quality of life in a supported speed treadmill training exercise program versus a strengthening program for children with cerebral palsy. *Journal of Pediatric Rehabilitation Medicine* 2012; 5: 75-88.
41. Katz-Leurer M, Rotem H, Keren O, Meyer S. The effects of a 'home-based' task-oriented exercise programme on motor and balance performance in children with spastic cerebral palsy and severe traumatic brain injury. *Clinical Rehabilitation* 2009; 23: 714-724.
42. Maher CA, Williams MT, Olds T, Lane AE. An internet-based physical activity intervention for adolescents with cerebral palsy: A randomized controlled trial. *Developmental Medicine & Child Neurology* 2010; 52: 448-455.
43. Scholtes VA, Becher JG, Comuth A, et al. Effectiveness of functional progressive resistance exercise strength training on muscle strength and mobility in children with cerebral palsy: A randomized controlled trial. *Developmental Medicine & Child Neurology* 2010; 52: e107-13.
44. Scholtes VA, Becher JG, Janssen-Potten YJ, et al. Effectiveness of functional progressive resistance exercise training on walking ability in children with cerebral palsy: A randomized controlled trial. *Research in Developmental Disabilities* 2012; 33: 181-188.

45. Slaman J, Roebroek M, Dallmijer A, et al. Can a lifestyle intervention programme improve physical behaviour among adolescents and young adults with spastic cerebral palsy? A randomized controlled trial [with consumer summary]. *Developmental Medicine and Child Neurology* 2015; 57: 159-166.
46. Verschuren O, Ketelaar M, Gorter JW, et al. Exercise training program in children and adolescents with cerebral palsy. *Archives of Pediatrics & Adolescent Medicine* 2007; 161: 1075-1081.
47. Wang T, Peng Y, Chen Y, et al. A home-based program using patterned sensory enhancement improves resistance exercise effects for children with cerebral palsy: A randomized controlled trial. *Neurorehabilitation and Neural Repair* 2013; 27: 684-694.
48. Blundell SW, Sheperd RB, Dean CM, et al. Functional strength training in cerebral palsy: A pilot study of a group circuit training class for children aged 4-8 years. *Clinical Rehabilitation* 2003; 17: 48-57.
49. Damiano DL, Vaughan CL, Abel MF. Muscle response to heavy resistance exercise in children with spastic cerebral palsy. *Developmental Medicine & Child Neurology* 1995; 37: 731-739.
50. Chen C, Hong W, Cheng H, et al. Muscle strength enhancement following home-based virtual cycling training in ambulatory children with cerebral palsy. *Research in Developmental Disabilities* 2012; 33: 1087-1094.
51. Crompton J, Imms C, McCoy AT, et al. Group-based task-related training for children with cerebral palsy: A pilot study. *Physical & Occupational Therapy in Pediatrics* 2007; 27: 43-65.
52. Kelly M, Legg D. On-land community-based aerobic and strength training program for children with cerebral palsy. *European Journal of Adapted Physical Activity* 2009; 2: 7-20.
53. Lancioni GE, Singh NN, O'Reilly MF, et al. Use of simple exercise tools by students with multiple disabilities: Impact of automatically delivered stimulation on activity level and mood. *Journal of Developmental and Physical Disabilities* 2004; 16: 171-178.
54. Shinohara TA, Suzuki N, Oba M, et al. Effect of exercise at the AT point for children with cerebral palsy. *Bulletin of the NYU Hospital for Joint Diseases* 2002; 61: 63-67.

55. Carter B, Grey J, McWilliams E, Clair Z, Blake K, & Byatt R. 'Just kids playing sport (in a chair)': Experiences of children, families and stakeholders attending a wheelchair sports club. *Disability & Society* 2014; 29: 938-952.
56. Faigenbaum AD, Kraemer WJ, Blimkie CJ, *et al.* Youth resistance training: updated position statement paper from the National Strength and Conditioning Association. *Journal of Strength and Conditioning Research* 2009; 23: S60–S79.
57. Bray N, Noyes J, Edwards RT, Harris N. Wheelchair interventions, services and provision for disabled children: a mixed-method systematic review and conceptual framework. *BMC Health Serv Res* 2014; 14: 309
58. CONSolidated Standards of Reporting Trials (CONSORT). <http://www.consort-statement.org/consort-2010> 2010; accessed January 2014.
59. Medical Research Council. A framework for the development and evaluation of RCTs for complex interventions to improve health. <http://www.mrc.ac.uk/Utilities/Documentrecord/index.htm?d=MRC003372> 2000; accessed December 2012.
60. Medical Research Council. Developing and evaluating complex interventions: new guidance. <http://www.mrc.ac.uk/Utilities/Documentrecord/index.htm?d=MRC004871> 2008; accessed November 2012.

FIGURE LEGENDS

Figure 1. Flow of article screening and inclusion in this review.

Figure 2. Harvest plot of intervention effects on fitness and exercise capacity. Each box represents the findings of a single article, and contains the article number, which is underlined for studies with undefined wheelchair use, and the risk of bias rating (High, Unclear or Low). Article 2 measured aerobic (Aero) and anaerobic (Ana) fitness, these are distinguished below article number.

Figure 3. Harvest plot of intervention effects on quality of life and attitudes. Each box represents the findings of a single article, and contains the article number, which is underlined for studies with undefined wheelchair use, and the risk of bias rating (High, Unclear or Low).

Figure 4. Harvest plot of intervention effects on gross motor function. Each box represents the findings of a single article, and contains the article number, which is underlined for studies with undefined wheelchair use, and the risk of bias rating (High, Unclear or Low).

Figure 5. Harvest plot of intervention effects on ambulation and mobility. Each box represents the findings of a single article, and contains the article number, which is underlined for studies with undefined wheelchair use, and the risk of bias rating (High, Unclear or Low).

Figure 6. Harvest plot of intervention effects on muscle strength. Each box represents the findings of a single article, and contains the article number, which is underlined for studies with undefined wheelchair use, and the risk of bias rating (High, Unclear or Low).

Figure 7. Harvest plot of intervention effects on muscle physical activity levels. Each box represents the findings of a single article, and contains the article number, which is underlined for studies with undefined wheelchair use, and the risk of bias rating (High, Unclear or Low). Articles 4 and 21 measured physical activity objectively (Obj) and self-reported (Self) by parents and children, these are distinguished below article number.