Powered mobility as an intervention for children with cerebral palsy: A systematic review; The smart wheelchair: Is it an appropriate mobility training tool for children with cerebral palsy?

Sarah McGarry

Edith Cowan University

Recommended Citation


This Thesis is posted at Research Online.
https://ro.ecu.edu.au/theses_hons/1212
Edith Cowan University

Copyright Warning

You may print or download ONE copy of this document for the purpose of your own research or study.

The University does not authorize you to copy, communicate or otherwise make available electronically to any other person any copyright material contained on this site.

You are reminded of the following:

- Copyright owners are entitled to take legal action against persons who infringe their copyright.

- A reproduction of material that is protected by copyright may be a copyright infringement. Where the reproduction of such material is done without attribution of authorship, with false attribution of authorship or the authorship is treated in a derogatory manner, this may be a breach of the author’s moral rights contained in Part IX of the Copyright Act 1968 (Cth).

- Courts have the power to impose a wide range of civil and criminal sanctions for infringement of copyright, infringement of moral rights and other offences under the Copyright Act 1968 (Cth). Higher penalties may apply, and higher damages may be awarded, for offences and infringements involving the conversion of material into digital or electronic form.
Powered mobility as an intervention for children with cerebral palsy: A systematic review

AND

The Smart Wheelchair: Is it an appropriate mobility training tool for children with cerebral palsy?

Sarah McGarry

A report submitted in Partial Fulfilment of the Requirements for the Award of Bachelor of Science (Occupational Therapy)(Honours), Faculty of Computing, Health and Science, Edith Cowan University.

Submitted (September, 2009)

"I declare that this written assignment is my own work and does not include:

(i) Material form published sources used without proper acknowledgement; or
(ii) Material copied from the work of other students”.

Signed

Dated 18.12.2009
USE OF THESIS

The Use of Thesis statement is not included in this version of the thesis.
COPYRIGHT AND ACCESS DECLARATION

I certify that this thesis does not, to the best of my knowledge and belief:

(i) Incorporate without acknowledgment any material previously submitted for a degree or diploma in any institution of higher degree or diploma in any institution of higher education;

(ii) Contain any material previously published or written by another person except where due reference is made in the text of this thesis; or

(iii) Contain any defamatory material.

(iv) Contain any data that has not been collected in a manner consistent with ethics approval.

The Ethics Committee may refer any incidents involving requests for ethics approval after data collection to the relevant Faculty for action.

Signed.. 

Date.....18.12.2009..................
Powered mobility as an intervention for children with cerebral palsy: A systematic review

AND

The Smart Wheelchair: Is it an appropriate mobility training tool for children with cerebral palsy?

Sarah McGarry

A report submitted in Partial Fulfilment of the Requirements for the Award of Bachelor of Science (Occupational Therapy)(Honours), Faculty of Computing, Health and Science, Edith Cowan University.

Submitted (September, 2009)

"I declare that this written assignment is my own work and does not include:

(i) Material form published sources used without proper acknowledgement; or

(ii) Material copied from the work of other students”.

Signed

Dated 18.12.2009
Declaration

I certify that this literature review and research project does not incorporate, without my acknowledgement, any material previously submitted for a degree or diploma in any institution of higher education and that, to the best of my knowledge and belief, it does not contain any material previously published or written by another person except where due reference is made in the text.

Signed

Dated 18.12.2009
ACKNOWLEDGEMENTS

I would like to express my sincere gratitude to all those who supported me in the completion of this study.

Firstly, to the children and parents who participated in this study. Thank you for volunteering your time and sharing this experience with me. Sincere thanks to The Centre for Cerebral Palsy for their support and for making this research possible.

I would like to especially thank my supervisors Dr Sonya Girdler and Lois Moir for their constant support, guidance and encouragement. I really appreciate all the time and effort you both spent on this project and, I am thankful to have the experience of working with two inspiring and motivational occupational therapists.

Finally, thank you to my family for their constant support.
Table of Contents

LITERATURE REVIEW

1. Abstract ................................................................. 2
2. Introduction ............................................................. 3
3. Methods ................................................................. 4
4. Results ................................................................. 6
5. Discussion ............................................................. 9
6. References ........................................................... 13
7. Guidelines for contributions by authors ......................... 17

RESEARCH REPORT

8. Abstract ............................................................... 28
9. Introduction ........................................................... 29
10. Materials and methods ............................................. 31
11. Findings ............................................................... 34
12. Discussion ........................................................... 41
13. Conclusion .......................................................... 44
14. Graphs ............................................................... 46
15. References ......................................................................................47

APPENDICES

Appendix A: Intervention protocol ..............................................................52
Appendix B: Driving Skills Assessment .......................................................55
Appendix C: Semi structured interview guide ................................................56
Appendix D: Description of included studies .................................................57
Guidelines for contributions by authors .......................................................60
Powered mobility as an intervention for children with cerebral palsy: A systematic review

Sarah McGarry
Powered mobility as an intervention for children with cerebral palsy: A systematic review

Abstract

Purpose. To systematically review the effectiveness of powered mobility training interventions in improving driving skills and psychosocial outcomes in children with cerebral palsy.

Method. Electronic searches of CINAHL, Medline and Meditext were conducted. Using two assessment tools, two independent reviewers assessed the quality of selected studies and level of evidence. Studies were included if a powered mobility training intervention was described, at least one participant had cerebral palsy and if the majority of participants were aged between 18 months and 18 years old. A narrative analysis was conducted.

Results. Seven articles were eligible for inclusion. Intervention protocols included computer simulator training, mobility training on a powered riding toy, and a powered wheelchair. The quality of the studies ranged from strong to limited, with six out of seven studies rated as level IV evidence and one study rated as level III evidence as defined by the National Health and Medical Research Council level of evidence table.

Conclusions. Although research is limited observational investigations suggest that powered mobility training programs are potentially beneficial in the development of driving skills for children with cerebral palsy and may have a positive impact on psychosocial outcomes. Methodological weaknesses of the studies were small sample sizes, absence of control groups and largely descriptive data analysis. There is a need for future, more rigorous research which addresses these weaknesses and contributes to an understanding of the utility of powered mobility both as a method of access and as a therapeutic intervention tool.

Key Words: powered wheelchairs, electric wheelchairs, mobility interventions, children with disabilities

Sarah McGarry,
Dr Sonya Girdler,
Powered mobility as an intervention for children with cerebral palsy: A systematic review

Introduction

Cerebral palsy (CP) is the most common physical childhood disability [1-3], accounting for 600 to 700 Australian births each year [1]. Improvements and advances in perinatal care are associated with declining rates of infant mortality among preterm infants and increasing numbers of infants born with CP [4-5]. It is estimated that 33,797 Australians have CP and this number is expected to increase to 47,601 by 2050 [1]. In 2007 the financial cost of CP in Australia was estimated to be $1.47 billion [1]. This value increases to $2.4 billion when considering the value of lost wellbeing, due to disability and premature death [1]. The per annum financial cost of CP is $43,431 per person. Including the value of lost wellbeing the cost increases to $115,000 per annum, per person [1]. With increasing numbers of children born with CP, there is a need to understand effective therapeutic interventions as demands on services will increase.

Cerebral palsy is a neurological condition referring to a group of permanent disorders affecting the development of movement and posture [6]. In addition many individuals experience limitations in cognition, communication, sensation and perception [6]. The severity of CP can vary, with motor and cognitive impairments ranging from minimal to profound [7,8]. The functional limitations that result from CP mean that children are dependent on their carers for many aspects of activities of daily living, instrumental activities of daily living and mobility [2,7,8]. It is estimated that 35 percent of the population with CP are also severely cognitively impaired, leading to reduced ability to care for themselves and others [9]. Restrictions in social interactions increase progressively with impairment type and severity level of CP [10,11]. These functional limitations may impact participation in meaningful activities and in the fulfilment of social roles.

Developmental theorists propose that early experiences of movement impact on physical and psychological development [12,13]. Piaget [14] believed independent movement to be an important
building block for the development of knowledge. Locomotion encourages a child’s understanding of space, objects, self awareness, and principles of cause and effect [14,15]. In addition, independent movement allows a child to gain a sense of independence and competence, as well as providing opportunities to learn and socialise [15,16]. Supporting this stance, research has demonstrated that for children, gains in mobility are linked with improvements in cognitive, social-emotional, perceptual and motor functioning [13,17,18].

Many children with CP are unable to independently mobilise due to cognitive and motor impairments [8,19]. These children rely on caregivers to move them from place to place, and consequently miss many crucial learning opportunities. This impacts negatively on their skill development in the areas of cognition, behaviour, emotion and communication [20-22]. Secondary developmental problems can result from a child’s increasing frustration at their inability to control the world around them [23]. This often results in a cycle of deprivation, and reduced motivation [24].

Powered wheelchairs can provide children with CP the opportunity to explore their environments through play and locomotion [19] and decrease their dependency on others [19,25]. Despite the importance of independent mobility in child development and the wide use of powered wheelchairs in clinical settings for children with disabilities, little attention has been given to the evidence base underpinning powered mobility interventions. The purpose of this systematic literature review was to examine the effectiveness of powered mobility training interventions in improving driving skills and psychosocial outcomes in young children with CP.

Methods
Principles and techniques of systematic reviews were used to review the literature and locate relevant studies for inclusion. Electronic searches of CINAHL, Medline and Meditext were conducted. Each database was searched from its earliest records to most recent (CINAHL 1937-2009, Medline 1948-2009,
The main search terms included a combination of both participant (children, young children and cerebral palsy) and intervention terms (powered mobility, powered wheelchairs, powered mobility training and mobility aids). All search terms were truncated, exploded and adjusted to match the identified databases with assistance from a librarian. Reference lists of all retrieved articles were manually searched for relevant articles. The literature search was limited to peer reviewed (English language) published data only, and excluded grey literature and expert opinion.

A priori criteria for inclusion of studies were applied to identify abstracts and then full text articles. Studies were included if a powered mobility training intervention was described. The mode, length and frequency of the interventions were not limited. Due to the paucity of studies describing powered mobility interventions with children with CP, studies were included if at least one participant had CP. Studies were limited to those with the majority of participants aged 18 months - 18 years old. Outcomes of interest included driving skills and psychosocial outcomes related to powered mobility, for example improvements in confidence and independence.

Assessment of methodological quality
Using an assessment tool developed by Kmet, Lee and Cook [26] two independent reviewers assessed the quality of the selected studies. The assessment consists of 14 questions and a scoring system for methodological quality. Using this system each study was categorised as strong (score of >80%), good (70-80%), adequate (50-70%) or limited (<50%) [27].

Data extraction
Two reviewers used methods set out in Section 7 of the Cochrane Handbook of Systematic Reviews of Interventions [28] to extract relevant data from the studies. Data extracted included: study objective(s), study design, method of participant selection, characteristics of the participants, random allocation, masking of the investigators, masking of the participants, outcomes, sample size, analytical methods,
estimate of variance, confounding controls, results and conclusions. Once data was extracted the level of
evvidence was determined based on the National Health and Medical Research Council guidelines
(NHMRC) [29]. Discrepancies between reviewers were resolved through discussion.

Data synthesis and analysis

Due to the use of non traditional analysis of methods such as parent interviews and video footage, and
small sample sizes in the articles, meta-analysis was not possible. A narrative review was conducted to
summarise findings and provide an assessment of methodological issues.

Results

Electronic searches located 21 articles from CINAHL, five from Meditext and 22 from Medline. A total of
39 potential articles were located after duplicates were accounted for. Using the stated inclusion criteria,
abstracts and full texts were reviewed, resulting in seven articles judged as eligible for inclusion. Excluded
papers included expert opinion, descriptions of the use of powered wheelchairs with adults and children,
discussions on types of wheelchairs and assessment practices, and descriptive data about the perspectives
of parents and providers.

Articles meeting the selection criteria ranged in publication date from 1983 to 2003. Of the seven articles
selected two were written by the same authors, and one author is responsible for three papers. These three
papers were published in 1983, 1984 and 1986, an average of 25 years ago. One article was published in
1998, and the remaining three articles were published in 2001, 2002 and 2003, a minimum of six years
ago. All studies investigated a powered mobility training intervention with a total of 83 participants.
Intervention protocols of these articles included computer simulator training to improve powered
wheelchair driving, mobility training on a powered riding toy and mobility training on a powered
wheelchair.
Quality assessment of studies

Assessment of methodological quality of the research ranged from strong to limited (see appendix D). A study investigating powered wheelchair training in the home was rated as methodologically limited (27%) [30], four separate studies reporting on a mobility toy, the use of a computer simulator to increase driving skills and powered mobility training in the home, were rated as adequate (50-68%) [31-34], and two studies examining the concept of driving to learn and powered wheelchairs for independence were rated strong [35-36]. Overall, methodological weaknesses of the studies were small sample sizes, absence of control groups, lack of blinding, and largely descriptive data analysis. In clinical research that requires the active involvement of participants these weaknesses are common and can be difficult to overcome. Authors did not report effect size or describe controls used for confounding, and three authors did not report estimates of variance for the main results. Apart from one study, psychometric properties of the outcome measures used were not described. The NHMRC [29] level of evidence table was used to rank studies according to four levels. Level I is the strongest and most rigorous, accounting for systematic reviews of all relevant randomised control trials. Level IV is the least rigorous, consisting of evidence obtained from case series, either post test, or pre test and post test [29]. Six out of seven studies included in this review were rated as level IV evidence and one study was rated as level III evidence, due to the inclusion of a comparison group (see appendix D).

Interventions

The reviewed interventions were powered mobility training programmes for children with CP and other disabilities. Three studies assessed the participant’s ability to learn to drive a powered wheelchair at home under parent supervision [30-32]. Another study evaluated power wheelchair training in a clinic setting and at home [36]. In a cohort design study using a computer simulator, children navigated a game of mazes with a joystick [34]. Lastly, a powered mobility riding toy was used as an intervention in a single subject withdrawal design study of two children aged five years old [33]. One study failed to provide
details about the powered mobility program [35] and no studies described explicitly the strategies used to promote intervention fidelity.

Several studies attempted to employ control strategies to provide a baseline of comparison, to support the interpretation of the effect of the intervention [37]. The use of a comparison group was evident in only one study [34]. This study investigated whether training on a computer simulator would increase the driving skills of children who had never used a powered wheelchair in comparison to children who were experienced drivers. A single subject design with multiple baselines was described as successful in using the participants as their own control [31]. Whereas a study using a case history survey design argued that the use of a control group would be unethical as they believed it would be detrimental to the participants to delay provision of a powered wheelchair [35].

Outcomes

In five of the seven studies powered wheelchair driving skills were measured as the primary dependant variable [30,32,34,-36]. One study described frequency of self initiated physical interaction with objects, communications with care giver and changes in location in space as dependent variables [31]. Similarly, Diatz and colleagues [33] reported child initiated movement, child initiated contact with others, and affect as dependent variables. Psychosocial outcomes were measured in five of the seven studies; outcomes included social, emotional and intellectual behaviours, level of independence, and development of initiative and exploratory behaviours [30,31,33,35,36]. Video footage, a seven skill driving checklist, Powered Mobility Program assessment battery [38], and a computer game were measures used to assess participant's driving skills. Five out of the seven studies utilised parent interviews to enrich their quantitative findings [30,32,33,35,36]. To investigate the effect of early provision of powered wheelchairs Bottos and colleagues [35] employed standardised measures, including Impact of Childhood Illness Scale, Gross Motor Functional Measure, the Canadian Occupational Performance Measure, Performance IQ
score of the Leiter International Performance Scale, and Verbal IQ score of the Peabody Developmental Verbal Scale.

Collectively the results of the five studies measuring the driving skills of participants suggested that children and young adults aged from 23 months to 22 years old with physical and cognitive impairments can learn to drive a powered wheelchair independently, with a minimum training time of 16 days and within a maximum training time of one year [30,32,34-36]. Five studies evaluating psychosocial outcomes found independent powered mobility promoted increased social interaction, increased motivation to move and interact with the environment, and increased independence [30,31,33,35,36]. Independent powered mobility did not affect motor impairment, Intelligence Quotient or quality of life [35]. However, powered mobility was linked to an increase in exploratory behaviours, initiative, and ability to react to external stimuli [36]. Qualitative findings supported the conclusions that powered mobility stimulated the child’s understanding of cause and effect relationships, encouraged limited use of participant’s arms and hands, as well as stimulated the child’s level of alertness [36].

Three studies reported statistically significant changes following the powered mobility intervention. Participant’s driving skills improved significantly (p<0.01) in two studies [34,35]. An increase in self initiated change in location (p<0.05) was found to be associated with independent driving of a powered wheelchair [31], as was changes in occupational performance, measured by the Canadian Occupational Performance Model (p<0.00001) [34,35].

Discussion

Overall this systematic review found a paucity of evidence to support powered mobility as an intervention for children with CP. Findings from level III and IV evidence suggest that young children with physical and cognitive disabilities can learn to drive a powered wheelchair and a powered riding toy independently. In addition, improvements in psychosocial outcomes were found to be associated with independent
powered mobility. However, the reviewed research has many methodological limitations, being based on small sample sizes and largely descriptive data analysis. Many of the studies excluded from the review were published by clinicians and consisted of ‘expert opinion’. Although papers in this category are valid contributions to literature in this area, they are viewed as the least rigorous in terms of evidence and are susceptible to bias [39,29]. Evidence gained from this literature review was limited as the majority of the studies were of adequate quality and were ranked level IV evidence, which can only allude to a causal relationship between the independent and dependent variables [29].

The studies reviewed varied considerably in intervention, study design, and outcome measures making it difficult to draw collective conclusions from the results. In depth descriptions of intervention protocols were absent and intervention fidelity was not accounted for in the studies. However, a number of studies used both qualitative and quantitative methods, an approach which has been described as having merit in this field of research [39]. Four studies used a single subject design, limiting the inference of outcomes to groups or situations other than those directly involved in the study [37]. Further, the reliability and validity of the findings are questionable given that authors did not discuss the psychometric properties of the outcome measures and only one study used standardised assessments [35]. When evaluating the impact of powered mobility devices for children with CP there is clearly a need for higher quality, more rigorous research involving larger sample groups, a comparison group, and standardised measures to produce adequate evidence for therapists and families.

Over the past three decades, the development of new assistive technologies for individuals with disabilities has increased [40,41]. Using the latest technologies and robotics, new models of wheelchairs such as the CALL Centre Smart Wheelchair and the Collaborative Wheelchair Assistant have been developed [42,43]. This systematic review found that there is a very scarce evidence base for the use of technologically advanced models of powered wheelchairs with children with CP. Of the seven articles reviewed three were at least 25 years old, with the most recent study in the review being six years old.
This suggests that this evidence is from powered wheelchairs that are since outdated and possibly no longer prescribed to children. There is a clear need for current research using the latest most advanced models of powered wheelchairs which may be suitable to a wider population of children with various disabilities.

The notion of ‘driving to learn’ was discussed in a study evaluating powered mobility for children with profound cognitive disabilities [36]. This method of training is based on the notion that mobility can be used as a therapeutic tool, rather than purely a means of gaining access between one place and another. It is proposed that the sensory stimulation and change of body positions resulting from powered mobility can raise the individual’s alertness, enhancing their ability to understand simple causal relationships [36]. This method of training may be useful for uncovering skills and abilities previously unrecognised in children with severe and multiple disabilities. Further the full mobility potential of children who may have previously been excluded from powered wheelchair training may be revealed. Given the potential of driving to learn for many children and their families, there is a great need to further investigate this concept.

Findings from this systematic review highlight the need for future research to evaluate the relationship between powered mobility and the components of the World Health Organization’s International Classification of Functioning, Disability and Health framework [44]. The development of targeted interventions to enhance independent powered mobility for children with CP will depend on a clearer understanding of the link between body structures and functions, activity, and their impact on participation [39]. In doing so the impact of contextual factors such as environment and personal factors on powered mobility interventions can be explored enabling an understanding of the length of training required, potential discontinuation of mobility devices, and a much broader perspective of the impact powered mobility has on an individual’s life [25].
Although a systematic search strategy was applied, the authors acknowledge that conclusions may be affected if relevant studies were unintentionally overlooked. Another limitation of this paper is that many studies in the field of powered mobility are not published in peer reviewed journals but appear in conference proceedings. The selection criteria excluded grey literature therefore these studies were not reviewed.

Although many experts in the field have advocated for early provision of powered wheelchairs for young children, the gap between research and practice continues. The existing evidence base for effective powered mobility interventions is limited however, the weight of the evidence suggests a positive relationship between powered mobility and child development. Limitations in the current body of knowledge outline directions for future research. Future studies should incorporate current clinical practices and examine intervention effectiveness not just for improvement of isolated mobility skills, but to gain a greater understanding of interventions that can contribute to the development of children with cerebral palsy.
References


Guidelines for contributions by authors

Disability and Rehabilitation: Assistive Technology is an international, multidisciplinary, peer-reviewed journal devoted specifically to the broad range of technological developments and related supports and issues which enhance the rehabilitation process. The journal seeks to encourage a better understanding of all aspects of disability, and to promote the rehabilitation process. New submissions on any aspect of technology, disability and rehabilitation are encouraged and the journal welcomes contributions from a wide range of professional groups, including medical practitioners, occupational therapists, physiotherapists, speech and language therapists, clinical psychologists and those involved in nursing, education, ergonomics, and engineering.

Disability and Rehabilitation: Assistive Technology is organised into sections: Literature Reviews; Research Papers, Case Studies, Clinical Commentaries; reports on Rehabilitation in Practice, Products and Devices/Rehabilitation Engineering, Occasional Special Issues and specific sections on contemporary themes of interest to the Journal's readership are published.

Disability and Rehabilitation: Assistive Technology will be of interest to a wide range of professional groups, including medical practitioners, occupational therapists, physiotherapists, speech and language therapists, clinical psychologists and those involved in nursing, education, ergonomics and engineering.

Editor

Marcia J. Scherer, PhD, MPH, FACRM
Institute for Matching Person & Technology
486 Lake Road
Webster, NY 14580 USA
585-671-3461 (phone/fax)
Email: IMPT97@aol.com

Submissions

All submissions should be made online at Disability and Rehabilitation: Assistive Technology's Manuscript Central site. New users should first create an account. Once a user is logged onto the site submissions should be made via the Author Centre.

Papers should be submitted with any tables, figures, or photographs, all of which should be of high quality suitable for reproduction. Submissions should be in English in double line spacing.

The submission should include a separate title page with the name(s) and affiliation(s) of the author(s) and the name and address for offprint requests with a telephone, fax number (including country and area codes), and electronic mail address.

Submissions should be accompanied by a covering letter stating that all authors have read the paper and agree to its submission and should include where appropriate a formal statement that ethical consent for the work to be carried out has been given. Photographs of individuals should be avoided, but if essential individual consent in writing must accompany the manuscript. It is not sufficient to mask identity by covering the person's eyes.

Refereeing will be carried out anonymously unless otherwise requested, and manuscripts should be prepared accordingly.
There are no page charges for papers published in the journal. Three complimentary copies of the issue in which your article appears will be sent to the principal or sole author or articles. Larger quantities may be ordered at a special discount price. An order form will accompany the proof which must be completed and returned, irrespective of whether you require additional copies.

Writing a paper for Disability and Rehabilitation: Assistive Technology

For all manuscripts, non-discriminatory and person-first language is mandatory. Sexist or racist terms should not be used.

Structured abstracts of around 200 words are required for all papers submitted and should precede the text of a paper. There are no lower or upper word limits for papers submitted to the Journal.

Manuscripts should be typed double-spaced throughout, including the reference section.

Submitted manuscripts must cite relevant original sources, not only review articles or articles published subsequently by the same or different author(s).

In writing your paper, you are encouraged to review articles in the area you are addressing which have been previously published in the journal, and where you feel appropriate, to reference them. This will enhance context, coherence, and continuity for our readers.

Structure of Paper

An introductory section should state the purpose of the paper and give a brief account of previous work. New techniques and modifications should be described concisely but in sufficient detail to permit their evaluation; standard methods should simply be referenced. Experimental results should be presented in the most appropriate form, with sufficient explanation to assist their interpretation; their discussion should form a distinct section. Extensive tabulations will not be accepted unless their inclusion is essential.

Abstracts

Structured abstracts are required for all papers, and should be submitted as detailed below, following the title and author's name and address, preceding the main text.

Intending contributors should note that Disability and Rehabilitation: Assistive Technology is changing the format of the summary of each paper to that of a structured abstract. The change will operate with immediate effect. There is clear evidence that structured abstracts contain more accessible information than summaries and are therefore of more use to the readership.

All papers submitted to the Journal of Disability and Rehabilitation: Assistive Technology should have a 'structured abstract' of no more than 200 words. The following headings should be used, following the title, author's name and address, and preceding the main text:

Purpose State the main aims and objectives of the paper.
Method Describe the design, and methodological procedures adopted.
Results Present the main results.
Conclusions State the conclusions that have been drawn and their relevance to the study of disability and rehabilitation.

Nomenclature and Units
All abbreviations and units should conform to SI practice. Drugs should be referred to by generic names; trade names of substances, their sources, and details of manufacturers of scientific instruments should be given only if the information is important to the evaluation of the experimental data.

Copyright Permission

Contributors are required to secure permission for the reproduction of any figure, table, or extensive (more than fifty word) extract from the text, from a source which is copyrighted - or owned - by a party other than Informa UK Ltd or the contributor.

This applies both to direct reproduction or 'derivative reproduction' - when the contributor has created a new figure or table which derives substantially from a copyrighted source.

The following form of words can be used in seeking permission:

Dear [COPYRIGHT HOLDER]

I/we are preparing for publication an article entitled

[STATE TITLE]

to be published by Informa UK Ltd in Disability and Rehabilitation: Assistive Technology.

I/we should be grateful if you would grant us permission to include the following materials:

[STATE FIGURE NUMBER AND ORIGINAL SOURCE]

We are requesting non-exclusive rights in this edition and in all forms. It is understood, of course, that full acknowledgement will be given to the source.

Please note that Informa UK Ltd are signatories of and respect the spirit of the STM Agreement regarding the free sharing and dissemination of scholarly information.

Your prompt consideration of this request would be greatly appreciated.

Yours faithfully

Code of Experimental Ethics and Practice

Contributors are required to follow the procedures in force in their countries which govern the ethics of work done with human or animal subjects. The Code of Ethics of the World Medical Association (Declaration of Helsinki) represents a minimal requirement.

When experimental animals are used, state the species, strain, number used, and other pertinent descriptive characteristics.

For human subjects or patients, describe their characteristics.

For human participants in a research survey, secure the consent for data and other material - verbatim quotations from interviews, etc. - to be used.

When describing surgical procedures on animals, identify the pre-anaesthetic and anaesthetic agents used
and state the amount of concentration and the route and frequency of administration for each. The use of paralytic agents, such as curare or succinylcholine, is not an acceptable substitute for anaesthetics. For other invasive procedures on animals, report the analgesic or tranquilizing drugs used; if none were used, provide justification for such exclusion.

When reporting studies on unanaesthetized animals or on humans, indicate that the procedures followed were in accordance with institutional guidelines.

Specific permission for facial photographs of patients is required. A letter of consent must accompany the photographs of patients in which a possibility of identification exists. It is not sufficient to cover the eyes to mask identity.

Clinical Trials Registry

Disability and Rehabilitation: Assistive Technology requests, as a consideration of publication, that clinical trials are registered in a public repository at their inception and prior to patient enrolment.

The registry must be accessible to the public at no charge, be open to all prospective registrants and managed by a not-for-profit organization. For a list of registries that meet all of these requirements, please see the WHO International Clinical Trials Registry Platform (ICTRP) http://www.who.int/ictrp/en/. This is in accordance with the guidelines published by the International Committee of Medical Journal Editors (ICMJE). For more information, see ICMJE Uniform Requirements for Manuscripts Submitted to Biomedical Journals http://www.icmje.org

The registration of all clinical trials facilitates the dissemination of information among clinicians, researchers and patients, and enhances public confidence in the research enterprise.

Offprints and Reprints

Offprints and reprints of articles published in this journal can be purchased once the article has been published online. Corresponding authors will receive free online access to their article through our website (www.informaworld.com). Reprints of articles published in this journal can be purchased through the Publisher when proofs are received. Copies of the Journal can be purchased separately at the author's preferential rate of £15.00/$25.00 per copy.

Colour Figures

a. Any figure submitted as a colour original will appear in colour in the journal's online edition free of charge and can be downloaded.

b. Paper copy colour reproduction will only be considered on condition that authors contribute to the associated costs. Charges are: £500/US$1030 for the first colour page and £250/US$515 for each colour page after per article. (Colour costs will be waived for invited Review Articles)

Electronic Processing

The following procedures which will assure we can process your article efficiently.

1. Authors using standard word-processing software packages

For the main text of your article, most standard PC or Mac word-processing software packages are acceptable, although we prefer Microsoft Word in a PC format.
Word-processed files should be prepared according to the journal style.

Avoid the use of embedded footnotes. For numbered tables, use the table function provided with the word-processing package.

All text should be saved in one file with the complete text (including the title page, abstract, all sections of the body of the paper, references), followed by numbered tables and the figure captions.

2. Authors using LaTeX mathematical software packages

Authors who wish to prepare their articles using the LaTeX document preparation system are advised to use article.sty (for LaTeX 2.09) or article.cls (for LaTeX2e).

The use of macros should be kept to an absolute minimum but if any are used they should be gathered together in the file, just before the \begin{document} command.

Articles prepared using LaTeX should be converted to PDF documents (incorporating the illustrations) and these should be submitted online in addition to the associated LaTeX source and graphics files.

The files you send must be text-only (often called an ASCII file), with no system-dependent control codes.

3. A guide for authors using graphics software packages

We welcome figures, but care and attention to these guidelines is essential, as importing graphics packages can often be problematic.

Figures must be saved separately from the text.

Avoid the use of colour and tints for aesthetic reasons. Figures should be produced as near to the finished size as possible.

All figures must be numbered in the order in which they occur (e.g. figure 1, figure 2 etc.). In multi-part figures, each part should be labelled (e.g. figure 1 (a), figure 1 (b) etc.)

The figure captions must be saved as a separate file with the text and numbered correspondingly.

The filename for the graphic should be descriptive of the graphic e.g. Figure1, Figure2a.

Files should be saved as TIFF (tagged image file format), PostScript or EPS (encapsulated PostScript), containing all the necessary font information and the source file of the application (e.g., CorelDraw/Mac, CorelDraw/PC).

Notes on Style

All authors are asked to take account of the diverse audience of the journal. Clearly explain, or avoid the use of, terms that might be meaningful only to a local or national audience.

Some specific points of style for the text of articles, research reports, case studies, reports, essay reviews, and reviews follow:

1. We prefer US to 'American', USA to 'United States', and UK to 'United Kingdom'.

2. We use conservative (British, not US, spelling, i.e. colour not color; behaviour (behavioural) not behavior; [school] programme not program; [he] practises not practices; centre not center; organization not organisation; analyse not analyze, etc.

3. Single 'quotes' are used for quotations rather than double "quotes", unless the 'quote is "within" another quote'.

4. Punctuation should follow the British style, e.g. 'quotes precede punctuation'.

5. Punctuation of common abbreviations should follow the following conventions: e.g. i.e. cf. Note that such abbreviations are not followed by a comma or a (double) point/period.

6. Dashes (M-dash) should be clearly indicated in manuscripts by way of either a clear dash (-) or a double hyphen (--).

7. We are sparing in our use of the upper case in headings and references, e.g. only the first word in paper titles and all subheads is in upper case; titles of papers from journals in the references and other places are not in upper case.

8. Apostrophes should be used sparingly. Thus, decades should be referred to as follows: 'The 1980s [not the 1980's] saw ...'. Possessives associated with acronyms (e.g. PA), should be written as follows: 'The APU's findings that ...', but, NB, the plural is APUs.

9. All acronyms for national agencies, examinations, etc., should be spelled out the first time they are introduced in text or references. Thereafter the acronym can be used if appropriate, e.g. 'The work of the Assessment of Performance Unit (APU) in the early 1980s ...'. Subsequently, 'The APU studies of achievement ...', in a reference ... (Department of Education and Science [DES] 1989a).

10. Brief biographical details of significant national figures should be outlined in the text unless it is quite clear that the person concerned would be known internationally. Some suggested editorial emendations to a 'typical' text are indicated in the following with square brackets: 'From the time of H. E. Armstrong [in the 19th century] to the curriculum development work associated with the Nuffield Foundation [in the 1960s], there has been a shift from heurism to constructivism in the design of [British] science courses'.

11. The preferred local (national) usage for ethnic and other minorities should be used in all papers. For the USA, 'African-American', 'Hispanic' and 'Native American' are used, e.g. 'The African American presidential candidate, Jesse Jackson ... '; for the UK, 'Afro-Caribbean' (not 'West Indian'), etc.

12. Material to be emphasized (italicized in the printed version) should be underlined in the typescript rather than italicized. Please use such emphasis sparingly.

Mathematics

Special care should be taken with mathematical scripts, especially subscripts and superscripts and differentiation between the letter 'ell' and the figure one, and the letter 'oh' and the figure zero. If your keyboard does not have the characters you need, it is preferable to use longhand, in which case it is important to differentiate between capital and small letters, K, k and x and other similar groups of letters. Special symbols should be highlighted in the text and explained in the margin. In some cases it is helpful to supply annotated lists of symbols for the guidance of the sub-editor and the typesetter, and/or a 'Nomenclature' section preceding the 'Introduction'.

For simple fractions in the text, the solidus / should be used instead of a horizontal line, care being taken to insert parentheses where necessary to avoid ambiguity, for example, $1/(n-1)$. Exceptions are the proper fractions available as single type on a keyboard.

Full formulae or equations should be displayed, that is, written on a separate line. Horizontal lines are preferable to solidi, for example:

$$\frac{61+5h+q}{3n+3yz^2}$$

But: $a/b + c/d + a/d$
\[ P = (a^2 - b^2)(c^2 + d^2) \]

The solidus is not generally used for units: \( \text{ms}^{-1} \) not m/s, but note electrons/s, counts/channel, etc.

Displayed equations referred to in the text should be numbered serially (1, 2, etc.) on the right hand side of the page. Short expressions not referred to by any number will usually be incorporated in the text.

Symbols should not be underlined to indicate fonts except for tensors, vectors and matrices, which are indicated with a wavy line in the manuscript (not with a straight arrow or arrow above) and rendered in heavy type in print: upright sans serif \( \mathbf{r} \) (tensor), sloping serif \( \mathbf{r} \) (vector) upright serif \( \mathbf{r} \) (matrix).

Typographical requirements must be clearly indicated at their first occurrence, e.g. Greek, Roman, script, sans serif, bold, italic. Authors will be charged for corrections at proof stage resulting from a failure to do so.

Braces, brackets and parentheses are used in the order \{[( )]&reuben;/, except where mathematical convention dictates otherwise (i.e. square brackets for commutators and anticommutators).

Citations in Text

We prefer that references are cited using the numerical system (e.g. \[3\], \[5-9\]). They should be listed separately at the end of the paper in the order in which they appear in the text.

Notes on Tables and Figures

1. Tables and figures should be valuable, relevant, and visually attractive. Tables and figures must be referred to in the text and numbered in order of their appearance. Each table and figure should have a complete, descriptive title; and each table column an appropriate heading.

   Tables and figures should be referred to in text as follows: figure 1, table 1, i.e. lower case. 'As seen in table [or figure] 1 ...' (not Tab., fig. or Fig).

2. The place at which a table or figure is to be inserted in the printed text should be indicated clearly on a manuscript:
   
   [Insert table 2 about here ]

3. Each table and/or figure must have a title that explains its purpose without reference to the text.

4. All figures and tables must be on separate sheets and not embedded in the text. Original copies of figures should be supplied. All figures should allow for reduction to column width (130 mm) or page width (160mm). Please avoid figures that would require landscape reproduction, i.e., reading from bottom to top of the page. Photographs may be sent as glossy prints or negatives.

   Do not type the caption to a figure on that figure; the legends to any illustrations must be typed separately following the main text and should be grouped together.

Acknowledgements

Any acknowledgements authors wish to make should be included in a separate headed section at the end of the manuscript. Please do not incorporate these into the bionote or notes.
Declaration of interest

It is the policy of all Informa Healthcare to adhere in principle to the Conflict of Interest policy recommended by the International Committee of Medical Journal Editors (ICMJE, http://www.icmje.org/index.html#conflict).

All authors must disclose any financial and personal relationships with other people or organisations that could inappropriately influence (bias) their work. It is the sole responsibility of authors to disclose any affiliation with any organisation with a financial interest, direct or indirect, in the subject matter or materials discussed in the manuscript (such as consultancies, employment, paid expert testimony, honoraria, speakers' bureaus, retainers, stock options or ownership, patents or patent applications or travel grants) that may affect the conduct or reporting of the work submitted. All sources of funding for research are to be explicitly stated. If uncertain as to what might be considered a potential conflict of interest, authors should err on the side of full disclosure.

All submissions to the journal must include full disclosure of all relationships that could be viewed as presenting a potential conflict of interest. If there are no conflicts of interest, authors should state that there are none. This must be stated at the point of submission (within the manuscript after the main text under a subheading "Declaration of interest" and, where available, within the appropriate field on the journal's Manuscript Central site). This may be made available to reviewers and will appear in the published article at the discretion of the Editors or Publisher.

If no conflict is declared, the following statement will be attached to all articles:

Declaration of interest: The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

The intent of this policy is not to prevent authors with these relationships from publishing work, but rather to adopt transparency such that readers can make objective judgements on conclusions drawn.

Plagiarism

Informa Healthcare has a strict policy against plagiarism. We define plagiarism as the use of extracts from another person's work that are not placed in quotation marks, without the permission of that person, and without acknowledgement to that person (using the appropriate reference style), with the result that your article presents these extracts as original to you. By submitting your work to an Informa Healthcare journal, you warrant that it is your original work, and that you have secured the necessary written permission from the appropriate copyright owner or authority for the reproduction of any text, illustration, or other material.

If any article submitted to an Informa Healthcare journal is found to have breached any of these conditions, Informa Healthcare reserves the right to reject that article and any others submitted by the same authors. Informa Healthcare may also contact the authors' affiliated institutions to inform them of its findings.

References

References should follow the CBE Citation & Sequence format. Only works actually cited in the text should be included in the references. Indicate in the text with Arabic numbers inside square brackets. Spelling in the reference list should follow the original. References should then be listed in numerical order at the end of the article. Examples are provided as follows:
Journal article:

Book chapter:

Conference proceedings:

Dissertations or Thesis:

Journal article on internet:

Webpage:

Internet databases:

Further examples and information can be found in the CBE style manual Scientific Style and Format, sixth edition.

NIH Public Access Policy

In consideration of the National Institutes of Health (NIH) Public Access Policy, Informa Healthcare acknowledges that the broad and open dissemination of NIH-funded-research results may benefit future scientific and medical research. Because we value the current and future contributions our journals make to the scientific body of knowledge, we have made certain that our policies accommodate those authors who wish to submit to PubMed Central.

Informa Healthcare's position with respect to public access to NIH-funded work published in Informa Healthcare journals is as follows:

- Informa Healthcare authors may voluntarily submit their funded work to PubMed Central after a 12-month embargo period;
"funded work" shall be defined as the final, peer-reviewed manuscript that is accepted by the Editor in Chief of the journal. This manuscript must not be altered by Publisher's copyediting and typesetting services; and

this embargo period begins the day the work is published online at www.informaworld.com.
The Smart Wheelchair: Is it an appropriate mobility training tool for children with cerebral palsy?

Sarah McGarry
The Smart Wheelchair: Is it an appropriate mobility training tool for children with cerebral palsy?

Abstract

Purpose: To describe the impact of a mobility training program using the Smart Wheelchair on the driving skills and psychosocial outcomes of children with cerebral palsy.

Method: A multiple case study design using mixed methods was used. Four children with cerebral palsy were recruited for the study. The intervention was a Smart Wheelchair mobility training program. Data was collected using a quantitative driving skills assessment, field notes and qualitative parent interviews.

Results: Three out of four children gained independence in at least three driving skills or more, whilst one child was competent with verbal prompts. Three out of four mothers reported positive changes in their child’s confidence, motivation and affect.

Conclusions: The Smart Wheelchair has the ability to uncover learning potential and facilitate the recognition of abilities in children with cerebral palsy previously excluded from access to independent mobility. Given the significant limitation that restrictions in mobility pose to participation for children with cerebral palsy, occupational therapists must begin to understand the effectiveness of interventions such as the Smart Wheelchair. The descriptive findings of this study allow for future, more rigorous research, to be conducted on the effectiveness of the Smart Wheelchair as a mobility training tool.

Key Words: powered wheelchairs, intelligent wheelchairs, mobility interventions, children, cerebral palsy

Sarah McGarry,
Lois Moir,
Dr Sonya Girdler,
September, 2009
The Smart Wheelchair: Is It an Appropriate Mobility Training Tool For Children with Cerebral Palsy?

Introduction

Cerebral palsy (CP) is the most common physical childhood disability [1-3], accounting for 600 to 700 Australian births each year [1]. Advances in perinatal care have been associated with declining rates of infant mortality among pre-term infants and an increase in the number of infants born with CP [4,5]. It was estimated in 2007 that 33,797 Australians have CP and this number is expected to increase to 47,601 by 2050 [1]. This increase in absolute numbers of Australians living with CP will result in a corresponding increase in demand for effective therapeutic interventions.

Many children with CP are unable to independently mobilise and consequently miss many crucial learning opportunities. This impacts negatively on their skill development in the areas of cognition, behaviour, emotion and communication [6-8]. Secondary developmental problems can result from a child’s increasing frustration at their inability to control the world around them [9]. Mobility impairment therefore often results in a cycle of deprivation, and reduced motivation [10].

Many children with CP are unable to operate a powered wheelchair (PWC) due to difficulties in motor and cognitive domains [11]. Children with CP have often been excluded from powered mobility research as they typically have higher learning needs compared to other children with developmental disabilities [12]. Clinically, therapists are confronted with the challenge of assessing whether a child is eligible for powered mobility training, and if they are not, what further options are available for mobility [13]. In 2005 a national survey of providers of paediatric PWCs in the USA, found that cognitive, physical, and behavioural factors were the three main reasons therapists did not refer children with disabilities for a PWC [14].

Occupational therapists assist individuals to perform meaningful and functional activities in order to enhance occupational performance and play an important role in facilitating the learning process for
children to develop motor skills [15]. Motor learning theory can be used by occupational therapists to conduct occupation-based interventions that are meaningful to individuals [16]. Many researchers suggest that programs enhancing functional abilities for children with developmental delays should include motor learning theory [16-18]. Motor learning theory facilitates skill acquisition as a result of practice and experience [19]. Feedback and repetition are the two key strategies used to facilitate skill acquisition [19], and it has been noted that children with CP respond well to this approach [18]. The use of feedback allows the therapist to be a responsive partner in therapy whilst maintaining a child-centred approach [17,20,21]. Occupational therapists can use motor learning theory to develop mobility training programs for children with disabilities.

The Smart Wheelchair is a mobility option for children who are considered to not have the prerequisite skills required to drive a PWC. The Smart Wheelchair is an augmentative mobility aid equipped with specialised sensors and a computerised control system. Functions of the chair include collision sensors and a line follower, allowing the user to navigate difficult environments with the use of a single capacity switch [22,23]. The aim of the Smart Wheelchair is not to replace the user’s abilities with technology, but to complement and expand the user’s skills [22]. The Smart Wheelchair can be viewed as a therapeutic tool as its objective is not only to allow the child to move from one area to another, but to learn new skills and progress developmentally. It has been proposed that using the Smart Wheelchair in combination with a clinical assessment tool, therapists are able to evaluate how far along the mobility continuum a child has progressed [22].

There is a need for new technologies such as the Smart Wheelchair to be evaluated. To date the inventors of the CALL Centre Smart Wheelchair are the only researchers to publish a study investigating the use of this wheelchair with children with motor and cognitive disabilities [10]. Given that the number of children living with CP is expected to increase in coming years and the significant impact that restrictions in mobility has on development, research must begin to evaluate the effectiveness of interventions such as
the Smart Wheelchair. Understanding the effectiveness of technologically advanced PWCs in maximising the mobility of children with CP has the potential to reduce the burden of disability associated with CP. There is a need for therapists and families to gain an understanding of the utility of the Smart Wheelchair, both as a method of access and as a therapeutic intervention tool. The purpose of this study was to describe the impact of a mobility training program using the Smart Wheelchair on the driving skills and psychosocial outcomes of children with CP.

**Materials and Methods**

This study employed a multiple case study design using mixed methods. Case studies are a suitable approach to explore, describe and explain complex phenomena within their natural context [24,25]. Case studies are appropriate designs to evaluate interventions in health science research because of their rigor, flexibility [24] and direct applicability to patient care [26,27]. Evidence from multiple case studies is often considered to be more robust and compelling compared to single case studies due to the replication of patterns and trends [25].

A mixed method approach to data collection was utilised in this study as it facilitates the collection of information from multiple perspectives, allowing a greater understanding of the research topic [28,29]. The purpose of using mixed methods was complementarity which allowed facets of the research phenomenon to be measured, facilitating an enriched and elaborated understanding of the phenomenon [28]. Ethical approval was obtained from the Edith Cowan University Human Research Ethics Committee and the Centre for Cerebral Palsy (TCCP).

Children in this study were recruited through TCCP. TCCP is a not for profit charitable organisation which provides services and support to more than 1000 individuals with CP and their families. Literal replication logic was used to select participants. This permitted the researcher to select cases that were similar allowing patterns in the data to emerge, enhancing the transferability of the evidence collected.
Four participants were deemed adequate to allow for replication within case studies [25]. Informed consent was gained from all participant’s legal guardians and all participants were: 1) aged between four and fourteen years; 2) diagnosed with cerebral palsy; 3) classified as level V on the Gross Motor Functioning Classification Scale [31]; 4) had minimal to extensive experience using a switch; and, 6) had never been exposed to a Smart Wheelchair.

**Intervention**

Each participant took part in the Smart Wheelchair Mobility Training Program (SWMTP). The intervention protocol for this program is outlined in appendix A. The SWMTP was conducted at TCCP and consisted of two one hour sessions per week (on consecutive days), for eight weeks. The researcher, a fourth year occupational therapy (honours) student conducted the SWMTP with supervision and guidance from an experienced occupational therapist. Activities in each session were aimed at encouraging the driving skills of the children, and were based on motor learning theory [19]. Motor learning theory supported a focus on motor skill acquisition and the learning process of these skills. An important component of motor learning theory is the therapist’s use of feedback [32]. To facilitate skill acquisition the therapist provided extrinsic feedback both verbal and physical, combined with structured practice to illicit an intrinsic feedback response in each child [32]. For four weeks, the sessions were held in a large room with toys to increase the child’s motivation and basic mobility skills. As the child’s abilities progressed sessions were held in the school environment, and outside areas to enhance functional driving skills.

**Driving skills**

Driving skills were measured using the Powered Mobility Program assessment battery, adapted (with permission) to reflect the use of a switch control rather than a joystick in basic mobility skills [33]. The adapted assessment was piloted with a child (not enrolled in the study) meeting the inclusion criteria [25]. In administering the assessment the researcher observed each participant driving the Smart Wheelchair for
a minimum of 45 minutes and scored the assessment according to the highest observed level of skill. For example a score of one is given if a child requires maximal hands on assistance compared to a score of 5 which equates to the child conducting the skill independently with age appropriate supervision. Skills observed included basic cause and effect association, contacting the switch to engage the wheelchair in motion, maintaining contact with the switch for continuous driving, looking in the direction of movement, turning in one or two directions and stopping on command and spontaneously to avoid objects (see appendix B). During each intervention session the researcher recorded field notes which included observations of changes in the child’s behaviour, important events and the physical setting of the environment [34].

Psychosocial outcomes
In order to describe psychosocial changes for each child semi-structured interviews were conducted with parents. Five open ended questions addressing the impact of the SWMTP on changes in communication, behaviours, motivation, affect, and initiation formed the interview guide, which while allowing for flexibility helped to centre the interview on the area of inquiry (see appendix C) [35]. The interview guide was piloted with a parent of a child with CP participating in a Smart Wheelchair clinic but not enrolled in the present study [36]. Parent interviews were conducted by the researcher after the completion of the SWMTP. Each interview was face-to-face in nature and conducted at TCCP or the parent’s home. The interviews were recorded using a voice recorder.

In order to describe changes in driving skills, data was entered into Microsoft Excel to produce line graphs [26]. For each participant a total average score was calculated for the first and last session [26]. Inductive analysis and reflexive interpretation of the field notes occurred at the completion of data collection in order to search for meaning and patterns within the data [34,37].
Parent interviews were transcribed verbatim onto a secure computer and compared with field notes taken during each interview. Individual interviews were analysed using narrative analysis to reduce the data and develop a summary of the phenomenon researched [35,38]. Narrative analysis can assist occupational therapists to gain an enhanced understanding of the participant's experience of the therapeutic process [38]. The credibility of the interview findings was enhanced by member checking with one parent to verify data and interpretation [39].

The trustworthiness of this study was enhanced by triangulation of data collection, literal replication logic to select participants, peer debriefing, member checks, field notes, an audit trail, and piloting of data collection tools [39,40].

Findings

Four children meeting the selection criteria were recruited. The participants in this study represent children typically excluded from powered wheelchair access. Participants ages ranged from five to 13 years. All participants lived at home with their parents and three of the children attended respite care regularly. Case studies are presented in the form of vignettes and pseudonyms were used to maintain confidentiality.

David

David was a 13 year old boy with West Syndrome, bilateral astigmatism, central hypotonia with mild peripherally increased tone, and an intellectual disability. David was dependent on his parents and carers for all activities of daily living (ADL) and communicated with vocalisations. He attended a WA Department of Education Support School. David experienced regular seizures which impacted on his level of alertness and it was unknown whether David understood cause and effect relationships.
In the past David’s switching had been unreliable and his therapist was unable to recommend a consistent switch site. Due to ill health David attended 11 of the 16 SWMTP sessions. David’s switching improved with a large plate switch covered in tinfoil to incorporate sensory stimulation. He was encouraged to use his right hand to reach forward to activate the switch with his fingers. David required hand-over-hand prompting in the first session and became very excited, shouting loudly and banging his head. In the second session of the SWMTP David responded well to the Smart Wheelchair, he was alert and activated the switch independently, with verbal prompting, three times. For the next five sessions David required mostly hand-over-hand prompting to activate the switch, and encouragement through verbal feedback. During the sessions David was alert and responsive to his environment. In the last four sessions with constant repetition, motivational toys, and intrinsic and extrinsic feedback, David switched independently, looked in the direction of movement and demonstrated an understanding of cause and effect. For example, David was able to activate the switch three times to reach his favourite toy. In each of the categories presented in graph 1, David improved from a score of 0 (not attempting the task) to 4 (verbal cuing only). In the first session David’s total average score on the driving skills assessment was 0.09 compared to the last session where he scored 1.09. David used the line following function for all sessions of the SWMTP.

In the interview following the SWMTP David’s mother discussed that she was pleased that David was switching independently and eager for David to transfer these skills to the classroom. She felt that it was hard to directly attribute participation in the SWMTP with positive changes in his behaviour such as increased eye contact, but she did feel that David demonstrated an increase in positive affect:

"Few occasions when I have said to my husband ‘oh Daniel is really happy today’. And he can be but, [has been a] few more times than usual."

David’s mother had also observed changes in his motor patterns:

"Recently looking for toys and reaching around [his wheelchair] in the last couple of months. More use of his right arm".
Sandra

Sandra was a 12 year old girl with spastic quadriplegia and dyskinetic CP. She was dependent on her parents and carers for all ADLs and attended a WA Department of Education Support School. She communicated ‘yes’ and ‘no’ through eye movements and was fed through a gastronomy feeding tube. Switching had been trialled previously with Sandra, but due to her dyskinetic movements, her switch access was often unreliable. Two switches were trialled with Sandra, the most successful being a large plate switch, which was placed on her tray. Sandra made contact with this switch with her fingers, back of her hand or her wrist. Due to ill health Sandra was able to attend seven of the 16 SWMTP sessions.

Sandra’s increased motivation to mobilise was evident through her high level of alertness and determination to overcome dysfunctional movement synergies. She self regulated her fatigue by taking regular rest breaks. Sandra understood cause and effect in relation to driving the Smart Wheelchair and she looked in the direction of movement at all times. Sandra used the line following function for all sessions of the SWMTP. During the first five SWMTP sessions, Sandra required stand-by assistance to activate the switch (see graph 2). In her last two sessions, the switch was changed to one with a larger surface area which assisted her to dampen her dyskinetic movements. Repetition facilitated intrinsic feedback for Sandra to control her movement synergies and energy expenditure. In session six Sandra began to independently switch (see graph 2). As described by Sandra’s mother this represented an important milestone:

“We have been trying to get Sandra to switch for years, she is 12 and we started when she was four. We haven’t had much progress, because she really gets too frustrated and in the end she just gives up. Where as something like that she wants to do it because she wants to move forward...
Sandra’s [Teacher] saw Sandra in [the Smart Wheelchair] she said she just did so well, it was voluntary, she was accurate in her switching, she stopped when she wanted to stop, she obviously feels more comfortable.”
In the first session Sandra’s total average score on the driving skills assessment was 0.63 compared to the last session where she scored 1.36. Sandra’s improved switching abilities also had a positive impact on her performance at school. As her mother explained:

“As far as school is concerned, they definitely see improvement.”

Following participation in the SWMTP Sandra’s mother observed an increase in her daughter’s confidence and affect:

“She was really proud of herself because it was something she achieved on her own...It was almost like the penny dropped with her to physically be able to control the switch and that is a massive achievement for Sandra to do something like that...When they do something right, they become confident. It is nice to see when we mention something that she has achieved, she lights up and she thinks she is pretty good, it is so nice, because you don’t get that very often...There are not many things that are available for these kids to make them feel good about themselves.”

Sandra’s mother summarised the impact the SWMTP had on Sandra and her perceptions of the Smart Wheelchair as an intervention tool:

“It would be really nice to have it ongoing, it would improve a lot of things in their life, it is an incentive, they do it on their own. They need something that they want to do independently. Everything gets done for them but this is something they can do on their own. It’s a great thing, I’m really pleased that it is out there.”

Blake

Blake was a five year old boy with spastic quadriplegia and epilepsy. He was dependent on his parents and carers for all ADLs and communicated with vocalisations and facial expressions. He attended conductive education classes at a WA Department of Education Support School. Blake had poor head control and was fed through a naso-gastric tube. He had previously been encouraged to switch, however his switching performance was unreliable. Due to ill health Blake attended 11 of the 16 SWMTP sessions
A large button switch was placed on Blake’s tray, with the intention that he would reach forward with his left arm to activate the switch with his fingers. To begin with Blake required hand over hand prompting to press the switch, however he clearly understood the causal relationship between activating the switch and driving forward. After six sessions Blake was independently activating the switch to drive forward (see graph 3). Once Blake achieved this skill the switch was adapted from timed, where movement is set to a specific amount of time, to momentary, where Blake determined the amount of time spent moving by maintaining constant pressure on the switch. Blake’s hand control gradually improved and he was capable of independently holding his hand on the switch to drive for a minimum of ten seconds (see graph 3). By the eleventh session Blake was able to stop on command with verbal prompting (see graph 3). In the first session Blake’s total average score on the driving skills assessment was 0.18 compared to the last session where he scored 3.27. Blake progressed from the line following function to using the collision sensors.

One key improvement observed over the course of the SWMTP was Blake’s ability to look in the direction of movement. For the first five sessions Blake held his head down and looked only at the switch and laterally out of the corner of his eye. With verbal prompting, Blake began to gradually lift his head to look in the direction of movement as well as to notice his environment. This was particularly apparent when Blake began driving outside. This change in physical context engaged Blake and facilitated an improvement in his performance. Blake looked in the direction of movement for the last three sessions (see graph 3).

In the post intervention interview Blake’s mother described the process she felt her son undertook to establish the correct timing and sequencing of his movement synergies:

"He was head up looking around for the first time... as opposed to the weeks before hand, where he was just looking down at the switch to know where it was, but it is as if that the message to the arm has become automated so he knows how to move his arm to reach the switch, he knows where
the switch is going to be, and now he has given himself stage two, I am now able to do that and look where I am going."

Following the SWTMP, Blake demonstrated skills he had previously not had the opportunity or motivation to develop:

"Talking with [the therapist], we think he might be able to work on, not just cause and effect type activities on the computer but now scanning and choice... which then translates to communication devices and more inclusion and accessibility to other people and being involved in that way. But that is a stage that he wasn't ready for at all before, we just hadn't put him in that category yet, but he seems to be more improved now."

Blake’s mother observed several psychosocial improvements following the SWMTP. She reported that Blake had demonstrated more initiative to attempt tasks, was more motivated to participate in activities, and was demonstrating increased positive affect. Blake was vocalising more and his mother reported he was communicating with new vocalisations. In addition, Blake’s use of his left arm had increased:

"He reached out a bit more to play with a toy...or to knock down blocks. A little bit more use of arm and reaching movement...Huge [change in vocalisations], much more vocal and even different sounds...He is wanting to be part of a group’s activities vocally and that's a difference...More initiative, willingness and some more enthusiasm for trying and motivation, rather than giving up."

Blake’s mother observed the motivational impact independent mobility had on her child:

"It has come at the right time, because even emotionally and mentally he was at a giving up stage, we hadn’t done hardly any programs for a long time, so now we did this and it is as if he was given a chance to prove that he can do something new, he is happier, more vocal and more enthusiastic."

Pauline
Pauline was a six year old girl with spastic diplegia involving both spasticity and dystonia. Pauline attended a main stream school and was capable of assisting her parents with some ADLs such as dressing and bathing. Pauline independently ate soft foods and could follow simple instructions as well as verbalise her needs. Pauline was a very social girl and was capable of basic communication in both English and Swedish. Due to ill health Pauline attended 11 of the 16 SWMTP sessions.

Pauline demonstrated poor hand control due to her tonal pattern and initially required a soft switch. Pauline began the SWMTP using the line follower function, then progressed to collision sensors, and finally was capable of driving without the use of sensors. As Pauline progressed she began to use three switches set on momentary, to manoeuvre forward, left and right. Through constant repetition and intrinsic feedback, Pauline developed the necessary postural control and movement synergies required to activate a flat button switch. Pauline’s problem solving, motor planning, spatial understanding and bilateral coordination significantly improved from requiring verbal prompts to navigate and stop, to independently driving through doorways and turning corners (see graph 4). In the first session Pauline’s total average score on the driving skills assessment was 1.90 compared to the last session were she scored 5.

Pauline consistently demonstrated positive affect whilst driving by laughing and smiling. Pauline continually vocalised her ability to drive by stating “I can” proudly. She also confirmed her independence by repeating frequently “on my own”. In the last three SWMTP sessions Pauline began to incorporate imaginary play, stopping for pretend trucks to pass, and asking an imaginary person for directions. This demonstrated significant developmental progression in play.

Pauline’s mother observed the positive impact the SWMTP had on her child’s confidence, independence and initiative to perform tasks:
“Because she sees she can do the driving on her own and she’s controlling that, I think she now realises that she can control a few things and is more willing to try to do that... She is more confident... that she can do things, amongst them thinking that she can tell me what to do while we are in the car, because she has been doing her driving lessons. Definitely more content to do things on her own. More wanting to try things on her own.”

Pauline’s mother reported that she was previously reluctant for Pauline to use a PWC however, Pauline’s increased independence and positive affect were all factors facilitating her to change her point of view:

“Personally, so excited to see her being independent, being able to do something herself, and able to control something herself in her environment that she can’t really always do otherwise. And the excitement, she has been so happy doing it and absolutely loving it... I’ve always been very hesitant about a motorised wheelchair for lots of different reasons, but seeing her do this and that she is able to do it, I think has changed my view of the whole thing.”

As a result of the SWMTP Pauline’s therapist and family are applying for funding for a PWC, a piece of equipment her family had not previously considered.

Discussion

The aim of this study was to describe the impact of a mobility training program using the Smart Wheelchair on the driving skills and psychosocial outcomes of children with CP. Following the SWMTP three out of four children gained independence using the Smart Wheelchair in at least three driving skills or more, whilst one child was competent with verbal prompts. Three out of four mothers reported positive changes in their child’s confidence, motivation and affect. Although the evidence is descriptive in nature, these findings suggest that the Smart Wheelchair has the potential to improve the driving skills and psychosocial outcomes of children with CP, who have previously been excluded from powered mobility training.
Findings from this study highlighted the need to look beyond a child’s limitations in order to uncover potential abilities. The Smart Wheelchair was a useful tool for uncovering skills and abilities previously unrecognised in all of the children. The importance of powered mobility as a therapeutic tool has been discussed by therapists working with children with multiple and complex disabilities [41,42]. This approach is based on the premise that sensory stimulation and change in body positions resulting from powered mobility can raise an individual’s alertness, enhancing their ability to understand simple causal relationships [41]. Findings from the present study suggest that powered mobility not only has the potential to help develop understanding of causal relationships, but to increase motor performance, postural control and influence positive psychosocial changes.

The SWMTP was underpinned by motor learning theory [19]. Previously, children in this study were believed to lack the pre-requisite skills required for powered mobility. Application of this theory allowed the children to receive constant intrinsic and extrinsic feedback through repetition and practice, consequently facilitating skill acquisition [19]. Children were able to reinforce the development of learnt skills on consecutive days and, repetition and practice were key factors in the acquisition of driving skills in all four children. This is in line with findings which suggest repetition over time is essential for skill development in children with multiple and complex disabilities [17,41]. This study also confirms that removing pre-determined criteria and rigid protocols when providing powered mobility training, enhances the development of skill learning specific to each individual child [20]. These findings support the idea that the emphasis in therapy should be placed on the therapist as a responsive partner who can elicit new learning through providing responsive feedback rather than commanding instructions [17,20]. Research suggests that this method reduces the child’s dependency on adults and encourages children to take more initiative for their own learning [43].

Parents perceived that the SWMTP facilitated an increase in their child’s motivation, affect, and initiative to participate in play and therapeutic activities. These changes in psychosocial outcomes are important
achievements for children with multiple and complex disabilities as they have fewer opportunities for contingency learning and often develop learned helplessness as a secondary effect [9,44]. For these reasons it is imperative that children with multiple and complex disabilities are provided with as many opportunities as possible to explore their environment. The Smart Wheelchair has the potential to break this cycle of learnt helplessness. The findings of this study are consistent with findings from a systematic literature review of powered mobility interventions for children with CP [45]. Five studies in the review evaluating psychosocial outcomes, found independent powered mobility promoted increased social interaction, increased motivation to move and interact with the environment, and increased independence [41,46-49].

The development of new assistive technologies for individuals with disabilities has increased over the past three decades [50,51]. New models of wheelchairs such as the CALL Centre Smart Wheelchair and the Collaborative Wheelchair Assistant have been developed using advanced technologies and robotics [22,52]. Current evidence suggests that technologically advanced wheelchairs can increase driving skills in adults and children with disabilities [10,52,53]. However, there is a paucity of evidence to describe how effective these new technologies are for children with CP. This study described how the Smart Wheelchair can be used as a safe therapeutic tool for children with CP. The Smart Wheelchair allows the therapist to tailor the level of driver control, to suit the child’s needs. As a child’s skills progress, the Smart Wheelchair can be adapted to complement and expand their skills [10]. During the SWMTP it became apparent that this capability was crucial in allowing children with multiple and complex disabilities the opportunity to explore their abilities. The Smart Wheelchair appeared to be a reliable and efficient therapeutic tool, however, there is a clear need for future research to further explore the effectiveness of the Smart Wheelchair.

Occupational therapists are concerned with enabling the achievement of occupational performance for all individuals [15]. When considering powered mobility in the context of occupational performance, Smart
Wheelchairs are clearly a valuable intervention tool. Independent mobility has the ability to enhance occupational performance and participation in valued life roles, enhancing feelings of self efficacy and competence [54]. Children with multiple and complex disabilities are generally dependent for all ADLs and instrumental ADLs. The Smart Wheelchair has the capacity to provide these children with an activity they can control independently. There is an increased demand for clinical interventions such as the Smart Wheelchair as there are growing numbers of children with CP. Wheelchairs using advanced technology are expensive and, purchase of an inappropriate PWC can lead to technology abandonment [55]. The Smart Wheelchair allows for a complete assessment of abilities and helps to ensure that the most beneficial equipment is purchased, reducing the financial burden on families and service providers. However, an evidence base supporting the use of Smart Wheelchairs with children with CP must first be established in order for occupational therapists to support evidence based practice.

These findings must be interpreted with caution due to the following limitations. The small number of participants and the short duration of the SWMTP limits the transferability of the research findings [39]. Internal validity was weakened due to the effects of extraneous factors such as therapies received outside of the SWMTP and school programs [26]. The broad nature of the intervention protocol permitted variations in therapy for each child, limiting the intervention fidelity [56]. This research was descriptive in nature, and for this reason a causal relationship between the SWMTP and improvements in driving skills and psychosocial outcomes cannot be assumed [26]. Children with multiple and complex disabilities often have high levels of co morbidity and ill health. All four children were unwell several times during the SWMTP disrupting the continuity of the intervention however, this was expected when conducting clinical research with children of this nature.

**Conclusion**

This study described how the SWMTP improved driving skills and impacted positively on psychosocial outcomes in four children with CP. The Smart Wheelchair has the ability to uncover learning potentials
and facilitate the recognition of abilities in children with CP. Advances in technology are providing more individuals access to mobility opportunities previously inaccessible. These technologies have the potential to maximise the quality of life and independence of children with CP. Clinical research involving the clinical application of the Smart Wheelchair and its therapeutic benefits should be conducted. There is also a need for future, more rigorous research to determine causal relationships between the Smart Wheelchair and mobility outcomes. This study is descriptive in nature however, these findings contribute valuable knowledge in the field of powered mobility for children previously excluded from access to independent mobility.
References


22. Odor P, Watson M. Learning through Smart Wheelchairs: A formative evaluation of the effective use of the CALL Centre's Smart Wheelchairs as part of children's emerging mobility,


Appendix A
Intervention protocol for weekly sessions

Set up child ready to switch and drive the chair

Say “Sue, we’re going to do some (more) driving.

Sue, when you press (push, hit, squeeze) your switch, you can make the chair go.

Sue, you press (hit, squeeze) your switch to make the chair go.

SUCCESS.
Say “Wow, you’ve made the chair go.”
If child appears to be responding to stimulus (chair movement) stay reasonably quiet.
If not attending to stimulus, attempt to get their attention on the movement of the chair.
Remember you are a response, not the stimulus ie your response does not need to be the focus.

Wait 5 seconds after stimulus switches off.
Sue, press your switch again to make the chair move again.
Give short response – “great, you’ve made the chair go.”

As child begins attending to stimulus, reduce prompting to get child to be more aware of the mobility.

After 6 successful switching attempts, increase waiting time (10 - 15 seconds) before verbally cueing. Also reduce verbal praise: let the stimulus be the reward.

NO SUCCESS
* In 20 seconds, say “Sue I will help you.”
“When you push, (press, hit) your switch like this, see what happens. We made the chair move.”
If attending to the stimulus, stay quiet.
If not attending to stimulus, attempt to get their attention on the movement of the chair.
Don’t over do it and be more of a response than the stimulus.

Sue, Now you press (hit, squeeze) your switch to make the chair move.

No response in 20 seconds, repeat

No response again in 20 seconds, say “Sue, I will help you” and repeat step *
After 6 more successful switching attempts, increase waiting time to 30 seconds before verbally cueing. Reduce verbal response to one/two words - “wow”, “great”, “good switching”, “you’re moving”.

If success reduces go back to sequence where child was successful and double number of times of success before moving on again.

Reduce verbal cues as success is maintained. “Sue, make the chair go”. Don’t just say “press your switch” – this does not help child to learn or reinforce cause and effect.

Provide verbal feedback, but allow the switched stimulus to provide the feedback.

Monitor child’s responses – child may work hard for 15 minutes then be tired. Note this and allow for altered levels of response.
Intervention Protocol

Weekly Sessions

Each session was 45 minutes in duration. This was shortened or modified if the children showed signs of fatigue or distress.

Chair Settings

The following Smart Wheelchair settings were used during the program to enhance the driver's abilities:
- Line following
- Collision sensors
- Driver control

Indoor Activities

Basic mobility sessions included the following meaningful and fun activities:
- Driving through a streamer curtain
- Driving through a sound beam
- Driving past mirrors
- Driving on a mat switch connected to a switch adapted toy

Outdoor Activities

Functional mobility sessions included the following meaningful and fun activities:
- Driving into the classroom
- Taking a message to the principal’s office
- Visiting the library
- Driving around the playground
Appendix B

Driving Skills Assessment

Assessment Scale:

0 – Task not attempted

1 – Maximal hands on assistance of switch with verbal cuing

2 – Minimal hands on assistance of switch with verbal cuing

3 – Direct stand-by guarding with verbal cuing

4 – Verbal cuing only

5 – Age appropriate supervision

<table>
<thead>
<tr>
<th>Basic Mobility Skills</th>
<th>1(^{st}) Session</th>
<th>2(^{nd}) Session</th>
<th>3(^{rd}) Session</th>
<th>4(^{th}) Session</th>
<th>5(^{th}) Session</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basic cause and effect association</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Contacts switch to engage wheelchair in motion</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maintains contact with the switch for a minimum of 2 seconds</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maintains contact with the switch for a minimum of 5 seconds</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maintains contact with the switch for a minimum of 5 seconds and stops on command</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Looks in the direction of movement</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maintains contact with switch for 10 seconds</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Maintains contact with switch for 10 seconds and stops on command</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stops spontaneously to avoid stationary objects</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Switches to turn in one direction</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Switches to turn in two directions</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Modified from Powered Mobility Program developed by Furumasu, Guertette & Tefft (1996)
Appendix C

Semi Structured Interview Guide

Today we are going to discuss the impact of the mobility training program on your child, but before we start could you tell me a little bit about your son/daughter?

- Can you tell me about any changes you have noticed in your child’s behaviour over the past eight weeks?
  
  Probe:
  - At home?
  - At school?
  - Self stimulating behaviour?

- Can you tell me about any changes you have noticed in your child’s communication with others?
  
  Probe:
  - Family?
  - School friends?

- Can you tell me about any changes you have noticed in your child’s initiative to do things?
  
  Probe:
  - Socialise?
  - Mobilise?
  - Play?

- What are the positive and negative experiences your child has had as a result of the mobility training program?

- Is there anything else you would like to share as part of this interview?
### Appendix D: Description of included studies

<table>
<thead>
<tr>
<th>Authors, year of publication</th>
<th>Design and sample</th>
<th>Intervention details</th>
<th>Outcome measures</th>
<th>Results</th>
<th>Methodological quality</th>
</tr>
</thead>
</table>
| Butler, Okamoto, McKay (1983) | Single case design (n = 9) children | - Children were provided with powered wheelchairs and learned to drive at home under parent supervision.  
- Intervention duration = 6 weeks | Driving skills  
- Seven skill driving checklist  
- Video recordings  
- Parent's perception  
- Parent interview | Driving skills  
8/9 children gained independent use of their power wheelchair (PWC).  
Parent's perception  
Parents noticed a positive effect on their child's social, emotional and intellectual behaviours, and an increased motivation in other forms of movement. | Limited quality (score = 6/22, 27%)  
Level IV evidence  
Conclusion was limited because of small sample size, no rigorous randomisation, and no blinding. |
| Butler, Okamoto, McKay (1984) | Single case design (n = 13) children | - Children were provided with PWCs and learned to drive at home under parent supervision.  
- Intervention duration = 1 month | Driving skills  
- Parents recorded their child's experience with the PWC including hours driving, time spent in the PWC and behaviours.  
- Seven skill driving checklist | Driving skills  
12/13 children learnt to competently drive a PWC in an average of 16.3 days with a median of 14 days.  
Children learnt 4-5 driving skills within a 1-5 day period.  
Cumulative time of wheelchair driving averaged 8.1 hours with a median of 6.0 hours (range 1.7-26.1 hours). The cumulative time spent sitting in the wheelchair averaged 34.4 hours with a median of 20 hours (range 6.6-168 hours). | Adequate quality (score = 11/22, 50%)  
Level IV evidence  
Conclusion was limited because of small sample size, no rigorous randomisation, and no blinding. |
| Butler, (1986) | Single subject time series design with multiple baselines. (n = 6) children | - Each child was provided with a PWC and learnt to use it to a level of proficiency (defined by 7 a priori criteria).  
- Intervention duration = 3 weeks. | Self initiated interaction with increased change in location.  
- Video footage  
Self initiated interaction with objects  
- Video footage  
Self initiated interaction in communication  
- Video footage | Self initiated interaction with increased change in location  
Clinically and statistically significant for all participants. T= 3.57, (p<0.05)  
Self initiated interaction with objects  
3 children increased in their self initiated interaction with objects. No consistent changes were recorded for the group. Variability in changes meant the results were not statistically reliable. T= 0.86, (p<0.21)  
Self initiated interaction in communication  
3 children increased in self initiated interaction in communication. No consistent changes were recorded for the group. Variability in changes means the results were not statistically reliable. T= 0.80, (p<0.22) | Adequate quality (score = 12/22, 54.5%)  
Level IV evidence  
Conclusion was limited because of small sample size, no rigorous randomisation, and no blinding. |
- 6-8 month treatment | Impairment  
- Performance IQ score of the Leiter International | Impairment  
Relationship between driving skills and Performance IQ score of the Leiter International Performance Scale or the Verbal IQ score of the | Strong quality (score = 19/22, 86%)  
Level IV evidence |
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>• Data for achievement/ non achievement of driving competence (n=27)</td>
<td></td>
</tr>
<tr>
<td>• Data for influence of PWC provision on outcome indexes (n=25)</td>
<td></td>
</tr>
<tr>
<td>period (PWC provision)</td>
<td>Performance Scale.</td>
</tr>
<tr>
<td></td>
<td>• Verbal IQ score of the Peabody Developmental Verbal Scale.</td>
</tr>
<tr>
<td></td>
<td><strong>Functional Limitation/Activity</strong></td>
</tr>
<tr>
<td></td>
<td>• Furumasu’s Driving Test</td>
</tr>
<tr>
<td></td>
<td>• Gross Motor Functional Measure</td>
</tr>
<tr>
<td></td>
<td>• Canadian Occupational Performance Measure</td>
</tr>
<tr>
<td></td>
<td><strong>Disability/Participation</strong></td>
</tr>
<tr>
<td></td>
<td>• Impact of Childhood Illness Scale</td>
</tr>
<tr>
<td></td>
<td><strong>Parental perceptions and acceptance</strong></td>
</tr>
<tr>
<td></td>
<td>• Parent interview</td>
</tr>
<tr>
<td>Peabody Developmental Verbal Scale was not statistically significant. <strong>Functional Limitation/Activity</strong> Significant change (p&lt;0.01), 21/27 children were competent driving a PWC. Relationship between Gross Motor and Functional Measure and driving skills was not statistically significant. Significant change between 2&lt;sup&gt;rd&lt;/sup&gt; and 3&lt;sup&gt;rd&lt;/sup&gt; assessment scores on the Canadian Occupational Performance measure (p&lt;0.0001). <strong>Disability/Participation</strong> No significant change (p= 0.027) on the Impact of Childhood Illness Scale. <strong>Parental perceptions and acceptance</strong> 21/25 parents were not in favour of PWC provision when the study started. After the study 23/25 parents expressed positive feelings about PWC provision. 25/25 parents reported increase independence in their child. 23/25 children expressed positive reactions from the beginning of the study, and did not change following the intervention.</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>• Powered mobility training sessions on a powered riding toy in child’s school environment. Intervention conducted by an occupational therapist and a researcher.</td>
<td></td>
</tr>
<tr>
<td>• Intervention duration = 14 weeks</td>
<td></td>
</tr>
<tr>
<td><strong>Self initiative movement</strong></td>
<td></td>
</tr>
<tr>
<td>• Video footage</td>
<td></td>
</tr>
<tr>
<td>• Manual counter (used on one participant only).</td>
<td></td>
</tr>
<tr>
<td><strong>Contact with others</strong></td>
<td></td>
</tr>
<tr>
<td>• Partial interval video recording.</td>
<td></td>
</tr>
<tr>
<td><strong>Affect</strong></td>
<td></td>
</tr>
<tr>
<td>• Momentary time sampling using video footage.</td>
<td></td>
</tr>
<tr>
<td><strong>Practicality of the powered riding toy</strong></td>
<td></td>
</tr>
<tr>
<td>• Researchers experiences and interviews with</td>
<td></td>
</tr>
<tr>
<td>• Self initiative movement Increase in number of self initiated movements. Participant 1 increased from 44 (baseline stage) to 87 (intervention stage) and participant 2 increased from 28 (baseline stage) to 65 (intervention stage). <strong>Contact with others</strong> Children made more positive contact with participant 2 during the intervention stage, and less positive contact with participant 1 during the intervention stage. Adults initiated more negative contact with participant 2 during the intervention stage and more positive contacts with participant 1 during intervention stage. Participant 2 had less positive initiations with</td>
<td></td>
</tr>
</tbody>
</table>

Conclusion was limited because of small sample size, no rigorous randomisation, and no blinding. Adequate quality (score = 16/24, 66 %) Level IV evidence Conclusion was limited because of small sample size, no rigorous randomisation, and no blinding.
school staff and one parent. adults during intervention stage and participant 1 had more positive initiations with others during intervention stage.

**Affect**
No clear impact on the amount of positive or negative affect could be determined.

**Practicality of the powered riding toy**
Looks like a toy, gains children's interest, easy to adapt, low cost, useful evaluation and early training tool and adjustable speed are all positive aspects.

Negative aspects include too loud, challenged by rough surfaces, impractical in small areas, and breaks down.

<table>
<thead>
<tr>
<th>Case study design</th>
<th>Driving skills</th>
<th>Participant behaviour</th>
<th>Abbreviations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hasdai, Jessel, Weiss, (1998)</td>
<td>Adequate quality (score = 15/26, 57.6%)</td>
<td>Level III evidence</td>
<td>PWC = power wheelchair</td>
</tr>
<tr>
<td>Nilsson, Nyberg, (2003)</td>
<td>Significant improvement in driving performance for intervention group (p&lt;0.01)</td>
<td>Level IV evidence</td>
<td></td>
</tr>
<tr>
<td>Cohort design (n = 22) children and young adults</td>
<td>Participants were divided into two groups, those with experience driving a PWC and those with no experience.</td>
<td>Strong quality (score = 17/20, 85%)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>After a driving assessment with a PWC the inexperienced drivers were trained on a joystick-controlled computer game in which they navigated mazes.</td>
<td>Conclusion was limited because of small sample size.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Both groups driving skills were evaluated using a PWC.</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Driving skills</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>- Functional evaluation rating scale</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>- Simulation program</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Participant behaviour</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>- Video recordings</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>- Field notes</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>- In depth parent interviews</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Participant behaviour</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Increased ability to react to external stimuli. Increased development of initiative and exploratory behaviours. Increase in awareness and alertness. Encouragement of simple cause and effect relationships. Increased use of child's arms and hands.</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Driving skills</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Significant improvement in driving performance for intervention group (p&lt;0.01)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Scores of the inexperienced PWC users were lower than the experienced PWC users (p&lt;0.01)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Driving skills</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Significant improvement in driving performance for intervention group (p&lt;0.01)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Scores of the inexperienced PWC users were lower than the experienced PWC users (p&lt;0.01)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>Driving skills</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Significant improvement in driving performance for intervention group (p&lt;0.01)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Scores of the inexperienced PWC users were lower than the experienced PWC users (p&lt;0.01)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Guidelines for contributions by authors

Disability and Rehabilitation: Assistive Technology is an international, multidisciplinary, peer-reviewed journal devoted specifically to the broad range of technological developments and related supports and issues which enhance the rehabilitation process. The journal seeks to encourage a better understanding of all aspects of disability, and to promote the rehabilitation process. New submissions on any aspect of technology, disability and rehabilitation are encouraged and the journal welcomes contributions from a wide range of professional groups, including medical practitioners, occupational therapists, physiotherapists, speech and language therapists, clinical psychologists and those involved in nursing, education, ergonomics, and engineering.

Disability and Rehabilitation: Assistive Technology is organised into sections: Literature Reviews; Research Papers, Case Studies, Clinical Commentaries; reports on Rehabilitation in Practice, Products and Devices/Rehabilitation Engineering, Occasional Special Issues and specific sections on contemporary themes of interest to the Journal's readership are published.

Disability and Rehabilitation: Assistive Technology will be of interest to a wide range of professional groups, including medical practitioners, occupational therapists, physiotherapists, speech and language therapists, clinical psychologists and those involved in nursing, education, ergonomics and engineering.

Editor

Marcia J. Scherer, PhD, MPH, FACRM
Institute for Matching Person & Technology
486 Lake Road
Webster, NY 14580 USA
585-671-3461 (phone/fax)
Email: IMPT97@aol.com

Submissions

All submissions should be made online at Disability and Rehabilitation: Assistive Technology's Manuscript Central site. New users should first create an account. Once a user is logged onto the site submissions should be made via the Author Centre.

Papers should be submitted with any tables, figures, or photographs, all of which should be of high quality suitable for reproduction. Submissions should be in English in double line spacing.

The submission should include a separate title page with the name(s) and affiliation(s) of the author(s) and the name and address for offprint requests with a telephone, fax number (including country and area codes), and electronic mail address.
Submissions should be accompanied by a covering letter stating that all authors have read the paper and agree to its submission and should include where appropriate a formal statement that ethical consent for the work to be carried out has been given. Photographs of individuals should be avoided, but if essential individual consent in writing must accompany the manuscript. It is not sufficient to mask identity by covering the person's eyes.

Refereeing will be carried out anonymously unless otherwise requested, and manuscripts should be prepared accordingly.

There are no page charges for papers published in the journal. Three complimentary copies of the issue in which your article appears will be sent to the principal or sole author or articles. Larger quantities may be ordered at a special discount price. An order form will accompany the proof which must be completed and returned, irrespective of whether you require additional copies.

**Writing a paper for Disability and Rehabilitation: Assistive Technology**

For all manuscripts, non-discriminatory and person-first language is mandatory. Sexist or racist terms should not be used.

Structured abstracts of around 200 words are required for all papers submitted and should precede the text of a paper. There are no lower or upper word limits for papers submitted to the Journal.

Manuscripts should be typed double-spaced throughout, including the reference section.

Submitted manuscripts must cite relevant original sources, not only review articles or articles published subsequently by the same or different author(s).

In writing your paper, you are encouraged to review articles in the area you are addressing which have been previously published in the journal, and where you feel appropriate, to reference them. This will enhance context, coherence, and continuity for our readers.

**Structure of Paper**

An introductory section should state the purpose of the paper and give a brief account of previous work. New techniques and modifications should be described concisely but in sufficient detail to permit their evaluation; standard methods should simply be referenced. Experimental results should be presented in the most appropriate form, with sufficient explanation to assist their interpretation; their discussion should form a distinct section. Extensive tabulations will not be accepted unless their inclusion is essential.

**Abstracts**
Structured abstracts are required for all papers, and should be submitted as detailed below, following the title and author's name and address, preceding the main text.

Intending contributors should note that *Disability and Rehabilitation: Assistive Technology* is changing the format of the summary of each paper to that of a structured abstract. The change will operate with immediate effect. There is clear evidence that structured abstracts contain more accessible information than summaries and are therefore of more use to the readership.

All papers submitted to the Journal of *Disability and Rehabilitation: Assistive Technology* should have a 'structured abstract' of no more than 200 words. The following headings should be used, following the title, author's name and address, and preceding the main text:

**Purpose** State the main aims and objectives of the paper.

**Method** Describe the design, and methodological procedures adopted.

**Results** Present the main results.

**Conclusions** State the conclusions that have been drawn and their relevance to the study of disability and rehabilitation.

**Nomenclature and Units**

All abbreviations and units should conform to SI practice. Drugs should be referred to by generic names; trade names of substances, their sources, and details of manufacturers of scientific instruments should be given only if the information is important to the evaluation of the experimental data.

**Copyright Permission**

Contributors are required to secure permission for the reproduction of any figure, table, or extensive (more than fifty word) extract from the text, from a source which is copyrighted - or owned - by a party other than Informa UK Ltd or the contributor.

This applies both to direct reproduction or 'derivative reproduction' - when the contributor has created a new figure or table which derives substantially from a copyrighted source.

The following form of words can be used in seeking permission:

Dear [COPYRIGHT HOLDER]

I/we are preparing for publication an article entitled

[STATE TITLE]
to be published by Informa UK Ltd in Disability and Rehabilitation: Assistive Technology.

I/we should be grateful if you would grant us permission to include the following materials:

[STATE FIGURE NUMBER AND ORIGINAL SOURCE]

We are requesting non-exclusive rights in this edition and in all forms. It is understood, of course, that full acknowledgement will be given to the source.

Please note that Informa UK Ltd are signatories of and respect the spirit of the STM Agreement regarding the free sharing and dissemination of scholarly information.

Your prompt consideration of this request would be greatly appreciated.

Yours faithfully

Code of Experimental Ethics and Practice

Contributors are required to follow the procedures in force in their countries which govern the ethics of work done with human or animal subjects. The Code of Ethics of the World Medical Association (Declaration of Helsinki) represents a minimal requirement.

When experimental animals are used, state the species, strain, number used, and other pertinent descriptive characteristics.

For human subjects or patients, describe their characteristics.

For human participants in a research survey, secure the consent for data and other material - verbatim quotations from interviews, etc. - to be used.

When describing surgical procedures on animals, identify the pre-anaesthetic and anaesthetic agents used and state the amount of concentration and the route and frequency of administration for each. The use of paralytic agents, such as curare or succinylcholine, is not an acceptable substitute for anaesthetics. For other invasive procedures on animals, report the analgesic or tranquilizing drugs used; if none were used, provide justification for such exclusion.

When reporting studies on unanaesthetized animals or on humans, indicate that the procedures followed were in accordance with institutional guidelines.

Specific permission for facial photographs of patients is required. A letter of consent must accompany the photographs of patients in which a possibility of identification exists. It is not sufficient to cover the eyes to mask identity.
Clinical Trials Registry

Disability and Rehabilitation: Assistive Technology requests, as a consideration of publication, that clinical trials are registered in a public repository at their inception and prior to patient enrolment.

The registry must be accessible to the public at no charge, be open to all prospective registrants and managed by a not-for-profit organization. For a list of registries that meet all of these requirements, please see the WHO International Clinical Trials Registry Platform (ICTRP) http://www.who.int/ictrp/en/. This is in accordance with the guidelines published by the International Committee of Medical Journal Editors (ICMJE). For more information, see ICMJE Uniform Requirements for Manuscripts Submitted to Biomedical Journals http://www.icmje.org

The registration of all clinical trials facilitates the dissemination of information among clinicians, researchers and patients, and enhances public confidence in the research enterprise.

Offprints and Reprints

Offprints and reprints of articles published in this journal can be purchased once the article has been published online. Corresponding authors will receive free online access to their article through our website (www.informaworld.com). Reprints of articles published in this journal can be purchased through the Publisher when proofs are received. Copies of the Journal can be purchased separately at the author's preferential rate of £15.00/$25.00 per copy.

Colour Figures

a. Any figure submitted as a colour original will appear in colour in the journal's online edition free of charge and can be downloaded.

b. Paper copy colour reproduction will only be considered on condition that authors contribute to the associated costs. Charges are: £500/US$1030 for the first colour page and £250/US$515 for each colour page after per article. (Colour costs will be waived for invited Review Articles)

Electronic Processing

The following procedures which will assure we can process your article efficiently.

1. Authors using standard word-processing software packages

For the main text of your article, most standard PC or Mac word-processing software packages are acceptable, although we prefer Microsoft Word in a PC format.

- Word-processed files should be prepared according to the journal style.
Avoid the use of embedded footnotes. For numbered tables, use the table function provided with the word-processing package.

All text should be saved in one file with the complete text (including the title page, abstract, all sections of the body of the paper, references), followed by numbered tables and the figure captions.

2. Authors using LaTeX mathematical software packages

Authors who wish to prepare their articles using the LaTeX document preparation system are advised to use article.sty (for LaTeX 2.09) or article.cls (for LaTeX2e).

The use of macros should be kept to an absolute minimum but if any are used they should be gathered together in the file, just before the \begin{document} command.

Articles prepared using LaTeX should be converted to PDF documents (incorporating the illustrations) and these should be submitted online in addition to the associated LaTeX source and graphics files.

The files you send must be text-only (often called an ASCII file), with no system-dependent control codes.

3. A guide for authors using graphics software packages

We welcome figures, but care and attention to these guidelines is essential, as importing graphics packages can often be problematic.

Figures must be saved separately from the text.

Avoid the use of colour and tints for aesthetic reasons. Figures should be produced as near to the finished size as possible.

All figures must be numbered in the order in which they occur (e.g. figure 1, figure 2 etc.). In multi-part figures, each part should be labelled (e.g. figure 1 (a), figure 1 (b) etc.)

The figure captions must be saved as a separate file with the text and numbered correspondingly.

The filename for the graphic should be descriptive of the graphic e.g. Figure1, Figure2a.

Files should be saved as TIFF (tagged image file format), PostScript or EPS (encapsulated PostScript), containing all the necessary font information and the source file of the application (e.g., CorelDraw/Mac, CorelDraw/PC).
Notes on Style

All authors are asked to take account of the diverse audience of the journal. Clearly explain, or avoid the use of, terms that might be meaningful only to a local or national audience.

Some specific points of style for the text of articles, research reports, case studies, reports, essay reviews, and reviews follow:

1. We prefer US to 'American', USA to 'United States', and UK to 'United Kingdom'.
2. We use conservative (British, not US, spelling, i.e. colour not color; behaviour (behavioural) not behavior; [school] programme not program; [he] practises not practices; centre not center; organization not organisation; analyse not analyze, etc.
3. Single 'quotes' are used for quotations rather than double "quotes", unless the 'quote is "within" another quote'.
4. Punctuation should follow the British style, e.g. 'quotes precede punctuation'.
5. Punctuation of common abbreviations should follow the following conventions: e.g. i.e. cf. Note that such abbreviations are not followed by a comma or a (double) point/period.
6. Dashes (M-dash) should be clearly indicated in manuscripts by way of either a clear dash (-) or a double hyphen (--).
7. We are sparing in our use of the upper case in headings and references, e.g. only the first word in paper titles and all subheads is in upper case; titles of papers from journals in the references and other places are not in upper case.
8. Apostrophes should be used sparingly. Thus, decades should be referred to as follows: 'The 1980s [not the 1980's] saw ...'. Possessives associated with acronyms (e.g. PA), should be written as follows: 'The APU's findings that ...', but, NB, the plural is APUs.
9. All acronyms for national agencies, examinations, etc., should be spelled out the first time they are introduced in text or references. Thereafter the acronym can be used if appropriate, e.g. 'The work of the Assessment of Performance Unit (APU) in the early 1980s ...'. Subsequently, 'The APU studies of achievement ...', in a reference ...
10. Brief biographical details of significant national figures should be outlined in the text unless it is quite clear that the person concerned would be known internationally. Some suggested editorial emendations to a 'typical' text are indicated in the following with square brackets: 'From the time of H. E. Armstrong [in the 19th century] to the curriculum development work associated with the Nuffield Foundation [in the 1960s], there has been a shift from heurism to constructivism in the design of [British] science courses'.
11. The preferred local (national) usage for ethnic and other minorities should be used in all papers. For the USA, 'African-American', 'Hispanic' and 'Native American' are used, e.g. 'The African American presidential candidate, Jesse Jackson...'; for the UK, 'Afro-Caribbean' (not 'West Indian'), etc.
12. Material to be emphasized (italicized in the printed version) should be underlined in the typescript rather than italicized. Please use such emphasis sparingly.
Mathematics

Special care should be taken with mathematical scripts, especially subscripts and superscripts and differentiation between the letter 'ell' and the figure one, and the letter 'oh' and the figure zero. If your keyboard does not have the characters you need, it is preferable to use longhand, in which case it is important to differentiate between capital and small letters, K, k and x and other similar groups of letters. Special symbols should be highlighted in the text and explained in the margin. In some cases it is helpful to supply annotated lists of symbols for the guidance of the sub-editor and the typesetter, and/or a 'Nomenclature' section preceding the 'Introduction'.

For simple fractions in the text, the solidus / should be used instead of a horizontal line, care being taken to insert parentheses where necessary to avoid ambiguity, for example, $\frac{1}{(n-1)}$. Exceptions are the proper fractions available as single type on a keyboard.

Full formulae or equations should be displayed, that is, written on a separate line. Horizontal lines are preferable to solidi, for example:

$$\frac{61 + 5h + g}{3n + 3y^2}$$

But: $a/b + c/d + a/d$

$$P = (a^2 - b^2)(c^2 + d^2)$$

The solidus is not generally used for units: ms$^{-1}$ not m/s, but note electrons/s, counts/channel, etc.

Displayed equations referred to in the text should be numbered serially (1, 2, etc.) on the right hand side of the page. Short expressions not referred to by any number will usually be incorporated in the text.

Symbols should not be underlined to indicate fonts except for tensors, vectors and matrices, which are indicated with a wavy line in the manuscript (not with a straight arrow or arrow above) and rendered in heavy type in print: upright sans serif r (tensor), sloping serif r (vector) upright serif r (matrix).

Typographical requirements must be clearly indicated at their first occurrence, e.g. Greek, Roman, script, sans serif, bold, italic. Authors will be charged for corrections at proof stage resulting from a failure to do so.

Braces, brackets and parentheses are used in the order \{ ( ) \}, except where mathematical convention dictates otherwise (i.e. square brackets for commutators and anticommutators)

Citations in Text
We prefer that references are cited using the numerical system (e.g. [3], [5-9]). They should be listed separately at the end of the paper in the order in which they appear in the text.

Notes on Tables and Figures

1. Tables and figures should be valuable, relevant, and visually attractive. Tables and figures must be referred to in the text and numbered in order of their appearance. Each table and figure should have a complete, descriptive title; and each table column an appropriate heading.

Tables and figures should be referred to in text as follows: figure 1, table 1, i.e. lower case. 'As seen in table [or figure] 1 ...' (not Tab., fig. or Fig).

2. The place at which a table or figure is to be inserted in the printed text should be indicated clearly on a manuscript:

   [Insert table 2 about here ]

3. Each table and/or figure must have a title that explains its purpose without reference to the text.

4. All figures and tables must be on separate sheets and not embedded in the text. Original copies of figures should be supplied. All figures should allow for reduction to column width (130 mm) or page width (160mm). Please avoid figures that would require landscape reproduction, i.e., reading from bottom to top of the page. Photographs may be sent as glossy prints or negatives.

Do not type the caption to a figure on that figure; the legends to any illustrations must be typed separately following the main text and should be grouped together.

Acknowledgements

Any acknowledgements authors wish to make should be included in a separate headed section at the end of the manuscript. Please do not incorporate these into the bionote or notes.

Declaration of interest

It is the policy of all Informa Healthcare to adhere in principle to the Conflict of Interest policy recommended by the International Committee of Medical Journal Editors (ICMJE, http://www.icmje.org/index.html#conflict).

All authors must disclose any financial and personal relationships with other people or organisations that could inappropriately influence (bias) their work. It is the sole
responsibility of authors to disclose any affiliation with any organisation with a financial interest, direct or indirect, in the subject matter or materials discussed in the manuscript (such as consultancies, employment, paid expert testimony, honoraria, speakers' bureaus, retainers, stock options or ownership, patents or patent applications or travel grants) that may affect the conduct or reporting of the work submitted. All sources of funding for research are to be explicitly stated. If uncertain as to what might be considered a potential conflict of interest, authors should err on the side of full disclosure.

All submissions to the journal must include full disclosure of all relationships that could be viewed as presenting a potential conflict of interest. If there are no conflicts of interest, authors should state that there are none. This must be stated at the point of submission (within the manuscript after the main text under a subheading "Declaration of interest" and, where available, within the appropriate field on the journal's Manuscript Central site). This may be made available to reviewers and will appear in the published article at the discretion of the Editors or Publisher.

If no conflict is declared, the following statement will be attached to all articles:

**Declaration of interest:** The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

The intent of this policy is not to prevent authors with these relationships from publishing work, but rather to adopt transparency such that readers can make objective judgements on conclusions drawn.

**Plagiarism**

Informa Healthcare has a strict policy against plagiarism. We define plagiarism as the use of extracts from another person's work that are not placed in quotation marks, without the permission of that person, and without acknowledgement to that person (using the appropriate reference style), with the result that your article presents these extracts as original to you. By submitting your work to an Informa Healthcare journal, you warrant that it is your original work, and that you have secured the necessary written permission from the appropriate copyright owner or authority for the reproduction of any text, illustration, or other material.

If any article submitted to an Informa Healthcare journal is found to have breached any of these conditions, Informa Healthcare reserves the right to reject that article and any others submitted by the same authors. Informa Healthcare may also contact the authors' affiliated institutions to inform them of its findings.

**References**

References should follow the CBE Citation & Sequence format. Only works actually cited in the text should be included in the references. Indicate in the text with Arabic numbers inside square brackets. Spelling in the reference list should follow the original.
References should then be listed in numerical order at the end of the article. Examples are provided as follows:

Journal article:

Book chapter:

Conference proceedings:

Dissertations or Thesis:

Journal article on internet:

Webpage:

Internet databases:

Further examples and information can be found in the CBE style manual Scientific Style and Format, sixth edition.

NIH Public Access Policy

In consideration of the National Institutes of Health (NIH) Public Access Policy, Informa Healthcare acknowledges that the broad and open dissemination of NIH-funded-research
results may benefit future scientific and medical research. Because we value the current and future contributions our journals make to the scientific body of knowledge, we have made certain that our policies accommodate those authors who wish to submit to PubMed Central.

Informa Healthcare's position with respect to public access to NIH-funded work published in Informa Healthcare journals is as follows:

- Informa Healthcare authors may voluntarily submit their funded work to PubMed Central after a 12-month embargo period;
- "funded work" shall be defined as the final, peer-reviewed manuscript that is accepted by the Editor in Chief of the journal. This manuscript must not be altered by Publisher’s copyediting and typesetting services; and
- this embargo period begins the day the work is published online at www.informaworld.com.