

Children's and adolescents' views on upper limb prostheses in relation to their daily occupations.

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Abstract:	Prosthetic treatment is widely used for children and adolescents with upper limb (UL) difference, for functional benefit, cosmetic restoration or performance of specific activities. Children and adolescents feel, however, that prostheses do not help function, are uncomfortable, are unreliable, are heavy and are not aesthetically acceptable. This research aims to develop an understanding of children's and adolescents' views on UL prostheses in relation to their daily occupations. Eight children and adolescents with UL difference participated in a focus group (n=4) or an individual interview (n=4). Data was analysed using thematic analysis. Participants described both positive and negative views of their prostheses. Participants described experiences where their prosthesis was helpful to them for social interactions, when playing and in self-care and productive activities. However, participants also described choosing not to use prostheses due to lack of functional benefit, causing a hindrance in certain activities and restriction of movement. Prostheses can be useful tools for occupational participation for children. General purpose functional prostheses for children can result in the prosthesis not being useful, becoming a hindrance and restricting movement. Task specific devices can overcome these issues to help children in social interactions, when playing and in carrying out self-care and productive activities.

SCHOLARONE™ Manuscripts Children's and adolescents' views on upper limb prostheses in relation to their daily occupations.

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Abstract

Introduction

Prosthetic treatment is widely used for children and adolescents with upper limb (UL) difference, for functional benefit, cosmetic restoration or performance of specific activities. Children and adolescents feel, however, that prostheses do not help function, are uncomfortable, are unreliable, are heavy and are not aesthetically acceptable. This research aims to develop an understanding of children's and adolescents' views on UL prostheses in relation to their daily occupations.

Method

Eight children and adolescents with UL difference participated in a focus group (n=4) or an individual interview (n=4). Data was analysed using thematic analysis.

Results

Participants described both positive and negative views of their prostheses. Participants described experiences where their prosthesis was helpful to them for social interactions, when

playing and in self-care and productive activities. However, participants also described choosing not to use prostheses due to lack of functional benefit, causing a hindrance in certain activities and restriction of movement.

Conclusion

Prostheses can be useful tools for occupational participation for children and adolescents.

General purpose functional prostheses for children and adolescents can result in the prosthesis not being useful, becoming a hindrance and restricting movement. Task specific devices can overcome these issues to help children and adolescents in social interactions, when playing and in carrying out self-care and productive activities.

Keywords: Paediatrics, Upper Limb, Prosthetics, Qualitative Research

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Introduction

Upper limb difference can have both a physical and psychological effect on a child (Smith, 2006). Physical impacts may include asymmetrical development of the spine, postural problems, back pain (Broomfield 2009) and overuse syndrome from increased workload for the remaining arm and compensatory movement strategies (Peterson 2011).

Developmentally, children may struggle with playing with objects in midline, sitting balance, crawling, walking and bimanual skills (Ibbotson 2002). Psychosocial impacts may include lower quality of social activities than peers (Michielsen et al 2011), greater behavioural and emotional problems, lower social competence and challenges to developing positive body image (Winchell 1996). Occupational therapy aims to limit the negative physical and psychological effects of upper limb difference for children and adolescents and promote

active and independent participation in daily occupations. Occupational therapy can include

adaptation of activities in the areas of self-care, work and leisure (Dillingham et al 2002), such as how to carry out activities using one-hand or with feet (Hanspal 2003), and the provision of upper limb prostheses for functional benefit, cosmetic restoration or performance of specific activities.

Non-use rates of upper limb prostheses amongst children vary widely throughout the literature but they have, in some studies globally, been reported to be as high as 50% (Shida-Tokeshi et al., 2005). Studies relating to children's general satisfaction with prostheses have found that prostheses do not help children's function, are uncomfortable, unreliable, too heavy and not aesthetically acceptable (Biddiss and Chau, 2007; Pylatiuk et al., 2007; Routhier et al., 2001; Wagner et al., 2007). Exploring and understanding children's and adolescents' views about, and use of, prostheses within the context of their daily occupations is needed to guide occupational therapy practice and ensure that provision of prostheses is an effective and positive intervention for children with upper limb difference.

Previous research has used parent proxy reports or quantitative methods to explore the views of children. A drawback of parent proxy reports is that they rely on adults having the ability to know what children think and feel, which is an outmoded concept (O'Kane 2008). Additionally, the reality of how disability and technology affect an individual may be subjective in nature. Sheffler et al (2009) compared self-report with parent proxy report of function and quality of life amongst children with limb difference and found that parents underestimated their children's physical and social function and overestimated their comfort.

Qualitative research that directly addresses children's experiences and viewpoints is needed.

Furthermore, from an ethical perspective, Article 12 of the United Nations Convention on the

Rights of the Child (1990) states that children have the right to say what they think about anything that affects them and that what they say must be listened to carefully. Using qualitative methods ensures that the factors and areas that are important to the participant are highlighted to the researcher, allowing participants to speak in their own voice, rather than being confined to categories imposed on them by others (Mack et al 2005).

This study aims to explore children's and adolescents' views on UL prostheses to develop an understanding of prostheses in relation to children's daily occupations.

Literature Review

Prosthetic treatment is widely used for children with upper limb difference, leading to prescription of prostheses for functional benefit, cosmetic restoration or for performance of particular activities. In current practice, there are three main types of prosthesis that may be prescribed for children - passive prostheses, body-powered prostheses and myoelectric prostheses (Broomfield, 2009).

A passive prosthesis is designed to look as natural and aesthetically pleasing as possible (Broomfield, 2009) and is shaped and coloured to match the unaffected arm (Watve et al., 2011). A passive prosthesis is usually the first prosthesis a child will be prescribed with (Watve et al., 2011). Passive prostheses are lightweight and do not usually contain moving parts (Broomfield, 2009). Despite the drawback of a lack of movement, a passive prosthesis can still be useful. The child can support objects with the prosthetic hand (Broomfield, 2009) and it may also perform a passive assistive role to the unaffected arm, such as steadying a piece of paper when writing (Watve et al., 2011).

A body-powered prosthesis, such as a voluntary opening/closing split hook or hand, contains moving parts and is, therefore, more practical for use in activities of daily living (ADLs) than a passive prosthesis (Broomfield, 2009). A body-powered prosthesis is an arm that utilises a harness and cabling to control a hand (also known as a terminal device) and an elbow unit (Muzumdar, 2004). The harness system is controlled by gross body movements (such as scapular abduction) (Muzumdar, 2004). It is simple in design, making it highly durable and suitable for use in tasks that involve water, dust, and other conditions that could damage an electric prosthesis (Muzumdar, 2004). Some users of body-powered prostheses report that the cable tension simulates proprioception, enabling enhanced spatial awareness and allowing the wearer improved feedback on the position of the prosthesis (Muzumdar, 2004). Body-powered prostheses are heavier than passive prostheses, which may cause physical strain to be put on a child's body, resulting in discomfort (Biddiss and Andrysek, 2011).

Myoelectric prostheses offer the advantage of combining looks and functionality as they can have cosmeses, which have the appearance of a real hand, but are more useful than passive prostheses when it comes to performing ADLs (Egermann et al., 2009). A myoelectric prosthesis uses electromyographic signals from voluntarily contracted muscles within a person's residual limb (Watve et al., 2011). These signals are detected by sensors on the surface of the skin and control the movements of the prosthesis. Myoelectric prostheses tend to be heavier than other prosthetic options and are susceptible to damage when exposed to moisture (Muzumdar, 2004).

In addition to commercially available prostheses for children and adolescents, there may also be a need for custom-made task specific devices (Broomfield, 2009). Much research has suggested that task specific devices are most useful for children as they can be used for

specific activities, making them most appropriate for the desired use (Buffart et al., 2007; Egermann et al., 2009; James et al., 2006). This has been evidenced in the recent provision by the UK Department of Health (DoH) (2017) for the funding of sports prostheses for children in England who have experienced limb loss or who were born with a limb difference, to enable them to engage in physical activity and sports.

Different types of prostheses offer different levels of functional and cosmetic benefit to their users. For over 30 years it has been recognised that a child is likely to go through different stages of prosthesis use, just as they will go through different developmental stages (Biddiss, 2010; Celikyol, 1984; National Academies of Sciences, Engineering, and Medicine 2017; Patton, 2004; Sims, 2018). It is suggested that children of primary school age, having previously been accepting of prostheses, may opt not to use them as they may become a hindrance during this very physically active developmental stage (Patton, 2004). During adolescence, however, both aesthetics and functionality may become more important to the young person as appearance becomes a more valued attribute and leisure or work activities become more demanding bimanually, requiring fine prehension (Patton, 2004).

Clinical decisions regarding which type of prosthesis is most suitable for a child are likely to be guided by the child's age and proposed use for the device, as well as the cost of the device and the funds available, leading to variations in treatment (Broomfield, 2009). Furthermore, since Celikyol's (1984) guidance was published, huge technological advances have occurred both in prosthesis development and in the environments with which children interact.

Current research to explore children's and adolescents' views about, and use of, upper limb prostheses in daily occupations is needed to provide additional evidence to guide

occupational therapy practice. This research aims to develop an understanding of children's and adolescents' views on upper limb prostheses in relation to their daily activities.

Method

Ethical approval for the study was granted by the University of Southampton's Faculty of Health Sciences' Ethics Committee in June 2011 (FoHS-ETHICS-2011-056).

Methodology

This study formed part of a larger programme of research which used an emancipatory lens (Mertens, 2015) to carry out the BRuger Involvering i Design, GEntænkt (BRIDGE) (English: User Involvement in Design, Revised) (Iversen and Brodersen, 2008) method of participatory design to develop new upper limb prostheses with children and adolescents, resulting in the production of several prototypes of cost-effective task-specific prosthetic devices (Sims et al, 2017).

Participants

Eight children and adolescents with below elbow upper limb difference participated in the study. They were recruited through a national charitable organisation (Reach Charity Ltd: http://reach.org.uk/). The charity sent participant information packs to their members and three months later followed up with an email reminder about the study. The packs contained: an introductory letter, two participant information sheets (one for parents and older children and a separate, simpler information sheet for children aged 7-11 years) and a reply slip which included some basic screening information about the respondents. The parents of eight participants contacted the researcher after receiving the information pack or reminder email and their children were recruited to participate in the study.

The eight participants of this study constitutes a subsample of the 34 participants (eight children/adolescents, nine parents, eight prosthetists and nine occupational therapists) from the larger programme of research, who contributed to the development of new devices (for further information, see Sims et al, 2017). Table 1 provides an outline of the demographic characteristics of the participants. Written informed consent was given by parents of participants, and participants provided written informed assent.

[Insert table 1 here]

Pseudonyms are used throughout the results to protect anonymity, but actual ages are provided.

Design

Participants were offered the choice of participating in a focus group or individual interview. The research combined focus groups and interviews for both pragmatic and confirmatory reasons. The focus group and interviews used the same questioning guide and included questions about what the participants liked or did not like about their prostheses, the activities they could or could not do with their prostheses and what their ideal prostheses would be like. Areas for discussion were based on previous work by Donovan-Hall (2010). See appendix for interview and focus group guide.

Four participants chose to take part in a focus group, lasting approximately 90 minutes and four participants chose to participate in an individual interview lasting approximately 60 minutes. The focus group was carried out in a central London location and interviews were

carried out in participants' homes. No financial incentive was offered for participation, but participants were reimbursed expenses and provided with a certificate of participation.

Data was transcribed verbatim and analysed using thematic analysis (Braun and Clarke, 2006). The stages of thematic analysis, as outlined by Braun and Clarke (2006), were followed: transcription and immersion; generating initial codes; searching for themes; reviewing themes; and defining and naming themes. Analysis was conducted on each transcript individually before combining to develop the themes.

Trustworthiness and credibility

Combining two data collection methods (focus groups and interviews) enhances credibility of the findings through triangulation of methods (Guba and Lincoln, 1985; Shenton, 2004). Credibility was further enhanced by conducting a pilot focus group (not included in the study findings) in order to improve skills in focus group facilitation, as it is acknowledged that the facilitator's role is crucial to the quality of data collected (Kennedy et al., 2001). Further details on the pilot focus group can be found in Sims (2017).

Throughout the research process, reflections and ideas were kept in a research journal to consciously acknowledge any biases. Meetings with an advisory group were also used as a way to broaden influences on the study.

Results

Participants described both positive and negative views of their prostheses, with some participants displaying a seemingly ambivalent relationship with their prosthesis, recognizing that it is helpful in some activities yet is "more hassle than it is worth" (Anna, aged 14) at

other times. The following section describes two overarching themes – "Prosthesis as a help" and "Prosthesis as a hindrance". Figure 1 shows the breakdown of themes into subthemes.

[Insert figure 1 here]

Theme one: "Prosthesis as a help":

Participants described experiences where their prosthesis has been helpful to them for social interactions, when playing and for carrying out self-care and productive activities.

In social interactions

The participants described finding their prostheses useful in activities which involved an element of social interaction. The participants in the study used their prostheses to deal with unwanted attention in different settings, such as when at school and when out in public, particularly in situations where they would encounter people they did not know. Becky (aged 8) described how the assumptions people make about limb difference, and the things they say, can be hurtful:

"If they can see and they come up to me and say "Hey, you've got one hand" that might make me feel in a way sad. But, then if they didn't like, if they saw and then didn't make an opinion on it I wouldn't mind that."

The unwanted attention the participants described included staring, asking questions and teasing:

"Say if I've had a bad day at school I normally wear it the next day. I don't know why. Like, people are teasing me or something" (Becky, aged 8).

Inherent in this desire to deflect unwanted attention was a feeling that the participants were viewed by other people as "different" as a result of their limb difference, and that this could cause embarrassment and isolation for them. Anna (aged 14), for example, remembered a time when she was younger and was required to wear gloves as part of a costume: "We had this dancing thing, where we had to wear, everyone had to wear, two blue gloves, and so I had one, a cosmetic hand, so that I could wear two blue gloves as well. I was

embarrassed that everyone else had two and I only had one".

Conversely, some participants reported choosing not to wear their prosthesis in social situations because they found it led to more unwanted attention than when not wearing it. For example,

"I think I feel more, not as much embarrassed, but more self-conscious if I do wear one [a prosthesis], especially if I have short sleeves. Then I feel more self-conscious when I'm out and about" (Anna, aged 14).

In play

Another positive element of prosthesis use described by the participants in the study was having fun with their prosthesis and using it in games with friends or for practical jokes, and some of them even viewed it as a toy, storing it in their toy box. For example:

"For a joke at home I say to my sisters 'you wanna piece of me? You wanna piece of me?' and I take my [prosthetic] arm off and give it to them" (Becky, aged 8).

"When we played games of tag, I used to use a prosthetic arm to actually tag the person or grab their t-shirt" (Gareth, aged 15).

Using the prosthesis for play, fun and humour were also seen as a way of celebrating their difference and appreciating something that is individual about them.

For self-care and productivity

Participants talked about using their prostheses to increase their ability to perform functional tasks. Some participants talked about using a prosthesis as a general aid for day-to-day tasks, whereas others talked about using one for specific activities.

"I used to use it for holding paper and stuff [...] It was useful when I was younger for cutting and stuff" (Lauren, aged 14).

"I might wear it if I was cleaning the bathroom because then I wouldn't hurt my little arm when I'm cleaning or something like that" (Emma, aged 9).

Some participants talked about having prostheses that are specifically designed to aid with particular activities that are important to them. Specific activities that the participants used their prostheses for generally involved taking part in sports / Physical Education at school: "When I play hockey it helps me hold my hockey stick because you have to have your right arm at the bottom and I obviously can't reach" (Emma, aged 9).

Participants appeared to find prostheses that were specifically designed for a task more useful for sports:

"I find the one that I've got at the moment, which I use for riding a bike, I find it really helpful when I'm riding a bike because that's the only thing I use it for" (Chris, aged 12).

As well as having prostheses for use specifically in sports, which included prostheses for cricket, cycling, golf and press-ups, the participants talked about having prostheses to help with typing and playing musical instruments:

"It's a thing that goes on my arm so that I can attach a plectrum to it so that I can play the guitar" (Anna, aged 14).

Participants described using their prostheses at times when they wanted to blend in and avoid attention from others. However, at other times they were used to enhance play and humour with friends. Prostheses were also seen to have practical functional benefit for some self-care and productive activities, particularly when they had been designed for use specifically in those activities.

Theme two: "Prosthesis as a hindrance":

Participants described choosing not to use prostheses as they found they provided no functional benefit. In some instances (particularly sport) they were even reported to inhibit participation and restrict the wearer's range of movement.

No functional benefit

Several of the participants reported having periods of not wearing a prosthesis because it did not provide any functional benefit. Lauren (14) reported not using a prosthesis because it did not help with activities:

"I used to wear it because I thought it would always help me with things but then after a while I thought it's not really helping me to do anything I couldn't already. You can't do anything with it that you can't do without it".

In sporting activities

Not only did the participants express the view that prostheses do not help function, but that they can actually be a hindrance in certain activities or due to certain design features.

"You couldn't hold the club because it was sliding around" (Gareth, aged 15, talking about playing golf).

"When you're doing sports you always feel like you're trailing it along [...] Just think it'd kind of slow me down a bit. I wore it for football once but just kinda gave up after a while just because [...] I just felt like I was trailing it around" (Gareth, 15).

In these instances, the participants were referring to trying to use general purpose passive or body-powered prostheses for sports, rather than task-specific devices.

Restricting movement

Lauren (aged 14) talked about her prosthesis restricting her elbow movement:

"It blocks you from using your elbow so you can't use it. So it's there but you can't do much stuff with it [...] it was kind of blocking how to pick up things and stuff like that. I found it made it harder a bit".

The weight of the prosthesis, combined with limited movement, was also identified as a factor in the prosthesis hindering function in daily tasks:

"It was more hassle than it was worth because all it did was that [demonstrates opening and closing of grip] and it was really heavy so it ended up being less convenient than without" (Lauren, aged 14).

The results show prostheses will not be used if they do not provide some practical functional benefit, particularly if issues with fit and weight result in restricted movement and difficulties with participating in activities.

Discussion and implications

The participants in the current study evidently find prostheses useful at some times, and in some activities, but a burden in others. Participants of the current study made much reference to the responses of others to limb difference and the impact of this on prosthesis use. Being viewed as different as a result of their impairment could result in feelings of embarrassment and isolation for the participants, leading them to choose to use prostheses to blend in or conform. Participants discussed finding their prostheses useful in social situations for preventing unwanted attention. Conversely, they sometimes chose to not use their prostheses when others around them accepted their differences, and the children and adolescents in our study suggested that in these social contexts use of a prosthesis only highlighted their differences. This reinforces the notion of disability being a socially constructed phenomenon (Shakespeare and Watson, 2002), as unwanted attention is an extrinsic societal factor as opposed to an impairment-based issue. The participants in our study chose whether or not to wear their prostheses to conform to societal regulations rather than for making an activity easier to perform, highlighting the prominence of socially constructed disability.

Participants reported using their prostheses for self-care and productivity. They also described having fun with their prostheses and using them for play, which supports our understanding of play as an important occupation for children (Case-Smith and O'Brien, 2015). Play is recognised as a central occupation of childhood, an essential part of children's development (Larson 2004). It has been argued that play in childhood is equivalent to work in

adult life, in its capacity to encourage personal development and provide opportunities for formative experiences (Ginsburg 2007). It is generally recognized that the two main functions of a prosthesis are to improve a person's appearance and to increase their ability to perform functional tasks (Spires et al., 2000). By directly exploring the views of children and adolescents, however, it has been revealed that there is a third important function of prostheses for this client group, which is as something to have fun with and include in play with friends and family. It is interesting that, when with friends and family, participants described using prostheses to enhance playful interactions, whereas with unfamiliar people prostheses were used to conceal limb difference. Using prostheses in a fun and humorous way may be an appropriate coping strategy for children and adolescents in managing stressful events (Dowling 2002). It can enable them to view an upsetting or stressful event from a different perspective and reappraise it as less threatening, lessening feelings of anxiety, fear, anger and frustration and helping them to gain control of the situation (Bennett 2003).

In parallel to previous research by Wagner et al. (2007), several of the participants reported having periods of not wearing a prosthesis because it did not provide any functional benefit. Furthermore, they also reported finding them to be a hindrance in certain activities, due to design features or restriction of movement, with particular emphasis on the weight of prostheses. This supports previous findings that prostheses do not always help function and are often too heavy for children (Biddiss and Chau, 2007; Pylatiuk et al., 2007; Routhier et al., 2001; Wagner et al., 2007).

The prostheses that were described most positively by participants were those that had been designed for a specific use, rather than general purpose functional devices. This supports previous research that children and adolescents should be provided with prostheses to serve

specific purposes and for use in particular activities so that they can select the device that will be most appropriate for the desired use (Buffart et al., 2007; Egermann et al., 2009; James et al., 2006). Prescribing general purpose functional prostheses implies the prosthesis is aimed at 'replacing' some function or cosmesis which is perceived to be 'lacking' or attempting to replicate a human hand. This strongly suggests an impairment-based approach to therapy, through treatment of the medical condition rather than adaptation of the social, cultural and environmental factors that may be leading to disability. Approaching prosthetic treatment as providing a 'replacement' for limb difference can be detrimental to ongoing rehabilitation and personal independence (Gallop, 2012). The current study suggests that prostheses as a general replacement for limb difference is largely unnecessary, with participants reporting that they feel comfortable with their difference and are able to complete their daily activities, as they have developed their own methods for performing the activities in a functional way. Our study suggests prescribing a device to assist with specific tasks may be more beneficial. Prosthesis development and prescription for children and adolescents may be more beneficial if it focused on what they wish to gain from the use of a device (i.e. their activity goal) as opposed to trying to 'treat' their 'deficiency' or replace the function they are (perceived to be) lacking. By providing appropriate tools for completing particular activities (rather than a general purpose prosthesis) emphasis is placed on adapting the physical demands of the environment to enable a young person with limb difference to participate in the task.

Limitations

The findings have indicated that prostheses can be valuable tools for participation in occupations, although, due to the relatively small sample size, conclusions should be tentative. Furthermore, the environmental and task demands of performing these occupations

require further study. An ethnographic study of prosthesis use in children's and adolescents' daily occupations could provide this insight.

Conclusion

Children and adolescents recognise that prostheses can serve particular functions for them and that they are useful in certain circumstances, but that prostheses may also have many drawbacks that counter the perceived benefits. Prostheses can be useful tools for occupational participation for children and adolescents providing that the right tool is provided for the task. Prescribing general purpose functional prostheses for children can result in the prosthesis not being useful or, worse, becoming a hindrance through restriction of movement and increased weight. Task specific devices can overcome these issues, as well as reflecting a less impairment-based approach to limb difference, which can be detrimental to developing independence.

Key findings

- Paediatric prostheses are used in social, productive and leisure occupations, including play;
- General purpose prostheses can be a hindrance in activities and represent an impairment-based approach to therapy;
- Task-specific devices can help to overcome some of these issues.

What the study has added

Upper limb prostheses can be useful to children and adolescents for participating in occupations. However, they can also be a hindrance. Prosthesis prescription for children and

adolescents should focus on what they wish to gain from the use of a device as opposed to trying to 'treat' their 'deficiency'.

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Research Ethics

Ethical approval was obtained from the University of Southampton's Faculty of Health Sciences' Ethics Committee (FoHS-ETHICS-2011-056; June 2011).

Consent

All participants provided written informed assent to be interviewed or participate in a focus group for the study. Their parents provided written informed consent for their children to participate.

Declaration of Conflicting Interests

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Contributorship

Tara Sims researched literature. All authors contributed to applying for ethical approval, developing methodology and carrying out data collection. Tara Sims carried out the data analysis. Tara Sims wrote the first draft of the manuscript. All authors reviewed and edited the manuscript and approved the final version.

References

Bennett HJ (2003) Humour in medicine. Southern Medical Journal 96(12): 1257–1261.

Biddiss E (2010) Need-directed design of prostheses and enabling resources. In: Murray C (ed) *Amputation, prosthesis use and phantom limb pain: an interdisciplinary perspective*. London, Springer, pp. 7-22.

Biddiss E and Andrysek J (2011) Upper and Lower Extremity Prosthetics for Children and Youth. In: Chau T and Fairley J (eds) *Paediatric Rehabilitation Engineering: From Disability to Possibility*. Boca Raton, Florida, CRC Press, pp. 189-240.

Biddiss EA and Chau TT (2007) Upper limb prosthesis use and abandonment, a survey of the last 25 years. *Prosthetics and Orthotics International* 31(3): 236-57.

Braun V and Clarke V (2006) Using thematic analysis in psychology. *Qualitative Research in Psychology* 3(2): 77-101.

Broomfield M (2009) Guide to Artificial Limbs. Tavistock, Devon: Reach Charity Ltd.

Buffart LM, Roebroeck ME, van Heijningen VG, Pesch-Batenburg JM and Stam HJ (2007) Evaluation of arm and prosthetic functioning in children with a congenital transverse reduction deficiency of the upper limb. *Journal of rehabilitation Medicine* 39(5): 379-386.

Case-Smith J and O'Brien JC (2015) *Occupational Therapy for Children and Adolescents* (7th Edition). London: Mosby.

Celikyol F (1984) Prostheses, Equipment, Adapted Performance, Reflections on these choices for the Training of Amputees. *Occupational Therapy in Health Care* 1(4): 89-115.

Department of Health (2017) Children given sports prostheses to help them get active.

Available at: https://www.gov.uk/government/news/children-given-sports-prostheses-to-help-them-get-active (accessed 25 July 2017).

Dillingham TR, Pezzin LE and MacKenzie EJ (2002) Limb amputation and limb deficiency, Epidemiology and recent trends in the United States. *Southern Medical Journal* 95(8): 875–883.

Donovan-Hall M (2010) Supporting the positive development and well-being of children and young people with prostheses: the influence of the appearance of the prosthesis and individual choice. *Big Lottery Final Report (RG1/010120846)*.

Dowling JS (2002) Humor: A coping strategy for pediatric patients. *Pediatric Nursing* 28(2): 123-131.

Egermann M, Kasten P and Thomsen M (2009) Myoelectric hand prostheses in very young children. *International Orthopaedics* 33(4): 1101-1105.

Gallop S (2012) Limb Loss of Personal Independence: Conflicts between artificial limb wearing and quality of life. Available at:

http://www.dysnet.org/images/files/limb_loss_personal_independence_final-1.pdf (accessed 6 April 2017).

Ginsburg KR (2007) The importance of play in promoting healthy child development and maintaining strong parent-child bonds. *Pediatrics* 119(1): 182–191.

Guba EG and Lincoln YS (1985) Naturalistic Inquiry. Thousand Oaks, CA: Sage

Hanspal R (2003) *Amputee & Prosthetic Rehabilitation-standards & Guidelines*. British Society of Rehabilitation Medicine, London.

Ibbotson V (2002) Upper limb amputees and limb-deficient children. In: Turner A, Foster M and Johnson S (eds) *Occupational therapy and physical dysfunction – principles, skills and practice* (5th edition). London, Elsevier, pp. 379-393.

Iversen OS and Brodersen C (2008) Bridging the Gap between users and children - A socio-cultural approach to designing with children. *Cognition, Technology and Work* 10(2): 83-93.

James MA, Bagley AM, Brasington K, Lutz C, McConnell S and Molitor F (2006) Impact of Prostheses on Function and Quality of Life for Children with Unilateral Congenital Below-the-Elbow Deficiency. *The Journal of Bone and Joint Surgery* 88(11): 2356–2365.

Kennedy C, Kools S and Kruger R (2001) Methodological considerations in children's focus groups. *Nursing Research* 50(3): 84–87.

Larson EA (2004) Children's work, the less considered occupation. *American Journal of Occupational Therapy 58*: 369-379.

Mack N, Woodsong C, MacQueen K, Guest G and Namey E (2005) *Qualitative research methods: A data collector's field guide*. North Carolina: Family Health International.

Mertens DM (2015) Research and evaluation in education and psychology: Integrating diversity with quantitative, qualitative, and mixed methods (4th edition). Thousand Oaks, CA: Sage.

Michielsen A, van Wijk I and Ketelaar M (2011) Participation and health-related quality of life of Dutch children and adolescents with congenital lower limb deficiencies. *Journal of Rehabilitation Medicine* 43: 584–589.

Muzumdar A (2004) *Powered Upper Limb Prostheses, Control, Implementation and Clinical Application*. Berlin, Germany: Springer.

National Academies of Sciences, Engineering, and Medicine (2017) *The Promise of Assistive Technology to Enhance Activity and Work Participation*. Washington, DC, The National Academies Press.

O'Kane C (2008) The Development of Participatory Techniques. In: Christensen PM and James A (eds.) *Research with Children, Perspectives and Practices*. Oxford, Routledge, pp.125-155.

Patton JG (2004) Training the child with a unilateral upper-extremity prosthesis. In: Meier RH and Atkins DG (eds) *Functional restoration of adults and children with upper extremity amputation*. New York, Demos Medical Publishing, pp. 297-316.

Peterson J (2011) The Prosthetic Habilitation of a Congenital, Transradial Limb Deficient Child, A Case Study Analyzing the Functional Effectiveness and the Benefits of Early Prosthetic Fitting, Appropriate Prosthetic Equipment, and Consistent Caregiver Follow up. *MEC 11 Raising the Standard, Proceedings of the 2011 MyoElectric Controls/Powered Prosthetics Symposium* Fredericton, New Brunswick, Canada.

Pylatiuk C, Schulz S and Doderlein L (2007) Results of an Internet survey of myoelectric prosthetic hand users. *Prosthetics and Orthotics International* 31(4): 362-70.

Routhier F, Vincent C, Morissette MJ and Desaulniers L (2001) Clinical results of an investigation of paediatric upper limb myoelectric prosthesis fitting at the Quebec Rehabilitation Institute. *Prosthetics & Orthotics International* 25(2): 119-131.

Shakespeare T and Watson N (2002) The social model of disability: An outdated ideology? *Research in Social Science and Disability* 2: 9-28.

Sheffler LC, Hanley C, Bagley A, Molitor F and James MA (2009) Comparison of Self-Reports and Parent Proxy-Reports of Function and Quality of Life of Children with Below-the-Elbow Deficiency. *Journal of Bone and Joint Surgery* 91: 2852-2859.

Shenton AK (2004) Strategies for ensuring trustworthiness in qualitative research projects. *Education for Information* 22: 63–75.

Shida-Tokeshi J, Bagley A, Molitor F, Tomhave W, Liberatore J, Brasington K and Montpetit K (2005) Predictors of Continued Prosthetic Wear in Children With Upper Extremity Prostheses. *Journal of Prosthetics and Orthotics* 17(4): 119-124.

Sims T (2018) Prosthesis Use in Times of Transition for Children and Adolescents with Upper Limb Difference. *Children, Young People and Families Occupational Therapy Journal* 22(1): 15-18.

Sims T (2017) *Piloting the Focus Group Method: Preparing for a Participatory Design*Study with Children and Young People. Sage Research Methods Cases Part 2 [Edited Collection]. Available at: http://methods.sagepub.com/case/focus-group-method-preparing-participatory-design-children-young-people (accessed 25 July 2017).

Sims T, Cranny A, Metcalf C, Chappell P and Donovan-Hall M (2017) Participatory Design of Paediatric Upper Limb Prostheses: qualitative methods and prototyping. *International Journal of Technology Assessment in Health Care* 33(6): 629-637.

Smith DJ (2006) Congenital Limb Deficiencies and Acquired Amputations in Childhood Part 3. *In Motion* 16(3): 56-61.

Spires MC, Miner L, and Colwell MO (2000) Upper-Extremity Amputation and Prosthetic Rehabilitation. In: Grabois M, Garrison SJ, Hart KA and Lehmkuhl LD (eds) *Physical Medicine and Rehabilitation: The Complete Approach*. Malden, Massachusetts, Blackwell Science, pp. 549-582.

United Nations (1990) *Convention on the Rights of the Child*. Available at: http://treaties.un.org/Pages/Treaties.aspx?id=4&subid=A&lang=en (accessed 21 July 2018).

Wagner LV, Bagley AM and James MA (2007) Reasons for prosthetic rejection by children with unilateral congenital transverse forearm total deficiency. *Journal of Prosthetics and Orthotics* 19(2): 51-54.

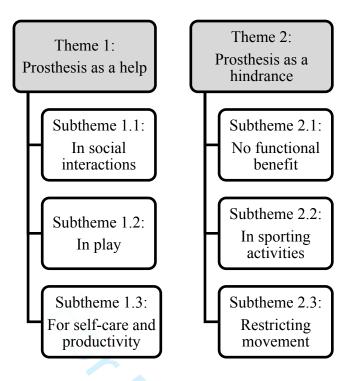
Watve S, Dodd G, MacDonald R and Stoppard ER (2011) Upper limb prosthetic rehabilitation. *Orthopaedics and Trauma* 25(2): 135-142.

Winchell E (1996) Body Image, Acceptance is Step One. *UK Limb Loss Information Centre Journal*, 6 (4): 1-3.

Table 1: Characteristics of participants

Participant	Gender	Age	Aetiology	Side of	Experience	Part of UK
(pseudonym)				limb	of	
				difference	prosthesis	
					use (daily,	
					occasional,	
					previous)	
Becky	Female	8	Congenital	Left	Occasional	England
Anna	Female	14	Congenital	Left	Occasional	England
Gareth	Male	15	Congenital	Right	Previous	Scotland
Lauren	Female	14	Congenital	Right	Previous	Wales
Emma	Female	9	Congenital	Right	Previous	England
Chris	Male	12	Acquired	Right	Previous	England
David	Male	10	Congenital	Right	Daily	England
James	Male	8	Congenital	Left	Daily	England

Figure 1: Themes and subthemes



Appendix: Questioning Route for interviews and focus group

For inclusion in focus group, interview or both	
Focus Group	Opening Questions Tell us your name and where you live Tell us your first name and your favourite colour
Both	 Introductory Questions Think about the first prosthetic arm you can remember having, what was it like? What were the things you liked about it? What were the things you didn't like about it?

Both	Transition Questions				
	What has been the best prosthetic arm you've had? – What made				
	it the best?				
	Thinking about all the different prosthetic arms you've had				
	• What's been good or bad about the way they've looked?				
	What's been good or bad about the things you could use				
	them for/what you could do with them?				
	• What's been good or bad about the way they felt?				
	Were they comfortable? What did or didn't help them to				
	be comfortable?				
Both	Key Questions				
	I want you to imagine that I have a magic wand and can conjure				
	up your ideal prosthetic arm:				
	What would it look like?				
	• Colour				
	Special features				
	Feel/texture of the surface				
	What activities would it help you to do?				
	• at home				
	• at school				
	• with friends				
	How would you feel about it?				

	What would other people think about it?
	How would it feel when you're wearing it?
	Ending Questions
Interview	Is there anything else that we haven't already talked about that
	you would like to tell me about?
Focus	Is there anything else that we haven't already talked about
Group	that anyone would like to add about their ideal prosthetic UL?