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# Facilitators and Barriers of Physical Activity Participation in Children with a Single Ventricle Physiology: a Mixed Methods Study

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## ABSTRACT

### Background

The present study focuses on assessing the physical activity level of children with Fontan circulation for Hypoplastic Left Heart Syndrome, and identifying potential barriers and facilitators toward their participation in physical activity.

### Patients and Methods

Seven children aged 5 to 16 years (mean(SD) 8.8 (3.7) years) with a Fontan procedure for HLHS, their parents (n=7) and siblings (n=1) were recruited. Data were collected using a mixed-methods approach: **(i)** children wore an activity monitor for 7 days to record physical activity, with sedentary time and level of activity calculated from accelerometer data; **(ii)** children completed a bespoke questionnaire recording limitations in physical activity; **(iii)** parents completed a semi-structured interview discussing perceptions about their child's physical activity participation. Qualitative data were analysed using thematic analysis.

### Results

Activity monitors data recorded highly active children with a mean(SD) of 153 (36) minutes/day spent in moderate-to-vigorous physical activity. Time spent in sedentary behaviour was also high (57.5% of total accelerometer wearing time). Four key themes relating to parental perceptions of physical activity were identified: **(i)** A new lease of life – post Fontan; **(ii)** Setting limits – managing limitations; **(iii)** The wider world – how others set limits; and **(iv)** “I fear the future” – parental concerns.

### Conclusion

Following completion of the Fontan circulation, children engaged in higher levels of physical activity in comparison to the national average. However, more than half their time was spent

in sedentary behaviour. Fears and anxiety from parents and teachers may act as a barrier toward physical activity participation.

**Keywords:** Congenital heart disease; hypoplastic left heart syndrome; mixed-methods; physical activity.

## INTRODUCTION

Congenital heart disease (CHD) is a broad term that defines a range of conditions involving the structure of the heart present at birth<sup>1</sup>. CHD is the most common type of birth defect, affecting around 8/1,000 babies born in the UK<sup>2</sup>. Hypoplastic Left Heart Syndrome is a severe defect, where the left-sided structures of the heart are severely underdeveloped, leaving the right ventricle as the only functional pumping chamber of the heart. Hypoplastic Left Heart Syndrome, accounts for 2-3% of all CHD and if left untreated leads to heart failure and death in the first weeks of life<sup>3</sup>. After birth, the neonate will present with cyanosis, failure to thrive, tachypnoea and lethargy<sup>4</sup>. Treatment involves a series of staged surgical interventions aiming to use the only functioning ventricle (right) to support the systemic circulation, while the systemic venous return is redirected to the lungs, bypassing the heart<sup>5</sup>. This is achieved with the Norwood sequence (involving three stages): Stage 1, uses the Right Ventricle to pump blood systemically via a common outlet provided by a reconstructed aorta together with the pulmonary artery, while pulmonary blood flow is provided by a systemic to pulmonary shunt; Stage 2 occurs at around 3-4 months of age when the systemic to pulmonary shunt is exchanged for a Glenn Shunt, where the superior vena cava is redirected to the pulmonary artery; and Stage 3, the Fontan procedure, to redirect blood from the inferior vena cava to the pulmonary artery via a conduit. Thus, the systemic venous return is separated from the pulmonary venous return avoiding intracardiac mixing of blood and cyanosis. Despite advancements in treatment and surgical techniques during the last 30 years<sup>6</sup>, children and adults with Hypoplastic Left Heart Syndrome still report several physical limitations during daily activities, a reduced exercise capacity, and a poorer quality of life overall<sup>7</sup>.

Physical activity plays a fundamental role in childhood development, contributing to healthy growth<sup>8</sup>; improved cardiovascular fitness; and quality of life. For children with CHD, these benefits may be heightened<sup>9</sup>. However, children diagnosed with CHD (irrespective of the complexity) usually report more sedentary behaviours compared to their peers, and reduced physical activity participation<sup>10</sup>. This worsens with age and disproportionately affects girls, and those from areas of higher deprivation<sup>10</sup>. Several barriers to physical activity participation, such as social stigma and parental overprotection, have been identified in children with CHD<sup>11</sup>. Currently, there is no consensus on what constitutes optimal physical activity levels. However, based on evidence from other chronic conditions, an individualised approach to physical activity, with adaptation over the life-course is likely to be beneficial for this cohort<sup>12</sup>.

The aim of this study was to explore the engagement in physical activity and activity levels in these children, and how family social functioning/lifestyle acts as barriers or enablers.

## PATIENTS AND METHODS

Data were collated via a combination of activity monitors, questionnaires and semi-structured interviews.

### Participants

Children with a Fontan circulation for HLHS, aged 5-16 years were identified via a research database at Alder Hey Children's Hospital, and social media between June 2021 and March 2022. Each participant received information explaining the study. Written consent was recorded from participants aged 16 years and older, with assent from those <16 years.

Inclusion criteria comprised diagnosis of a single ventricle congenital heart pathology in the child, and subsequent Fontan circulation.

### Data Collection

Data collection comprised three stages: **(i)** children wore an activity monitor for  $\geq 7$  days; **(ii)** children and siblings (if any) completed a bespoke questionnaire (10-15 minutes to complete); and **(iii)** parents completed a semi-structured interview (20-30 minutes).

### Activity Monitors

Physical activity was monitored using a tri-axial accelerometer (Actigraph wGT3x-BT) worn on the right hip for 7 consecutive days, with removal for sleeping and water-based activities. Times the device was worn were recorded manually. Accelerometer non-wear time was defined as 60 consecutive minutes of zero counts/min<sup>-1</sup><sup>13</sup>. Data were considered for analysis if there was  $>7$  hours of wear time per day<sup>14</sup>, for a minimum of 4 days, including at least one weekend day<sup>15</sup>.

### Activity Monitors' Data Analysis

The ActiLife software, version 6.2 (ActiGraph, Florida) was used to download the data and perform scoring and wear-time validation analysis. Raw acceleration data were converted to 60s-epoch activity count data (counts.min-1). Physical activity intensity was based on the following cut points (counts.min-1)<sup>16</sup>: light ( $\geq 150$ ), moderate ( $\geq 500$ ), and vigorous ( $\geq 4000$ ). Total time (minutes) spent in the above thresholds was calculated (Figure 1).

### Children/Siblings questionnaire

Two questionnaires were developed: one for children with CHD, and one for siblings. Questionnaires comprised two sections: **(i)** multiple choices questions based on "emoji scores" <sup>17</sup> and **(ii)** free narrative section where the child could describe daily routines. For

children with CHD, the questions (Supplementary Table S1) related to aspects of daily life, providing insight into barriers/facilitators in physical activity. The sibling questionnaire, enabled the children to record the impact of their siblings' condition on their life. Questionnaires were developed and piloted with a dedicated Patient and Public Involvement group.

#### Children/Siblings' Questionnaire Analysis

The results of the questionnaires were reported using descriptive statistics. No formal statistical significance testing was undertaken due to small numbers.

#### Parents' interviews

The interviews explored parental experiences, particularly the impact of CHD on family activities, daily activities, sport, and physical activity. A semi-structured interview format was utilised, with each interview lasting ~30 minutes.

#### Parents' interview analysis

Interviews were digitally recorded and transcribed verbatim. Data was analysed in accordance with the 6-step thematic analysis<sup>18</sup>. All transcripts were read by three researchers (DGL/DAL/RRL) and discussed in dedicated meetings.



## Results

### Participants

Fifteen participants were recruited, including 7 parents and 7 children (3 girls/4 boys), ranging from 5-16 years (mean (SD) 8.8 (3.7) years; one aged >11 years). Only one sibling was recruited and therefore this data was excluded to ensure confidentiality.

### Activity Monitors

All children wore the activity monitors for  $\geq 7$  hours per day. Mean daily sedentary time (ST) over 7 days (including two weekend days) was higher (57.5% of wear time) than mean daily total activity (TA) time (42.5% of wear time). Mean (SD) daily time spent in moderate-to-vigorous activity was 153 (36) minutes (26.5% of the total wear time) (Figure 1). No difference in the amount of activity was observed between weekdays and weekend days (Table 1).

### Children/Siblings questionnaires

Results from children's questionnaires are presented in Figure 2. Physical activities of interest suggested by the children (H.7 – Supplementary Table S1) were dance (33%), short walks (17%), and climbing (17%).

### Parents' interviews

Data generated from six interviews (one interview undertaken with both parents together) were analysed. Four key themes were identified. *A new lease of life – post Fontan* highlights the impact of the surgery on physical activity engagement by children following completion of the Fontan circulation. *Setting limits – managing limitations* explores the different mechanisms through which families managed physical activity with a child with a single

ventricle. *The wider world – how others set limits* examines the way in which the community affects engagement in physical activity in children with a single ventricle. Finally, *I fear the future – parental concerns* examining fears and anxiety that parents experience in relation to their child's condition.

#### *A new lease of life – post Fontan*

All the parents spoke about the positive impact of the Fontan circulation on their child's ability to engage in physical activity. Post-surgery, parents reported that their child was less exhausted and had fewer limitations in physical activity participation:

*"Before his operation he was taking Wednesday off as he was exhausted.*

*After the operation he is in full time school, he keeps up with his peers. I'm*

*thrilled about it" [05 Mother]*

Prior to completion of the Fontan circulation, cyanosis on exertion was common. Parents frequently described these changes as visible indicators of the child's limitations, providing a measure of how much a child could do.

*"He would join PE, but his teachers are really aware of his condition, so if his*

*lips get blue or if he gets breathless, then they make sure he sits down and*

*rest." [09 Mother]*

However, these external markers disappeared post-surgery, and changed the way the child was perceived.

*"Since he started school, it shocked the people I tell it to, they are like "has*

*he?!". This because he is doing just fine... he doesn't look like he has got a*

*heart condition. People are very shocked" [05 Mother]*

For some parents, this raised concerns that their child risked being pushed too hard when it came to engaging in physical activity:

*“He had some swimming lessons at a sport facility and the instructor was like ‘ok, ok, I got it’ [referring to #03’s condition] and then one day she [the instructor] pushes (#03) too much and (#03) is like ‘I am not coming again’... because she [the instructor] just did not get it...” [03 Mother]*

### ***Setting limits – managing limitations***

Parents managed their child’s condition in different ways, influencing the way in which limits were imposed and the way the child was included in their activities as a family. In general, the parents interviewed recognised the importance of physical activity to improve their child’s fitness and cardiovascular health:

*“When she got 6 months old I would let her participate in ‘water babies’, which I thought was a good respiratory training... and from there I always tried to keep her active.” [01 Mother]*

For some parents, this meant supported the child to effectively judge when to push themselves, as well as when to stop and have a quick break:

*“He self-manages himself very well. So, he understands his limitations and knows when he has to stop, ... he does self-manage very well.” [03 Father]*

On occasions this could mean pushing their own concerns away and allowing the child to make their own decisions.

*"If he want to do something we completely let him do that, even if sometimes we caught us thinking 'uhm... not so sure about that... it could be dangerous...' " [03 Mother]*

However, other parents reported a more risk-averse approach, stepping in when they perceived their child was overdoing things:

*"He just keeps going, to the point I have to tell him 'come and get your drink, you are getting very hot' ". [05 Mother]*

All the parents discussed their attempts not to treat their child differently to their siblings. However, for some families, this meant adjusting activities to enable the child to participate:

*"We are very active, really...But when #03 was 3 he was riding a quad bike because there was a significant difference in mobility... he couldn't keep up with his brother and his cousins ..." [03 Mother]*

Conversely, others reported adapting family activities to their child's condition, thereby avoiding activities that could potentially exhaust their child:

*"We don't tend to do anything in the evening as a family because of #09 limitations. What we don't want to do is to tire him out to the point where he can't attend school the next day." [09 Mother]*

Additional limitations to engaging in physical activity were associated with environmental factors, such as temperature:

*"He struggled a little when he joined the football team, and it [the training] was outside and was in the winter... and he got too cold even if he had thermal clothes on, we had to stop him from continuing with the training." [08 Mother]*

### *The wider world – how others set limits*

Parents spoke about a spectrum of approaches to managing engagement in physical activity at school. For some, it was treated as a risky activity by the school, resulting in limitations placed on the child.

*"School, obviously there was always the risk assessment... I mean, why shouldn't she do PE? You know, she can sit down if she needs to rest... I don't see the point to limit her participation i." [01 Mother]*

For others, whilst the child was included in activities, they were identified as 'disabled' by the teacher:

*"At school they were letting them have a race and were trying to make it even for everyone [...] and they had to wear a head-band to let the other boys know they had to give them a chance in the race, but he literally outraced everyone" [05 Mother]*

In contrast, other children were trusted by their parents and teachers to self-manage.

*"They know he has a problem, but I don't think it worries them at all [in limiting what he can do], so I don't think they are limiting him but as soon as he says 'I would like to sit down for a minute' they are quite happy to allow him to step back." [03 Father]*

### *Fear for the future – parental concerns*

Uncertainty about the future, led all the interviewed parents to express fear, with most of them preferring to live in the present and not talk/think about the future.

*"I think it is quite hard to know what to plan for the future not knowing how much time we have got with them [their children]..." [09 Mother]*

## DISCUSSION

The children recruited for this study reported an average daily physical activity higher than the minimum recommended by the national guidelines, but also a high proportion spent in sedentary behaviours. The interviews highlighted different mechanisms for managing their child's level of physical activity. Whilst some parents supported the child in self-managing activities, others felt it necessary to impose limitations to protect their child. This was confounded by the spectrum across which the wider community perceived their child, either adding to restrictions, or being unaware of the child's limitations. Whilst the children were all much more active following completion of the Fontan circulation, parents had ongoing concerns over the future.

Evidence examining engagement and perception of physical activity in children with a Fontan circulation is limited. This study demonstrated an average daily moderate-to-vigorous activity more than 2.5 times the recommended 60 minutes for children and adolescents<sup>19</sup>. Other observational studies measuring physical activity in healthy children in England<sup>20,21</sup> suggest that only 31% of boys and 22% of girls aged 4-15 years in England<sup>20</sup>, and 27% of children living in the North-West<sup>21</sup> meet recommended levels of daily physical activity. Moderate-to-

vigorous activity has been shown to decrease with age, with children aged 4-7 averaging 124 minutes per day (boys) and 101 minutes (girls), dropping to 52 (boys) and 28 (girls) minutes in those aged 12-15 years<sup>20</sup>. Data derived from the CHD population is variable, with some evidence of reduced physical activity levels and increased sedentary behaviour<sup>22</sup>, while others have shown that despite a CHD diagnosis, children are meeting physical activity recommendations<sup>23</sup> with no significant differences in sedentary time<sup>24</sup>. Sedentary behaviour in children has been identified as a risk factor for obesity and cardio-metabolic diseases<sup>25</sup> which may disproportionately impact those with CHD<sup>26</sup>, further reducing physical fitness and negatively impacting quality of life<sup>27</sup>. Physical activity interventions have shown to increase the submaximal cardiorespiratory fitness in children with CHD<sup>28</sup>. However, children with HLHS represent the highest risk group among CHD and further studies are required to assess the effects of physical activity/exercise intervention in this population before making appropriate recommendations.

Parental attitudes to physical activity were predominantly positive, although approaches to moderating exertion differed. The impact of parental fears and anxiety on physical activity are well documented,<sup>29</sup> with a positive parental attitude toward physical activity important in promoting self-efficacy in CHD children.<sup>30</sup> Parental perceptions of staff attitudes, suggests teachers often adopt a precautionary approach to physical activity in children with CHD. Since children spend a considerable proportion of their waking time at school<sup>31</sup>, limiting physical education and other play activities may significantly impact their overall physical activity levels. Perceived risk of physical activity in children with CHD therefore needs to be more widely addressed, with effective support in place for schools and families.

## Study limitations

The sample size is small, although acceptable as a pilot study. Geographic diversity of the sample was limited, and participation was likely to be biased towards families who had a more positive attitude towards physical activity. Additionally, activity level of the children in the sample may have been modified by observation (Hawthorne effect).

## PRACTICAL IMPLICATIONS

Although the level of sedentary behaviour of children in this study was similar to their healthy peers, the children spent a significant time sedentary. Reducing sedentary behaviour is crucial in preventing co-morbidities and further reducing quality of life in this cohort<sup>26</sup>. Children with Hypoplastic Left Heart Syndrome (and CHD in general) have been shown to have a reduced exercise capacity<sup>32</sup>, which has been linked to reduced neurodevelopment<sup>33</sup>. Physical activity increases exercise capacity for children with single ventricle heart disease<sup>32</sup>, and has also been shown to potentially reduce NT-proBNP levels (markers of cardiac stress) in these children<sup>34</sup>. Despite this, physical activity participation is still restricted in these children due to healthcare professionals and parents' concerns regarding safety<sup>35</sup>. Further studies that assess the long-term effects of physical activity in children with CHD are required<sup>28</sup>, with appropriate assessment of cardiorespiratory fitness pre- and post-intervention. Engaging the whole family in physical activity<sup>36</sup>, and reducing barriers through increased awareness of the beneficial effects, and safety of physical activity, are the first steps to improving physical activity participation.



## CONCLUSION

Children with a Fontan circulation appear to participate in physical activity commensurate with national recommendations, with similar time spent engaged in sedentary behaviours. Parents' support and external influences (e.g., teachers/relatives) play an important role in supporting engagement in physical activity. Increased awareness of the safety and effectiveness of physical activity in children with CHD amongst parents, supported by educational materials around physical activity for the wider community are needed to promote greater engagement. Further studies to investigate interventions that promote physical activity participation and the long-term effects of physical activity in this population are required.

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## CONFLICTS OF INTEREST

DGL, MR, AAL and RRL declare no conflict of interest. DAL has received investigator-initiated educational grants from Bristol Myers Squibb (BMS); been a speaker for Boehringer Ingelheim, Bayer, and BMS/Pfizer and consulted for Boehringer Ingelheim, Bayer, and BMS/Pfizer; all outside the submitted work.

## ETHICAL STANDARDS

The authors assert that all procedures contributing to this work comply with the ethical standards of the relevant national guidelines on human experimentation (UK HRA and NHS ethics) and with the Helsinki Declaration of 1975, as revised in 2008, and has been approved by the Wales Research Ethics Committee 4 (ref. 20/WA/0144).

## REFERENCES

1. Dolk H, Loane M, Garne E and European Surveillance of Congenital Anomalies Working G. Congenital heart defects in Europe: prevalence and perinatal mortality, 2000 to 2005. *Circulation* 2011; 123: 841-849.
2. Dadvand P, Rankin J, Shirley MD, Rushton S and Pless-Mullooli T. Descriptive epidemiology of congenital heart disease in Northern England. *Paediatric and perinatal epidemiology* 2009; 23: 58-65.
3. Liu L, Oza S, Hogan D et al. Global, regional, and national causes of child mortality in 2000–13, with projections to inform post-2015 priorities: an updated systematic analysis. *The Lancet* 2015; 385: 430-440.
4. Moodie D. *Clinical Management of Congenital Heart Disease from Infancy to Adulthood*. Cardiotext Publishing, 2013.
5. Gobergs R, Salputra E and Lubaua I. Hypoplastic left heart syndrome: a review. *Acta medica Lituanica* 2016; 23: 86.
6. Kutty S, Jacobs ML, Thompson WR and Danford DA. Fontan circulation of the next generation: why it's necessary, what it might look like. *Journal of the American Heart Association* 2020; 9: e013691.
7. Uzark K, Zak V, Shrader P et al. Assessment of quality of life in young patients with single ventricle after the Fontan operation. *The Journal of pediatrics* 2016; 170: 166-172. e161.
8. Malina RM. Physical activity and fitness: pathways from childhood to adulthood. *American Journal of Human Biology: The Official Journal of the Human Biology Association* 2001; 13: 162-172.
9. Tran DL, Gibson H, Maiorana AJ et al. Exercise Intolerance, Benefits, and Prescription for People Living With a Fontan Circulation: The Fontan Fitness Intervention Trial (F-FIT)—Rationale and Design. *Frontiers in Pediatrics* 2021; 9.
10. Voss C, Duncombe SL, Dean PH, de Souza AM and Harris KC. Physical activity and sedentary behavior in children with congenital heart disease. *Journal of the American Heart Association* 2017; 6: e004665.
11. Moola F, McCrindle BW and Longmuir PE. Physical activity participation in youth with surgically corrected congenital heart disease: devising guidelines so Johnny can participate. *Paediatrics & child health* 2009; 14: 167-170.
12. Anderson E and Durstine JL. Physical activity, exercise, and chronic diseases: A brief review. *Sports Medicine and Health Science* 2019; 1: 3-10.
13. Troiano RP. Large-scale applications of accelerometers: new frontiers and new questions. *Medicine and science in sports and exercise* 2007; 39: 1501-1501.
14. Corder K, Ekelund U, Steele RM, Wareham NJ and Brage S. Assessment of physical activity in youth. *Journal of applied physiology* 2008; 105: 977-987.
15. Trost SG, Loprinzi PD, Moore R and Pfeiffer KA. Comparison of accelerometer cut points for predicting activity intensity in youth. *Med Sci Sports Exerc* 2011; 43: 1360-1368.
16. Freedson P, Pober D and Janz KF. Calibration of accelerometer output for children. *Medicine and science in sports and exercise* 2005; 37: S523.
17. Leo DG, Murphy R, Gambling T, Long A, Jones H and Perry DC. Perspectives on the social, physical, and emotional impact of living with Perthes' Disease in children and their family: A mixed methods study. *Global pediatric health* 2019; 6: 2333794X19835235.
18. Clarke V and Braun V. *Thematic analysis: a practical guide*. Thematic Analysis 2021: 1-100.
19. Oja P and Titze S. Physical activity recommendations for public health: development and policy context. *EPMA Journal* 2011; 2: 253-259.
20. Craig R, Mindell J and Hirani V. *Health survey for England 2008: physical activity and fitness*. 2009.
21. Ramirez-Rico E, Hilland TA, Foweather L, Fernandez-Garcia E and Fairclough SJ. Weekday and weekend patterns of physical activity and sedentary time among Liverpool and Madrid youth. *European Journal of Sport Science* 2014; 14: 287-293.

22. McCrindle BW, Williams RV, Mital S et al. Physical activity levels in children and adolescents are reduced after the Fontan procedure, independent of exercise capacity, and are associated with lower perceived general health. *Archives of Disease in Childhood* 2007; 92: 509.
23. Brudy L, Hock J, Häcker A-L et al. Children with Congenital Heart Disease Are Active but Need to Keep Moving: A Cross-Sectional Study Using Wrist-Worn Physical Activity Trackers. *The Journal of Pediatrics* 2020; 217: 13-19.
24. Ewalt LA, Danduran MJ, Strath SJ, Moerchen V and Swartz AM. Objectively assessed physical activity and sedentary behaviour does not differ between children and adolescents with and without a congenital heart defect: a pilot examination. *Cardiology in the Young* 2012; 22: 34-41.
25. Saunders TJ, Chaput J-P and Tremblay MS. Sedentary Behaviour as an Emerging Risk Factor for Cardiometabolic Diseases in Children and Youth. *Canadian Journal of Diabetes* 2014; 38: 53-61.
26. Andonian C, Langer F, Beckmann J et al. Overweight and obesity: an emerging problem in patients with congenital heart disease. *Cardiovasc Diagn Ther* 2019; 9: S360-S368.
27. Gauthier N, Curran T, O'Neill JA, Alexander ME and Rhodes J. Establishing a Comprehensive Pediatric Cardiac Fitness and Rehabilitation Program for Congenital Heart Disease. *Pediatric Cardiology* 2020; 41: 1569-1579.
28. Williams CA, Wadey C, Pielles G, Stuart G, Taylor RS and Long L. Physical activity interventions for people with congenital heart disease. *Cochrane Database Syst Rev* 2020; 10: Cd013400.
29. Bennett E, Voss C, Faulkner G and Harris K. From 'it makes me feel free' to 'they won't let me play': The body and physical activity-related perceptions and experiences of children with congenital heart disease and their parents. *Qualitative Research in Sport, Exercise and Health* 2021; 13: 325-341.
30. Moola F, Faulkner GE, Kirsh JA and Kilburn J. Physical activity and sport participation in youth with congenital heart disease: perceptions of children and parents. *Adapted Physical Activity Quarterly* 2008; 25.
31. Long R. House of Commons Library: Briefing Paper: Number 07148, 24 September 2021: The School Day and Year (England). 2021.
32. Haley JE and Davis C. Exercising with a Single Ventricle: Limitations and Therapies. *Journal of Cardiovascular Development and Disease* 2022; 9: 167.
33. Cooney SJ, Campbell K, Wolfe K, DiMaria MV and Rausch CM. Is Neurodevelopment Related to Exercise Capacity in Single Ventricle Patients Who Have Undergone Fontan Palliation? *Pediatr Cardiol* 2021; 42: 408-416.
34. Perrone MA, Santilli A, De Zorzi A et al. The effects of physical activity in children with hypoplastic left heart syndrome after complete palliation with Fontan procedure. *Med Sport* 2020; 73: 526-533.
35. Caterini JE, Campisi ES and Cifra B. Physical Activity Promotion in Pediatric Congenital Heart Disease: Are We Running Late? *Canadian Journal of Cardiology* 2020; 36: 1406-1416.
36. Brown HE, Atkin AJ, Panter J, Wong G, Chinapaw MJM and van Sluijs EMF. Family-based interventions to increase physical activity in children: a systematic review, meta-analysis and realist synthesis. *Obesity Reviews* 2016; 17: 345-360.

## LIST OF TABLES AND FIGURES

**Table 1:** Differences in activity time and sedentary time between week days and weekend days in children with Hypoplastic Left Heart Syndrome who received a Fontan procedure.

**Figure 1:** Activity and sedentary time in children with hypoplastic left heart syndrome. Figure shows the data from the activity monitors.

**Figure 2a:** Results from the children questionnaire: Limits imposed by parents in physical activity participation in children with hypoplastic left heart syndrome (A), and the way these child feels about engagement in physical activity (B).

**Figure 2b:** Results from the children questionnaire: Limitations reported by children with hypoplastic left heart syndrome when doing physical activity (A and B).

**Figure 2c:** Results from the children questionnaire: Considerations on participation in physical activity in children with hypoplastic left heart syndrome in their free time (A); and possibilities of these children to engage in physical activities (B).

**Figure 2d:** Results from the children questionnaire: Interaction between children with hypoplastic left heart syndrome and others during physical activity (A); and activities that these children like (B).