Effects of a therapeutic exercise program in children with non-cystic fibrosis bronchiectasis: A pilot randomized controlled trial

Barbara Joschtel1, Sjaan R. Gomersall1,2, Sean Tweedy1, Helen Petsky3,4, Anne B. Chang3,4,5 and Stewart G. Trost1,3*

1School of Human Movement and Nutrition Sciences, The University of Queensland, Brisbane, QLD, Australia, 2School of Health and Rehabilitation Sciences, The University of Queensland, Brisbane, QLD, Australia, 3Faculty of Health, Centre for Children’s Health Research, Queensland University of Technology, Brisbane, QLD, Australia, 4Department of Respiratory and Sleep Medicine, Queensland Children’s Hospital, Children’s Health Queensland, Brisbane, QLD, Australia, 5Child Health Division, Menzies School of Health Research, Charles Darwin University, Tiwi, NT, Australia

Background: In the absence of randomized controlled trials (RCTs) on the benefits of therapeutic exercise programs involving children with bronchiectasis, we undertook a pilot RCT to evaluate the effects of a play-based therapeutic exercise program on fundamental movement skill (FMS) proficiency. The effects of the program on cardiorespiratory fitness, perceived competence, and health-related quality of life (HR-QoL) were examined as secondary outcomes.

Methods: Children [median (IQR) age: 6.8 (5.3–8.8) years] with bronchiectasis unrelated to cystic fibrosis were randomized to a 7-week therapeutic exercise program (n = 11) or wait-list control (n = 10). The exercise program comprised 7 × 60-min weekly sessions and was supplemented by a home-based program 2-days/week. Participants were assessed on: FMS (locomotor and object control skills) using the Test of Gross Motor Development 2nd Edition (TGMD-2); cardiovascular fitness by calculating the percent change in heart rate (%ΔHR) from rest to completion of the first stage of a submaximal treadmill test; perceived competence using Harter’s athletic competence subscale; and QoL with the PedsQL.

Results: Significant group by time interactions were observed for locomotor and object control skills. Children completing the therapeutic exercise program exhibited significant improvements in both locomotor (pre 29.0 ± 2.0, post 35.2 ± 2.2, p = 0.01) and object control (pre 35.2 ± 2.0, post 35.5 ± 2.2, p = 0.01) skills, with no significant change in controls (pre 31.6 ± 2.1, post 31.8 ± 2.3 and pre 31.0 ± 2.1, post 32.3 ± 2.3, respectively). Among children completing the program, %ΔHR declined by 6% points, while %ΔHR declined only marginally among controls (0.9% points), but the group by time interaction was not statistically significant. The program had a small
positive impact on competence perceptions (Cohen’s $d = 0.2$) and HR-QoL (Cohen’s $d = 0.3$).

**Conclusion:** This pilot RCT provides preliminary evidence for the efficacy of a play-based therapeutic exercise program to improve proficiency in FMS and fitness in children with bronchiectasis. The results are sufficiently positive to warrant conducting a larger RCT testing the efficacy of the exercise program in children with bronchiectasis and/or other chronic respiratory conditions.

**KEYWORDS**

physical activity, pediatrics, clinical trial, bronchiectasis, fundamental movement skills, respiratory disease, motor competence, fitness

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**Introduction**

Globally bronchiectasis is recognized as a major cause of respiratory morbidity, mortality and health-care utilization (1). It is part of the chronic suppurative lung disease spectrum and caused by chronic airway inflammation resulting in abnormal dilatation of the bronchial wall (1, 2). Typical clinical features are recurrent or persistent wet cough (2, 3) and they regularly experience exacerbations of their condition resulting in frequent acute visits to the doctor, hospitalization, further deterioration of lung tissue, and decreased quality of life (QoL) (4, 5). The prevalence of bronchiectasis ranges from 0.2 to 15 cases per 100,000 children (2). Of particular concern, the prevalence is substantially higher among socially disadvantaged populations such as the indigenous communities of Australia, New Zealand, Pacific Islands, Alaska, and Canada (2, 3). For example, a study of indigenous children residing in central Australia reported a prevalence of 1,470 cases per 100,000 (2, 6).

Current guidelines for the treatment and management of bronchiectasis call for regular exercise as a means of improving cardiovascular fitness and QoL. (3, 7). However, to date, the health benefits resulting from therapeutic exercise have not been systematically investigated in children with bronchiectasis. A recent study involving 39 children and adolescents with non-CF bronchiectasis reported active video gaming to improve aerobic fitness and muscular strength; however, because exercise was combined with an extensive home-based physiotherapy program comprising ventilatory muscle training and postural drainage with percussion, the independent effects of exercise training on the fitness and respiratory outcomes could not be delineated (8). We have previously shown that children with bronchiectasis are insufficiently active for health benefit (9) and exhibit significant developmental delays in fundamental movement skill (FMS) proficiency (10). Further, children with bronchiectasis who achieve their age equivalency for FMS have higher levels of physical activity (10). On the basis of these findings, and the consistent evidence that exercise training improves cardiorespiratory fitness and QoL in children with other respiratory conditions (11), we hypothesized that a play-based, therapeutic exercise program designed to improve FMS would be beneficial to the health and well-being of children with bronchiectasis. Therefore, the primary aim of this pilot randomized controlled trial (RCT) was to evaluate the efficacy of a play-based, therapeutic exercise program to increase proficiency in FMS in children with bronchiectasis. A secondary aim was to determine the effects of the program on cardiorespiratory fitness, perceived movement competence, and QoL.

**Materials and methods**

**Participant recruitment**

Children meeting the following inclusion criteria were eligible to participate in the study: diagnosed with bronchiectasis, aged between 4 and 13 years, and able to complete an exercise program. Bronchiectasis was diagnosed according to the then current guideline published by the Thoracic Society of Australia and New Zealand (3). Exclusion criteria were unstable medical condition, unstable emotional or behavioral status, or recent musculoskeletal injury (e.g., sprains, fracture, and muscle strains).

Participants were recruited through the Department of Respiratory and Sleep Medicine at the Royal Children’s Hospital (now called Queensland Children’s Hospital), Brisbane, Australia, between March 2015 and February 2017. Clinicians were provided a detailed study description, along with the inclusion and exclusion criteria, and asked to identify children who met the criteria and would benefit from participating in a therapeutic exercise program addressing movement skills and aerobic fitness. Parents of potential participants were subsequently contacted by the primary investigator (BJ) who formally assessed eligibility and provided details about the study. After discussing the study, parents with children who were eligible and interested in participating provided written
which consisted of two games from their most recent hourly

Children completed a home-based program 2 days per week

supervised sessions consisted of one 60-min session per week,

physiologist (BJ). To supplement each supervised session,

week, conducted over 7-weeks, led by a clinical exercise

FMS and cardiorespiratory fitness. The program comprised

assessments.

Therapeutic exercise program

The therapeutic exercise program aimed to improve

FMS and cardiorespiratory fitness. The program comprised

a combination of supervised and unsupervised exercise. The

supervised sessions consisted of one 60-min session per

week, conducted over 7-weeks, led by a clinical exercise

physiologist (BJ). To supplement each supervised session,

children completed a home-based program 2 days per week

which consisted of two games from their most recent hourly

session. During each 60-min session, children rotated through

six activity stations where they completed developmentally

appropriate games targeting specific FMS. A description of

each game is provided in Table 1. The activity stations were

sequenced so that a high intensity game was always followed

by a low intensity game. Depending on the age of the

participant, two to three new games were introduced each

session. Activities were tailored to the child's fitness and skill

level by modifying the intensity and duration of each session,

and/or modifying the equipment. All sessions started with a 5-

min warm up and finished with a 5-min cool down consisting

of stretching and relaxation exercises. Exercise intensity during

each program session was monitored continuously with a heart

rate monitor (PolarV800; Polar Electro Oy, Kempele, Finland).

To accommodate families who had limited time and resources

for travel to a research facility, participants were given the

option of completing the supervised sessions at the CCHR or

at home. Approximately half (52%) of the exercise sessions

were completed at the CCHR, with the remaining half (48%)

completed at the participant's home.

Measures

Parental questionnaire

Parents completed a brief questionnaire providing

information about their child's health history, duration of
cough, type of cough, current medication, frequency of doctor
visits during the last 12 months, and family demographics (i.e.,
family structure, parental age, parental smoking status, and
household income).

Fundamental movement skills

Fundamental movement skills were assessed using the Test
of Gross Motor Development 2nd Edition (TGMD-2) (14).
The TGMD-2 evaluates 12 movement skills representing two
dimensions of gross motor performance – locomotor and object
control (14). To assess children, a skill was demonstrated
once, after which the child had one practice trial and two
test trials, with both test scores being recorded. Scores were
based on ratings of movement process characteristics for six
locomotion (i.e., run, gallop, hop, leap, horizontal jump, and
slide) and six object control skills (i.e., striking a stationary ball,
stationary dribble, catch, kick, throw, and roll). For each skill,
the nominated process characteristics were rated as “1” (present)
or “0” (absent). Ratings were summed to derive a raw score for
the locomotor and object control skill dimensions. The TGMD-
2 is a reliable measure of proficiency in FMS in children with
test-retest reliability ranging from 0.86 to 0.96 (14).

Cardiorespiratory fitness

Cardiorespiratory fitness was measured using a submaximal
graded treadmill test following the Balke protocol for low
active children (15). For ethical and safety reasons, only children aged 6 years and older completed the test. After a 2-min familiarization period, participants walked at a speed of 5.2 km/h at 6% grade, with the grade increasing by 2% every 3 min thereafter. The test was terminated at the end of the stage in which the participant’s heart rate exceeded 170 bpm. The percentage change in heart rate (%ΔHR) from rest to the completion of the first stage of the test was used as an indicator of cardiorespiratory fitness, with a decrease in %ΔHR indicative of an endurance training effect (15).

**Perceived competence**

Perceived competence was assessed using the athletic competence subscale from Harter’s Self-Perception Profile for
children (16). This subscale consists of six items measuring children’s perception of their ability to do well in sports and outdoor games. To offset children’s tendency to give socially desirable responses, the Self-Perception Profile utilizes a structured alternative response format. For each item, the child is presented with a description of two children who differ on their ability to play sport or outdoor games. The child is asked to select which of the two children he or she is more like, and to indicate if the description is “really true for me” or “sort of true for me.” An example of an item is: “Some kids wish they could be a lot better at sports BUT other kids feel they are good enough at sports.” Responses were assigned a score of 1 to 4, with a score of 1 indicating the lowest perceived competence and a score of 4 indicating the highest level of perceived competence. Scores for each item were averaged to provide an overall score for perceived competence. Children aged 4–7 years completed the pictorial scale version of the scale (17). Children aged 8–13 years completed the written version of the test (16). The perceived athletic competence subscale is a reliable measure with internal consistency ranging from 0.84 to 0.91 (18). Internal consistency in the current sample was 0.54.

**Quality of life**

The Pediatric Quality of Life Inventory (PedsQL) 4.0 was utilized to assess QoL (19). The PedsQL consists of 23 items independently completed by parents and children. The domains covered include physical functioning, emotional functioning, social functioning, and school functioning. Items address functioning over the past month and responses are recorded using a 5-point Likert scale: 0 (never a problem) to 4 (almost always a problem). This measure has demonstrated reliability and validity in a diverse sample of healthy children and pediatric patients with acute or chronic health conditions (19), and within the sample, internal consistency for the total score was 0.85 for the child report and 0.90 for the parent report.

**Parent-proxy cough-specific quality of life**

The parent-proxy cough-specific quality of life (PC-QoL-8) (20) consists of eight items, exploring how their child’s cough affected their life during the previous week. Items were scored on a 7-point Likert scale, with 1 being “all the time” and 7 “none of the time.” The internal consistency of the scale within the current sample was 0.89.
Statistical analysis

Means, SDs and frequencies were used as descriptive statistics for the sample characteristics and study outcomes. A two-way (group × time) ANOVA was used to assess the effect of the therapeutic movement program on FMS, cardiovascular fitness, perceived competence, QoL and PCQoL. Data were analyzed according to the intention to treat principle, with missing values imputed using the last observation being carried forward method (21). Cohen’s d was used to describe effect sizes. A Cohen’s d of 0.2 corresponded to a small effect size, 0.5 to a medium effect size and 0.8 to a large effect size (22). All analyses were conducted using SPSS software version 26 (IBM Corp., Armonk, NY, USA) and statistical significance was set at alpha level of 0.05.

Results

In total, 21 children were enrolled in the study, with 11 randomized to the exercise group and 10 to the control group. The flow of the participants through the study is shown in Figure 1. Three children in the intervention group dropped out of the study, one due to an injury not related to participation in the exercise program, and two for personal reasons.

Descriptive statistics for the exercise group and control group are reported in Table 2. In both groups, a majority were currently on medication and reported ≥5 doctor visits in the previous year. Of the 11 children randomized to the exercise group, 8 (72.7%) completed all seven supervised exercise sessions. Of the three children who dropped out, two completed six sessions, and one completed two sessions. The mean ± SD heart rate during the supervised exercise sessions was 137 ± 12 beats per minute (bpm). On average, children’s heart rate was above 120 bpm for 81% of the session time and above 140 bpm for 47% of the session time.

Table 3 presents baseline and post-test scores for object control skills, locomotor skills, perceived competence, PedsQL, and PCQoL. There was a significant group by time interaction for both locomotor (F(1,18) = 7.6, p = 0.01) and object control skills (F(1,18) = 8.3, p = 0.01). Children randomized to the exercise group exhibited statistically significant and substantial improvements in both locomotion and object control movement skills, while the control group exhibited little change. The associated effect sizes indicated that the exercise group had a large positive effect on locomotor and object control skills. No significant group by time interactions were observed for scores on the Harter’s perceived athletic competence scale, PedsQL and PCQoL.

Figure 2 illustrates the effects of the exercise group on cardiorespiratory fitness. Among children completing the exercise program, %ΔHR declined 6% points while %ΔHR declined only marginally among control participants (0.9% points). However, due to the small sample size, the group by time interaction was not statistically significant (F(1,13) = 0.9, p = 0.4). Expressed as an effect size, the exercise program had a moderate positive effect on cardiorespiratory fitness (Cohen’s d = 0.5).

Discussion

This pilot RCT evaluated the efficacy of a therapeutic exercise program on FMS, cardiovascular fitness, perceived competence and QoL in children with bronchiectasis. The program significantly improved proficiency in FMS, with a 21% increase in locomotion skills and a 31% increase in object control. The effect sizes associated with these improvements were, by convention, large. In addition, the program had a moderately positive effect on cardiorespiratory fitness. Among children completing the exercise program, %ΔHR declined by 6% points, while children in the control group exhibited minimal change. Although the small sample size precluded statistical significance, the observed effect size for %ΔHR (d = 0.5) was commensurate with that reported for improvements in aerobic fitness in asthmatic children completing exercise training (23). The program had no significant impact on perceived competence or QoL.

Table 2. Descriptive statistics of children aged 4–13 years with non-cystic fibrosis bronchiectasis randomized to the exercise intervention (n = 11) or control (n = 10).  

|                          | Exercise (n = 11) | Control (n = 10) |
|--------------------------|-------------------|------------------|
| Age (years)a             | 6.8 (4.3–8.8)     | 6.9 (5.7–8.9)    |
| Male                     | 73.0%             | 80.0%            |
| Currently on medicationb | 8 (72.7)          | 6 (60.0)         |
| Inhaled steroids          | 1 (9.1)           | 3 (30.0)         |
| Bronchodilators           | 1 (9.1)           | 2 (20.0)         |
| Oral steroids             | 0 (0.0)           | 1 (10.0)         |
| Antibiotics               | 6 (54.5)          | 5 (50.0)         |

Number of doctor visits in the last 12 months

|                      | Exercise (n = 11) | Control (n = 10) |
|----------------------|-------------------|------------------|
| <5 visits            | 18.2%             | 10.0%            |
| 5–9 visits           | 54.5%             | 60.0%            |
| 10–14 visits         | 18.2%             | 20.0%            |
| 15–20 visits         | 9.1%              | 10.0%            |
| Children with a single parent | 9.1% | 20.0% |

Average household incomec

|                      | Exercise (n = 11) | Control (n = 10) |
|----------------------|-------------------|------------------|
| <$25,000 AUD         | 0.0%              | 30.0%            |
| $26,000–$50,000 AUD  | 18.2%             | 20.0%            |
| $51,000–$75,000 AUD  | 9.1%              | 0.0%             |
| >$76,000 AUD         | 72.7%             | 50.0%            |
| Families with a smoker | 18.2% | 30.0% |

AUD, Australian dollar.

aData are expressed as median (interquartile range).
bData are expressed as n and percentage. Some children were on more than 1 medication so percentage exceeds 100%.
cPoverty line <$46,551 AUD/year (37).


| Exercise Control |  |  |  |  |  |
|------------------|---|---|---|---|---|
|                  | Mean | SEM | Mean | SEM | df | F   | P-value | ES |
| **FMS**          |      |     |      |     |    |     |         |    |
| Locomotor pre    | 29.0 | 2.0 | 31.6 | 2.1 |    |     |         |    |
| Locomotor post   | 35.2 | 2.2 | 31.8 | 2.3 | 1.19 | 7.6 | 0.01     | 1.2 |
| Object control pre | 27.0 | 2.0 | 31.0 | 2.1 |    |     |         |    |
| Object control post | 35.5 | 2.2 | 32.3 | 2.3 | 1.19 | 8.3 | 0.01     | 1.3 |
| **Perceived competence** |      |     |      |     |    |     |         |    |
| PAC pre          | 3.0  | 0.2 | 3.0  | 0.2 |    |     |         |    |
| PAC post         | 2.8  | 0.2 | 3.0  | 0.2 | 1.19 | 0.2 | 0.63     | 0.2 |
| **Quality of life** |      |     |      |     |    |     |         |    |
| PQL physical pre | 74.1 | 5.6 | 67.2 | 5.6 |    |     |         |    |
| PQL physical post | 78.4 | 4.7 | 72.7 | 4.7 | 1.19 | 0.7 | 0.80     | 0.4 |
| PQL social pre | 73.5 | 6.1 | 70.0 | 6.1 |    |     |         |    |
| PQL social post | 70.5 | 7.1 | 68.0 | 7.0 | 1.19 | 0.04 | 0.84     | 0.1 |
| PQL school pre | 76.0 | 5.3 | 59.5 | 5.5 |    |     |         |    |
| PQL school post | 75.0 | 5.1 | 63.0 | 5.6 | 1.19 | 1.2 | 0.28     | 0.5 |
| PC-QoL-8 pre | 4.5  | 0.4 | 4.2  | 0.3 |    |     |         |    |
| PC-QoL-8 post | 4.5  | 0.5 | 4.6  | 0.5 | 1.19 | 0.4 | 0.54     | 0.3 |

SEM, standard error of measurement; ES, effect size; PAC, perceived athletic competence; PQL, PedsQL 4.0 quality of life; PCQoL-8, parent-proxy cough-specific quality of life-8.

The results indicate that a play-based therapeutic exercise program can improve FMS proficiency and fitness in children with bronchiectasis and that a larger trial is warranted to test the efficacy of the program.

Previous studies have shown exercise training to be effective in improving cardiorespiratory fitness and QoL in children with asthma and CF (11, 24–26). However, to our knowledge, this is the first study to evaluate the effects of a therapeutic exercise program on proficiency in FMS in children with chronic respiratory disease. Improvements in FMS resulting from the seven-session program were not only statistically significant, but clinically important. Improving FMS is important as such improvements have been shown to contribute positively to children's health and well-being. Children with FMS proficiency have higher levels of cardiorespiratory fitness (27, 28), perceived competence (29, 30) and self-esteem (31). Additionally, children with FMS proficiency are more likely to be physically active (32, 33) and less likely to be overweight and obese (34). Thus, by improving FMS, the capacity for children with bronchiectasis to engage in sport and other physical activities is potentially enhanced, which may mitigate the risk of disabling secondary conditions such as obesity and depression. In support of this assertion, we had undertaken a cross-sectional study which found that children with bronchiectasis who achieve their age equivalency for FMS have higher levels of physical activity, greater perceived competence, and better health-related QoL (10). Future trials evaluating the effectiveness of programs to improve FMS in children with chronic respiratory conditions such as bronchiectasis should therefore include a follow-up period of 6–12 months to determine the extent to which improvements in FMS lead to increases in physical activity performance and improvements in health outcomes such as weight status, cardiovascular fitness, and lung function.

Quality of life is an important multi-dimensional health outcome in children with chronic health conditions (19). In a systematic review and meta-analysis, it was concluded that exercise training significantly improves QoL in children with asthma and CF (26). However, our seven-session therapeutic movement program did not have a significant impact on perceived competence or QoL. This may be due to our small sample size, but other reasons include insufficient duration and intensity of our program to change these outcomes and a lag in time is required before one would expect to see improvements in QoL.

The current study had several strengths. It is the first study to examine the effects of a therapeutic movement program
to improve FMS proficiency in children with bronchiectasis. Further, the therapeutic exercise program was game-based and designed to match the naturally intermittent activity patterns of children (35). Finally, the program is highly scalable. It does not require expensive equipment and can be delivered in variety of settings, including the participant's home.

There were, however, a number of limitations that need to be considered. This was a pilot study providing initial data on the effects of a therapeutic exercise program. We therefore recruited a relatively small number of participants, resulting in low statistical power for some of the study outcomes. However, the findings are promising and provide empirical estimates for effect size and variance of the primary outcome measure for larger trials testing the efficacy of the program on FMS proficiency. Second, due to the nature of exercise intervention studies, it is not possible to blind participants to group allocation, and in this trial, assessor blinding also was not possible as the principal investigator (BJ) delivered the program and completed the assessments. While assessor bias cannot be ruled out, it is important to note that the assessments of cardiorespiratory fitness, perceived competence and QoL were instrument based and therefore did not require interpretation from the assessor. While locomotion and object control scores were based on assessor ratings, the TGMD-2 assessment protocol mitigates against observer bias by clearly delineating the critical features for each skill, which are coded as being present or absent. Third, the exercise program comprised of seven supervised 60-min weekly sessions, with a supplementary home-based program completed twice a week, which is a relatively low dose for an exercise intervention (36). Despite eliciting large positive improvements in FMS proficiency, a longer program may be required to favorably impact perceived competence and QoL. Fourth, despite randomization, there was an imbalance between the two groups in relation to the proportion of participants with a household income of greater than $76,000 AUD and families with a smoker. However, in absolute terms, the differences equated to just 1 or 2 participants per group. Fifth, as noted above, the internal consistency for Harter’s athletic competence subscale was low, potentially indicating problems with children’s understanding of the structured alternative format scale. Thus, the findings related to perceived competence need to be interpreted with considerable caution. Sixth, compliance to the home exercise component was not adequately monitored. At each supervised session, children would verbally self-report their homework from the previous week, but no data were recorded. Therefore, the full dose of the program is unknown.

Conclusion

In conclusion, this pilot RCT provides preliminary evidence for the efficacy of a seven-session play-based therapeutic movement program to improve proficiency in FMS in children with bronchiectasis. The program had a moderate positive effect on cardiorespiratory fitness, but no improvements were noted for perceived competence and QoL. The results are sufficiently
positive to warrant conducting a larger RCT testing the efficacy of the exercise program in children with bronchiectasis and/or other chronic respiratory conditions. Based on the observed effect sizes for the impact of the program on locomotor skills (Cohen $d = 1.2$) and object control skills (Cohen $d = 1.3$) and an estimated SD of 6.8 for locomotor and object control standard scores (based on the observed SD and the recommended inflation factor of 1.099), a sample size of 24 participants will be required. To offset a projected dropout rate of up to 25%, the larger trial should recruit and enroll 15 participants per group, providing a total trial sample size of $N = 30$. This calculation assumes a power level of 80% and a two-sided alpha level of 0.05. In addition, a future RCT should consider implementing the program over a longer duration (8–10-weeks) and employ accelerometer-based motion sensors to monitor the supplemental home-based exercise sessions. Where possible, future studies should be carefully designed to minimize bias (e.g., blinded assessors), and include a follow-up period to determine the extent to which improvements in FMS proficiency are maintained over time and contribute to positive changes in other health outcomes such as physical activity performance, cardiorespiratory fitness, and health-related QoL.

Author contributions

BJ and SGT designed the study. BJ collected and analyzed the data and wrote the manuscript. SGT was a major contributor in designing the study, analyzing data, and writing the manuscript. SG and ST helped designing the study and writing the manuscript. HP and AC helped with recruiting the participants and writing the manuscript. All authors read and approved the final manuscript.

Funding

AC was supported by an Australian National Health and Medical Research Council (NHMRC) Practitioner Fellowship (Grant APP1058213).

Acknowledgments

We want to thank all the children and families who gave their valuable time to participate in our study.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

Ethics statement

The studies involving human participants were reviewed and approved by the Children’s Health Queensland Hospital and Health Service (HREC/14/QRCH/136) and the Human Research Ethics Committee at the University of Queensland (2014001176). Written informed consent to participate in this study was provided by the participants’ legal guardian/next of kin.

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