Longitudinal Myocardial Deformation Does Not Predict Single Ventricle Ejection Fraction Assessed by Cardiac Magnetic Resonance Imaging in Children with a Total Cavopulmonary Connection

L. P. Koopman1 · L. M. Geerdink2,3 · S. S. M. Bossers1 · N. Duppen1 · I. M. Kuipers4 · A. D. ten Harkel5 · G. van Iperen6 · G. Weijers2 · C. de Korte2 · W. A. Helbing1,7 · L. Kapusta2,8

Received: 23 June 2017 / Accepted: 13 October 2017 / Published online: 25 October 2017
© The Author(s) 2017. This article is an open access publication

Abstract Survival of children with single ventricle heart defects after the total cavopulmonary connection (TCPC) has improved, but impaired cardiac function remains a major cause of morbidity and mortality. Cardiac magnetic resonance imaging (cMRI) is the gold standard in assessing single ventricle volume and function, but high costs and limited availability hamper its routine use. A cheaper and more available alternative is echocardiography. Myocardial function can be studied in more detail using speckle tracking echocardiography (STE). The purpose of the study was to describe the association between myocardial deformation assessed by speckle tracking echocardiography (STE) and single ventricle function assessed by cMRI and to evaluate differences in myocardial deformation in children with single left and single right ventricular morphology. Cross-sectional, multicenter study in 77 children after TCPC was conducted. STE segmental and global longitudinal peak strain and systolic strain rate (SR) of the dominant ventricle were measured. Impaired SV function by cMRI was defined as ejection fraction (EF) < 45%. Mean age was 11.8 (range 9.7–14.3) years. Pearson R for cMRI EF versus global longitudinal strain and SR was \( -0.25 (p = 0.06) \) and \( -0.03 (p = 0.82) \), respectively. Global single ventricle longitudinal strain and SR was similar in patients after TCPC with single left and single right ventricular morphology (\( -19.0 \pm 3.1\% \) vs \( 19.2 \pm 3.2\% \), \( p = 0.94 \)). STE myocardial deformation parameters do not correlate with single ventricle ejection fraction assessed by cMRI.

Keywords Fontan · Total cavopulmonary connection · Cardiac MRI · Speckle tracking echocardiography

Introduction

The Fontan circulation is a palliative surgical strategy for children with congenital heart disease who cannot be offered a biventricular repair [1]. The Fontan circulation in the current era is achieved using a staged approach by connecting the superior caval vein and inferior caval vein (either by an extra cardiac conduit [ECC] or intra-atrial lateral tunnel [ILT]) directly to the pulmonary arterial circulation, resulting in a total cavopulmonary connection (TCPC) [2].

While the short- and median-term survival in children palliated with a TCPC has improved considerably during the last decades [3], it is well recognized that cardiac...
function and exercise capacity in children with a TCPC is reduced as compared to healthy children and may deteriorate over time [4].

Cardiac magnetic resonance imaging (cMRI) is considered to be the gold standard for the non-invasive assessment of single ventricle volume and function, although its routine use is hampered by high costs, long analysis time, lack of availability, need for sedation in small children, and inability to use in most patients with pacemaker devices [5]. Echocardiography is cheaper, faster, widely available, and can be used in non-sedated children and patients with pacemaker devices. However, standard 2-dimensional echocardiographic measurements of ventricular volume and function such as ejection fraction (EF) by biplane Simpson and M-mode shortening fraction cannot be used in the single ventricle due to its different geometry as compared to the normal left ventricle (LV). Speckle tracking echocardiography (STE)-derived myocardial deformation [strain and strain rate (SR)] is a less geometry-dependent technique to assess myocardial function compared to standard echocardiographic methods [6]. Various, yet rather small studies in children with a TCPC, have been published, suggesting that measurement of myocardial deformation is feasible and reproducible, and that lower strain and SR values are measured in patients versus controls as well as in TCPC patients with single right ventricular (RV) versus single LV morphology [7–9]. It is unknown whether myocardial strain and SR parameters provide additional information on single ventricular function in patients with a TCPC assessed by cMRI.

The purpose of this study is to describe the association between longitudinal STE-derived myocardial deformation parameters and single ventricle function assessed by cMRI in a relatively large cohort of contemporary pediatric TCPC patients and to evaluate differences in myocardial deformation in patients with single right versus single left ventricular morphology.

Methods

Patients

We performed a cross-sectional multicenter study in single ventricle patients with a TCPC. Details of the recruitment procedures and in- and exclusion criteria are described elsewhere [10]. In short, a total of 77 patients after the TCPC procedure were included. Inclusion criteria were as follows: having undergone a TCPC through a staged approach with a partial cavopulmonary connection procedure prior to the completion of the TCPC according to a current technique (ILT or ECC); completion of the TCPC before the age of 7 years and inclusion in the study at an age of at least 8 years or older; and a minimum of 3 years of follow-up since completion of the TCPC. Patients with mental retardation were excluded from this study. Patients were recruited from four tertiary referral centers in the Netherlands that used the same ultrasound equipment (center that used another ultrasound system was excluded). The study was approved by the institutional medical ethical review boards of the participating centers. Written informed consent was obtained from all patients and/or their parents. All patients underwent routine physical examination, including body weight, height, blood pressure.

Echocardiographic Assessment

All patients underwent a detailed transthoracic echocardiographic examination performed by an experienced technician following the recommendations of the American Society of Echocardiography [11]. All images were obtained with an appropriate transducer on locally available machines (GE Vivid7; GE Vingmed Ultrasound AS, Horten, Norway) according to a standardized protocol that was used in all participating centers. Images were reviewed and supervised by an experienced cardiologist (LK) in the core lab situated in one location.

Conventional Echocardiographic Measurements

Analyses were performed offline using EchoPAC version 11.0 (GE Healthcare, USA). All measurements were performed using 3 different cardiac cycles and the values were averaged. Pulsed wave Doppler flow measurements were obtained at the right upper pulmonary vein. Continuous wave Doppler flow velocities of the ascending (neo) aorta and descending aorta were obtained. Pulsed wave Tissue Doppler measurements were obtained at the basal segment of the lateral wall of the dominant ventricle. Annular plane systolic excursion (APSE) was assessed using M-mode interrogation of the lateral side of the dominant AV valve.

Speckle Tracking Echocardiography (STE) Image Acquisition and Off-Line Analysis

Apical grayscale images were acquired at frame rates of 55–90 frames/s. The images were analyzed by a single investigator (LMG) using EchoPAC (version 11.2, GE Vingmed Ultrasound, Horten, Norway) according to the vendor’s instructions and as previously described by our group [12]. In short, endomyocardial borders of the dominant ventricle were manually traced within the end-systolic frame. Secondly, epicardial tracing was automatically generated by the computer algorithm and, when necessary,
manually adjusted to cover the whole myocardial wall. The tracking algorithm then followed the myocardial speckles during the cardiac cycle. Tracking was accepted only if both visual inspection as well as the software indicated adequate tracking and strain curves. The software automatically divided the apical chamber views in 6 segments (3 septal and 3 lateral). Strain curves of 3 separate cardiac cycles were averaged after equalizing cycle lengths, by padding zeros behind the diastolic phases of the shorter cycles [13]. To determine global longitudinal strain and SR, the average of the six segments was calculated and at least 4 out of 6 valid lateral segments were needed. To determine septal or lateral wall strain and strain rate, at least 2/3 segments should be included.

**Cardiac Magnetic Resonance Imaging (cMRI)**

The participants underwent a cMRI study on the same day as the echocardiographic study. Patients with pacemaker devices were excluded for undergoing cMRI. The ventricular volumes were imaged using a multi-slice, multi-phase, steady-state, free precession sequence as previously reported [10]. In short, the series were acquired at rest and volumes were analyzed using an Advanced Windows workstation (GE Medical Systems, Milwaukee, Wis) and MASS software (Medis Medical Imaging Systems, Leiden, The Netherlands). The endo- and epicardial contours were manually drawn in the end-diastolic and end-systolic phases. The volumes, EF, and mass of the LV and RV were added to calculate the single ventricular indices to compare the different cardiac configurations possible. Cardiac index (CI) was calculated by multiplying stroke volume with heart rate. Abnormal single ventricle EF was defined as <45%.

**Reproducibility**

Inter-observer and intra-observer variability of global longitudinal strain measurements were assessed in 10 randomly selected studies. To assess intra-observer variability, the same observer who performed all the measurements re-measured the same segments at least 2 weeks later to avoid recall bias (LMG). To assess inter-observer variability, deformation parameters were measured by one independent and blinded observer (LK). Reproducibility was expressed as bias (limits of agreement) and the coefficient of variation, COV (the standard deviation of the difference of paired samples divided by the average of the paired samples times 100).

**Statistical Analysis**

Statistical analysis was performed using SPSS 21.0. (IBM corporation, New-York, USA). Normal distribution of the data was tested using Shapiro-Wilk test. Data are expressed as frequencies, mean (standard deviation) in case of normal distribution, or median (interquartile range) in case of non-normal distribution. Comparisons between groups were made using the independent T test (all comparisons were normally distributed). Categorical data are presented as counts and percentages. Pearson’s correlations and their statistical significance were calculated as measures of raw associations between measurements. Two-sided p values ≤ 0.05 were considered to be statistically significant.

**Results**

**General Characteristics**

Table 1 shows the general characteristics and conventional echocardiographic parameters of the whole study population:

**Table 1** General characteristics of the study population (n = 77)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender male</td>
<td>43 (55.8%)</td>
</tr>
<tr>
<td>Age at study (years)</td>
<td>11.8 (9.7–14.3)</td>
</tr>
<tr>
<td>Age at TCPC (years)</td>
<td>3.3 (2.5–3.9)</td>
</tr>
<tr>
<td>BSA (m²)</td>
<td>1.27 (1.09–1.53)</td>
</tr>
<tr>
<td>Resting HR (beats/min)</td>
<td>76 (68–88)</td>
</tr>
<tr>
<td>Resting SaO₂ (%)</td>
<td>96 (94–97)</td>
</tr>
<tr>
<td>SBP (mm Hg)</td>
<td>112 (104–118)</td>
</tr>
<tr>
<td>DBP (mm Hg)</td>
<td>67 ± 9</td>
</tr>
<tr>
<td>Dominant LV/RV morphology</td>
<td>55/22</td>
</tr>
<tr>
<td>Single ventricle diagnosis</td>
<td></td>
</tr>
<tr>
<td>HLHS</td>
<td>11 (14.3%)</td>
</tr>
<tr>
<td>RV hypoplasia with TA or PA</td>
<td>32 (41.6%)</td>
</tr>
<tr>
<td>DILV</td>
<td>8 (10.4%)</td>
</tr>
<tr>
<td>DORV. no TGA</td>
<td>6 (7.8%)</td>
</tr>
<tr>
<td>DORV. with TGA</td>
<td>2 (2.6%)</td>
</tr>
<tr>
<td>Other</td>
<td>18 (23.4%)</td>
</tr>
<tr>
<td>TCPC type (ECC/ILT)</td>
<td>11/66</td>
</tr>
<tr>
<td>Conventional echo parameters of dominant ventricle</td>
<td></td>
</tr>
<tr>
<td>S’ velocity (cm/s)</td>
<td>5.9 (4.9–6.6)</td>
</tr>
<tr>
<td>APSE (mm)</td>
<td>11.5 ± 2.4</td>
</tr>
</tbody>
</table>

APSE annular plane systolic excursion, BSA body surface area, DBP diastolic blood pressure, DILV double inlet left ventricle, DORV Double outlet right ventricle, ECC extra cardiac conduit, HLHS hypoplastic left heart syndrome, HR heart rate, ILT intra-atrial lateral tunnel, LV left ventricle, PA pulmonary atresia, RV right ventricle, S’ systolic velocity, pulsed Tissue Doppler, SaO₂ oxygen saturation, SBP systolic blood pressure, TA tricuspid atresia, TCPC total cavopulmonary connection, TGA transposition of great arteries
population. No significant Fontan baffle leakage was observed in our study population which was also represented by the high median oxygen saturation of 96%. Five participants had an epicardial pacemaker implanted due to heart block or sinus node dysfunction. Four children were diagnosed with right atrial isomerism and 1 child with left atrial isomerism. Two participants had a mildly increased flow velocity in the ascending aorta (1.6 and 2.1 m/s, respectively) and 3 participants had a mildly increased flow velocity in the descending aorta (1.9, 1.9, and 2.1 m/s, respectively). No patient had signs of clinically significant pulmonary vein stenosis at the time of assessment.

Feasibility and Reproducibility

Feasibility of STE was highest for mid and apical septal segments and lowest for the basal septal segment (due to frequent absence of the basal septum in the setting of a large ventricular septal defect), Fig. 1. Reproducibility data for longitudinal myocardial deformation parameters of the lateral wall are provided in Table 2. COVs of myocardial deformation were higher when individual segments were analyzed compared to global parameters (data not shown).

The Association Between Myocardial Deformation, cMRI and Conventional Echo Parameters

In Fig. 2, the associations between single ventricular function assessed by cMRI EF and longitudinal myocardial deformation parameters are shown. There was a trend towards lower (less negative) longitudinal strain values in children with lower EF, but none of the associations were statistically significant. No association was found between cMRI EF and STE strain rate (Fig. 1d–f). Figure 3 shows the results for lateral wall and global longitudinal strain and strain rate when children with a TCPC were dichotomized in a group with normal EF and abnormal EF by cMRI. Complete data on echocardiography and cMRI were available in 50 to 57 out of 77 participants and no statistically significant differences in myocardial deformation were found between children with normal and low single ventricle ejection fraction. Also, no association was found between global longitudinal strain by STE and cMRI SV volume (Fig. 4a), more conventional echocardiographic function parameters SV APSE (4B), SV S’ by pulsed wave TDI (4C) and other cMRI and conventional echocardiographic parameters (Table 3). Longitudinal systolic strain rate was unrelated to SV volumes by cMRI and echocardiographic parameters (Table 3). When septal or lateral wall longitudinal strain or strain rate was correlated to the parameters shown in Table 3 similar results were found (data not shown). A weak correlation was found between cMRI EF and SV APSE (Pearson R 0.34, p = 0.01). No association was found between cMRI EF and SV S’ (Pearson R 0.17, p = 0.19).

Single Ventricle Morphology and Myocardial Deformation

Figure 5 shows that global, lateral wall and septal longitudinal strain was similar in TCPC patients with a dominant right and left single ventricle morphology. Also, when segmental myocardial deformation was compared no differences are observed in children with left ventricular morphology and right ventricular morphology, except for longitudinal systolic strain rate of the basal lateral segment (Table 4).

Discussion

This multicenter study in a relatively large cohort of children and young adults with a TCPC shows that myocardial deformation parameters assessed by STE are not related to single ventricle function assessed by cMRI and conventional echocardiography (APSE). This study also shows that single ventricular morphology in TCPC patients is unrelated to longitudinal myocardial deformation parameters.
Myocardial Deformation and cMRI-Derived Cardiac Assessment

We found no correlation between longitudinal strain of the single ventricle assessed by echocardiography and single ventricle ejection fraction, single ventricle volume and cardiac index assessed by MRI. Khoo et al. performed a study in single RV patients before performing the bidirectional cavo-pulmonary connection (mean age 0.42 year) combining cMRI data with myocardial deformation data assessed by speckle tracking echocardiography [14]. In this small study, the authors found highly statistically significant correlation coefficients for global longitudinal deformation (strain and SR) versus cMRI-derived RV volumes and ejection fraction in the range of 0.57–0.78 [14]. It is difficult to explain why Khoo et al. found these strong correlations between cMRI-derived parameters and STE-derived myocardial deformation parameters compared to our results. A possible explanation might be that the range in ejection fraction seems large in the study by Khoo et al., ranging from 25 to 80%, while in our study ejection fraction by cMRI was generally well preserved and less widely ranged [10, 14]. When the range of values of one parameter is limited, Pearson correlation might not be the most appropriate test to investigate an association. Another possible explanation of the differences between our study and the study by Khoo et al. is that in the later study children were much younger and had less cardiac surgical procedures. Of interest is that when we dichotomized TCPC patients with relatively low and those with normal cMRI EF, no statistically significant differences in global and lateral wall longitudinal strain and systolic SR were found (Fig. 3), suggesting that at least at a relatively young age STE is not very helpful in predicting low EF assessed by cMRI.

Myocardial Deformation in Right Versus Left Dominant Ventricles

Conceptually, a single RV may be more likely to fail over time compared to a single LV, since a RV is not intrinsically designed to sustain systemic blood pressures. Studies in patients with a systemic RV due to transposition of the great arteries (TGA, either D-TGA after Mustard/Senning operation or congenitally corrected TGA without surgical intervention) show that short- and mid-term ventricular function is generally well preserved, but that the majority of patients develop clinical RV failure after the age of 25 years [15, 16]. Many studies have investigated differences in clinical outcome, exercise capacity, and single ventricular function in patients with single RV and LV morphology with a TCPC and found conflicting results [9]. We found that STE-derived global longitudinal myocardial deformation parameters were comparable in children with LV and RV morphology, which is in agreement with a relatively large study by Petko et al. in children with a mean age of 7.8 years [9]. However, Kaneko et al. [17] found in a small study (20 children with TCPC, mean age 3.4 years) that global peak longitudinal strain was $-16.6\%$ in children with RV morphology versus $-20.5\%$ in children with LV morphology ($p = 0.01$). Studies performed in TCPC patients with only one type of single ventricle morphology report global longitudinal strain values in RV morphology ranging from $-12.0$ to $-18.9\%$ [7, 18–21] and in LV morphology ranging from $-14.2$ to $-18.3\%$ [18, 22, 23]. However, these studies are difficult to compare, since they vary in important factors, including age of study participants, time after completion of the TCPC, type of surgical procedure, type of echocardiographic equipment used, and underlying specific single ventricle morphology. In a recently published study by our group in the same population as described in this study, we found a higher single ventricle EF assessed by cMRI in children with LV morphology compared to children with RV morphology ($55 \pm 8\%$ vs $49 \pm 9\%, p = 0.001$) [10], suggesting a slightly better preserved global systolic ventricular function in TCPC patients with LV morphology. We also found indications for less favorable diastolic ventricular performance in children with RV morphology compared to LV morphology. Further studies are required to obtain more insight into this topic.

Feasibility and Reproducibility

Longitudinal deformation measurements by STE in patients with a TCPC were feasible in 65–81% (segmental)
and > 80% (global) of the participants. We have previously reported feasibility figures for normal children in the range of 91–96% for segmental longitudinal deformation and 97–100% for global longitudinal deformation [6, 24]. Previous studies using STE in children and adults with single ventricle morphology report feasibility between 63 and 100% for longitudinal strain [9, 19, 25], although some studies do not provide information on feasibility [17, 22, 26]. Based on our own experience and publications by other groups, we conclude that feasibility of STE in patients with single ventricle morphology is modest, probably due to poor acoustic window as a result of abnormal anatomy (e.g., missing segments in the setting of 

Fig. 2 Association between single ventricle ejection fraction by cMRI and longitudinal peak strain of the septum (a), lateral wall (b), and global (c); longitudinal systolic strain rate of the septum (d), lateral wall (e), and global (f)
large ventricular septum defects) and/or various thoracotomies.

Reproducibility data are not reported in many studies using STE in children and adults with a TCPC [8, 17, 27]. Some studies use inter-class correlation coefficients to report on intra- and inter-observer reproducibility and found figures in the range of 0.8 for both longitudinal strain and SR [19, 26]. Others use COV which were found to be between 3 and 25% for both strain and SR [9, 18, 22]. Our data on reproducibility are comparable with these studies and confirm that performing STE is challenging in patients with single ventricle morphology. Singh et al. have clearly shown that reproducibility of STE is worse in children with a TCPC compared to subjects with anatomical normal hearts [23].

Strength and Limitations

The major strengths of this study are the relatively large number of TCPC patients included and the ability to compare a number of potentially relevant functional parameters, including extensive echocardiographic data (both conventional and innovative parameters such as strain and SR) as well as cMRI data. Other studies using myocardial deformation in single ventricle patients with a TCPC often included rather small numbers of patients [7, 8, 17–19, 21–23, 28] which might result in lack of statistical power.

A recent study by Ghelani et al. investigated the comparability of myocardial deformation assessed by echocardiography and myocardial deformation assessed by cMRI using vendor independent software [29]. They found that intermodality agreement was modest for circumferential strain and poor for longitudinal strain [29]. We decided not to use the vendor independent software, since this software package uses DICOM data with reduced frame rate, which hampers the analysis of strain rate, since this parameter is highly dependent on adequate frame rate. We however acknowledge that it would be interesting to compare myocardial strain assessed by vendor-dependent software with cMRI strain.

Fig. 3 Lateral wall longitudinal strain (a) and strain rate (c) and global longitudinal strain (b) and strain rate (d) in children with a TCPC with normal EF (≥ 45%) and abnormal EF (< 45%) by cMRI

Pediatr Cardiol (2018) 39:283–293
We decided to only include longitudinal deformation in our study. In contrast to the experience of Ghelani et al., who used circumferential strain and SR in a rather large study in young adults with a TCPC [30], we found it very difficult to measure circumferential strain in our study population. By definition, when studying a heterogeneous single ventricle population, many participants will (at least partly) miss their ventricular septum, making it very hard to study circumferential strain. We therefore believe that circumferential strain is an unlikely parameter to become of clinical use in following myocardial function in the majority of TCPC patients.

Finally, although sample size of our study is considerably larger compared to previously published studies, data were missing in some participants for echocardiographic or cMRI parameters due to poor echocardiographic acoustic

---

**Table 3** Correlation between global longitudinal peak strain or strain rate and cMRI parameters and conventional echocardiographic parameters

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Global longitudinal strain</th>
<th>Global longitudinal strain rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>SV end-diastolic volume/m²</td>
<td>$-0.08$</td>
<td>$0.06$</td>
</tr>
<tr>
<td>SV cardiac index</td>
<td>$-0.25$</td>
<td>$-0.23$</td>
</tr>
<tr>
<td>$S'$ velocity</td>
<td>$-0.08$</td>
<td>$-0.22$</td>
</tr>
<tr>
<td>APSE</td>
<td>$-0.17$</td>
<td>$-0.10$</td>
</tr>
</tbody>
</table>

APSE annular plane systolic excursion, $S'$ systolic velocity, pulsed Tissue Doppler
window, the presence of a pacemaker device or claustrophobia.

Clinical Implications

The present study emphasizes that echocardiography and cMRI are complementary diagnostic tests to evaluate the cardiac function of TCPC patient. Our results provide little support that myocardial deformation by STE might have a role in the assessment of this patient group, at least at this relatively young age. Since our study was of cross-sectional design, we cannot exclude that STE might have a role in predicting single ventricle failure in follow-up studies and further investigations on this topic are warranted.

Table 4  Segmental lateral longitudinal strain and systolic SR, depending on type of dominant ventricle (mean ± SD)

<table>
<thead>
<tr>
<th></th>
<th>N</th>
<th>Longitudinal peak strain (%)</th>
<th></th>
<th>Longitudinal systolic SR (1/s.)</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>LV</td>
<td>RV</td>
<td>p value</td>
<td>LV</td>
</tr>
<tr>
<td>Basal-septal</td>
<td>50</td>
<td>16.5 ± 6.5</td>
<td>-14.6 ± 5.9</td>
<td>0.33</td>
<td>-0.93 ± 0.28</td>
</tr>
<tr>
<td>Mid-septal</td>
<td>63</td>
<td>18.6 ± 5.5</td>
<td>-19.6 ± 4.3</td>
<td>0.45</td>
<td>-1.02 ± 0.28</td>
</tr>
<tr>
<td>Apical-septal</td>
<td>62</td>
<td>21.0 ± 5.8</td>
<td>-21.9 ± 5.9</td>
<td>0.55</td>
<td>-1.18 ± 0.41</td>
</tr>
<tr>
<td>Apical-lateral</td>
<td>58</td>
<td>19.7 ± 5.6</td>
<td>-19.3 ± 5.8</td>
<td>0.83</td>
<td>-1.16 ± 0.35</td>
</tr>
<tr>
<td>Mid-lateral</td>
<td>56</td>
<td>19.3 ± 4.1</td>
<td>-19.7 ± 4.9</td>
<td>0.80</td>
<td>-1.04 ± 0.28</td>
</tr>
<tr>
<td>Basal-lateral</td>
<td>59</td>
<td>18.3 ± 4.7</td>
<td>-18.5 ± 4.1</td>
<td>0.89</td>
<td>-1.19 ± 0.52</td>
</tr>
</tbody>
</table>

LV left ventricle, RV right ventricle, SR strain rate
Conclusions

Myocardial function in contemporary TCPC patients with univentricular hearts is on average well preserved in children and young adults. Longitudinal myocardial deformation by STE, a novel non-invasive technique yielding relatively low costs, does not predict the presence of low EF by cMRI. Clinical utility is also hampered by moderate feasibility and variability of the current available techniques. Based on STE-derived parameters, the myocardial function in morphological RVs is similar compared to morphological LVs.

Acknowledgements The authors would like to thank Prof. Dr. E. Boersma, Department of Cardiology, Thorax Center, Erasmus Medical Center, Rotterdam, the Netherlands for his advice regarding the statistical analysis. This work was financially supported by the Dutch Heart Foundation (Grant number 2008T037).

Compliance with Ethical Standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical Approval All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Informed Consent Informed consent was obtained from all individual participants and/or their legal parents included in the study.

Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References


