



# Vortex formation time as an index of left ventricular filling efficiency: comparison between children volunteers and patients with tetralogy of Fallot

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**Background:** Vortex formation time (VFT) had been considered a useful marker for assessing diastolic performance. The VFT assessment of diastolic function using four-dimensional (4D) flow cardiovascular magnetic resonance (CMR) has not been used in repair of tetralogy of Fallot (rTOF) patient. The aims of this study were as follows: (I) establish reference ranges for VFT measurements in healthy children and adolescents using 4D flow CMR imaging; and (II) analyze VFT parameters to assess diastole dysfunction in rTOF patients group.

**Methods:** We acquired the CMR data was of 62 healthy participants (aged 6–18 years; male: 40, female: 22) and 20 patients with rTOF (aged 10–13 years; male: 15, female: 5) using 4D flow and cine sequence in routine chamber view. The VFT was calculated based on comparison of different algorithms from cine measurements ( $VFT_{\text{volume}}$ ) and 4D flow measurements ( $VFT_{\text{blood}}$ ). Then, VFT measurements were compared to subject peak filling rate (PFR), age, and cardiac mass using simple linear regression and multiple regression analyses. Data were also categorized according to age for VFT and cardiac functional assessment comparisons between 3 age groups (Group 1: 6–9 years; Group 2: 10–13 years; Group 3: 14–18 years). The correlation of VFT and cardiac function parameters were analyzed in the rTOF group.

**Results:** Normal mean value of  $VFT_{\text{volume}}$  and  $VFT_{\text{blood}}$  were  $4.25 \pm 0.92$  and  $3.77 \pm 1.11$  in healthy children participants. The  $VFT_{\text{volume}}$  was correlated with  $VFT_{\text{blood}}$  ( $r=0.61$ ,  $P<0.001$ ). There was a moderately significant correlation between  $VFT_{\text{volume}}$  and PFR ( $r=0.46$ ,  $P<0.001$ ) and between  $VFT_{\text{blood}}$  and PFR ( $r=0.47$ ,  $P<0.001$ ), age ( $r=0.41$ ,  $P=0.002$ ) and left ventricular (LV) mass ( $r=0.48$ ,  $P<0.001$ ). Multiple regression analyses demonstrated that  $VFT_{\text{volume}}$  was independently associated with PFR ( $T=2.239$ ;  $P<0.05$ ) and  $VFT_{\text{blood}}$  ( $T=4.361$ ;  $P<0.001$ ). There was a significant difference in  $VFT_{\text{volume}}$  between healthy controls and rTOF patients ( $5.44 \pm 1.93$  vs.  $4.27 \pm 0.88$ ,  $P=0.018$ ).

**Conclusions:** The VFT measurements showed that the LV that had appropriate space to form the optimal vortex ring in normal children and adolescents aged 6–18 years old. The  $VFT_{\text{volume}}$  could potentially be helpful in improving our understanding of LV diastolic dysfunction in rTOF patients.

**Keywords:** Cardiovascular magnetic resonance (CMR); 4D flow; vortex formation time (VFT); vortex flow; intra-cardiac blood flow patterns

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## Introduction

Vortex formation time (VFT), as originally defined by Gharib *et al.*, has been applied as an important index to characterize propulsion efficiency in cardiovascular disease (1). A normal VFT implied optimal formation of the transmitral vortex, thus ensuring an effective transfer of circulation impulse energy for efficient blood flow (2). The vortex rings generated from starting jets stop forming and pinched off from the generating jet for sufficiently ventricular filling. The open distance of the mitral valve determined value of VFT following equation (1). In recent years, several studies have shown that VFT reflects the quality of ventricular filling and is correlated with transmitral thrust and mitral annulus recoil, and in particular, transmitral vortex formation has been shown to play an important role in diastolic function (2-4). Therefore, VFT has been considered a useful marker for assessing diastolic performance (5,6).

Cardiovascular magnetic resonance (CMR) has been accepted as the “gold standard” for the assessment of left ventricular (LV) morphology and systolic function, but CMR assessment of LV diastolic function has been limited (7). Four-dimensional flow (4D flow) CMR was developed to provide a comprehensive assessment of blood flow through the heart and large vessels. This unique technique enabled both visualization and quantification of blood flow in the heart (8). Several authors reported that tricuspid flow and regurgitation can be assessed with good reproducibility and consistency using 4D flow CMR and echocardiography (9). Automated 4D flow analysis has improved the accuracy of peak mitral inflow velocity measurements and enabled tracking of the mitral valve plane motion (10). These studies provided the basis of evaluating VFT by 4D flow CMR.

Tetralogy of Fallot (TOF) is the most common cyanotic congenital heart disease. Early surgical intervention has improved survival rates of TOF patients, but complications such as pulmonary regurgitation (PR), right ventricle hypertrophy, and structural myocardial changes have continued to occur (11,12). To the best of our knowledge, the VFT assessment of diastolic function using 4D flow CMR has not been used in repair of TOF (rTOF) (1). Our previous study highlighted the importance of vortical

blood flow patterns for kinetic energy preservation in right ventricle of rTOF patients (13). This was the first study for evaluating LV diastolic dysfunction by VFT measurements using 4D flow CMR in healthy children and rTOF patients. Thus, the aims of this study were to (I) establish reference ranges for VFT measurements in healthy children and adolescents using 4D flow CMR imaging; and (II) analyze VFT parameters to assess diastole dysfunction in a rTOF patient group. We present the following article in accordance with the MDAR reporting checklist (available at <https://tp.amegroups.com/article/view/10.21037/tp-22-67/rc>).

## Methods

### Participants

We acquired CMR datasets of 62 healthy Chinese children and adolescents (aged 6–18 years; male: 40; female: 22) and 20 patients with rTOF (aged 10–13 years; male: 15, female: 5) from studies performed between June 2017 and April 2021. We recruited 6–18 years old participants by the network platform ([https://h.eqxiu.com/s/Mgzk8jcR?eqrcode=1&from=singlemessage&share\\_level=1&from\\_user=2022032551a044a1&from\\_id=61704003-9&share\\_time=1648518974336](https://h.eqxiu.com/s/Mgzk8jcR?eqrcode=1&from=singlemessage&share_level=1&from_user=2022032551a044a1&from_id=61704003-9&share_time=1648518974336)). All volunteers were healthy children who had normal blood pressure and were free of cardiovascular disease as determined by echocardiography. Children with a family history of any other form of heart disease and obesity were excluded. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by research ethics committee board of Shanghai Children’s Medical Center (No. SCMCIRB-K2017062). Informed consent was taken from all the participants’ guardians. The study sample median age was 13.55 years, encompassing 3 different age groups: (Group 1, 6–under 10 years old, 18 cases; Group 2, over 10–under 14 years old: 30 cases; Group 3, 14–18 years old: 14 cases). The data of healthy controls were acquired without contrast-medium administration.

The inclusion criteria for 20 patients with rTOF were as follows: (I) surgery by pulmonary transannular patching; (II) diagnosis of impaired diastolic function (grade 1 or grade 2) according to the criteria described in the 2016 American Society of Echocardiography guidelines (14); (III) patients

with mild-moderate right ventricular (RV) dilation [RV end-diastolic volume (EDV) index  $<150 \text{ mL/m}^2$ ] and significant PR ( $>20\%$ ); and (IV) LV diastolic dysfunction [early and/or late peak diastolic mitral inflow velocity (E/A)  $<1$  or early peak mitral inflow velocity and/or mean mitral annular peak early diastolic velocity (E/E')  $>13$ ]. The exclusion criteria were as follows: rTOF patients with mitral regurgitation or left atrial enlargement (left atrial maximum volume index  $>34 \text{ mL/m}^2$ ).

Intravenous bolus injection of  $0.1 \text{ mmol/kg}$  gadopentetate dimeglumine was administered to rTOF patients within 15–20 minutes before 4D-flow imaging.

### CMR imaging

We conducted CMR on a 3.0 Tesla scanner (Discovery MR 750, GE, Chicago, IL, USA) using an 8-channel phased-array cardiac coil. Whole heart 4D flow CMR was performed [spatial resolution  $= (1.2\text{--}1.8) \times (1.2\text{--}1.8) \times (1.2\text{--}1.8) \text{ mm}^3$ , field of view (FOV)  $= 340\text{--}420 \text{ mm}^2$ , slab thickness  $= 60\text{--}85 \text{ mm}$ , temporal resolution  $= 18.2\text{--}34.4 \text{ ms}$ , echo time (TE)  $= 2.1 \text{ ms}$ , repetition time (TR)  $= 4.3 \text{ ms}$ , flip angle  $= 8\text{--}12^\circ$ , bandwidth  $= 62.5 \text{ Hz/pixel}$ , and velocity sensitivity  $= 120\text{--}160 \text{ cm/s}$ ] with acquisition volume covering the entire heart. The scan time was 6–8 minutes. We followed the parameter settings of protocol according to 4D flow CMR consensus statement (8). Due to the smaller FOV, higher spatial resolution was recommended in a pediatric population. Retrospective electrocardiogram (ECG)-gating was used and 20–30 cardiac phases were reconstructed to represent 1 average heartbeat. Free breathing was allowed and no respiratory motion compensation was used. We also acquired 2-, 3-, 4-chamber, and short-axis cine two-dimensional balanced steady state free precession (2D b-SSFP) images using standard clinical protocols.

### Image post-processing analysis

#### 2D velocity vector field visualization

The 4D flow data from the 62 volunteers were used for 2D velocity vector field visualization in LV outflow tract view using pre-process software (MASS Version 30 March 2020; Leiden University Medical Center, Leiden, The Netherlands) (Figure 1).

We used 2 different strategies for assessment of VFT. The first strategy (Strategy 1) was based on ventricular volume measurements performed using the 2D cine b-SSFP

CMR data. The second strategy (Strategy 2) used 4D flow CMR data for calculation of the VFT (15).

#### Strategy 1

The VFT was calculated by the following equation:

$$VFT_{\text{volume}} = \frac{4}{\pi} \cdot \alpha^3 \cdot EF \cdot \frac{EWW}{SV} = \frac{4}{\pi} \cdot \frac{EWW}{D^3} \quad [1]$$

where,  $\alpha$  was defined as:

$$\alpha = \frac{EDV^{\frac{1}{3}}}{D} \quad [2]$$

with the open distance of the mitral valve,  $D$ , calculated as the mean of the maximum distance between the leaflet tips during the rapid filling phase, perpendicular to the flow direction, in the three-chamber view, and the distance between the commissures on a short-axis slice through the mitral valve (Figure 2). The E-wave volume (EWW) was defined as the difference in volumes of the LV at the end of the E-wave and ESV.

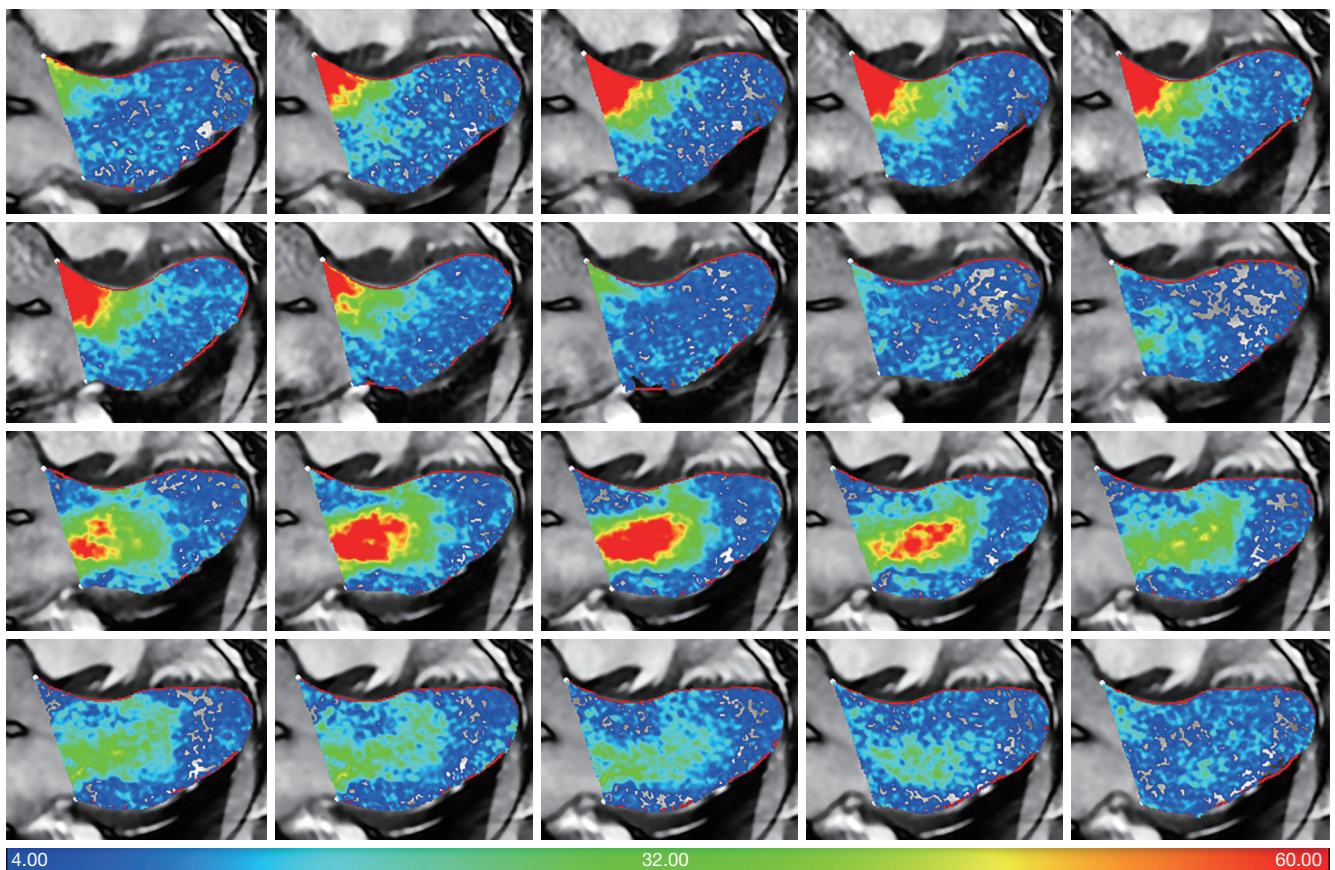
#### Strategy 2

The trans-mitral valve velocities and time to peak of E-wave and A-wave were determined from the 4D flow CMR data by semi-automatic segmentation using commercial postprocessing software (Version 5.1; Pie Medical Imaging, Maastricht, The Netherlands) by a single observer with over 3 years of experience in CMR. These measurements were verified by a radiologist with over 20 years of experience in CMR. Excursion of the mitral valve was identified on the LV 2-chamber, 4-chamber 2D cine b-SSFP images. By using the valve-tracking algorithms, measurement planes corrected for aliasing, velocity offset, and through-plane motion were obtained for mitral valve via multiplanar reformatting (Figure 3). Once obtained, these planes provided instantaneous velocity and time-averaged velocity measurements within the mitral valve.

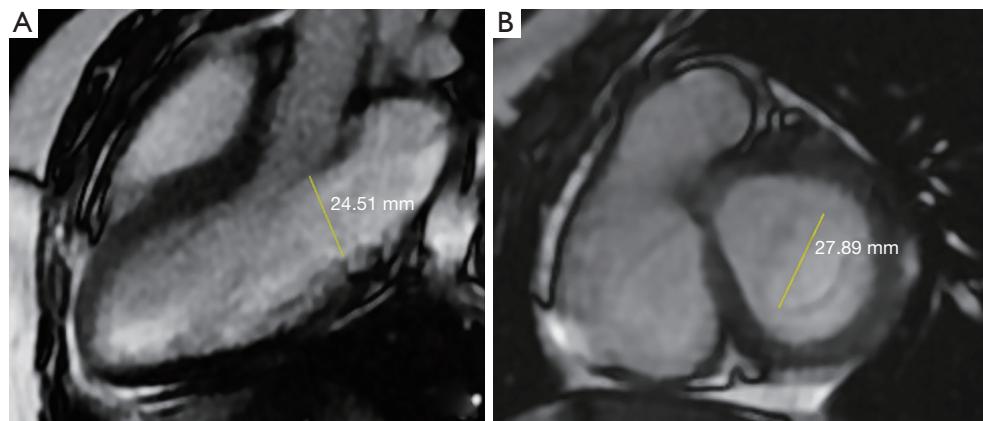
The peak filling rate (PFR), EDV, end-systolic volumes (ESV), stroke volume (SV), and ejection fraction (EF) of the LV were determined from short-axis 2D cine b-SSFP images using an automatic segmentation method based on a deep learning algorithm. Papillary muscles and trabeculations were included within the LV blood pool. We assessed ventricular volumes normalized to body surface area (BSA) and Z score based on multivariable models with the square of height, height, weight, and age (16).

We analyzed VFT parameters between age matched subgroup of healthy participants and the rTOF patients group (Table 1). Post-processing procedure of vortex ring

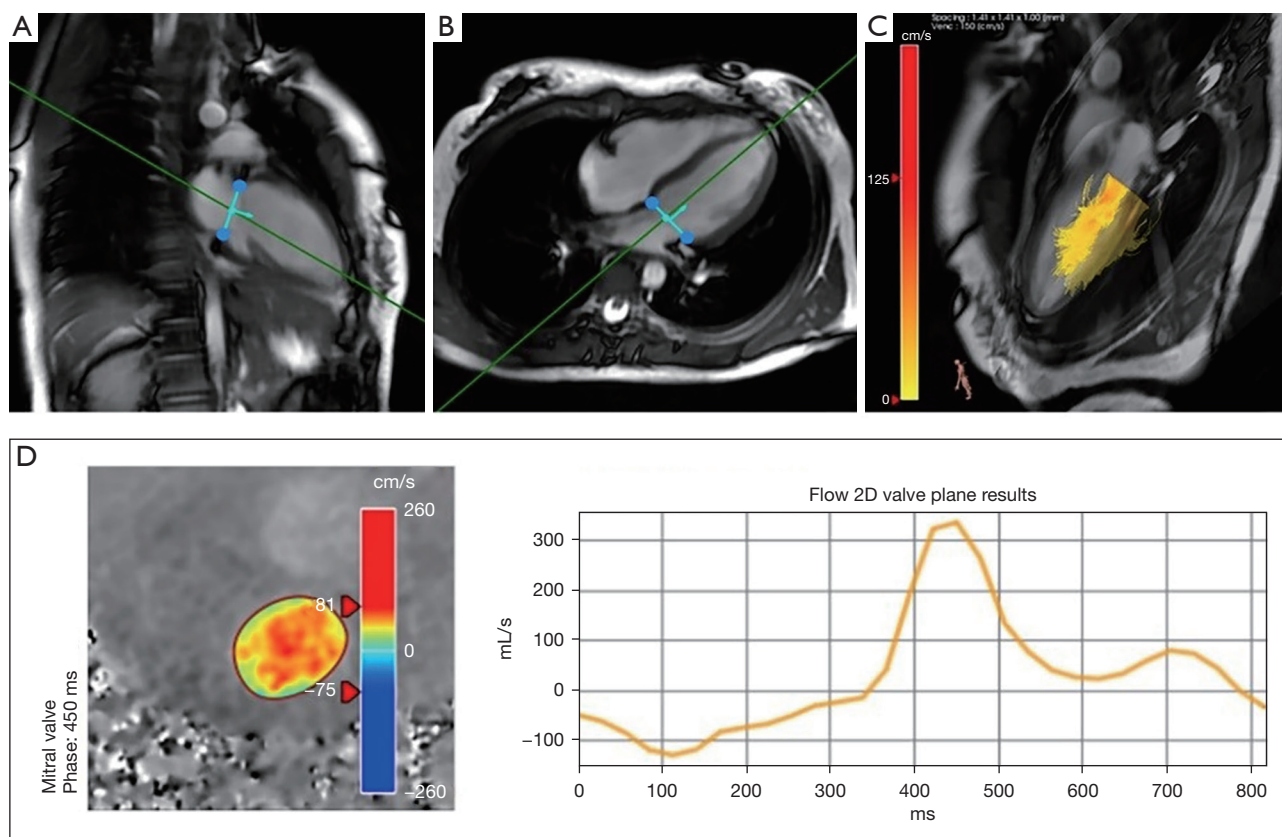




**Figure 1** Representative diastolic left ventricular visualizations in control using 4D CMR flow. The color bar represents 2D velocity (cm/s) vector field visualization. CMR, cardiovascular magnetic resonance.



**Figure 2** The mitral valve opening distance was calculated by the average of the distance between the commissures seen in the short axis slice (B) of the mitral valve in the 3-chamber view (A), in the rapid filling phase (E wave), perpendicular to the flow direction, the maximum observation distance between the leaflet tips (E wave), and in the short axis slice (B).



**Figure 3** Line graph showing blood flow through the mitral valve during a single cardiac cycle in healthy child volunteer measured using 4D CMR flow sequence. (A) Valve-tracking in two chamber view; (B) valve-tracking in four chamber view; (C) streamline map in the LV; (D) diastolic phase includes early filling, diastole and late filling. CMR, cardiovascular magnetic resonance; LV, left ventricular.

parameters are described in the additional file [Figure S1](#).

### Statistical analysis

Statistical analysis was performed using GraphPad Prism version 6.0 (GraphPad Inc., San Diego, CA, USA). Continuous data were reported as mean  $\pm$  standard deviation. Categorical data were reported as numbers with percentages. We calculated the correlation between VFT and age, heart rate, PFR, time-averaged velocity, and open distance of mitral orifice. Correlation was evaluated using simple linear regression analysis with Pearson  $r$ -values calculated for normally distributed data and Spearman  $r$ -values for non-normally distributed continuous variables. Correlations were categorized as follows: 0.95–0.80, strong; 0.80–0.60, good; 0.60–0.40, moderate; and less than 0.40, poor. Multiple regression analyses were performed between  $VFT_{\text{volume}}$  and age, heart rate, PFR, cardiac mass, and  $VFT_{\text{blood}}$ . A generic linear relationship was calculated

between  $VFT_{\text{volume}}$  and  $VFT_{\text{blood}}$ . Comparisons among 3 groups were performed using one-way analysis of variance (ANOVA) for normally distributed data and Kruskal-Wallis tests for non-normally distributed continuous variables. Comparison of continuous variables between the rTOF group and volunteer group was performed using an independent-samples  $t$ -test for normally distributed data, and Mann-Whitney U test for non-normally distributed data. Statistical significance was indicated by  $P < 0.05$ .

### Intra-observer and inter-observer reproducibility analysis

The only source of potential observer variability was the user segmentation of the mitral valve. The remaining steps for 2D cine b-SSFP and 4D flow CMR VFT estimations were fully automated. For evaluation of reproducibility, mitral valve segmentation for computation of VFT using the above-mentioned strategies was performed for a subgroup of 20 randomly selected participants, in a random



**Table 1** VFT analysis with respect to both their differences between age-matched healthy volunteers and patients

Parameter	rTOF group	Volunteer group	P value
Age (years)	11.85±1.27	12.01±1.27	0.585
N (case)	20	20	
Gender (male/female)	15/5	11/9	
Heart rate (beat/min)	76±11	75±13	0.823
LV EDV indexed to BSA (mL/m <sup>2</sup> )	85.18±22.13	80.08±12.19	0.499
LV ESV indexed to BSA (mL/m <sup>2</sup> )	35.83±9.16	30.12±6.03	0.026*
LV stroke volume indexed to BSA (mL/m <sup>2</sup> )	45.95±6.97	50.27±7.89	0.199
LV EF (%)	57.90±4.96	62.55±4.67	0.004*
VFT <sub>volume</sub>	5.44±1.93	4.27±0.88	0.018*
VFT <sub>blood</sub>	3.75±0.81	4.26±0.94	0.078

Data were expressed as mean ± standard deviation. \*, statistically significant differences. VFT, vortex formation time; rTOF, repair of tetralogy of Fallot; LV EDV, left ventricular end-diastole volume; BSA, body surface area; LV ESV, left ventricular end-systolic volume; EF, ejection fraction.

order by 2 independent operators (with 3 years' experience in CMR image processing), both blinded to participant's characteristics and quantitative results. Both the open distance of the mitral orifice and VFT measurements for the separate segmentations were compared with calculations of bias, intraclass correlation coefficient (ICC), and coefficient of variation (CoV).

## Results

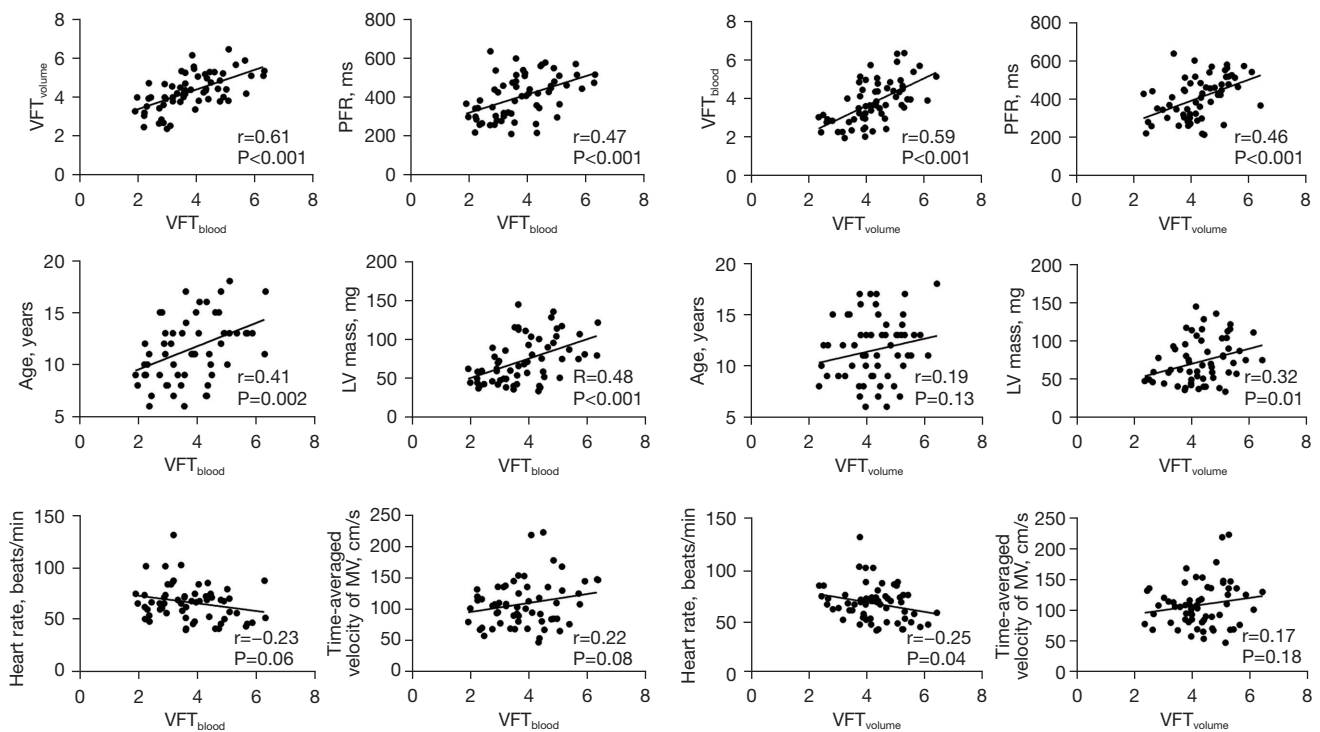
### *Comparisons between VFT measurements and clinical physiological parameters*

Normal mean value of VFT<sub>volume</sub> and VFT<sub>blood</sub> were 4.25±0.92 and 3.77±1.11 in healthy child participants. The VFT<sub>volume</sub> measurements demonstrated good correlation to VFT<sub>blood</sub> measurements ( $r=0.61$ ,  $P<0.001$ ). There was significant correlation between VFT<sub>volume</sub> and PFR ( $r=0.46$ ,  $P<0.001$ ), VFT<sub>volume</sub> and heart rate ( $r=-0.25$ ,  $P=0.04$ ), and VFT<sub>volume</sub> and LV mass ( $r=0.32$ ,  $P=0.01$ ). There was no significant correlation between VFT<sub>volume</sub> and age ( $r=0.19$ ,  $P=0.13$ ), or VFT<sub>volume</sub> and time-averaged velocity of mitral orifice ( $r=0.17$ ,  $P=0.18$ ). There were moderate but significant correlations between VFT<sub>blood</sub> and PFR ( $r=0.47$ ,  $P<0.001$ ), VFT<sub>blood</sub> and age ( $r=0.41$ ,  $P=0.002$ ), and VFT<sub>blood</sub> and LV mass ( $r=0.48$ ,  $P<0.001$ ). There was no significant correlation between VFT<sub>blood</sub> and heart rate ( $r=-0.23$ ,  $P=0.06$ ), and VFT<sub>blood</sub> and time-averaged velocity of MV

( $r=0.22$ ,  $P=0.08$ ) (Figure 4).

### *Age-dependence of cardiac function and VFT measurements*

The LV EDV, LV ESV, and LV SV were significantly increased with age in the 3 different age groups ( $P<0.001$ , Table 2). Normalized LV volumes and function by BSA and Z score were not significantly different with age in the 3 different age groups ( $P>0.05$ , Table 2). The E wave volume (EWW) also significantly increased with age (47.62±11.37 vs. 61.51±16.07 vs. 67.81±11.67 mL,  $P<0.001$ , Table 2). The open distance of the mitral orifice significantly increased with age (24.87±2.25 vs. 26.32±2.68 vs. 27.31±2.27 mm,  $P<0.05$ , Table 2). The average velocity of the mitral orifice also demonstrated significant differences between the 3 age groups (88.32±26.69 vs. 110.95±34.46 vs. 114.44±23.51 cm/s,  $P<0.05$ , Table 2). The VFT<sub>volume</sub> measurements were not significantly different between the age groups (3.96±0.77 vs. 4.34±1.02 vs. 4.32±0.99,  $P>0.05$ ). However, VFT<sub>blood</sub> measurements were significantly different between the 3 groups (3.23±0.84 vs. 3.93±1.24 vs. 4.15±0.95,  $P<0.05$ ) (Figure 5). The relationship was calculated between age and VFT by 2 methods (Figure 6). Multiple regression analyses demonstrated that VFT<sub>volume</sub> was independently associated with PFR ( $T=2.239$ ;  $P<0.05$ ) and VFT<sub>blood</sub> ( $T=4.361$ ;  $P<0.001$ ) in Table 3. A generic linear relationship was calculated between VFT<sub>volume</sub> and VFT<sub>blood</sub> ( $VFT_{volume}=0.49 VFT_{blood}+3.39$ ,  $P<0.001$ ) (Figure 7).



**Figure 4** Comparisons between VFT measurements and clinical physiological parameters. VFT, vortex formation time; PFR, peak filling rate; MV, mitral valve; LV, left ventricular.

### *VFT and MV vortex ring parameters in healthy and rTOF participants*

Both LV end systolic volume index (ESVi) and LV EF were significantly different with age among healthy controls and patients with rTOF ( $P < 0.05$ ). The  $VFT_{\text{volume}}$  was significantly different between healthy controls and patients with rTOF ( $5.44 \pm 1.93$  vs.  $4.27 \pm 0.88$ ,  $P = 0.018$ ). The  $VFT_{\text{blood}}$  was not significantly different between healthy controls and patients with rTOF ( $3.75 \pm 0.81$  vs.  $4.26 \pm 0.94$ ,  $P > 0.05$ , *Table 1*). The  $VFT_{\text{volume}}$  was significantly correlated with LV EDVi, LV ESV, and cardiac mass in the rTOF patients group ( $P > 0.05$ , *Table 4*).

The maximal vorticity of LV blood flow in the rTOF patient group was larger than that in the healthy volunteer group during E wave phase ( $29.3$  vs.  $25.9$   $s^{-1}$ ). The average vorticity of LV blood flow in the patient group was less than that in the volunteer group during E wave phase ( $11.7$  vs.  $14.9$   $s^{-1}$ ). The maximal vorticity of LV blood flow in the patient group was less than in the volunteer group during A wave phase ( $10.7$  vs.  $12.2$   $s^{-1}$ ). The average vorticity of LV blood flow in the patient group was less than in the volunteer group during A wave phase ( $2.1$  vs.  $2.3$   $s^{-1}$ ).

(*Figure S1* in supplementary material).

### *Reproducibility*

*Table 5* summarizes the results of the intra-observer and inter-observer analysis for mitral orifice open distance and VFT measurements. Intra-observer and inter-observer agreement were excellent for  $VFT_{\text{volume}}$ ,  $VFT_{\text{blood}}$  and open distance of mitral orifice (all ICCs  $> 96\%$ , CoV  $< 5.3$  for intra-observer variability; all ICCs  $> 88\%$ , CoV  $< 11.56$  for inter-observer variability).

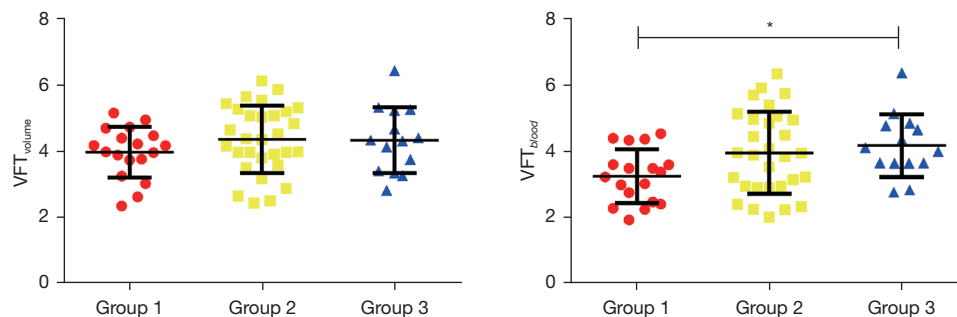
### **Discussion**

To the best of our knowledge, this was the first study to evaluate VFT measurements of mitral valve flow using 4D flow CMR in healthy children and rTOF patients. Previous investigators had highlighted the importance of vortical blood flow patterns for kinetic energy preservation (2). Other researchers (11,17) had reported 4D flow CMR for quantitative analysis of the LV vortex ring during early and late ventricular filling. We found LV vortex flow during

**Table 2** Cardiac physiology and VFT measurements

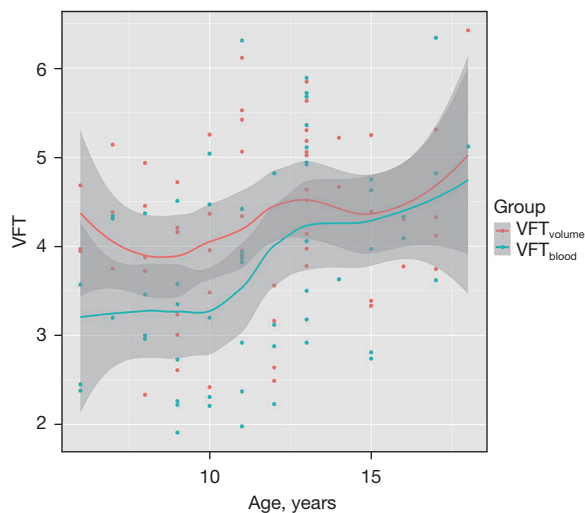
Parameter	Group 1	Group 2	Group 3	P value
Age (years)	6–9	10–13	14–18	
N (case)	18	30	14	
Gender (male/female)	15/3	14/16	9/5	
Heart rate (beat/min)	86±17	78±13	67±10	0.001*
Body surface area (m <sup>2</sup> )	1.08±0.20	1.39±0.19	1.65±0.94	<0.001*
LV EDV (mL/m <sup>2</sup> )	80.16±14.55	106.36±23.54	126.81±20.35	<0.001*
LV ESV (mL/m <sup>2</sup> )	30.47±8.27	39.63±10.18	51.58±12.73	<0.001*
LV stroke volume (mL/m <sup>2</sup> )	49.92±9.59	67.09±16.55	75.22±10.03	<0.001*
LV EDV indexed to BSA (mL/m <sup>2</sup> )	76.65±10.45	76.51±13.01	76.75±10.32	0.997
LV ESV indexed to BSA (mL/m <sup>2</sup> )	28.25±6.23	28.54±6.34	31.09±6.55	0.388
LV stroke volume indexed to BSA (mL/m <sup>2</sup> )	48.38±5.88	48.16±9.45	45.65±5.94	0.554
LV EDV indexed to Z score (mL/m <sup>2</sup> )	74.82±10.93	76.48±12.99	76.75±10.41	0.869
LV ESV indexed to Z score (mL/m <sup>2</sup> )	28.26±6.23	28.54±6.33	31.09±6.57	0.391
LV stroke volume indexed to Z score (mL/m <sup>2</sup> )	46.71±7.46	48.16±9.43	45.65±5.99	0.619
LV EF (%)	63.45±4.65	62.68±6.06	59.71±4.82	0.136
Open distance of mitral orifice (mm)	24.87±2.25	26.32±2.68	27.31±2.27	0.02*
Average velocity of mitral inflow (cm/s)	88.32±26.69	110.95±34.46	114.44±23.51	0.01*
EWV (mL)	47.62±11.37	61.51±16.07	67.81±11.67	<0.001*
VFT <sub>volume</sub>	3.96±0.77	4.34±1.02	4.32±0.99	0.19
VFT <sub>blood</sub>	3.23±0.84	3.93±1.24	4.15±0.95	0.03*

Data were expressed as mean ± standard deviation. \*, statistically significant differences denoted. Group 1: 6–9 years; Group 2: 10–13 years; Group 3: 14–18 years. VFT, vortex formation time; LV EDV, left ventricular end-diastole volume; LV ESV, left ventricular end-systolic volume; BSA, body surface area; EF, ejection fraction; EWV, E wave volume.



**Figure 5** VFT measurements performed with volume (A) and blood flow (B) based measurement strategies for each of the 3 different age groups (Group 1: 6–9 years; Group 2: 10–13 years; Group 3: 14–18 years). \*, indicates statically significant difference between VFT<sub>blood</sub> measurements. VFT, vortex formation time.



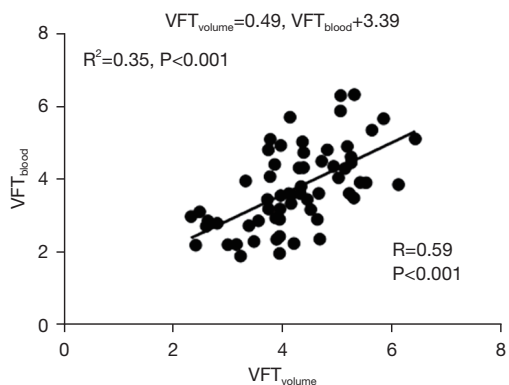


**Figure 6** The relationship between age and vortex formation time by 2 methods including 95% CI. VFT, vortex formation time; CI, confidence interval.

**Table 3** Multiple regression analyses of VFT and left cardiac function in healthy participants

Parameter	VFT <sub>volume</sub>	
	T value	P value
Age	-1.217	0.228
Heart rate	-1.157	0.252
Peak filling rate	2.239	0.03*
Cardiac mass	-0.377	0.708
VFT <sub>blood</sub>	4.361	<0.001*

\*, statistically significant differences. VFT, vortex formation time.



**Figure 7** A generic linear relationship (VFT<sub>volume</sub> and VFT<sub>blood</sub>) indicated by the solid black line. VFT, vortex formation time.

early and late ventricular filling in rTOF patients and control subjects can be imaging by 4D CMR flow and be visualized using postprocessing in software. In our study, VFT<sub>volume</sub> and VFT<sub>blood</sub> were calculated based upon CMR to offer the additional potential intracardiac markers (8).

**VFT as index of diastolic function**

Pinched-off vortex rings have been shown to be optimal for efficient blood flow within the LV and contribute to the distribution of flow-related stress loads. Previous studies had shown that vortex formation time could be affected by (I) time-dependent velocity of the fluid flow; (II) the Reynolds number of flow (18); and (III) the shape of the mitral orifice (19). In our study, VFT<sub>blood</sub> was calculated based on an equation that included the above factors. The VFT<sub>blood</sub> was found to be positively correlated with VFT<sub>volume</sub> measured based upon ventricular volume measurements. This positive correlation between VFT<sub>volume</sub> and VFT<sub>blood</sub> strongly supported the robustness of VFT measurement. We detected a generic linear relationship, as indicated by the solid black line in Figure 7. This formula was helpful to correct the variability of VFT<sub>blood</sub>. In a previous study, Kovács *et al.* (20) demonstrated that vortex formation was causally connected to rapid filling, which is a known consequence of overall diastolic function characterized by load, relaxation, and stiffness. The PFR was determined from ventricular time-volume curves obtained from CMR imaging and could be calculated as an indicator of diastolic function (21). In our study, multiple regression analyses demonstrated that VFT<sub>volume</sub> was independently associated with PFR and VFT<sub>blood</sub>, respectively. At present, the clinical significance of markers as surrogate is unknown for ventricular diastolic function and blood flow momentum (22). Matsuura *et al.* demonstrated the close relationship between vortex and intraventricular pressure difference and showed both of them can become new markers of the left ventricular relaxation property (23). Further research is needed to better understand intracardiac flow mechanisms, particularly in children and adolescents.

**VFT<sub>volume</sub> and VFT<sub>blood</sub> in children and adolescents**

Gharib *et al.* measured the VFT for healthy adults (>20 years old) and found an optimal value of VFT in a narrow range between 3.5 and 5.5. A negative correlation between age and VFT was shown in their

**Table 4** The correlation of VFT and left cardiac function in rTOF patients

LV function	VFT <sub>volume</sub>		VFT <sub>blood</sub>	
	r value	P value	r value	P value
LV EDV indexed to BSA (mL/m <sup>2</sup> )	0.546	0.012*	0.087	0.714
LV ESV indexed to BSA (mL/m <sup>2</sup> )	0.595	0.005*	-0.027	0.911
LV stroke volume indexed to BSA (mL/m <sup>2</sup> )	0.106	0.657	0.378	0.101
LV cardiac mass (g)	0.685	0.002*	-0.172	0.468

\*, statistical differences. VFT, vortex formation time; rTOF, repair of tetralogy of Fallot; LV EDV, left ventricular end-diastole volume; BSA, body surface area; LV ESV, left ventricular end-systolic volume.

**Table 5** Intra-observer and inter-observer variability for mitral orifice open distance and VFT measurements

Parameter	Intra-observer (n=20)			Inter-observer (n=20)		
	Bias (limits of agreement)	ICC	CoV	Bias (limits of agreement)	ICC	CoV
Mitral orifice open distance (mm)	0.17 (-0.91 to 1.26)	98.5%	2.21	0.33 (-1.15 to 1.81)	97.3%	3.15
VFT <sub>volume</sub>	-0.11 (-0.55 to 0.33)	96.8%	5.29	-0.03 (-0.69 to 0.63)	93%	7.56
VFT <sub>blood</sub>	0.11 (-0.26 to 0.47)	97.4%	5.30	0.19 (-0.56 to 0.93)	88.9%	11.56

VFT, vortex formation time; ICC, intraclass correlation coefficient; CoV, coefficient of variation.

measurements (1). Previous studies reported that age was associated with a reduction in LV filling efficiency quantified using VFT, and adverse or negative correlation was found between age and VFT using linear regression analysis (24-26). Based on this negative correlation, it could be speculated that the value of VFT should be between 6 and 8 for teenagers. However, our results were inconsistent with latter speculation. In our study, VFT<sub>volume</sub> did not exhibit significant differences between age groups and VFT<sub>volume</sub> approached the previously determined optimal VFT of 4.0. Nonetheless, our findings might be consistent with known processes of growth given that the reduction of the intraventricular pressure gradients occurs at the same time as the increase of mitral valve area (27). In theory, the vortex rings in the LV of teenagers should still satisfy the optimal principle of VFT with these vortex rings optimized for efficient fluid transport. Our findings based upon VFT<sub>volume</sub> also suggested that the LV had appropriate space to form the optimal vortex ring in normal children and adolescents from 6 to 18 years old.

By using VFT<sub>blood</sub> obtained by 4D flow CMR, the volunteer cohort data showed that VFT<sub>blood</sub> was in optimal range for Group 2 and Group 3, but below the previously determined optimal range in Group 1. This phenomenon

might imply that the blood transport in the LV for older children (>10 years old) was more efficient than for younger children (<10 years old). Our results suggested that the normal mean values of VFT<sub>volume</sub> and VFT<sub>blood</sub> in children are lower than the adults' normal range. Kulkarni *et al.* (28) indicated that VFT was attenuated in neonates and infants and increased to the adults' normal range by adolescence. Their explanation was that in children (<10 years old) LV has a low peak torsion, untwisting velocity, smaller dimensions, and faster heart rates compared to adults. In our study, VFT<sub>volume</sub> measurements were not significantly different between the age groups. Our results deviated from the literature due to the lack of neonates and infants, and further study is required. To sum up, the results from calculation of VFT<sub>volume</sub> and VFT<sub>blood</sub> presented that the value of vortex rings in the LV for children over 10 years old were consistent with previous determined unified optimal formation number (VFT ≈ 4).

#### **Clinical applications of VFT<sub>volume</sub> and VFT<sub>blood</sub> in rTOF**

Previous studies have reported that peak systolic LV kinetic energy was decreased in rTOF patients. Unlike

systolic kinetic energy, VFT is a more sensitive indicator for assessment of diastolic function (29). The abnormal flow has been associated with increased viscous energy loss and reduced LV function in pediatric TOF patients (30). We also quantified VFT in a pilot application in controls with rTOF patients. The  $VFT_{\text{volume}}$  of participants was statistically significant compared with the aged-matched subgroup of controls. We explained that right ventricular dilation leads to left ventricular diastolic insufficiency, the trans-mitral jet velocity was disproportionately increased, which leads to higher VFT. Diastolic dysfunction of LV disrupted the interaction between the vortex and ventricular wall. The trans-mitral vortex pinch-off tends to occur in infinity, or alternatively may never occur. In this situation, the blood flow momentum was mainly transferred through the jet that led to high-energy dissipation and incoherent flow structures. Previous investigations have indicated that there are significant reductions in systolic kinetic energy and altered distribution of hemodynamic forces in rTOF patients (31). In our study,  $VFT_{\text{volume}}$  was significantly correlated with cardiac mass in the rTOF patients group.

### **Reproducibility of VFT<sub>volume</sub> and VFT<sub>blood</sub>**

The ICC and CoV for  $VFT_{\text{volume}}$ ,  $VFT_{\text{blood}}$  and mitral orifice open distance are summarized in *Table 5* for the intra-observer and inter-observer variability study. Mitral orifice opening distance showed a higher ICC and CoV than  $VFT_{\text{volume}}$  and  $VFT_{\text{blood}}$ . However, there was still good reproducibility for both  $VFT_{\text{volume}}$  and  $VFT_{\text{blood}}$  in our study.

### **Limitations**

There were some limitations to this study. Firstly, this was a single-center prospective study performed at a large hospital. The inherent limitations of this study design cannot be avoided. Secondly, Rutkowski *et al.* (32) demonstrated a gender-based cardiac efficiency relationship between cardiac function and flow. Gender difference may also be one of the factors affecting VFT, which has not been reported. Our study did not consider the effect of gender upon VFT as a larger cohort would be needed to test this gender difference. Thirdly, the mitral orifice opening distances were determined from CMR data using semi-automatic segmentation. In general, mitral annulus morphology for size and shape was addressed adequately by as few as 6 radial slices (33). We calculated opening distance of the mitral valve only by 3-chamber view and short axis

view. Additional studies in the future may be required with direct tracing of the mitral orifice to further improve the accuracy of VFT calculations. Finally, according to our implemented VFT equation, the definition should be mass/energy supply ratio rather than a physical VFT. The total kinetic energy of a vortex ring consisted of not only the translational energy associated with impulse and velocity but also the rotational energy associated with vorticity distribution. Future prospective, large-scale, multicenter investigations are needed to verify the prognostic value of  $VFT_{\text{volume}}$  in rTOF patients.

### **Conclusions**

The VFT measurements showed that the LV had appropriate space to form the optimal vortex ring in normal children and adolescents aged 6 to 18 years old. The  $VFT_{\text{volume}}$  could potentially be helpful in improving our understanding of LV diastolic dysfunction in rTOF patients.

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### **Footnote**

*Reporting Checklist:* The authors have completed the MDAR reporting checklist. Available at <https://tp.amegroups.com/article/view/10.21037/tp-22-67/rc>

*Data Sharing Statement:* Available at <https://tp.amegroups.com/article/view/10.21037/tp-22-67/dss>

*Conflicts of Interest:* All authors have completed the ICMJE uniform disclosure form (available at <https://tp.amegroups.com/article/view/10.21037/tp-22-67/coif>). All authors report technical assistance from Gaston Vogel and Qiuchen Wang (Pie Medical Imaging Corporation Netherlands), and statistical assistance from Tingfan Wu (GE Healthcare). The authors have no other conflicts of interest to declare.

*Ethical Statement:* The authors are accountable for all

aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by research ethics committee board of Shanghai Children's Medical Center (No. SCMCIRB-K2017062). Informed consent was taken from all the participants' guardians.

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