

Original Article

Evaluation of right ventricular function in patients with tetralogy of Fallot using the myocardial performance index and isovolumic acceleration: a comparison with cardiac magnetic resonance imaging

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Abstract *Background:* Assessment of right ventricular function is a key point in the follow-up of operated patients with tetralogy of Fallot. Cardiac magnetic resonance assessment of right ventricular function is considered the gold standard. However, this technique is expensive, has limited availability, and requires significant expertise to acquire and interpret the images. Myocardial performance index and isovolumic acceleration have recently been studied for the assessment of right ventricular function and are shown to be simple yet powerful tools for assessing patients with right ventricular dysfunction of various origins. *Methods:* In this study, the integrity of myocardial performance index and isovolumic acceleration obtained by tissue Doppler imaging echocardiography to quantify right ventricular function was assessed in 31 patients operated for tetralogy of Fallot. Myocardial performance index and isovolumic acceleration measurements were compared with the parameters derived by cardiac magnetic resonance imaging. *Results:* In this study, a significant correlation has not been detected between cardiac magnetic resonance-originated right ventricular ejection fraction, pulmonary regurgitation fraction and myocardial performance index, isovolumic acceleration obtained by tissue Doppler imaging echocardiography from the lateral tricuspid annulus of the right ventricle. *Conclusion:* We have concluded that when evaluated separately, myocardial performance index and isovolumic acceleration obtained from tissue Doppler imaging echocardiography can be used in the long-term follow-up of patients who have been operated for tetralogy of Fallot, but that they do not show correlation with cardiac magnetic resonance-originated right ventricle ejection fraction and pulmonary regurgitation fraction.

Keywords: Tetralogy of Fallot; right ventricle function; tissue Doppler imaging; cardiac magnetic resonance; myocardial performance index; isovolumic acceleration

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TETRALOGY OF FALLOT IS THE MOST COMMON FORM of cyanotic congenital heart disease, with a prevalence of 0.26–0.8 per 1000 live births.¹ Total repair for tetralogy of Fallot has been available for 55 years with a favourable outcome in most

patients. Late survival has also improved, with reports showing a 20-year survival rate nearing 90%.^{2,3} Today, we are faced with an increasing number of adult patients who require regular follow-up for complications after initial correction of tetralogy of Fallot. These complications mostly consist of pulmonary regurgitation, residual or recurrent pulmonary stenosis, ventricular septal defect, or right ventricular outflow tract aneurysm.

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Echocardiography provides information on the presence of right ventricular dilatation and/or hypertrophy, the presence of pulmonary regurgitation, and by Doppler estimation of right ventricle systolic pressure. However, assessing the function of the morphological right ventricle is challenging because of its complex anatomy. Difficulties are compounded by irregularities in the ventricular cavities and abnormalities in wall motion in patients with congenital heart lesions. None of the geometric assumptions used to assess left ventricular function hold true for the systemic right ventricle. The complex shape of the right ventricle makes quantification difficult.⁴ Thus, in clinical scenarios, many centres rely on visual estimation of right ventricle systolic function, which is then subject to variability because of incomplete visualisation of the entire right ventricle and the experience of the observer. Therefore, cardiac magnetic resonance has evolved to be a better quantitative standard, especially for serial comparisons.⁵ However, this technique, which is contraindicated in some patients, is expensive, has limited availability, and requires significant resources and expertise to acquire and interpret the images.

The tissue Doppler imaging-derived systolic myocardial velocities are considered more useful compared with conventional echocardiography in the assessment of the right ventricle contractile function.⁶ In the literature, a limited number of studies have shown the potential usefulness of the tissue Doppler imaging-derived myocardial performance index (Tei index) and isovolumic acceleration in the evaluation of right ventricular function in post-operative tetralogy of Fallot patients.⁷ However, no research study has been performed yet to compare cardiac magnetic resonance and tissue Doppler imaging echocardiography in post-operative tetralogy of Fallot patients.

Thus, the aim of this study was to compare tissue Doppler imaging assessment of right ventricular function measuring the myocardial performance index and isovolumic acceleration values with cardiac magnetic resonance findings.

Methods

Study design and patients

This cross-sectional study was conducted at the Unit of Pediatric Cardiology of İstanbul University Cerrahpaşa Medical Faculty, İstanbul, Turkey. Before recruitment of patients and controls, the study protocol was reviewed and approved by the local ethics committee, in accordance with the ethical principles for human investigations, as

outlined by the Second Declaration of Helsinki, and written informed consents were obtained from all the patients. Between November, 2010 and January, 2011, we recruited 31 consecutive age- and gender-matched patients operated for tetralogy of Fallot and 36 healthy controls. No evidence of structural cardiovascular disease was detected by two-dimensional or Doppler echocardiography in the control group.

The study comprised two groups: group 1 (n = 31) consisted of patients operated for tetralogy of Fallot and group 2 (n = 36) consisted of healthy controls. Patients (mean ± standard deviations, 14.25 ± 3.61 years) who had undergone surgery for tetralogy of Fallot were selected according to the following exclusion criteria: resting significant arrhythmias by electrocardiographic Holter monitoring; echocardiogram of inadequate quality; any anamnestic and clinical evidence of heart failure; and treatment with digitalis, β-blockers, and antiarrhythmics. Patients were included in the study if (1) the echocardiographic assessment was made within 2 months of the cardiac magnetic resonance, and (2) they were clinically stable, in sinus rhythm. The symptoms of patients have been classified by using New York Heart Association Functional Classification. All patients were in New York Heart Association I–II and had regular sinus rhythm with complete right bundle branch block.

Cardiac magnetic resonance and tissue Doppler imaging echocardiography was performed in group 1. Only echocardiography was performed in group 2.

Baseline definitions and measurements

Height and weight were measured according to standardised protocols. Body surface area was calculated as the body surface area = (weight^{0.425} × height^{0.725}) × 0.007184 (Dubois formula) (m²). Blood pressure was measured using a mechanical sphygmomanometer in the medical office setting. In each individual, after 15 min of comfortably sitting, the average of three blood pressure measurements was calculated.

Echocardiography and Doppler measurements

A detailed echocardiographic evaluation, which included an M-mode, two-dimensional, and colour Doppler – continuous and pulsed wave – examination, was performed. Images were obtained on a Siemens Acuson CV-70 with a 4–2 MH transducer (East Flanders, Belgium). Tissue Doppler imaging echocardiography was performed by activating the tissue Doppler imaging functions in the same unit. The tissue Doppler imaging programme was set to the pulsed wave Doppler mode reported previously.^{8,9} Filters were

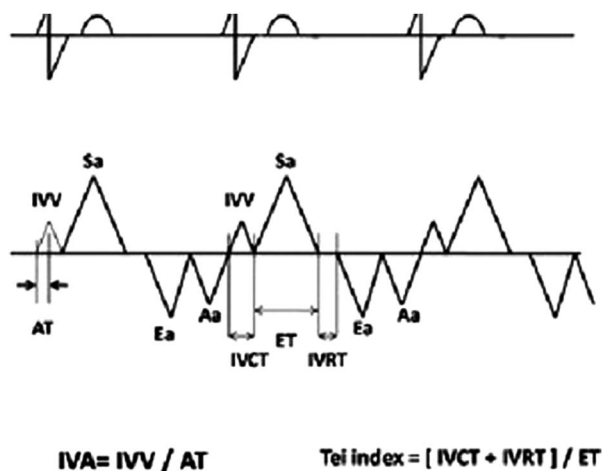


Figure 1.

Tissue Doppler-derived systolic velocities obtained at the lateral corner of tricuspid annulus. IVV begins before the R-wave on electrocardiogram and followed by peak systolic myocardial velocity (Sa). IVA was calculated by dividing the peak IVV by time interval from the onset of the wave during isovolumic contraction to the time at peak velocity of this wave (acceleration time). AT = acceleration time; ET = ejection time; IVA = isovolumic myocardial acceleration; IVCT = isovolumetric contraction time; IVRT = isovolumetric relaxation time; IVV = isovolumic velocity.

set to exclude high-frequency signals. Gains were minimised to allow a clear tissue signal with minimal background noise. Using the four-chamber view, a 2-mm sample volume was placed at the lateral corner of the tricuspid valve annulus, and peak systolic, early diastolic, and late diastolic myocardial velocities were obtained at a sweep speed of 100 mm/s and analysis was carried out as has been described before.⁸ Isovolumic contraction time was defined as the time period between the end of the late diastolic myocardial wave and the beginning of the peak systolic wave. Isovolumic relaxation time was defined as the time period between the end of the peak systolic wave and the beginning of the early diastolic myocardial wave. Ejection time was measured as the duration of ventricular outflow velocity pattern. Myocardial performance index was calculated according to the following equation: myocardial performance index = (isovolumic contraction time + isovolumic relaxation time)/ejection time (Fig 1). The myocardial acceleration during isovolumic contraction was measured by dividing the myocardial velocities during isovolumic contraction by the time interval from onset of the myocardial velocity during isovolumic contraction to the time at peak velocity of this wave.¹⁰ The mean values were recorded by averaging the results of three consecutive measurements.

Pulsed wave Doppler studies of right ventricular outflow tract were performed to assess the systolic

pressure gradient across the right ventricle and the presence and degree of pulmonary regurgitation. Mild pulmonary regurgitation was considered to be present if diastolic retrograde flow could be detected under the pulmonary valve. Moderate pulmonary regurgitation was diagnosed if the retrograde flow could be seen in the right ventricle further apically from the pulmonary valve and in the pulmonary trunk. Severe pulmonary regurgitation was diagnosed if abnormal retrograde diastolic flow could also be detected in the branch pulmonary arteries.¹¹

Cardiovascular magnetic resonance imaging protocol

Cardiac magnetic resonance were performed using a 1.5-T Philips Achieva magnetic resonance system (Rhineland-Palatinate, Germany) using SENSE XL Torso coil 16 elements. The timing of a typical protocol is approximately 35–40 min. After obtaining scout images and a reference scan, an axial stack of black blood turbo spin echo images is acquired to outline cardiac and non-cardiac anatomy. For analysis of valvular and ventricular function, we follow the axis of the heart rather than the axis of the body. For visualisation of valvular insufficiency, two- and four-chamber cines – steady-state free precession sequences – are acquired. To quantify right ventricular function, multiple cines are obtained in the short-axis orientation. An additional cine is aligned along the right ventricular outlet tract to visualise pulmonary insufficiency and right ventricular outlet tract enlargement. From the additional right ventricular outlet tract cine and a transversal black blood image, a velocity map across the pulmonary artery is acquired for calculation of pulmonary regurgitant volume. Quantification of regurgitation entails a post-processing technique in which through-plane velocity and area are measured by tracing the vessel border on sequential phase-contrast images obtained over one cardiac cycle.

For exact measurement of right ventricular volumes and ejection fraction, cardiac magnetic resonance is the most accurate imaging modality. Furthermore, cardiac magnetic resonance may give an unrestricted view of the right ventricular outlet tract, and an aneurysmatic enlargement can be observed. The right ventricular end-diastolic volume and right ventricular end-systolic volume are calculated by manually drawing endocardial contours at end diastole and end systole, respectively, on cine loops, oriented axial, or along the short axis of the right ventricle. A contrast-enhanced magnetic resonance angiography is used to visualise the pulmonary tree. As mentioned in the literature, pulmonary regurgitation fraction >40% was considered severe pulmonary regurgitation.⁷

Table 1. The distribution of the groups of variables.

	Statistics	p*
Age (years)	0.97	0.2
Heart rate (beats/min)	0.116	0.2
Systolic BP (mmHg)	0.179	0.032
Diastolic BP (mmHg)	0.239	0.001
BSA (m ²)	0.12	0.2
Ea (m/s)	0.157	0.097
Aa (m/s)	0.148	0.15
Ea/Aa	0.183	0.025
Sa (m/s)	0.155	0.111
IVRT (ms)	0.178	0.033
IVCT (ms)	0.143	0.179
ET (ms)	0.110	0.2
MPI	0.147	0.154
IVA (m/s ²)	0.233	0.001
RV-EF (%)	0.168	0.058
PRF (%)	0.128	0.2
RV-EDV (ml/m ²)	0.229	0.001
RV-ESV (ml/m ²)	0.127	0.2

Aa = late diastole; BP = blood pressure; BSA = body surface area; Ea = early diastole; ET = ejection time; IVA = isovolumetric acceleration; IVCT = isovolumetric contraction time; IVRT = isovolumetric relaxation time; MPI = myocardial performance index; PRF = pulmonary regurgitant fraction; RV-EDV = right ventricular end-diastolic volume; RV-EF = right ventricle ejection fraction; RV-ESV = right ventricular end-systolic volume; Sa = systole

*Kolmogorov–Smirnov

Cardiovascular magnetic resonance scans protocol:

- scout images single-phase steady-state free precession;
- parallel imaging reference scan;
- axial black blood turbo spin echocardiogram;
- 2-chamber multiphase steady-state free precession;
- 4-chamber multiphase steady-state free precession;
- right ventricular outlet tract cine multiphase steady-state free precession;
- multislice, multiphase short-axis steady-state free precession;
- magnetic resonance angiography pulmonary artery; and
- flow mapping pulmonary trunk (phase-contrast images).

Statistical analysis

All statistical analyses were performed using statistical package for the social sciences for Windows version 17.0 (SPSS, Chicago, Illinois, USA). Kolmogorov–Smirnov tests were used to test the normality of data distribution (Table 1). The data were expressed as arithmetic means and standard deviations. The χ^2 -test was used to compare the categorical variables between groups.

Table 2. Comparison of demographic and clinical characteristics of the study subjects.

	Group 1 (n = 31)	Group 2 (n = 36)	p
Gender (female)	9	10	ns
Age (years)	14.25 ± 3.61	13.72 ± 3.21	0.531
Heart rate (beats/min)	78.61 ± 12.91	81.72 ± 17.89	0.425
Systolic BP (mmHg)	104.03 ± 10.83	107.28 ± 10.93	0.292
Diastolic BP (mmHg)	68.87 ± 11.81	67.76 ± 11.01	0.06
BSA (m ²)	1.48 ± 0.3	1.46 ± 0.28	0.888

BP = blood pressure; BSA = body surface area; ns = non-significant
All measurable values were given with mean ± standard deviation

Independent sample *t*-test was used for comparison of continuous variables among the two homogeneous study groups. Mann–Whitney U-test was used for comparison of continuous variables among the two non-homogeneous study groups. Pearson's and Spearman's correlation analyses were used, respectively, to analyse continuous and categorical variables. Two-sided *p*-value <0.05 was considered statistically significant.

Results

Clinical characteristics of the study population

Clinical and demographic characteristics of controls and operated tetralogy of Fallot patients are presented in Table 2. There were no statistical differences with regard to gender, age, body surface area, heart rate, and both systolic and diastolic blood pressure between the controls and operated tetralogy of Fallot patients (*p* > 0.05 for all). By electrocardiographic analysis, all the patients were in sinus rhythm, and all patients with tetralogy of Fallot had a right bundle block. A total of 31 patients with a mean age of 11.98 ± 3.03 years at follow-up who underwent repair of tetralogy of Fallot at a mean age of 2.56 ± 1.61 years and 36 age- and sex-matched healthy children with a mean age of 13.72 ± 3.21 years were enrolled in this study. The 31 patients had transannular patch enlargement of the right ventricular outflow tract. In all, seven patients underwent Blalock–Taussig shunt before corrective surgery. Of the 31 patients, 24 were in New York Heart Association functional class I, whereas the remaining seven were in class II.

Standard echocardiographic examination

By Doppler criteria, severe pulmonary regurgitation was present in eight (25%) patients, moderate pulmonary regurgitation was present in 15 (50%) patients, and mild pulmonary regurgitation was present in eight (25%) patients.

Table 3. Tissue Doppler findings.

	Group 1 (n = 31)	Group 2 (n = 36)	p
Ea (m/s)	0.20 ± 0.03	0.26 ± 0.04	<0.0001
Aa (m/s)	0.15 ± 0.03	0.19 ± 0.03	<0.001
Ea/Aa	1.34 ± 0.39	1.43 ± 0.26	0.116
Sa (m/s)	0.19 ± 0.03	0.23 ± 0.03	<0.0001
IVCT (ms)	103.77 ± 27.78	71.91 ± 15.86	<0.001
IVRT (ms)	84.72 ± 21.13	69.91 ± 14.27	0.006
ET (ms)	232.63 ± 26.20	232.94 ± 23.37	0.96
IVA (m/s ²)	3.96 ± 1.20	5.77 ± 1.09	<0.0001
MPI	0.80 ± 0.18	0.60 ± 0.11	<0.0001

Aa = late diastole; Ea = early diastole; ET = ejection time; IVA = isovolumetric acceleration; IVCT = isovolumetric contraction time; IVRT = isovolumetric relaxation time; MPI = myocardial performance index; Sa = systole
All measurable values were given with mean ± standard deviation

Tissue Doppler imaging velocities and isovolumic time intervals in patients with tetralogy of Fallot and healthy children

Compared with those of normal children, tissue Doppler imaging velocities of tetralogy of Fallot patients showed decreases in early diastolic myocardial velocity (0.20 ± 0.03 versus 0.26 ± 0.04 cm/s, $p < 0.0001$, respectively), late diastolic myocardial velocity (0.15 ± 0.03 versus 0.19 ± 0.03 cm/s, $p < 0.001$, respectively), and peak systolic velocity (0.19 ± 0.03 versus 0.23 ± 0.03 cm/s, $p < 0.0001$, respectively). Both isovolumic contraction time and isovolumic relaxation time in tetralogy of Fallot patients were significantly longer than those in normal children (103.77 ± 27.78 versus 71.91 ± 15.86 and 84.72 ± 21.13 versus 69.91 ± 14.27 ms, $p < 0.001$ and 0.006 , respectively). Isovolumic acceleration value from the right ventricular tricuspid lateral annulus was significantly lower in patients than in controls (3.96 ± 1.20 versus 5.77 ± 1.09 m/s², $p < 0.0001$, respectively). However, myocardial performance index value from the right ventricular tricuspid lateral annulus was significantly longer in patients than in controls (0.80 ± 0.18 versus 0.60 ± 0.11 , $p < 0.0001$, respectively) (Table 3).

Cardiovascular magnetic resonance findings

The time interval between cardiac magnetic resonance scans and echocardiography was 2 months. Table 4 lists cardiac magnetic resonance parameters.

Derived from the right ventricle lateral tricuspid annulus myocardial performance index and isovolumic acceleration was not correlated with the right ventricle ejection fraction and pulmonary regurgitation fraction source from the cardiac magnetic resonance (Table 5).

Table 4. Cardiovascular magnetic resonance findings.

	Patients (n = 31)
Indexed right ventricle-EDV (ml/m ²)	102.3 ± 40
Indexed right ventricle-ESV (ml/m ²)	57 ± 17.5
Right ventricular-SV (ml/m ²)	53.2 ± 12.5
Right ventricular-EF (%)	39.5 ± 7.5
PRF (%)	38.6 ± 12.88

EDV = end-diastolic volume; EF = ejection fraction; ESV = end-systolic volume; PRF = pulmonary regurgitant fraction; SV = stroke volume

Table 5. Comparison of right ventricular tissue Doppler imaging parameters with cardiac magnetic resonance-derived parameters.

	RV-EF		PRF		RV-EDV		RV-ESV	
	r	p	r	p	r	p	r	p
Ea (m/s)	-0.04	0.80	0.3	0.11	0.37	0.43	0.14	0.44
Aa (m/s)	-0.06	0.72	0.13	0.48	-0.35	0.85	-0.05	0.77
Ea/Aa	0.02	0.91	0.15	0.42	0.17	0.36	0.19	0.31
Sa (ms)	0.03	0.85	0.24	0.20	0.15	0.42	-0.10	0.59
IVCT (ms)	-0.02	0.91	0.03	0.84	0.14	0.47	0.23	0.23
IVRT (ms)	0.02	0.88	-0.05	0.78	-0.28	0.13	-0.34	0.06
ET (ms)	-0.31	0.08	-0.03	0.98	0.23	0.22	0.20	0.26
MPI	0.19	0.33	-0.11	0.58	-0.28	0.15	-0.18	0.35
IVA (m/s ²)	0.12	0.50	0.20	0.28	0.09	0.06	0.91	0.87

Aa = late diastole; Ea = early diastole; EDV = end-diastolic volume; EF = ejection fraction; ESV = end-systolic volume; ET = ejection time; IVA = isovolumetric acceleration; IVCT = isovolumetric contraction time; IVRT = isovolumetric relaxation time; MPI = myocardial performance index; PRF = pulmonary regurgitant fraction; RV = right ventricle; Sa = systole; SV = stroke volume

Discussion

Most of the problems reported during the late follow-up of patients after repair of tetralogy of Fallot have been related to abnormal right ventricle physiology. Residual pulmonary incompetence, residual or recurrent pulmonary stenosis, exercise intolerance, residual ventricular septal defect, and malignant ventricular arrhythmias have been identified as predictors of unfavourable long-term outcome.¹² Long-term volume overload, in patients with moderate or severe pulmonary regurgitation, progressive right ventricle dysfunction takes place in time. For this reason, in patients with right ventricle dilatation, pulmonary valvular replacement should be done before irreversible changes take place.¹³ This finding signifies the importance of post-operative follow-up of the patients and the methods that should be used in the follow-up.

To the best of our knowledge, this is the first report to evaluate the relationship between right ventricle tissue Doppler imaging parameters and findings of cardiac magnetic resonance in patients

with operated tetralogy of Fallot. In this study, right ventricular function was assessed by tissue Doppler imaging and cardiac magnetic resonance. Cardiac magnetic resonance findings by comparing with myocardial performance index, isovolumic acceleration and tried to determine the prognostic values. However, in this study there was no correlation between myocardial performance index, isovolumic acceleration and right ventricular ejection fraction, pulmonary regurgitation fraction, and other results of cardiac magnetic resonance.

Although echocardiography, which has good correlation with radionuclide angiography,¹⁴ is used every day in clinical practice to quantitatively assess left ventricular function, it was previously considered an inaccurate tool to quantitatively assess right ventricular function because of a lack of an ideal geometric model for evaluation of ventricular volumes.¹⁵ In children, three-dimensional echocardiography has been found to have an excellent correlation with cardiac magnetic resonance in the assessment of right ventricular volumes and function.¹⁶ However, questions remain about the accuracy in adults because of inadequate windows and larger right ventricular volumes.

The myocardial performance index is a useful clinical index of global ventricular function for evaluating both systolic and diastolic function.^{17,18} The tissue Doppler imaging method of calculation may have advantages over the pulsed wave Doppler echocardiography calculation. The pulsed wave Doppler myocardial performance index has gained acceptance as a clinical examination for assessing cardiac function and is particularly useful as a predictor of clinical outcome in patients with cardiac disease.^{19–21} The main advantage of this index is that it appears to be independent of ventricular geometry and heart rate.²⁰ However, one of the main limitations of the pulsed wave Doppler myocardial performance index is that it cannot be calculated over a single cardiac cycle because the interval between the end and onset of tricuspid inflow and the ejection time is measured sequentially. By contrast, the tissue Doppler imaging can provide the timing elements necessary to calculate the myocardial performance index on a beat-to-beat basis.²² In addition, the tissue Doppler imaging derived from the tricuspid annulus velocities allows the determination of the isovolumic contraction and relaxation times and the ejection time over a single cardiac cycle in normal conditions.²² These results suggest that the pulsed wave Doppler myocardial performance index may be misinterpreted because it cannot be calculated over a single cardiac cycle, whereas the tissue Doppler imaging myocardial performance index should indicate right ventricular

global function. In our study, myocardial performance index value from the right ventricular tricuspid lateral annulus was significantly longer in patients than in controls.

Therefore, evaluation of right ventricular systolic function is important in this group of patients. Assessment of right ventricular function is difficult because of its asymmetrical shape and narrow acoustic window. In our study, we used tissue Doppler imaging-derived right ventricular isovolumic acceleration. It is a new parameter and has been validated to be a reliable and relatively load-independent measure of right ventricular systolic function.^{23,24} The main finding of our study is the evidence for its clinical use in assessing right ventricular systolic function to determine the severity of right ventricular dysfunction. In many studies, peak systolic velocity has also been shown to reflect right ventricular systolic function. This parameter was found to have a very good correlation with right ventricular fractional area and right ventricular ejection fraction assessed by radionuclide ventriculography.²⁵ Peak systolic velocity is significantly afterload dependent,⁶ whereas isovolumic acceleration reflects right ventricular systolic function during isovolumic contraction. In contrast to peak systolic velocity, isovolumic acceleration has the advantage of being relatively preload and afterload independent. This parameter has been successfully validated by both experimental and clinical studies. Vogel et al.²⁶ demonstrated that isovolumic acceleration was an accurate parameter to assess right ventricular systolic dysfunction and was able to measure the force-frequency relation. Harada et al.²⁴ showed that peak systolic velocity was lower in patients after repair of tetralogy of Fallot compared with the control group. Frigiola et al.²⁷ proposed that isovolumic acceleration was not related directly to right ventricular dilatation but could give information about global right ventricular function in a study composed of 124 patients. In another study, Toyono et al.²⁵ reported decreased right ventricular myocardial velocities and isovolumic acceleration after repair of tetralogy of Fallot.

Schwerzmann et al.²⁸ (mean operated age: 6.6 years, range: 2.1–19.9 years) in a series of 36 patients with tetralogy of Fallot found that inverse linear correlation between myocardial performance index and cardiac magnetic resonance right ventricular ejection fraction ($r = 0.73$, $p < 0.001$). In this study, myocardial performance index was measured using pulsed wave Doppler echocardiography. The follow-up period is longer than that in our study (median 18.2, range 3–37.1 years). Cheung et al.²⁹ (mean operated age 4 ± 1.8 years) in their study of 30 patients with tetralogy of Fallot found a correlation between myocardial performance index

and right ventricular ejection fraction, pulmonary regurgitation fraction (respectively, $r = -0.4$, $p = 0.028$ and $r = -0.4$, $p = 0.031$). In this study, myocardial performance index was measured using pulsed wave Doppler echocardiography. Again, in our patients the follow-up period was shorter than in the previous study.

Conclusion

In our study, a significant relationship has not been detected between the findings obtained from right ventricular cardiac magnetic resonance and myocardial performance index and isovolumic acceleration. Our results do not support the use of myocardial performance index and isovolumic acceleration as a sensitive or a specific tool in patients with operated tetralogy of Fallot.

We did not compare our results with findings of other modalities such as cardiac catheterisation and three-dimensional echocardiography. Cardiac catheterisation was not available for asymptomatic patients. The management of asymptomatic patients with an abnormal myocardial performance index and isovolumic acceleration is not clear because the prognostic importance of subclinical myocardial dysfunction as detected by the myocardial performance index and isovolumic acceleration is still uncertain. Further studies that would present comparative results with new diagnostic modalities are needed to evaluate the diagnostic value of right ventricular myocardial performance index and isovolumic acceleration in patients with operated tetralogy of Fallot. Therefore, in the future large prospective cohort studies are needed to address this issue.

Limitations of the study

When evaluating our findings, it should be considered that the post-operative follow-up period is not very long and that there were no patients older than the second decade and the extrapolation of these findings to long term may not be applicable.

The relatively small sample size and the cross-sectional design are limitations of the present study. The cross-sectional design of this study did not allow us to investigate whether tissue Doppler imaging measurements are predictive of clinical outcomes.

In this study, respiratory cycles were not recorded. Respiratory variations of preload and their effects on echocardiographic measurements were not assessed and may influence the relation of myocardial performance index and isovolumic acceleration to cardiac magnetic resonance-measured right ventricular function.

In addition, the non-simultaneous acquisition of the tissue Doppler imaging and cardiac magnetic resonance data might have introduced an error. However, the echocardiographic evaluation was made within 2 months of the cardiac magnetic resonance and truly simultaneous image acquisition was not feasible.

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None.

Conflicts of Interest

None.

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