

Review

Prevalence of left ventricular 'rigid body rotation', the near absence of left ventricular twist (insights from the MAGYAR studies)

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Abstract

Left ventricular (LV) twist is defined as the wringing motion of the LV around its long-axis during systole generated by rotation of the LV apex in a counterclockwise direction, as viewed from the apex, while the LV base moves in a clockwise direction. In several cases, the LV apex and base move in the same direction during ejection demonstrating a special condition called as LV 'rigid body rotation'. The present review aimed to summarize our knowledge about this rare but not fully understood entity demonstrating its theoretic pathophysiologic background, clinical significance, associated diseases, and reversibility based on available literature.

Keywords: Echocardiography; Left ventricular; Rotation; Twist; Speckle-tracking; Two-dimensional; Three-dimensional

1. The normal twisting motion of the heart

Left ventricular (LV) twist represents the wringing motion of the heart along its long-axis during systole generated by rotation of the LV apex in a counterclockwise (CCw) direction, as viewed from the apex, while the LV base moves in a clockwise (Cw) direction [1]. One of the main components of the mechanical efficacy of the heart is the peak systolic LV twist, resulting in 60% LV ejection fraction (EF) with only 15% fiber shortening [2]. This wringing motion is generated by the helical arrangement of myocardial fibers, subendocardially, the spiral holds a left-handed helix, while there is a right-handed helix in the subepicardium. The myocyte helix angle changes continuously from the subendocardium to the subepicardium, typically ranging from +60 degree at the subendocardium to -60 degree at the subepicardium [3]. The mid-myocardial circumferential fibers shorten simultaneously with the oblique fibers in the right- and left-handed helices during ejection providing a horizontal counterforce throughout ejection. The subepicardial fibers control the direction of the peak systolic LV twist, mainly due to their longer arm of movement. Therefore, the absence of the endocardial helix would result in increased Cw basal and CCw apical rotation, while the absence of the epicardial helix would cause CCw basal and Cw apical LV rotation [4].

2. Estimation of LV rotational mechanics

In earlier years, LV rotational mechanics could be examined only by invasive (microsonometry) or expensive (magnetic resonance imaging) methodologies [1]. Two-dimensional (2D) speckle-tracking echocardiography (STE) is a non-invasive opportunity to assess LV rotational mechanics. However, the LV is a three-dimensional (3D)

chamber, therefore its examination in 2D theoretically is associated with loss of information. This fact leads to guidelines not supporting 2DSTE in the quantification of LV rotational mechanics [5]. Moreover, significant differences between 2DSTE- and 3DSTE-derived LV rotational parameters could also be demonstrated [6]. 3DSTE is a novel method for 3D assessment of LV. Shortly, full volume 'echocloud' (3D echocardiographic dataset) is acquired using 3D capable matrix transducer at the beginning of 3DSTE examination in which analysis could be performed online or offline using a special software by creating a virtual 3D model of the LV. Volumetric changes in LV respecting the cardiac cycle, LV contractility represented by LV strains and LV rotational parameters could be calculated at the same time using this cast [7] (Fig. 1).

3. MAGYAR studies

The aim of our study conducted in our tertiary cardiovascular center was to assess the diagnostic and potentially prognostic values of 3DSTE-derived parameters including LV rotational parameters. LV apical and basal rotations and LV twist were measured in healthy subjects (MAGYAR-Healthy Study) and in several pathological states (MAGYAR-Path Study). LV rotational parameters were measured in elite athletes (MAGYAR-Sport Study), in twins (MAGYAR-Twin Study), in foetuses (MAGYAR-Fetus Study) and during stress protocols (MAGYAR-Stress Study) [7]. ('Magyar' means 'Hungarian' in Hungarian language).

4. Near absence of LV twist—LV 'rigid body rotation'

In special circumstances, LV 'rigid body rotation' (RBR), the near absence of LV twist may develop due to

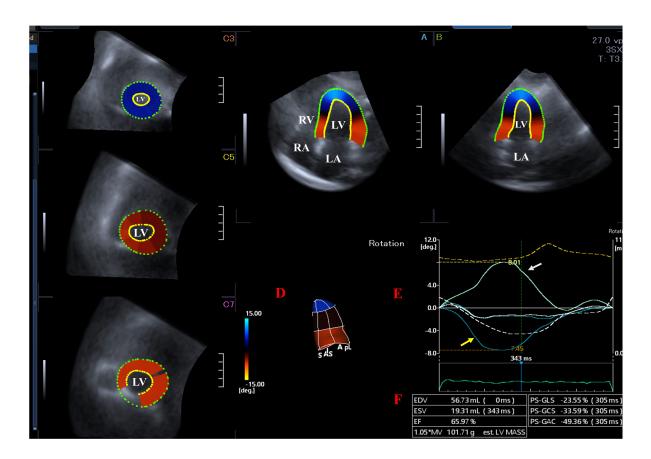


Fig. 1. Left ventricular (LV) rotational mechanics assessed by three-dimensional (3D) speckle-tracking echocardiography. Apical four-chamber (A) and two-chamber views (B) and basal (C3), midventricular (C5) and apical (C7) short-axis views are presented extracted from the acquired 3D volumetric dataset. LV cast (D), curves representing apical (white arrow) and basal (yellow arrow) LV rotations (E) and LV volumetric parameters (F) generated by the software are presented. LA, left atrium; LV, left ventricular; RA, right atrium; RV, right ventricle; EDV, end-diastolic volume; ESV, end-systolic volume; EF, ejection fraction; GLS, LV global longitudinal strain; GCS, LV global circumferential strain; GAC, LV global area strain (change).

changes in the direction of the movement of the myocardial fibers, the LV apex and LV base move in the same clockwise (Cw) or counterclockwise (CCw) direction [8] (Figs. 2,3). This phenomenon is considered to be normal in healthy neonates with an immature heart, LV-RBR has occurred due to the counterclockwise rotation of the basal and apical fibers [9]. However, later the normal LV rotational mechanics develops. In some special clinical circumstances, LV-RBR persists in adults, its real clinical significance and prevalence are not known. The present review aimed to summarize related information from known literature mainly based on the results of the MAGYAR Studies.

5. Cardiac diseases associated with LV-RBR

5.1 Noncompaction cardiomyopathy

LV noncompaction cardiomyopathy (NCCM) or spongiform cardiomyopathy is a rare congenital cardiomyopathy due to an intra-uterine arrest of compaction of the myocardial fibres during embryogenesis [10]. NCCM is characterized by a thin, compacted epicardium and a thick

non-compacted endocardium, with prominent trabeculation and deep intertrabecular recesses [10,11]. Characteristic complications of NCCM include chronic heart failure, lifethreatening ventricular arrhythmias and systemic embolic events [10,11].

According to the literature, NCCM is the most extensively examined LV-RBR-related disease, where its prevalence ranged from 26% to 100% (Table 1) [11–21]. van Dalen *et al.* [12] assessed patients with dilated cardiomyopathy (DCM), healthy subjects and cases with prominent trabeculations, of whom expert opinion in NCCM defined some of them having NCCM, and found that LV-RBR was an objective, quantitative and reproducible functional criterion, and had a good predictive value in the diagnosis of NCCM. Specificity and sensitivity of LV-RBR was 88% and 78% in differentiating NCCM from LV "hypertrabeculation", respectively [12]. Cw LV-RBR was detected in most NCCM cases [11,12,15,18–20], which was found to be related to LV-EF [17]. Almost one-third of the relatives of NCCM patients showed LV-RBR, the ratio of Cw and



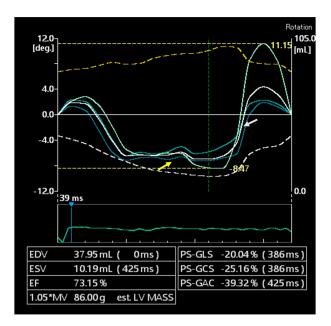


Fig. 2. Abnormal clockwise apical (white arrow) and normal clockwise basal (yellow arrow) left ventricular rotations (clockwise left ventricular 'rigid body rotation'). LV, left ventricular; EDV, end-diastolic volume; ESV, end-systolic volume; EF, ejection fraction; LV-GLS, LV global longitudinal strain; LV-GCS, LV global circumferential strain; LV-GAC, LV global area strain (change).

CCw direction was the same [18]. Ratio of LV-RBR was significantly different between NCCM children and subjects with hypertrabecularization [19]. In another study, reverse LV apical rotation (Cw LV-RBR) was present in 39% of NCCM children [20].

5.2 Dilated or nonischaemic cardiomyopathy

LV rotational mechanics was examined only in a limited number of studies in series of DCM or nonischaemic (NICM) cardiomyopathy patients. 88–100% of patients showed Cw LV-RBR [22,23]. Reversed apical rotation (Cw LV-RBR) and loss of LV torsion was found to be associated with significant LV remodelling, increased electrical dyssynchrony, reduced systolic function, and increased filling pressures in patients with DCM, indicating a more advanced stage of the disease [22]. The normal torsion pattern was observed more frequently in NICM patients without mid-wall fibrosis (MWF), and LV-RBR was more frequently observed in patients with MWF [24].

5.3 Hypertensive and other special cardiomyopathies

Hypertensive heart disease includes a number of complications due to high blood pressure including LV hypertrophy, heart failure, etc. In hypertensive cardiomyopathy patients with reduced LV-EF, 32% of subjects showed predominantly Cw LV-RBR suggesting compromised LV apical rotation similarly to NCCM/DCM/NICM patients [25].

Theoretically hypertension-related abnormalities, fibrosis, etc. could explain these findings. CCw LV-RBR was found in cases with peripartum and chloroquine cardiomyopathies [26,27]. In a small study with hypertrophic cardiomyopathy (HCM), only 1 case out of 18 showed Cw LV-RBR [21].

5.4 Infiltrative cardiac disorders

Regarding the literature, only positron emission tomography-proven cardiac lymphoma [28] and biopsy-proven cardiac amyloidosis (CA) [29] were found to be related to LV-RBR at this moment. Interestingly, CCw LV-RBR could be demonstrated in all cases suggesting compromised LV basal rotation [28,29]. Infiltration of the my-ocardial tissue by amyloid or lymphomatous tissue could lead to changes in LV twisting deformation [28,29].

5.5 Congenital heart diseases

LV-RBR was reported only in a few case reports in certain congenital heart diseases (CHDs) including hypoplastic right heart syndrome [30], univentricular heart [31] and Ebstein's anomaly [32]. Interestingly, hypoplastic right heart syndrome was associated with Cw LV-RBR, while univentricular heart and Ebstein's anomaly were associated with CCw LV-RBR in some individual patients. Abnormalities in the development of myocardial architecture during morphogenesis could be theorised to be responsible for LV-RBR in these diseases [30–32].

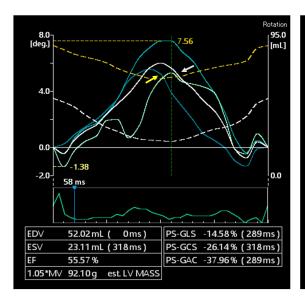
Tetralogy of Fallot (TOF) is a cardiac anomaly with combination of ventricular septal defect, overriding aorta, pulmonary stenosis and right ventricular hypertrophy [33–35]. In patients with corrected TOF from the MAGYAR-Path Study, 27% of subjects showed Cw LV-RBR, while 11% of patients had CCw LV-RBR [33]. In case of corrected TOF patients with early total reconstruction, the same values proved to be beneficial compared to those who had early palliation and late correction. In another earlier study, the values of the same parameters were 15% and 18%, respectively, with patients with almost zero LV rotations [34] (Fig. 4). In the study of Dragulescu *et al.* [35], 38% of corrected TOF patients showed CCw LV-RBR.

6. Disorders with LV-RBR without obvious cardiac involvement

6.1 Acromegaly

Acromegaly is a rare and chronic hormonal disorder that develops due to the hypersecretion of the human growth hormone (GH) and insulin-like growth factor-1 (IGF-1) in adults [36–38]. LV-RBR could be detected in 20% of the cases with normal LV-EF (without obvious signs of classic acromegalic cardiomyopathy), which did not depend on the presence or absence of diabetes mellitus [37,38]. LV-RBR proved to be CCw in 75% of cases, these abnormalities could be theorized to long-term effects of GH and IGF-1.





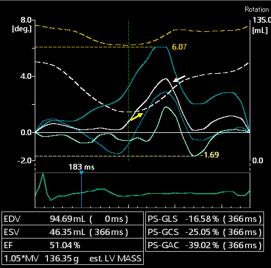


Fig. 3. Normal counterclockwise apical (white arrow) and abnormal counterclockwise basal (yellow arrow) left ventricular rotations (counterclockwise left ventricular 'rigid body rotation'). LV, left ventricular; EDV, end-diastolic volume; ESV, end-systolic volume; EF, ejection fraction; LV-GLS, LV global longitudinal strain; LV-GCS, LV global circumferential strain; LV-GAC, LV global area strain (change).

6.2 Hypopituitarism

Hypopituitarism is a rare and complex hormonal disease caused by the decreased secretion of one or more hormones in the pituitary gland. LV-RBR was present in 13% of patients, potentially due to abnormal secretion of certain hormones [39].

6.3 Lipedema and lymphedema

While lipedema is a feminine disorder with disproportional, bilateral and symmetrical obesity of unknown pathomechanism, excessive accumulation of lymphatic fluid could be detected due to certain factors in lymphedema [40]. In several cases, lipedema and lymphedema are difficult to be differentiated clinically. However, in both disorders, the ratio of LV-RBR was similar (14%) with typical CCw direction in lymphedema patients [40]. The use of compression medical stockings had significant effects on LV rotational mechanics both in lipedema [41] and lymphedema [42].

6.4 Hidradenitis suppurativa

Hidradenitis suppurativa is a dermatological disorder where chronic inflammation was suggested to be having effects on LV rotational mechanics [43].

6.5 Haemophilia

Haemophilia is an X-linked congenital coagulation factor deficiency. Although significant LV rotational abnormalities are present in haemophilia with reduced LV apical rotation and twist, ratio of patients with LV-RBR is not higher than in the normal population [44].

6.6 Hypereosinophilic syndrome

In hypereosinophilic syndrome (HES), peripheral eosinophilia and eosinophilic tissue/end-organ damage could be demonstrated [45]. In HES patients in the early necrotic stage, LV rotational abnormalities including presence of LV-RBR was found in 13% of cases.

6.7 Following kidney transplantation

In all patients with end-stage renal disease, kidney transplantation is the preferred treatment [46]. In post-transplant patients, ratio of LV-RBR was not significantly higher than in normal controls with CCw direction.

6.8 Twin-to-twin transfusion syndrome

In a recent case report, twin-to-twin transfusion syndrome was confirmed in the past medical history of an identical pair of twins who showed different patterns of LV-RBR, Cw LV-RBR was described in the recipient twin and CCw LV-RBR was found in the donor twin [47]. Similarly to CHDs, abnormalities in the development of myocardial architecture could lead to LV-RBR in twin-to-twin transfusion syndrome theoretically.

7. Healthy subjects and elite athletes

In a current study demonstrating normal reference values of 3DSTE-derived LV rotational parameters and their age- and gender-dependency, 6% of cases proved to have LV-RBR. At the time of 3DSTE, these subjects had no abnormality, pathologic state, drug use, symptoms, electrocardiographic or echocardiographic findings, which could explain LV-RBR. Theoretically, these subjects had subclin-



Table 1. Studies confirming left ventricular 'rigid body rotation' in different cardiac disorders.

	Publication	Method	No. of cases and ratio of	Cw/CCw
	Publication	Method	cases showing LV-RBR	LV-RBR
	C	Cardiomyopathi	es	
- Noncompaction CM	van Dalen et al. [11]	2D-STE	10 (100%)	Cw-70%
				CCw - 30%
	van Dalen et al. [12]	2D-STE	34 (88%)	Cw-38%
				initial CCw, then Cw – 41%
				CCw - 6%
	427 1 5107	4D 0000	4.4	initial Cw, then CCw – 3%
	*Nemes <i>et al</i> . [13]	3D-STE	1 (case report)	Cw
	*Kalapos et al. [14]	3D-STE	7 (100%)	Cw – 29% CCw – 71%
	Peters et al. [15]	2D-STE	60 (53%)	Cw-72%
				CCw - 28%
	Cortés et al. [16]	2D-STE	28 (57%)	-
	Szűcs et al. [17]	FT-MRI	with LVEF < 50%	Cw-92%
			31 (42%)	CCw - 8%
			with LVEF >50%	Cw – 12%
			31 (26%)	CCw - 88%
	Akhan <i>et al</i> . [18]	2DSTE	NCCM patients	Cw – 59%
			32 (53%)	CCw-41%
			patients' relatives 31 (30%)	Cw – 56% CCw – 44%
	Sobotino et al [10]	2DSTE	NCCM children	Cw - 100%
	Sabatino et al. [19]	2D81E	23 (56%)	CW - 100%
			hyper-trabecularization 24 (4%)	
	Nawaytou et al. [20]	2DSTE	NCCM children 28 (39%)	reverse apical rotation (Cw) - 100%
	Ashwal et al. [21]	2DSTE	12 (50%)	CCw - 100%
- Peripartum CM with noncompaction phenotype	Peters et al. [26]	2D-STE	1 (case report)	CCw
- Hypertrophic CM	Ashwal et al. [21]	2DSTE	18 (6%)	Cw
- Chloroquine-induced CM	*Nemes <i>et al</i> . [27]	3D-STE	1 (case report)	CCw
- Dilated CM with reduced EF	Popescu et al. [22]	2D-STE	50 (52%)	Cw - 100%
	Cortés et al. [16]	2D-STE	13 (15%)	-
	Setser et al. [23]	MRI-TT	21(76%)	Cw - 88%
			, ,	CCw - 12%
- Non-ischaemic CM	Taylor et al. [24]	FT-CMR	with MWF	-
			32 (64%)	
			without MWF	
			84 (28%)	
- Hypertensive CM with reduced EF	Maharaj et al. [25]	2D-STE	41 (32%)	Cw - 85%
				CCw – 15%



Table 1. Continued.

	Publication	Method	No. of cases and ratio of cases showing LV-RBR	Cw/CCw LV-RBR
	Infiltrative disorde	ers with cardiac in	nvolvement	
- Cardiac lymphoma	*Nemes <i>et al.</i> [28]	3D-STE	1 (case report)	CCw
- Cardiac amyloidosis	*Nemes et al. [29]	3D-STE	10 (60%)	CCw - 100%
	Congen	ital heart diseases	S	
- Hypoplastic right heart syn-	*Nemes <i>et al</i> . [30]	3D-STE	1 (case report)	Cw
drome				
- Univentricular heart	*Nemes <i>et al.</i> [31]	3D-STE	1 (case report)	CCw
- Ebstein' anomaly	*Nemes <i>et al.</i> [32]	3D-STE	1 (case report)	CCw
- Tetralogy of Fallot	*Nemes <i>et al</i> . [33]	3D-STE	26 (38%)	Cw - 70%
				CCw-30%
	Menting et al. [34]	2D-STE	82 (34%)	Cw - 46%
				CCw-54%
	Dragulescu et al. [35]	2D-STE	50 (38%)	CCw - 100%

2D, two-dimensional; 3D, three-dimensional; Cw, clockwise; CCw, counterclockwise; LV-RBR, left ventricular 'rigid body rotation'; CM, cardiomyopathy; LVEF, left ventricular ejection fraction; MWF, mid-wall fibrosis; NCCM, noncompaction cardiomyopathy; STE, speckle-tracking echocardiography; FT-CMR, feature-tracking cardiovascular magnetic resonance.

ical abnormalities or undiagnosed disorders which were not known at the time of 3DSTE [48]. In a study comparing LV rotational mechanics in NCCM patients and controls, 23% of healthy subjects showed CCw LV-RBR which higher rate could be partially explained by selection bias [17]. In contrast, none of the elite athletes doing high dynamic sports showed 3DSTE-proven LV-RBR in a recent study [49]. The above mentioned results suggest further examinations mainly focusing any diagnostic or prognostic impact of LV-RBR in subjects without symptoms or apparent abnormalities.

8. Special considerations regarding LV-RBR

8.1 Clockwise vs. counterclockwise LV-RBR

Normally, the LV base rotates clockwise, while the LV apex rotates in counterclockwise direction. In Cw LV-RBR, apical rotation of the LV has a clockwise direction (negative value on 3DSTE images, Fig. 2), while in CCw LV-RBR, basal rotation of the LV has a counterclockwise direction (positive value on 3DSTE images, Fig. 3). Direction of LV-RBR and their ratio in certain cardiac and non-cardiac disorders are presented in Tables 1 and 2 (Ref. [17,36–40,43–49]). Interestingly, while most cardiomyopathy patients showed Cw LV-RBR [11,12,15,18,19,22,23,25], patients with non-cardiac disorders without obvious cardiac involvement (lymphedema, acromegaly, following kidney transplantation) most likely had CCw LV-RBR [37,40,46].

8.2 Heart failure treatment, reversibility and LV-RBR

A recent report described a patient with evidence of reversible LV-RBR who met the clinical criteria for the diagnosis of both peripartum cardiomyopathy and LV noncompaction with reduced LV-EF. Following 6-months heart failure treatment, improvement in LV-EF could be observed together with normalization of LV rotational mechanics [26].

8.3 Vasodilation, stress test and LV-RBR

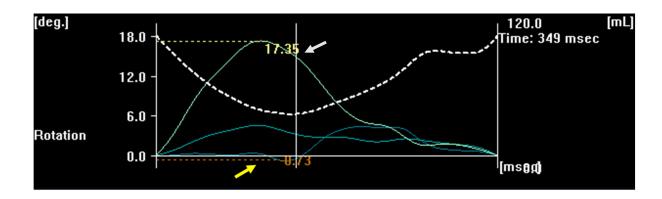
Vasodilation and stress-related LV-RBR is weakly assessed. In a patient with previous coronary stenting presented with stable angina and positive exercise treadmill test, LV-RBR could be demonstrated, its characteristics changed dramatically during dipyridamole-induced vasodilatation [50]. A case of abnormal CCw rotation of the LV base was reported in a patient with angina pectoris without significant coronary artery disease (CAD) at rest, the direction of the rotation of the LV base changed to Cw theoretically due to dipyridamole-induced vasodilation via improved blood supply at maximum hyperaemia [51]. At the recovery phase, LV-RBR could be detected again in this case. On the contrary, a CAD patient showed LV-RBR during dipyridamole stress, which normalized by the end of stress [52].

8.4 'Clear' LV-RBR vs. LV-RBR with apico-basal gradient

LV-RBR is the near absence of LV twist, when rotation at both apical and basal levels of the LV occur in the same Cw or CCw direction (Figs. 2,3,4). Although all segments move in the same direction, the amplitude of their motion



^{*} Results from the MAGYAR-Path Study.



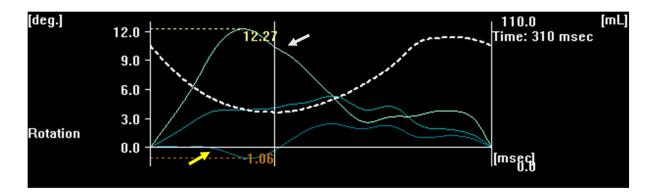


Fig. 4. Increased counterclockwise left ventricular apical rotation (white arrow) with almost zero left ventricular basal rotation (yellow arrow).

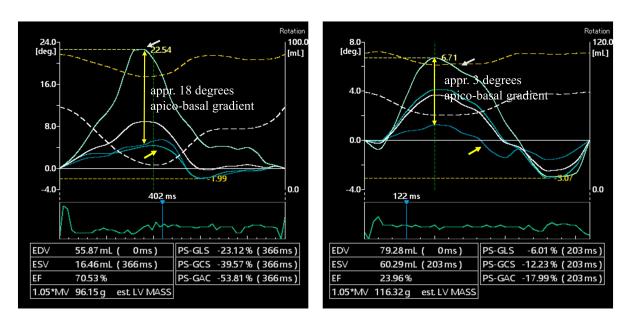


Fig. 5. Differences between apical and basal left ventricular rotations in the same counterclockwise direction demonstrating high (left panel) or low (right panel) left ventricular apico-basal rotational gradient. LV, left ventricular; EDV, end-diastolic volume; ESV, end-systolic volume; EF, ejection fraction; LV-GLS, LV global longitudinal strain; LV-GCS, LV global circumferential strain; LV-GAS, LV global area strain.

Table 2. Studies confirming left ventricular 'rigid body rotation' in different non-cardiac diseases without known cardiac involvement.

		m vor vement.		
	Publication Method	36.1.1	No. of cases and ratio of	Cw/CCw
		Method	cases showing LV-RBR	LV-RBR
	I	Healthy subject	S	
- Healthy subjects	*Kormányos et al. [48]	3D-STE	177 (6%)	-
	Szűcs et al. [17]	FT-MRI	31 (23%)	CCw - 100%
Sportsmen doing high dy-	*Gyenes <i>et al</i> . [49]	3D-STE	80 (0%)	-
namic sports				
		docrine disorde		
- Acromegaly	* Nemes <i>et al</i> . [36]	3D-STE	1 (case report)	Cw
	*Kormányos et al. [37]	3D-STE	20 (20%)	Cw-25%
				CCw - 75%
	*Nemes <i>et al.</i> [38]	3D-STE	with DM	-
			5 (20%)	
			without DM	
			19 (21%)	
Hypopituitarism	*Kormányos et al. [39]	3D-STE	31 (13%)	Cw-50%
				CCw - 50%
	Derm	natological diso	rders	
- Lipedema	*Nemes <i>et al</i> . [40]	3D-STE	22 (14%)	Cw - 33%
				CCw - 33%
				reversed twist
				− 33 %
Lymphedema	*Nemes et al. [40]	3D-STE	22 (14%)	CCw-100%
Hidradenitis suppurativa	*Nemes <i>et al</i> . [43]	3D-STE	1 (case report)	CCw
	Haen	natological disc	rders	
Haemophilia	*Nemes et al. [44]	3D-STE	14 (7%)	CCw
- Hypereosinophilic syndrome	*Nemes <i>et al</i> . [45]	3D-STE	11 (18%)	Cw - 50%
				CCw - 50%
		Other diseases		
Following kidney transplanta-	*Nemes <i>et al</i> . [46]	3D-STE	38 (8%)	CCw - 100%
ion				
Twin-to-twin transfusion syn-	*Nemes <i>et al.</i> [47]	3D-STE	2 (case report)	twin A – Cw
lrome				
				twin B – CCw

³D, three-dimensional; Cw, clockwise; CCw, counterclockwise; LV-RBR, left ventricular 'rigid body rotation'; STE, speckle-tracking echocardiography; FT-CMR, feature-tracking cardiovascular magnetic resonance.

could be different leading to regional differences between apical and basal LV rotations called as LV apico-basal gradient (Fig. 5). In some disorders, LV apico-basal gradient could be elevated like in case of CA [29]. Its clinical significance is not known but could be considered as an overcompensation due to absence of normally directed LV rotational mechanics. Similar changes could be detected during stress as well [50–52].

9. Prognostic value of LV-RBR

Limited information is available regarding prognostic impact of LV-RBR. In a recent study, reverse LV apical rotation (Cw LV-RBR) was not a sensitive but a specific indicator of complications in children with NCCM suggesting its prognostic rather than diagnostic value [20].



^{*} Results from the MAGYAR-Path, MAGYAR-Healthy and MAGYAR Sport Studies.

10. Possible pathophysiological explanations of LV-RBR

Theoretically, changes in LV rotational mechanics may occur due to the above mentioned special factors, as well as due to classic risk factors, such as hypertension, hypercholesterolaemia, diabetes mellitus, etc. or subclinical diseases (focal/local oedema, inflammation, fibrosis, etc.). The effect of CAD and related haemodynamic changes could not be excluded either.

11. The role of three-dimensional speckle-tracking (strain) echocardiography in current cardiology practice

Although 3DSTE is not a guideline-recommended technique at this moment, it has several advantages as compared to other echocardiographic methods. It sees the heart as it is: a 3D organ and all volumetric, functional (strain) and rotational parameters of heart chambers respecting the cardiac cycle can be calculated at the same time using the same virtual 3D models based on digitally acquired echocardiographic datasets. Moreover, data acquisition and analysis do not require significantly more time in practical hands than a routine echocardiographic examination. These facts could highlight its importance in clinical practice in more complicated cases, which require more detailed analysis (for instance in heart failure or following myocardial infarction). As demonstrated above, 3DSTE-derived parameters could help in differentiating NCCM from LV 'hypertrabecularization' or HCM from cardiac amyloidosis [12,53]. Moreover, prognostic impact of 3DSTE-derived ejection fraction and 3D strain parameters has also been demonstrated [54,55]. 3DSTE allows detailed assessment of the right ventricle and both atria, as well [53,55]. The role of 3DSTE in the assessment of atrioventricular valvular annuli has also been demonstrated [56].

Most of echocardiographic laboratories have access to 2D strain software and LV rotational parameters can be calculated with them, 3DSTE-derived ones are recommended according to the guidelines [5]. Although 2DSTE-derived global longitudinal strain is well known and a widely used parameter with a strong progmostic impact [57], limitations of 2DSTE should always be considered, when LV rotations are calculated.

12. Conclusions

LV-RBR is defined as the near absence of LV twist when LV apical and basal segments move in the same clockwise or counterclockwise direction. LV-RBR could be demonstrated in a number of diseases with larger prevalence in certain disorders with cardiac involvement. However, some non-cardiac diseases without overt cardiovascular involvement show limited prevalence of LV-RBR without obvious significance, which would require further investigations.

Author contributions

AN and ÁK designed the research study, performed the research and analyzed the data. All authors contributed to editorial changes in the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

All subjects gave their informed consent for inclusion before they participated in the MAGYAR studies. The study was conducted in accordance with the Declaration of Helsinki, and the protocol was approved by the Ethics Committee of University of Szeged (approval number: 71/2011).

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Conflict of interest

The authors declare no conflict of interest. Attila Nemes is serving as one of the Editorial Board members of this journal. We declare that Attila Nemes had no involvement in the peer review of this article and has no access to information regarding its peer review. Full responsibility for the editorial process for this article was delegated to Tasneem Z. Naqvi.

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