Three-dimensional speckle-tracking echocardiography detects different patterns of right atrial dysfunction in selected disorders: a short summary from the MAGYAR-Path Study

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Abstract: The right atrium (RA) is the most under-evaluated heart chamber in the clinical practice. Threedimensional (3D) speckle-tracking echocardiography (STE) merges the benefits of 3D echocardiography and STE: volumetric and strain assessment of the RA can be performed at the same time from the same digitally acquired 3D echocardiographic dataset with a special equipment using a virtual 3D model of the RA. A diagnostic focused study is under way at the Second Department of Medicine and Cardiology Center, University of Szeged, Hungary to assess 3D-STE-derived RA variables in different patient populations, which serves as a substudy of Motion Analysis of the heart and Great vessels bY three-dimensionAl speckletRacking echocardiography in Pathological cases (MAGYAR-Path) Study. The aim of the present review is to summarize these early results.

Keywords: Three-dimensional (3D); speckle-tracking echocardiography (STE); right atrium (RA)

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The right atrium (RA) is the most under-evaluated heart chamber in the clinical practice (1). Its role is to transit deoxygenated blood from the caval veins and coronary sinus to the right ventricle (RV) via the tricuspid valve, but it has effects on heart rhythm development and conductance, and endocrine and baroreceptor regulations as well (2). Similarly, to the left atrium (LA), the RA has a triple function during the cardiac cycle (2,3): (I) it acts as a "reservoir" storing blood arriving from the systemic venous circulation following closure of the tricuspid valve; (II) it acts as a "conduit" with passive filling from the caval veins to the RV after opening of the tricuspid valve; (III) it acts as a "booster pump" contributing to RV filling by atrial contraction in late diastole.

Routine transthoracic echocardiographic RA assessment has a lot of limitations (3). RA dimensions could be

directly assessed in apical 4-chamber view (AP4CH), with Doppler and tissue Doppler echocardiography tricuspid inflow and tricuspid annular velocities could be measured, respectively (3). With newer echocardiographic techniques, speckle-tracking echocardiography (STE) allows quantitative assessment of RA deformation, while threedimensional (3D) echocardiography enables accurate RA volumetric measurements (3,4). 3D-STE merges the benefits of 3D echocardiography and STE: volumetric and strain assessment of heart chambers could be performed at the same time from the same 3D echocardiographic dataset. According to recent practices 3D-STE analysis includes two steps: data acquisition and offline measurement (3,4). Acquisition of 3D echocardiographic dataset is performed with a specific echocardiographic equipment using a Quantitative Imaging in Medicine and Surgery, Vol 8, No 2 March 2018



Figure 1 3D-STE image of a typical patient: apical four-chamber (A), two-chamber views (B) and short-axis views at basal (C3), midatrial (C5) and superior (C7) levels of the RA. A 3D cast of the RA (D), volumetric RA data (E) and time-segmental strain curves (of a selected strain; coloured lines) together with time volume curve respecting the cardiac cycle (dashed line) (F) are also demonstrated. 3D, three-dimensional; STE, speckle-tracking echocardiography; RA, right atrium; EDV, end-diastolic volume; ESV, end-systolic volume; EF, ejection fraction, est; MASS, estimated mass.

matrix phased-array transducer with 3D capability. Using a dedicated software 3D cast of the RA could be created. Several apical and cross-sectional 2D views help this evaluation, on which endpoints of the tricuspid valve and endocardial surface of the RA superior region are set by the examiner (4-8) (*Figure 1*).

According to the literature, end-systolic largest RA volume (V_{max}), early diastolic RA volume before atrial contraction (V_{preA}) and late-diastolic smallest RA volume (V_{min}) are suggested to be measured. From these RA volumes, several volume-based functional properties respecting the cardiac cycle could be calculated which are presented in *Table 1* (3). Using the same 3D virtual RA cast, several unidirectional [radial (RS), longitudinal (LS) and circumferential (CS)] and multidirectional [area (AS) and 3D (3DS)] strains could be assessed with a typical twin peaks

curve (3). The first peak represents the systolic reservoir function and is called peak RA strain, while the second peak represents late diastolic booster pump phase and is called RA strain at atrial contraction. It is possible to calculate not only global and mean segmental, but segmental strains as well from which different regional strains (basal, midatrial and superior) could be measured for each patient.

Summarizing these data 3D-STE has an ability of detailed analysis of different phases of RA function as follows (5-8): (I) systolic reservoir phase with total atrial stroke volume (TASV) and emptying fraction (TAEF) together with peak RA strains; (II) early diastolic phase with passive atrial stroke volume (PASV) and emptying fraction (PAEF); (III) late diastolic phase with active atrial stroke volume (AASV) and emptying fraction (AAEF) together with RA strains at atrial contraction.

Table 1 Assessment of RA volume-based functional properties in different phases of the cardiac cycle

Parameters	Stroke volume (mL)	Emptying fraction (%)
Systolic reservoir function	$TASV = V_{max} - V_{min}$	$TAEF = TASV/V_{max} \times 100$
Early diastolic conduit function	$PASV = V_{max} - V_{preA}$	$PAEF = PASV/V_{max} \times 100$
Late-diastolic active contraction	$AASV = V_{\text{preA}} - V_{\text{min}}$	$AAEF = AASV/V_{preA} \times 100$

RA, right atrial; V_{max} , maximum RA volume; V_{min} , minimum RA volume; V_{preA} , RA volume before atrial contraction; TASV, total atrial stroke volume; PASV, passive atrial stroke volume; AASV, active atrial stroke volume; TAEF, total atrial emptying fraction; PAEF, passive atrial emptying fraction; AAEF, active atrial emptying fraction.

Table 2 3D-STE-derived RA volumes and volume-based functional properties in different disorders

Parameters	HES (5)	NCCM (6)	CA (7)	cTOF (8)
RA volumes (mL)				
V _{max}	↑	↑	1	↑
V _{min}	\uparrow	↑	\uparrow	↑
V _{preA}	1	1	\uparrow	↑
RA stroke volumes (mL)				
TASV	1	1	-	-
PASV	\uparrow	↑	-	-
AASV	-	-	-	-
RA emptying fraction (%)				
TAEF	-	-	\downarrow	\downarrow
PAEF	-	-	-	\downarrow
AAEF	_	_	Ţ	_

 \downarrow , decrease; \uparrow , increase; \neg , no changes as compared to age- and gender-matched healthy controls. 3D, three-dimensional; STE, speckletracking echocardiography; CA, cardiac amyloidosis; cTOF, corrected tetralogy of Fallot; HES, hypereosinophilic syndrome; NCCM, noncompaction cardiomyopathy; RA, right atrial; TASV, total atrial stroke volume; PASV, passive atrial stroke volume; AASV, active atrial stroke volume; TAEF, total atrial emptying fraction; PAEF, passive atrial emptying fraction; AAEF, active atrial emptying fraction; V_{max}, maximum RA volume; V_{min}, minimum RA volume; V_{preA}, RA volume before atrial contraction.

A particular diagnostic substudy is under way at the Second Department of Medicine and Cardiology Center, University of Szeged, Hungary to assess the above mentioned 3D-STEderived RA variables in different patient populations using a Toshiba ArtidaTM echocardiography device with a PST-30SBP phased-array transducer (5-8). These studies are parts of Motion Analysis of the heart and Great vessels bY three-dimensionAl speckle-tRacking echocardiography in Pathological cases (MAGYAR-Path) Study, where "Magyar" means "Hungarian" in Hungarian language.

RA volumes, volume-based functional properties and strains were found to have specific patterns in different

disorders as compared to age- and gender-matched healthy subjects as a part of the MAGYAR-Path Study (*Tables 2-4*) (5-8). Similar ability of 3D-STE for LA has already been confirmed (9).

In these early results increased RA volumes respecting cardiac cycles could be demonstrated in hypereosinophilic syndrome (HES) (5), noncompaction cardiomyopathy with LV involvement (NCCM) (6), cardiac amyloidosis (CA) (7) and corrected tetralogy of Fallot (cTOF) (8). Interestingly, HES and NCCM patients showed similar pattern of mild RA dysfunction affecting systolic reservoir and early diastolic conduit functions (increased TASV and PASV). No changes

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Table 5 5D-51 E-derived global KA peak strains and strains at atrial contraction in different disorders				
Parameters	HES (5)	NCCM (6)	CA (7)	cTOF (8)
RA global peak strains (%)				
Radial strain	-	-	-	\downarrow
Circumferential strain	-	-	-	-
Longitudinal strain	-	-	-	\downarrow
3D strain	-	-	-	-
Area strain	-	-	\downarrow	\downarrow
RA global strains at atrial contraction (%)				
Radial strain	-	-	-	-
Circumferential strain	-	-	-	\downarrow
Longitudinal strain	-	-	\downarrow	_
3D strain	-	-	-	\downarrow
Area strain	-	-	\downarrow	_

Table 3 3D-STE-derived global RA peak strains and strains at atrial contraction in different disorders

↓, decrease; –, no changes as compared to age- and gender-matched healthy controls. 3D, three-dimensional; STE, speckle-tracking echocardiography; CA, cardiac amyloidosis; cTOF, corrected tetralogy of Fallot; HES, hypereosinophilic syndrome; NCCM, noncompaction cardiomyopathy; RA, right atrial.

Table 4 Different patterns of RA dysfunction as assessed by 3D-STE in different disorders

Parameters	HES (5)	NCCM (6)	CA (7)	cTOF (8)
Systolic reservoir function	+	+	+	+
Diastolic conduit function	+	+	-	+
Diastolic active contraction	-	_	+	+

+, alterations; –, no alterations as compared to age- and gender-matched healthy controls. 3D, three-dimensional; STE, speckle-tracking echocardiography; CA, cardiac amyloidosis; cTOF, corrected tetralogy of Fallot; HES, hypereosinophilic syndrome; NCCM, noncompaction cardiomyopathy; RA, right atrial.

in RA emptying fractions or strains could be detected. These alterations were present without any clinical signs of right heart involvement in HES and NCCM suggesting subclinical changes (5,6). As expected CA and cTOF were associated with more severe alterations: reductions in RA emptying fractions (TAEF and AAEF) with preserved stroke volumes and impairment of RA strains featuring both systolic reservoir and late-diastolic booster pump functions could be detected (7,8). However, RA conduit function seemed to be intact in CA (7). cTOF showed a different pattern: all RA functions were found to be affected (reduced TAEF, PAEF and strains) (8).

These disease-related alterations could be theoretically explained by RA volume and pressure overloads, filling and emptying features, tissue wall characteristics. Similar alterations of the caval veins, RV and pulmonary artery could also explain the results, but presence and degree of tricuspid regurgitation, LA-RA interactions, conductance and congenital abnormalities, hormonal discrepancies, effects of associated cardiovascular risk factors, lung diseases, different degree left heart failure, autonomic neuropathy, etc. could also have a role. Moreover, differences could be interpreted whether a particular disease affects only left or right heart or both (1-3). Normal values of 3D-STE-derived RA volumetric and functional properties are not defined yet therefore selection of control subjects could also affect comparisons. These factors and their ratio in the development of above mentioned RA morphologic and functional changes theorized to have

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different effects which could highlight our attention whether pattern of these alterations is specific for certain disorders (or group of diseases), have any differential diagnostic or prognostic value or whether these are age- or gender-related. Therefore, large volume studies with significant number of patients with different disorders are warranted.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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