

Assessment of Aortic Valve Area Combining Echocardiography and Magnetic Resonance Imaging

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Abstract

Background: Transthoracic echocardiography (TTE) is routinely used to calculate aortic valve area (AVA) by continuity equation (CE). However, accurate measurement of the left ventricular outflow tract (LVOT) can be difficult and flow acceleration in the LVOT may lead to miscalculation of the AVA.

Objective: The aim of our study was to compare AVA measurements by standard TTE, cardiac magnetic resonance imaging (MRI) and a hybrid approach combining both techniques.

Methods: AVA was calculated in 38 patients (age 73 ± 9 years) with standard TTE, cine-MRI planimetry and a hybrid approach: Hybrid Method 1: TTE-derived LVOT measurement in the CE numerator was replaced by the MRI assessment of the LVOT and AVA was calculated: $(LVOT_{MRI} \cdot LVOT-VTI_{TTE}) / \text{transaortic-}VTI_{TTE}$. Method 2: We replaced the SV in the numerator by the MRI-derived SV and calculated $AVA = SV_{MRI} / \text{transaortic-}VTI_{TTE}$.

Results: Mean AVA derived by TTE was $0.86 \text{ cm}^2 \pm 0.23 \text{ cm}^2$ and $0.83 \text{ cm}^2 \pm 0.3 \text{ cm}^2$ by MRI- planimetry, respectively. The mean absolute difference in AVA was 0.03 cm^2 for TTE vs. MRI planimetry. AVA calculated with method 1 and method 2 was $1.23 \text{ cm}^2 \pm 0.4 \text{ cm}^2$ and $0.92 \text{ cm}^2 \pm 0.32 \text{ cm}^2$, respectively. The mean absolute difference between TTE and method 1 and method 2 was 0.37 cm^2 and 0.06 cm^2 , respectively ($p < 0.001$).

Conclusion: MRI-planimetry of AVA and hybrid method 2 are accurate and showed a good agreement with standard TTE measurements. Therefore, hybrid method 2 is a reasonable alternative if poor acoustic windows or LVOT flow accelerations limit the accuracy of TTE, particularly in patients at high risk for an invasive hemodynamic study. (Arq Bras Cardiol 2012;98(3):234-242)

Keywords: Aortic valve stenosis; echocardiography; magnetic resonance spectroscopy.

Introduction

Therapeutic decisions in aortic stenosis (AS) are based on disease-specific symptoms and on hemodynamic severity, measured by using the aortic valve area (AVA). Thus, an accurate evaluation of these two issues is mandatory. Transthoracic echocardiogram (TTE) with Doppler interrogation of the aortic valve serves as the mainstay of diagnosis^{1,2}. The parameters referred to in current guidelines¹ for grading the severity of AS are the $AVA \leq 1 \text{ cm}^2$, mean pressure gradient $\geq 40 \text{ mmHg}$ (Pmean) and peak flow velocity $\geq 400 \text{ cm/s}$ (Vmax). In patients with normal left ventricular ejection fraction (EF), these parameters should allow a dependable classification of the severity of AS as mild, moderate or severe. In clinical practise, a good TTE study can provide all the parameters necessary to assess the severity of stenosis.

Nevertheless, doppler-derived calculation of AVA using the continuity equation (CE) can be difficult, due to poor echocardiographic conditions such as a limited acoustic window, heavy calcifications of the aortic valve or flow acceleration in the LVOT due to a septal bulge, a common finding in AS. When these factors question a proper orifice area quantification, we have to rely on alternative methods including transesophageal planimetry of the AVA, replacing the Doppler-derived stroke volume (SV) in the CE by SV measured Swan-Ganz Catheter or invasive quantification by cardiac catheterization and applying the Gorlin formula³. However, the invasive nature and the increased stroke risk⁴ with retrograde catheterisation of the aortic valve (AV) limits its clinical use. Hence, there is still a need for an alternative approach to the standard CE and invasive procedures in selected patients. Cardiovascular magnetic resonance imaging (MRI) is an important non-invasive tool for the diagnosis and treatment evaluations of patients with cardiovascular disease. Measurements of both blood flow and flow volume qualify MRI as an ideal technique for the evaluation of cardiac function and volumes. Therefore, the purpose of our study was to compare AVA measurements by standard TTE, cardiac magnetic resonance imaging (MRI) and a hybrid approach combining both techniques.

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Methods

Patients

A cohort of 38 consecutive patients referred to the Department of Cardiology with a diagnosis of moderate to severe AS was recruited into the study. The study protocol was approved by the regional ethics committee. Exclusion criteria were concomitant mitral valve disease, severe low gradient AS, EF < 35%, hypertrophic obstructive cardiomyopathy, uncontrolled hypertension, severe arrhythmias, and general exclusion criteria for MRI.

Transthoracic Echocardiography

All patients underwent transthoracic echocardiography using standard 2-D, CW-, PW- and colour Doppler techniques (Vingmed Vivid 7, General Electric, Milwaukee, Wisconsin). LV EF was quantified according to the Simpson method. LV dimensions and left ventricular muscle mass (LVM) were measured in the parasternal long axis view according to the American Society of Echocardiography guidelines² and indexed for body surface area (BSA). Doppler assessment of AS included measurement of Pmean, Pmax and Vmax. AVA was calculated by means of the CE ($AVA = (LVOT_{\text{diameter}}/2)^2 \cdot \pi \cdot [LVOT\text{-}VTI/\text{transaortic-VTI}]$) and indexed for body surface area. The pressure recovery adjusted aortic valve area (i.e. energy loss index (ELI)) was calculated by a previously validated formula⁵. Severe AS was defined as $AVA < 1 \text{ cm}^2$ with a $P_{\text{mean}} \geq 40 \text{ mmHg}$. Tissue Doppler velocity imaging and 2D strain analysis were performed as previously described⁶.

Magnetic resonance imaging

A 1.5-T Achieva MRI scanner (Philips Medical Systems, Netherlands) equipped with a 5-element cardiac synergy coil was used. Cine-Images were acquired in breath hold SSFP sequences (TE 3.43, TR 1.72). Images were evaluated with the cmr42 research edition toolkit (circle cardiovascular imaging, Calgary, Canada) combining long and short axis views. Non-myocardial structures and papillary muscles were carefully excluded and endocardial as well as epicardial borders were outlined manually. Endsystolic and enddiastolic volumes were calculated by the program software. AVA planimetry was performed in the cardiac phase where the maximal opening area of the leaflets was observed. A three-chamber view and LVOT view were performed with cine imaging using fast-field echo sequence. The planimetric area of the LVOT cross sectional area ($LVOT_{\text{CSA}}$) was measured mid-systolic. Additionally, the diameter of the LVOT was measured in the three-chamber view (as normally done with TTE), and the LVOT area was calculated according to $LVOT_{\text{CSA}} = (LVOT_{\text{diameter}}/2)^2 \cdot \pi$. MRI velocity measurements in phase contrast technique of the aortic valve and the LVOT were performed as described before⁷.

Hybrid approach

Method 1: The $LVOT_{\text{CSA}}$ value determined by 2D echocardiography in the CE numerator was replaced by the

MRI planimetry of LVOT and calculated $AVA = (LVOT_{\text{CSA}} \text{ MRI} \cdot LVOT\text{-}VTI \text{ Echo})/\text{transaortic-VTI Echo}$.

Method 2: The CE numerator (stroke volume, SV) was determined by MRI and the denominator (velocity time integral through the aortic valve) was measured by echocardiography. We calculated $AVA = SV \text{ MRI}/\text{transaortic-VTI (echocardiography)}$.

Statistics

All analyses were performed using SPSS statistical software (SPSS 17.0, Chicago, IL). The data are presented as mean \pm SD unless otherwise specified. A p value < 0.05 was considered statistically significant and adjusted for multiple comparisons using Bonferroni adjustment. Comparison between two or more groups was assessed by one-way analysis of variance (ANOVA) or t- Test when appropriate.

When normality and/or equal variance testing conditions were not met, the Kruskal-Wallis rank test was used. Pearson's linear correlation coefficients were calculated for continuous variables. Stepwise multiple variable regression models including variable most predictive for the dependent variables were built.

Results

Study Population

A total number of 38 patients with moderate to severe AS (16 women, 22 men, mean age 73 ± 9 years) were included in the study. Baseline demographics, clinical characteristics and key echocardiographic and MRI measurements are highlighted in table 1a. Concomitant coronary artery disease (CAD) was present in 66% of patients.

Echocardiography

According to the AVA, 26 (68%) patients were classified as having severe aortic stenosis ($AVA_{\text{index}} < 1.0 \text{ cm}^2$) and 12 individuals (32%) were classified as moderate AS, respectively. The AVA ranged $0.43\text{-}1.35 \text{ cm}^2$ (mean $0.86 \pm 0.23 \text{ cm}^2$, mean $AVA_{\text{index}} 0.47 \pm 0.12 \text{ cm}^2/\text{m}^2$, table 1b). The mean pressure gradient was $45 \pm 18 \text{ mmHg}$, and Vmax was $434 \pm 71 \text{ cm/s}$. Mean left ventricular ejection fraction (EF) was $64 \pm 12\%$. The $LVOT_{\text{CSA}}$ ranged from $3.1\text{-}4.9 \text{ cm}^2$ (mean $3.62 \pm 0.62 \text{ cm}^2$) and the LVOT diameter from $2.0\text{-}2.5 \text{ cm}$ (mean $2.14 \pm 0.1818 \text{ cm}$). The mean stroke volume (SV) was $80 \pm 15 \text{ ml/beat}$, respectively.

Magnetic resonance imaging

All examinations were diagnostic and well-tolerated by the patients. No complications occurred during any of the investigations. MRI allowed the planimetry of the AVA and $LVOT_{\text{CSA}}$ as well as measuring the diameter of LVOT in 35 individuals. Mean heart rate was 65 ± 14 beats/min. Mean AVA measured by planimetry was $0.82 \pm 0.3 \text{ cm}^2$ ($AVA_{\text{index}} 0.46 \pm 0.16 \text{ cm}^2/\text{m}^2$), while the planimetry of the $LVOT_{\text{CSA}}$ ranged from $3.0\text{-}7.5 \text{ cm}^2$ (mean $5.01 \pm 1.08 \text{ cm}^2$). Mean LVOT diameter measured in the three-chamber view was $2.36 \pm 0.47 \text{ cm}$, and the $LVOT_{\text{CSA}}$ calculated from the LVOT

Table 1 - a) Demographics and clinical data b) Echocardiographic and MRI measurements in individuals with aortic valve stenosis summarized

Demographics Variable	n = 38	Echocardiography/MRI Variable	n = 38
Age (mean ±SD)	73±9	Vmax (cm/s)	434±71
Woman (n / %)	16/42	Pmax (mmHg)	77±26
CAD (n / %)	25/66	Pmean (mmHg)	45±18
Hx of MI (n / %)	4/10	AVA (cm ²)	0.86±0.23
Hx of CABG (n / %)	2/5	AVA index (cm ² /m ² BSA)	0.47±0.12
Diabetes mellitus (n / %)	13/34	ELI (cm ² /m ² BSA. mean ± SD)	0.54±0.16
Hypertension (n / %)	33/87	E/A	1.1±0.8
Hyperlipidemia (n / %)	20/53	Smax (cm/s)	4.8±1.3
Smoking (n / %)	6/16	E' (cm/s)	4.5±1.2
ACE-inhibitor (n / %)	26/66	E/E' (mean ± SD)	20.5±8.6
β-Blockers (n / %)	29/76	CO (Echo. l/min)	5.0±2.1
Statins (n / %)	19/50	SV (MRI. ml/beat)	86±22
CRP (mg/dl)	0.94	SV (Echo. ml/beat)	80±15
Hemoglobin (g/dl)	13.3±1.6	EF (Echo. %)	64±12
Creatinine (mg/dl)	1.08	EF (MRI. %)	68±28
Blood pressure (mmHg)	143/77	LVM (Echo. g/m ² BSA)	121±36
Euroscore (%)	11.1	GLS Baseline (%)	-15.2±3.6
Euroscore additive	7.1± 0.3	Aortic root (mm)	31±3.1
NYHA	2.61±0.97	LVOT diameter (MRI. cm)	2.4±0.5
NT-proBNP (pg/ml)	2939±4211	LVOT diameter (Echo. cm)	2.1±0.18

CAD - coronary artery disease; Hx - history of; LVM - left ventricular muscle mass index; MI - myocardial infarction; SD - standard deviation; AVA - aortic valve area; AVR - aortic valve replacement; BSA - body surface area; CO - cardiac output; ELI - energy loss index; EF - ejection fraction; GLS - global longitudinal strain; LVOT - left ventricular outflow tract; LVM - left ventricular mass index; MRI - magnetic resonance imaging; Smax - peak systolic tissue doppler velocity at the septal mitral annulus; SV - stroke volume; Continues variables are presented as mean ± SD.

diameter was 4.45±1.63cm², respectively. Mean SV was 86±22ml/beat.

Table 2 shows a comparison between echocardiographic and MRI measurement. The LVOT_{CSA} calculated from the LVOT diameter with MRI techniques was significantly different compared to the standard echocardiographic determination of the LVOT_{CSA} (mean difference 0.94cm²±1.54cm², CI 0.43-1.45, p=0,001). Likewise, the results from LVOT diameter measurement with MRI or TTE techniques were significantly different (mean difference 0.22±0.45cm, CI 0.07-0.37, p=0.04). In addition, the

planimetry of the LVOT_{CSA} using MRI reveals significantly larger LVOT_{CSA} in comparison to TTE determination (mean difference 1.39±0.88cm², CI 1.09-1.67, p<0.001). Stroke volume measurements based on MRI and TTE showed a moderate correlation (r=0.56, p<0.001, mean difference 6.8±18.6ml/b, CI 0.7-12.9, p = 0.03).

Comparing the hybrid approach with standard continuity equation

Results for calculation of AVA are summarized in Table 3. There was no statistically significant difference for mean

Table 2 - Comparison of LVOT measurement with TTE and MRI

Measurement	LVOT diameter (cm)	LVOT _{CSA} calculated (cm ²)	LVOT _{CSA} Planimetry (cm ²)
TTE (mean ± SD. cm)	2.14 ± 0.18*	3.62 ± 0.62**	-
MRI (mean ± SD. cm ²)	2.36 ± 0.47*	4.45 ± 1.63*	5.01 ± 1.08*
p-value	*0.004	*0.001	*<0.001

CSA - cross sectional area; TTE - transthoracic echocardiography; MRI - magnetic resonance imaging; LVOT - left ventricular outflow tract; SD - standard deviation.

Table 3 - Results for calculation of AVA based on standard CE (TTE) and the hybrid approach

Measurement	AVA CE TTE	AVA CE Method 1	AVA CE Method 2	AVA MRI planimetry
AVA (cm ²)	0.86±0.23	1.23±0.41	0.92±0.32	0.83±0.30
AVAindex (cm ² /m ²)	0.47±0.12 ^{*1,2,3}	0.68±0.21 ^{*1}	0.50±0.16 ^{*2}	0.46±0.16 ^{*3}
p-value	-	^{*1} <0.001 ⁺	^{*2} =0.11 n.s.	^{*3} =0.14 n.s.

AVA - Aortic valve area; CE - continuity equation; MRI - magnetic resonance imaging; n.s. - not significant; TTE - transthoracic echocardiography. + = significant p<0.05.

indexed AVA between standard CE and hybrid method 2 (mean difference 0.03±0.11cm²/m², CI = -0.007-0.069, p = 0.11) and between standard CE and MRI planimetry of the AVA (mean difference 0.04±0.15cm²/m², CI -0.09-0.012, p = 0.14). On the contrary, AVA measurements by hybrid approach 1 and standard CE revealed a significant difference (mean difference 0.21±0.14cm²/m², CI -0.17-0.25, p < 0.001, Fig. 1). Correlation between standard CE and the hybrid techniques were significant (indexed AVA) for the hybrid method 1, 2 and AVA planimetry (p<0,001, figure. 2a-c), respectively. Bland-Altman plots are shown in Figure 3 a-c. Table 4 illustrates the frequency of the severity of AS by the different techniques categorized into mild, moderate and severe.

The correlation between standard CE and hybrid techniques for the non-indexed AVA was also significant for the hybrid method 1, 2 and AVA planimetry (r = 0.556, r = 0.800 and r = 0.710 respectively, all p < 0.001).

Discussion

In individuals with severe aortic stenosis, a modified continuity equation combining MRI and echocardiography revealed a very good agreement compared to standard continuity equation. The comparison of both hybrid method 2 and MRI planimetry of the AVA to standard TTE by Bland Altman analysis showed a mean difference near zero, a spread within two standard deviations (SD) and very similar limits of agreement. Furthermore, the vast

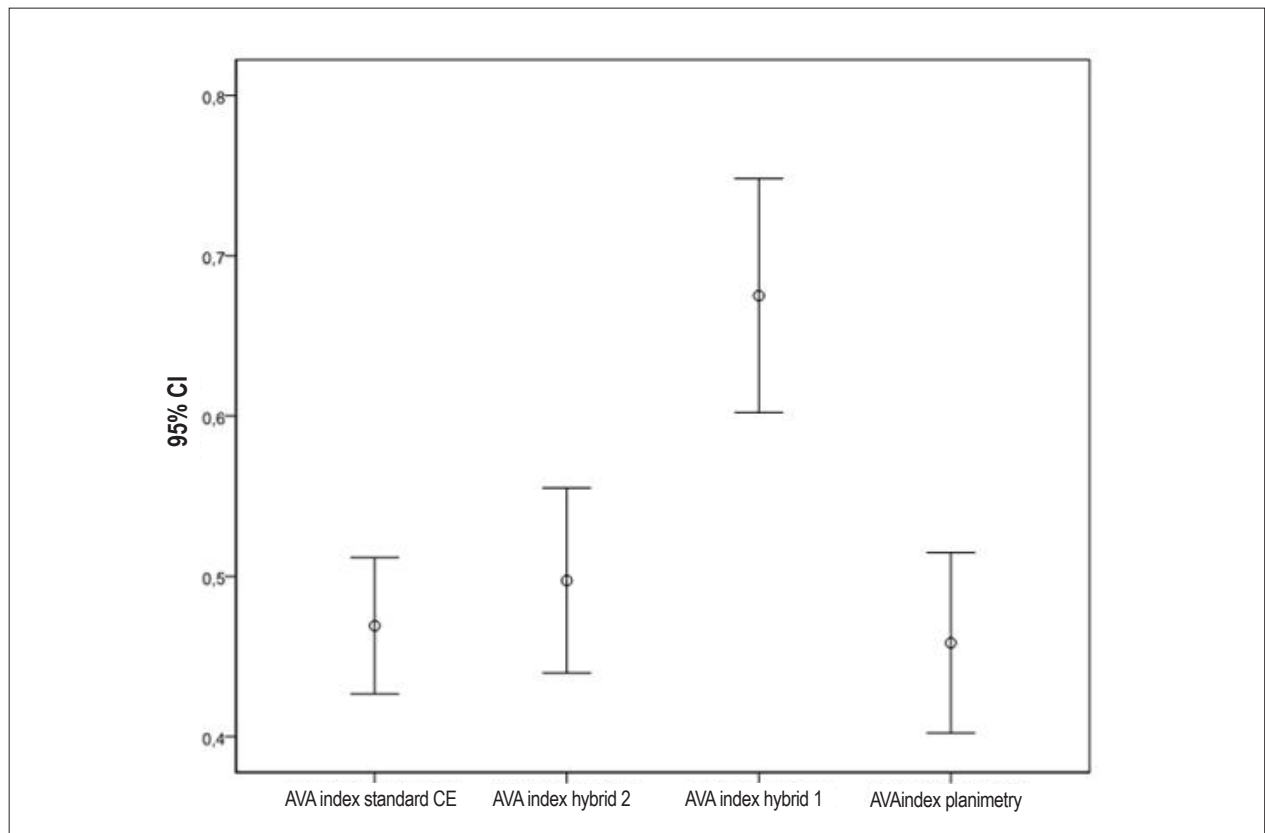


Figure 1 - Indexed aortic valve area (cm²/m²) determined by standard continuity equation and hybrid techniques; Hybrid method 1 reveals significant larger aortic valve areas indexed for body surface area (AVA index) compared to standard continuity equation based on echocardiography. There was no significant difference between standard echocardiography assessment of AVA index and hybrid method 2 and AVA planimetry, respectively. AVA - Aortic valve area

Fig. 2a

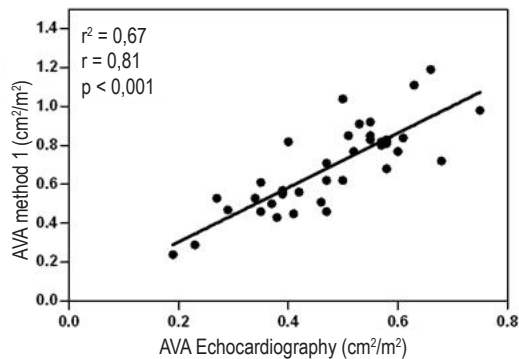


Fig. 2b

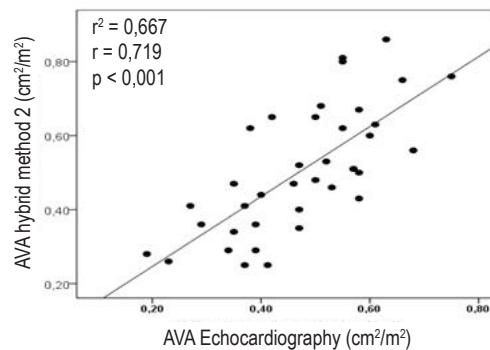


Fig. 2c

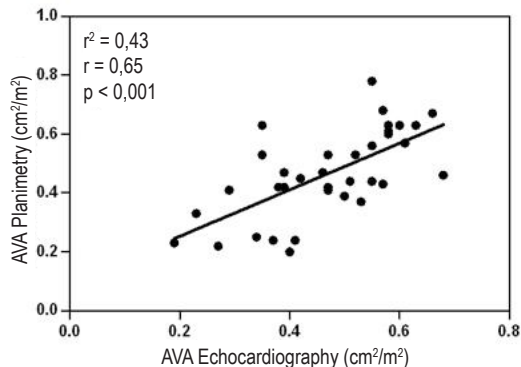


Figure 2 - Correlation of standard AVA index measurement and hybrid approach 1(a), 2(b) and MRI planimetry (c)

Figure 2a - Linear regression analysis demonstrates a significant correlation between AVA-index (cm²/m² BSA) derived by standard CE (TTE) and hybrid method 1.

Figure 2b - Linear regression analysis demonstrates a significant correlation between AVA-index (cm²/m² BSA) derived by standard CE (TTE) and hybrid method 2.

Figure 2c - Linear regression analysis demonstrates a significant correlation between AVA-index (cm²/m² BSA) derived by standard CE (TTE) and MRI planimetry. AVA - Aortic valve area; BSA - Body surface area; CE - continuity equation; TTE - transthoracic echocardiography.

majority of patients (92%) were classified as having severe AS with standard CE, hybrid method 2 and AVA planimetry. Since the ability to identify individuals with definite severe AS may have therapeutic implications; these findings are of great importance.

Calculation of AVA by means of Doppler echocardiography is a widely used standard for evaluation of aortic valve stenosis in clinical practice. Transthoracic echocardiography (TTE) plays an important role in management decisions including indications for surgery and scheduling follow-ups in individuals with AS. TTE is non invasive, easily to perform, readily available and inexpensive and offers an acceptable intraobserver and interobserver reproducibility. Therefore, a full hemodynamic study with retrograde catheterisation of the AV is no longer recommended if non invasive assessment of the severity of AS is adequate to assess valve hemodynamics.

Echocardiographic valve area calculation is based on the CE, which assumes that the flow (F) on both sides of the aortic valve orifice is equal³. Flow is defined as cross sectional area (CSA) multiplied by flow velocity (V). Across

the aortic valve (AV), the flow is equal to stroke volume (SV). Because SV is the product of the CSA and time velocity integral (VTI), the CE can be arranged to yield: $AVA = (LVOT_{VTI} \times LVOT_{CSA}) / AV_{VTI}$, in which $LVOT_{VTI}$ is the velocity time integral at the level of the stenotic valve and CSA is the cross sectional area of the LVOT.

In most echocardiographic studies, reproducibility of LVOT measurement is sufficient. Nevertheless, significant variability can occur⁸. Potential factors that may contribute to measurement errors include image quality, annular calcifications, a noncircular annulus and general failure in measuring the actual area. LVOT diameter measurement is squared; therefore any significant LVOT diameter measurement error will also be squared, producing a potentially unacceptable error in the AVA calculation. Accordingly, accurate measurement of the LVOT is critical. Previous studies have shown that the mean interobserver and intraobserver measurement variability is 5% to 8%, resulting in a variability of calculated valve area of about 0.15cm² for a borderline AS with an AVA of 1cm². Furthermore, flow acceleration in the LVOT may lead to overestimation of the Doppler-derived SV and therefore misrepresentation of

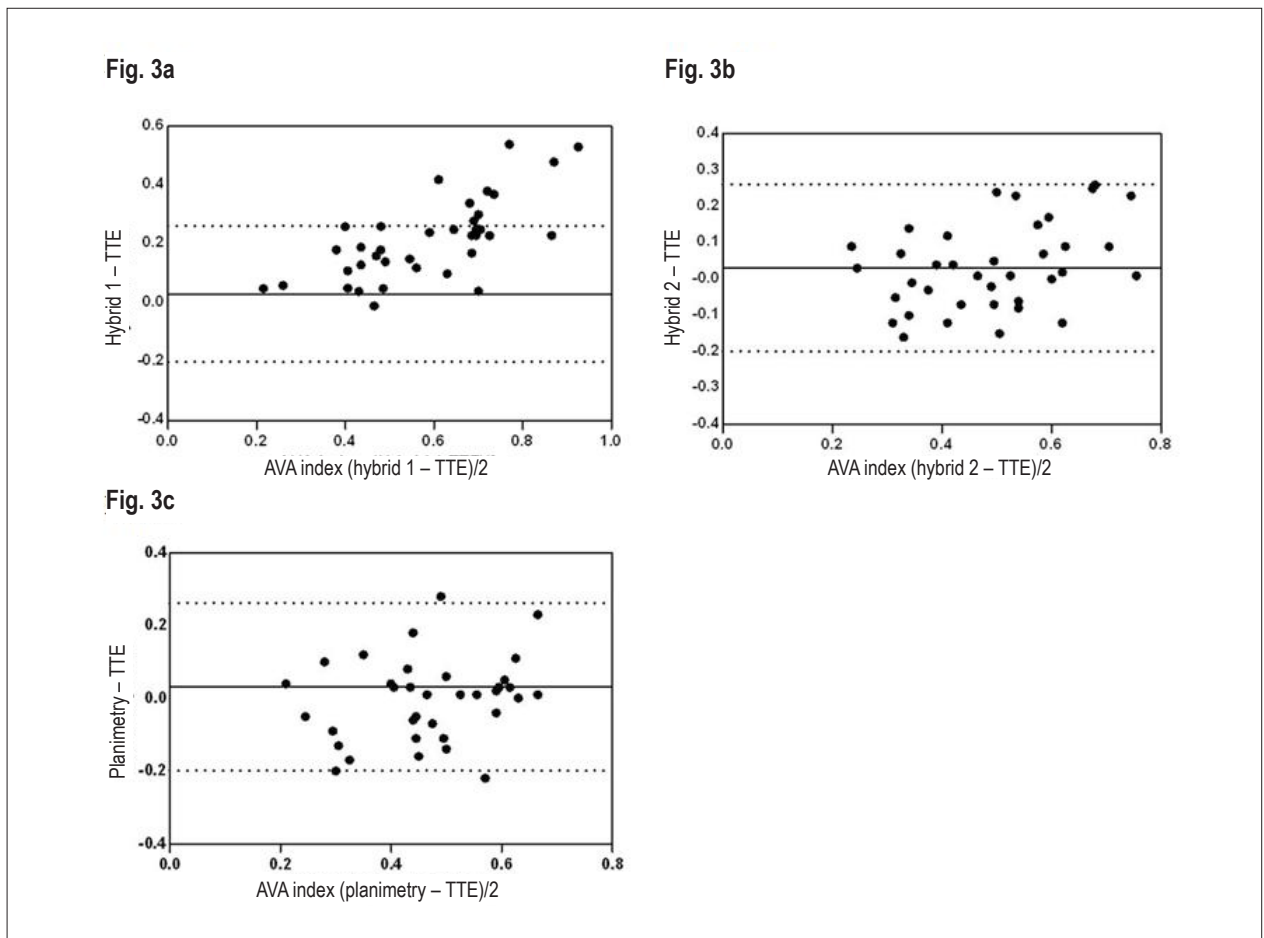


Figure 3 - Bland-Altman analyses illustrating the agreement between standard AVA-assessment and hybrid-approach 1(a), 2(b) and MRI planimetry(c).

Figure 3a - Agreement between AVA measured by standard CE (TTE) indexed for BSA (cm^2/m^2) and hybrid method 1. The continuous line represents the mean difference, and the dashed lines represent limits of agreement.

Figure 3b - Agreement between AVA measured by standard CE (TTE) indexed for BSA (cm^2/m^2) and hybrid method 2. The continuous line represents the mean difference, and the dashed lines represent limits of agreement.

Figure 3c - Agreement between AVA measured by standard CE (TTE) indexed for BSA (cm^2/m^2) and MRI planimetry of the AVE. The continuous line represents the mean difference, and the dashed lines represent limits of agreement.

AVA - Aortic valve area; BSA - Body surface area; CE - continuity equation; TTE - transthoracic echocardiography.

Table 4 - Frequency of the severity of AS by 4 different imaging techniques

Classification	TTE	Hybrid Method 1	Hybrid Method 2	AVA MRI planimetry
Mild ($> 1.5\text{cm}^2$)	-	26%	3%	-
moderate ($1.0\text{-}1.5\text{cm}^2$)	34%	40%	34%	32%
Severe ($< 1.0\text{cm}^2$)	66%	34%	63%	68%

Values are presented as percent. Aortic stenosis is categorized as mild, moderate and severe based on aortic valve area (cm^2); AVA - Aortic valve area; TTE - transthoracic echocardiography; MRI - magnetic resonance imaging.

the AVA. Hence, assessing the severity of AS is not always straightly apparent.

When echocardiographic measurements are inconsistent, invasive hemodynamic remain the gold standard. However, aortic atheroma is a common

finding in individuals with AS and is a potential source of ischaemic stroke, especially during cardiac catheterisation or surgery⁴. Therefore, a non-invasive alternative particularly in patients at high risk of procedure-associated complications is needed.

Our results demonstrate that the standard CE can be modified when necessary: the modification is simple and merely consists of replacing the Doppler-derived stroke volume by a MRI determination of stroke volume (hybrid method 2). Previous studies have demonstrated MRI as a very useful tool for evaluation of ventricular flow as it is independent of geometric assumptions, as well as accurate and reproducible⁹. On the contrary, the echocardiographic simplified Bernoulli equation ignores the flow acceleration proximal to the stenosis, which is an acceptable assumption as long as the transvalvular flow velocity is significantly greater than the proximal flow velocity. However, in the presence of accelerated flow or less significant stenosis, the simplified Bernoulli equation is less accurate.

Flow acceleration in the LVOT can affect AVA calculation by overestimating the actual AVA¹⁰. By contrast, SV calculation by MRI is not affected by flow acceleration in the LVOT. Therefore, replacing the SV in the CE by MRI derived SV measurement may be more accurate in situations where significant flow acceleration is present in the LVOT. On the other hand, Doppler-derived gradients correlate well with invasively measured pressure gradients as it has been demonstrated in the experimental and in the clinical setting^{11,12}. Therefore, combining measurements of SV by MRI with measurements of VTI across the aortic valve by echocardiography in hybrid approach 2 takes advantages of both methods. It yields results with very small differences compared to standard CE and accurately identifies the vast majority of severe AS. Using a hybrid approach bypasses the need for accurate measurement of the LVOT diameter and LVOT flow and therefore may be an acceptable alternative when measuring the LVOT is not possible.

Furthermore, planimetric-derived AVA by MRI is accurate and correlates well with the one obtained by echocardiography. These findings are congruent with the literature published concerning this matter¹³. Although attractive, direct planimetry of the AVA by MRI is nonetheless a technically demanding method. It requires precise measurement at the valve tips during their maximal opening. This can be challenging as there may be heavy calcifications of the tips leading to image artefacts, and a precise view of the aortic valve leaflets, since partial volume effects may be difficult. In addition, MRI planimetry represents a single time point measurement of AVA at maximal systole, while CE is measured as average over the whole ejection period. The AVA, however, may vary substantially during the ejection period¹⁴.

In contrast to planimetry and hybrid method 2, hybrid method 1 failed to demonstrate a good agreement compared to standard CE. The MRI images in our study reveal that the LVOT geometry resembles a more elliptical rather than a circular shape and therefore a larger LVOT_{CSA} was determined. Replacing the LVOT_{CSA} measurement in the standard CE with MRI planimetry of the LVOT, it remains doubtful whether the standard CE permits an accurate calculation of the AVA, because standard CE assumes a circular shape of the LVOT_{CSA}. Accordingly, whether LVOT is obtained with echocardiography or MRI, these two approaches cannot be used interchangeably.

In general, despite the fact that MRI overcomes some of the methodological limitations of echocardiography mentioned above, it seems unlikely for MRI to replace TTE as the image modality of first choice in evaluating individuals

with aortic stenosis. However, our data are consistent with previously published studies, suggesting that a hybrid approach combining TTE and MRI as a promising alternative diagnostic tool for evaluating the severity of AS noninvasively when echocardiography is inconclusive.

Limitations

The cohort of 38 individuals included in this pilot study is relatively small and does not allow performing a subgroup analysis. Therefore, we must remain cautious in the interpretation of these data considering the relatively small number of patients. However, these preliminary results encouraged us to investigate this issue in a larger study group.

Furthermore, only a small proportion of patients had invasive measurement of AVA by cardiac catheterization. Therefore, the results comparing standard CE and the hybrid approach should be analysed with caution because the “gold standard” in determining the severity of AS is not available, therefore new methods should be judged. Nevertheless, invasive measurement of AS is based on the “Gorlin formula” which has many well-recognized limitations¹⁵, whereas the continuity equation method for calculation of AVA has the advantage of being non-invasive and of not containing an empirical constant¹⁶.

Another limitation of our study was a well preserved EF in most patients included in the study ($64 \pm 12\%$). However, especially in patients with severely impaired EF (low flow, low gradient aortic stenosis), the accurate assessment of AVA using Doppler TTE is challenged by the limitations of Doppler flow parameters¹⁷.

Conclusions

The calculation of AVA by a non-invasive hybrid approach between echocardiography and MRI is a good alternative to standard CE and correlates well with the standard method. The hybrid approach is more time consuming, not widely available and more cost intensive than standard CE and will certainly not replace TTE in clinical routine. However, this method could be used as a non-invasive alternative in patients in whom echocardiographic results are inconclusive or in whom conflicting results require further clarifications, particularly in patients with a high risk for an invasive hemodynamic study.

Potential Conflict of Interest

No potential conflict of interest relevant to this article was reported.

Sources of Funding

There were no external funding sources for this study.

Study Association

This study is not associated with any post-graduation program.

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