Sonomicrometry-derived 3-dimensional geometry of the human tricuspid annulus

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ABSTRACT

Objectives: Surgical correction of functional tricuspid regurgitation is focused on prosthetic reduction and remodeling of the tricuspid annulus. We set out to investigate the precise geometry of the human tricuspid annulus to better guide surgical therapy.

Methods: Eleven human donor hearts with normal right ventricular function and without tricuspid regurgitation that were rejected for clinical transplantation were harvested. Sonomicrometry crystals were sewn around the tricuspid annulus and pressure sensors placed in the right ventricle and right atrium. The hearts were studied in the TransMedics Organ Care System (Andover, Mass) ex vivo perfusion apparatus in the right heart working model. Data were acquired at baseline and before and after bolus calcium infusion. Annular height, dimensions, strain, and curvature were calculated based on 3-dimensional crystal coordinates.

Results: Maximal annular area was 997 ± 258 mm² and minimal 902 ± 257 mm² with contraction of 10% ± 5% at baseline and 19% ± 6% after calcium (P = .007). Segmental contractility of anterior, posterior, and septal annular regions was 7% ± 5%, 6% ± 4%, and 6% ± 3%, respectively. Only anterior region had increased contractility after calcium infusion (to 15% ± 5%; P = .023). Annulus had its high points at anteroseptal commissure and the midposterior region and lowest point in the midseptal region with maximal and minimal height of 5.0 ± 1.1 mm and 4.0 ± 1.1 mm, respectively. The greatest curvature responsible for out of plane annular bending was observed at annular high points.

Conclusions: The human tricuspid annulus is a complex 3-dimensional dynamic structure with its high points and maximal degree of bending at the anteroseptal commissure and midposterior annulus. These detailed geometric data may aid the design of more physiologic annular prostheses and surgical reparative techniques. (J Thorac Cardiovasc Surg 2019;157:1452-61)
knowledge gap is fundamental to the advancement of current surgical therapy of FTR and to the reduction of recurrent disease.2

Several clinical studies have described human tricuspid annulus (TA) geometry utilizing magnetic resonance imaging,3 computed tomography,4,5 and 2-dimensional6 or 3-dimensional (3D) echocardiography.7-9 However, these advanced clinical imaging modalities cannot reliably assess small changes in valvular 3D geometry nor follow discreet annular or subvalvular points in time. Due to morphology of the right ventricle (RV) and complexity of the TV, clinical imaging does not allow for the detailed analysis of individual segments of the TA or the subvalvular apparatus.10 3D echocardiography is limited by low frame rate acquisition and inability to record the entire undisturbed heart cycle,9 whereas use of magnetic resonance imaging in the 3D assessment of valvular structures is possible only after multiplane reconstructions.3 Experimental methodology such as crystal sonomicrometry is much better suited to capture the subtle geometric alterations that are associated with cardiac chamber and valvular remodeling.11 and this technology has been used to mark specific sites on the TV complex, and elucidate detailed valvular geometry and dynamics in animal models.12-14 However, sonomicrometry requires surgical implantation of miniature crystals and a wired connection, and therefore cannot be used to study human hearts in normal clinical settings. The aim of the current study was to describe the detailed 3D geometry and dynamics of the human TA utilizing sonomicrometry crystal technology in isolated, working human hearts rejected for clinical transplantation.

METHODS
The study was evaluated by our local Spectrum Health Internal Review Board and determined as nonhuman research as defined by Department of Health and Human Services and Food and Drug Administration regulations (Spectrum Health Institutional Review Board No. 2016-248).

A total of 11 donor hearts rejected for clinical transplantation with normal RV function and without tricuspid regurgitation were studied. Hearts were harvested from donors with a mean age of 50 ± 13 years and a mean weight of 78 ± 19 kg. Six donors were men, and the mean left ventricular ejection fraction was 57% ± 4%. None of hearts had more than mild tricuspid regurgitation or right ventricular dysfunction. The average heart cold ischemia time was 118 ± 67 minutes. The mean coronary sinus lactate level at the time of data acquisition was 3.7 ± 1.1 mmol/L with arterial level at 3.9 ± 1.0 mmol/L indicative of cardiac lactic acid absorption and adequate physiologic resuscitation.15 No increase in tricuspid regurgitation from preoperative state was observed in the right heart working mode.

Surgical Preparation
All hearts were harvested using the normal clinical procurement protocol. The hearts were rejected for clinical transplantation due to infection concerns (hepatitis B and C), mild to moderate left system disease on coronary angiography, or mild left ventricle dysfunction. At least 1200 mL heparinized donor blood was collected in a blood storage system immediately before aortic crossclamping. Subsequently, the aorta was crossclamped and 2000 mL cold HTK-Custodiol solution (Essential Pharmaceuticals, Durham, NC) was delivered in the aortic root for myocardial preservation. After excision, the heart was transported on ice to the experimental site, and subsequently the right atrium was opened and 6 sonomicrometry crystals (Sonometrics Inc, London, Ontario, Canada) were sewn around the TA with 5-0 polypropylene suture (Figure 1); 4 along the mid-RV free wall and 1 at the apex. A solid-state pressure transducer (PA4.5-X6; Konigsberg Instruments Inc, Pasadena, Calif) was placed through the RV apex for RV pressure measurements and 1 in the right atrium for right atrial pressure measurements. The right atriotomy was closed and both superior and inferior vena cava oversewn. The aorta was sized and cannulated with a connector and the pulmonary artery was instrumented with a 30F cannula in preparation for the connection to the Organ Care System (OCS) (TransMedics, Andover, Mass).

The OCS is a portable ex vivo organ perfusion platform used in clinical transplantation and designed to perfuse the beating, unloaded donor heart with warm, oxygenated blood during transport to the transplant center (Figure 1). The OCS device was primed with donor blood in accordance to the clinical protocol and the heart mounted on the device, defibrillated if spontaneous rhythm did not return, ventricularily paced at 80 bpm, and perfused under normothermic conditions. The heart was perfused with a mean arterial blood pressure of 75 mm Hg in the clinical unloaded mode. The pumped blood to the aorta constituted the inflow of the oxygenated blood to the heart. Through the coronary arteries it reached the coronary sinus and served as the inflow to the right heart. The blood was subsequently ejected by the RV to the pulmonary artery outflow cannula and collected in the venous reservoir. Arterial (aortic inflow) and coronary sinus venous (pulmonary artery cannula) blood gases, including lactate levels, were collected every 15 minutes to assure stability of the experimental preparation.15

Data Acquisition
After 60 minutes of reperfusion, the outflow pulmonary artery cannula was partially occluded allowing the right heart to be filled with blood. The pulmonary artery was narrowed to achieve maximal RV pressure of 30 mm Hg, thus converting to the right heart working mode. TV competence was assessed with epicardial echocardiography utilizing a Vivid S6 machine (GE Healthcare, Chicago, Ill).

Hemodynamic and sonomicrometry data were acquired simultaneously under working right heart conditions (baseline n = 11). The last 5 hearts in the study were additionally tested before (precalcium) and after (calcium) 500 mg calcium chloride bolus infusion to assess the changes in annular geometry and strain under contractile stimulation.

Sonomicrometric Data
All sonomicrometry data were acquired using a Sonometrics Digital Ultrasonic Measurement System DS3 (Sonometrics Inc) as
Previously described, data from 3 consecutive cardiac cycles during hemodynamically stable conditions (Table E1) were averaged for each heart. Given that all 3 cardiac cycles were recorded in immediate succession, there were no physiologic or external reasons for changes in the signal. Therefore, averaging consecutive cardiac cycles increased the signal to noise ratio and made data more reliable. Data were acquired at 128 Hz with simultaneous RV pressure and right atrial pressure, and electrocardiogram recordings. The time-resolved 3D coordinates of the crystals were obtained from the SonoSoft software (Sonometrics Inc) and analyzed in Matlab (MathWorks, Natic, Mass) using custom-designed software. Least-squares cubic splines were fit to the discrete sonomicrometry annular crystal data to create approximations of the TA at each time point and for each heart. Spline-based annular metrics add to the standard distance-based sonomicrometry analysis by allowing to spatially resolve the configurational changes of the annulus as it undergoes complex in-plane and out-of-plane deformations during the cardiac cycle. This spatially continuous mathematical description was used to obtain complete height profile of the TA. Moreover, the annular 3D geometry was also expressed as the maximal displacement of annular crystals from annular best-fit plane. Annular height to commissural width ratio (AHCWR) was calculated as the ratio of maximal displacement of annular crystals from annular plane (height) to its anteroposterior (A-P) diameter. Annular area and perimeter were calculated based on the least-squares cubic splines. The septolateral (S-L) annular dimension was calculated as the distance between crystal 11 and the midpoint between crystals 7 and 8; the A-P annular dimension was calculated as the distance between crystal 1 and the midpoint between crystals 6 and 9. The eccentricity index was calculated as the A-P to S-L cubic splines. The septolateral (S-L) annular dimension was calculated as the distance between crystal 11 and the midpoint between crystals 7 and 8; the A-P annular dimension was calculated as the distance between crystal 1 and the midpoint between crystals 6 and 9. The eccentricity index was calculated as the A-P to S-L distance ratio. Intercommisural distances were calculated as the respective distances between crystals 6, 8, and 10. Annular contraction was computed as the difference between the maximal and minimal TA area ([Amax – Amin]/Amin × 100%). Sonomicrometry-derived RV contraction (ie, fractional area change) was calculated as the difference between the maximal and minimal area enclosed by crystals 1 to 4 ([AED – AES]/AED × 100%) and served as a surrogate of RV performance. RV volume was assessed using convex hull method. All values were calculated at end-diastole (ED), end-systole (ES), end of isovolumic contraction, end of isovolumic relaxation, maximal valve opening (defined as the time when annular area was the largest), minimal valve opening (the time when annular area was the smallest), ED was defined as the time of the beginning of the positive deflection in electrocardiogram voltage (R wave), whereas ES was determined to be the time of maximum negative dP/dt of the RV pressure. End of isovolumic contraction and end of isovolumic relaxation were determined as end points of the linear RV pressure segments during isovolumetric contraction and relaxation, respectively.

**Annular Strain and Curvature Analysis**

TA strain was calculated based on the first spatial derivative of the spline fits using the method previously described. Strain, as a relative measure of displacement and thus annular deformation, is calculated relative to a reference configuration. Baseline at ED and at maximal valve opening chosen as the reference configurations for other time points within the cardiac cycle. For the last 5 hearts in the study, we also compared strains after calcium stimulation with precalcium configuration used as the reference for the same time points during calcium infusion. Therefore, cardiac strain first reflects the deformation of the annulus in baseline conditions and intervention strain reflects the deformation of the annulus induced after calcium. Specifically, Green-Lagrange strain was calculated along the annulus for each heart and later displayed on a spline representation of the population-averaged annulus. The global and regional strain patterns were calculated for the entire annular circumference (crystals 6-11) and the anterior (crystals 6-8), posterior (crystals 8-10), and septal (crystals 10-6) regions separately. Negative or compressive strains imply that tissue is contracted, whereas positive or tensile strains imply that tissue is stretched. The absolute strains have positive values only and are indicative of annular deformation. Local annular curvature was calculated based on the principles previously presented in detail.

**Statistical Analysis**

Data are presented as mean ± standard deviation unless otherwise stated. The baseline data shown are predominantly descriptive. The within heart cycle comparisons and the precalcium and calcium conditions were tested with the use of Student t test for dependent observations or Wilcoxon
signed-rank test for repeated measurements when the normality test failed (SigmaPlot 12.5; Systat, San Jose, Calif).

RESULTS

Hemodynamic Parameters

During baseline conditions, mean heart rate was 80 ± 2 bpm, maximal RV pressure was 28 ± 4 mm Hg, RV end-diastolic pressure was 8 ± 3 mm Hg, and right atrial pressure was 6 ± 3 mm Hg. Hemodynamic variables before and after calcium infusion are presented in Table E2. The expected increase in the RV fractional area change was noted after stimulation with calcium with no difference in other measured parameters.

TA Size

The maximal area of the TA in diastole was 997 ± 258 mm² and minimal area during systole was 902 ± 257 mm². The maximal perimeter was 115 ± 14 mm and minimal perimeter was 110 ± 14 mm. The S-L distance at the maximal valve opening was 30.8 ± 5.7 mm and A-P diameter was 39.4 ± 6.6 mm. At minimal valve opening, S-L diameter was 28.7 ± 5.9 mm and A-P diameter was 36.4 ± 6.9 mm. Annular area and dimensions throughout the cardiac cycle at Baseline are shown in Figure 2. Values for annular distances before and after calcium infusion are illustrated in Table 1. Of note is the disproportionate diastolic enlargement between A-P and S-L diameter. The eccentricity index increased from 0.76 ± 0.19 in diastole to 0.85 ± 0.19 during systole at baseline (P = .007). There was no change in annular eccentricity after calcium, and within-heart cycle dynamics remained unaffected: 0.78 ± 0.21 during diastole and 0.92 ± 0.20 in systole (P = .029).

Annular Geometry

The 3D structure of the reconstructed annulus revealed its high points at the anteroseptal commissure and the midposterior annulus, whereas low points were observed in the septal and the A-P region as demonstrated in Figure 3. Annular height changed dynamically during the heart cycle by 20% ± 10% with maximal height of 5.0 ± 1.1 mm and minimal height of 4.0 ± 1.1 mm. Annular height and curvature maps at baseline are presented in Figure 4. Video 1 illustrates the annular height during 3 recorded heart cycles at baseline. Calcium stimulation did not alter annular height, maintaining the maximal height at 5.4 ± 1.3 mm and minimal height at 4.0 ± 1.5 mm (P = .9 and P = .51 vs precalcium, respectively). AHCWR at baseline was 12% ± 5% at maximal valve opening and 13% ± 6% at its minimum. No significant effect on AHCWR was observed with calcium at either maximal and minimal valve opening.
TA Dynamics

The TA dynamically changed its dimensions during the heart cycle with an area change of 10% ± 5% and perimeter change of 5% ± 2% at baseline, and 19% ± 6% and 8% ± 3% after calcium, respectively. The annulus presented uniform dynamics across all its segments at baseline. From diastole to systole, the anterior annulus contracted by 7% ± 5% from 49.5 ± 5.6 mm to 46.1 ± 6.7 mm (P < .001), the posterior annulus by 6% ± 4% from 28.3 ± 6.9 mm to 26.6 ± 7.1 mm (P < .001), and the septal annulus by 6% ± 3% from 39.4 ± 8.5 mm to 36.8 ± 7.5 mm (P < .001). Regional annular segment lengths and intercommisural distances before and after calcium stimulation are presented in Table 1.

Annular Strains

The TA showed significant deformation during the heart cycle with mean annular strain of −4% ± 2% at TV closure (P < .001 vs opening). All annular segments presented with compression during valve closure with largest at anterior annulus (−5% ± 4%) and posterior annulus (−6% ± 3%) and smallest at the septal annulus (−2% ± 2%) (all P values < .05 vs valve opening). Annular strain at baseline during 3 consecutive heart cycles is presented in Video 2. Absolute annular strains showing annular deformation in relation to ED across all regions are shown in Figure 5. The strain pattern with calcium infusion revealed heterogenous strain along the annulus with the minimal compression in the septal portion and the largest contraction in the anterior annulus close to the anteroseptal commissure (Figure 6).

DISCUSSION

The TA is the main therapeutic target in the surgical treatment of FTR, and its detailed morphologic mapping...
is critical to the design of annuloplastic prostheses that may better re-establish the physiologic geometry of the diseased valve. In the current study, we describe for the first time the detailed 3D geometry of the human TA using sonomicrometry crystals to mark discreet annular sites. We found the TA to be a complex 3D structure with its high points near the anteroseptal commissure and the midposterior annulus and low points in the midanterior and midseptal segments. The area of highest curvature was observed at the anteroseptal commissure and midposterior annulus. This complex spatial configuration was not altered by contractile stimulation with calcium.

Initial human cadaver studies of the TA described it as a flat structure. In 2004, Hiro and colleagues implanted sonomicrometry crystals onto the TV complex in sheep and showed the annulus to be a saddle-shaped structure with local maxima at the posterior annulus, the anteroseptal commissure, and the anterior annulus and minima at A-P and posteroseptal commissures. Together with its intricate geometry, the annulus was described to have an elliptical shape, becoming more circular during maximal valve opening. These results were later corroborated using the same experimental technology in pigs, and are consistent with findings reported by our laboratory and other investigators. The current human data support these experimental findings and suggest preservation of the TA shape across mammalian physiology.

Using clinical imaging, the human TA has also been described as nonplanar structure with an oval shape with multiple studies reporting the geometry of the TA using 3D echocardiography, magnetic resonance

**FIGURE 4.** Top panel, Tricuspid annular height map at baseline (n = 11). Bottom panel, Curvature map at baseline (n = 11). The metrics are shown on the average annular spline. Height is presented as the distance from the best-fit tricuspid annular plane. Crystals 6, 8, and 10 represent anteroseptal, anteroposterior, and posteroseptal commissures, respectively. ED, End-diastole; EIVC, end of isovolumic contraction; ES, end-systole; EIVR, end of isovolumic relaxation; Max, maximal valve opening; Min, minimal valve opening.

**VIDEO 1.** Tricuspid annular height at baseline. Height is expressed as the maximal displacement from the annular best-fit plane. Video available at: https://www.jtcvs.org/article/S0022-5223(18)32505-4/fulltext.

**VIDEO 2.** Tricuspid annular strain at baseline. Video available at: https://www.jtcvs.org/article/S0022-5223(18)32505-4/fulltext.
imaging, and computed tomography. The details of the saddle-like shape of the annulus vary between reports. There appears to be a consensus that, similar to current findings, the highest annular point is at the anteroseptal commissure, but some reports present the highest point at the posterior annulus only. The saddle geometry of the human TA mirrors the well-investigated shape of the mitral annulus, which was demonstrated to be potentially beneficial in reducing mitral leaflet stress/strain. However, the TV is more complex and only direct in vivo leaflet strains measurements can provide a definite answer to this important question. The TA undergoes dynamic changes during the heart cycle with maximal area and perimeter in diastole and minimal area and perimeter in midsystole, and we observed a similar pattern for tricuspid leaflet stretch with either a flat or a saddle-shaped annulus. AHCWR calculated in the current study for the human TA (around 10%–15%) was at a level where a 5% to 13% peak leaflet strain reduction could be expected if one were to extrapolate available mitral data. However, the TV is more complex and only direct in vivo leaflet strains measurements can provide a definite answer to this important question. The TA undergoes dynamic changes during the heart cycle with maximal area and perimeter in diastole and minimal area and perimeter in midsystole, and we observed a similar pattern for.

FIGURE 5. Regional absolute average annular cardiac strain. The strain is calculated relative to end-diastole at baseline (n = 11). The absolute strains are positive in value and do not reflect stretching or shortening. Crystal numbers defining annular segments as presented in Figure 1. A, Anterior annulus. B, Posterior annulus. C, Septal annulus. The upper and lower borders of the box represent the upper and lower quartile. The middle horizontal line represents the median. The upper and lower whiskers represent the maximum and minimum values. ES, End-systole; EIVC, end of isovolumic contraction; EIVR, end of isovolumic relaxation; MAX, time of maximal annular area; MIN, time of minimal annular area. *P < .05 versus end-diastole.
annular area and perimeter change in our study. However, the magnitude of annular area reduction during the cardiac cycle of approximately 10% was lower than previously reported. Tsakiris and colleagues demonstrated canine tricuspid annular reduction of 39% during the cardiac cycle, whereas sonomicrometry-derived annular area contraction of 21.9% and 21% has been reported in sheep and pigs, respectively. Clinically, tricuspid annular area was shown to decrease 29.6% to 35% during the heart cycle as measured by transthoracic echocardiography. The lower annular contraction observed in our study may be attributed to the experimental preparation of isolated, denervated hearts subjected to cold ischemic storage and without normal physiologic flow. However, the hearts used in our study remained responsive to contractile stimulation with calcium, and biochemical analysis suggested lactate absorption indicative of adequate myocardial resuscitation. Our results are comparable to the 3D transesophageal echocardiography study of Owais and colleagues in anesthetized patients undergoing coronary artery bypass surgery who reported tricuspid annular area contraction of 10%. Furthermore, the degree of TA contraction seen in the current study is similar to clinical and experimental area reduction of the mitral annulus.

The most responsive regions of the TA in isolated human hearts were anterior and midposterior segments. These portions of the annulus coincided with locations of the highest annular curvature, presenting as the sites of out-of-plane annular folding. In-plane diameter analysis revealed that changes in the A-P dimension were the primary drivers of the annular ellipse becoming more circular during the cardiac cycle, as reported previously by others. Limited S-L dynamics may be explained by the characteristics of the septal annulus and the more distensible RV free wall encircling most of the posterior annulus. The conformation changes in both in- (planar dimensions) and out-of- (height, curvature) plane human TA geometry were very similar to the mechanics of the ovine annulus reported by our group previously.

The least responsive portion of the TA to inotropic stimulation was the septal region, with strain of only –2% after calcium infusion. These findings may be explained by high stiffness of this segment due to its dense collagen content, although the septal annulus has previously been shown to be equally susceptible to annular dilation as other regions. Because the left ventricle was unloaded in our experimental preparation, it is feasible that septal contribution to annular dynamics was reduced by diminished interventricular interaction. However, we have previously shown in the ovine model that unloading the left ventricle in vivo has minimal effect of annular geometry or dynamics.

**Clinical Significance**

The results of our study may have important clinical implications. Currently used tricuspid annuloplasty prostheses greatly vary in their geometric shape yet lack clear scientific basis to support their design. Most commonly implanted 3D annuloplasty rings incorporate axial heights from 9 to 15 mm, which is significantly greater than the approximate 5- to 6-mm annular height of the normal human TAs reported herein. Detailed geometrical analysis of a human TA along with stress/strain mapping may therefore drive a more rational design of the shape and material properties of more physiologic annular prostheses. Specifically, strain and curvature data may aid in a detailed, spatially resolved understanding of annular dynamics. Strain and curvature changes throughout the cardiac cycle could thus inspire devices with differential stiffness to accommodate the native motion of the valve. This is of utmost importance for improved clinical results and solidifying the concept of annular remodeling in the treatment of functional tricuspid regurgitation.

**Limitations of the Study**

The results of our study must be interpreted in the context of important limitations. This study was performed in a small number of isolated human hearts, and the RV working mode was induced by increased RV afterload. Moreover, the use of ex vivo retrograde perfusion system keeps the left
ventricle unloaded throughout the entire experiment and excludes the influence of interventricular dependence. However, we have demonstrated previously that unloading the left ventricle in vivo does not alter the geometry of the TAs.26 Denervation of the heart and need for ventricular pacing may have influenced the dynamic behavior of the TA but would be unlikely to affect annular geometry, which was the main focus of this study. Similarly, lack of cardiac output at the physiologic level through the RV may be expected to influence annular contraction but have a limited effect on its 3D shape. Because of protocol change during the course of the study, only the 5 last hearts were tested before and after calcium stimulation. We chose to analyze 3 consecutive heart cycles to provide more reliable averaged data for all presented parameters. Although this is a short data acquisition time, we believe the results are reliable due to the stable condition of the experimental setup (Table E1). The same number of heart cycles was previously shown to be sufficient in a similar mitral valve study.17 The current study analyzed only the TA and did not investigate other key components of the valvular complex such as the valve leaflets and the subvalvular apparatus.

CONCLUSIONS
Using crystal sonomicrometry in isolated beating hearts, we found the human TA to be a complex nonplanar structure with distinct high points around the anteroseptal commissure and midposterior annulus with low points in the septal and the A-P regions. The TA maintained a saddle-like shape with out-of-plane bending and highest curvature at the anteroseptal commissure. The 3D geometry of the annulus was stable during the cardiac cycle and not influenced by inotropic stimulation. The planar shape of the annulus at ED resembled an ellipse that became more circular during ES. These geometric findings (shown in the Graphical Abstract) should be considered in the design of more physiologic TA prostheses.

Conflict of Interest Statement
Authors have nothing to disclose with regard to commercial support.

TransMedics Inc (Andover, Mass) provided the Organ Care System for research use and technical support during the experiments.

References
23. Spinner EM, Buice D, Yap CH, Yoganathan AP. The effects of a three-dimensional, saddle-shaped annulus on anterior and posterior leaflet...


Key Words: tricuspid valve, tricuspid annulus, strain, sonomicrometry
### Table E2. Hemodynamic parameters after calcium administration (n = 5)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Precalcium (mean ± SD)</th>
<th>Calcium (mean ± SD)</th>
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<td>Heart rate (bpm)</td>
<td>8.0 ± 1</td>
<td>8.2 ± 3</td>
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<td>RVP max (mm Hg)</td>
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<td>RAP (mm Hg)</td>
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<td>4 ± 2</td>
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<tr>
<td>RV EDV (mL)</td>
<td>97 ± 26</td>
<td>80 ± 14</td>
<td>.12</td>
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</table>

Data are presented as mean ± SD. RVP, Right ventricular pressure; RV EDP, right ventricular end diastolic pressure; RV FAC, right ventricular fractional area change, RAP, right atrial pressure; RV EDV, right ventricular end diastolic volume. *Data obtained from sonomicrometry.