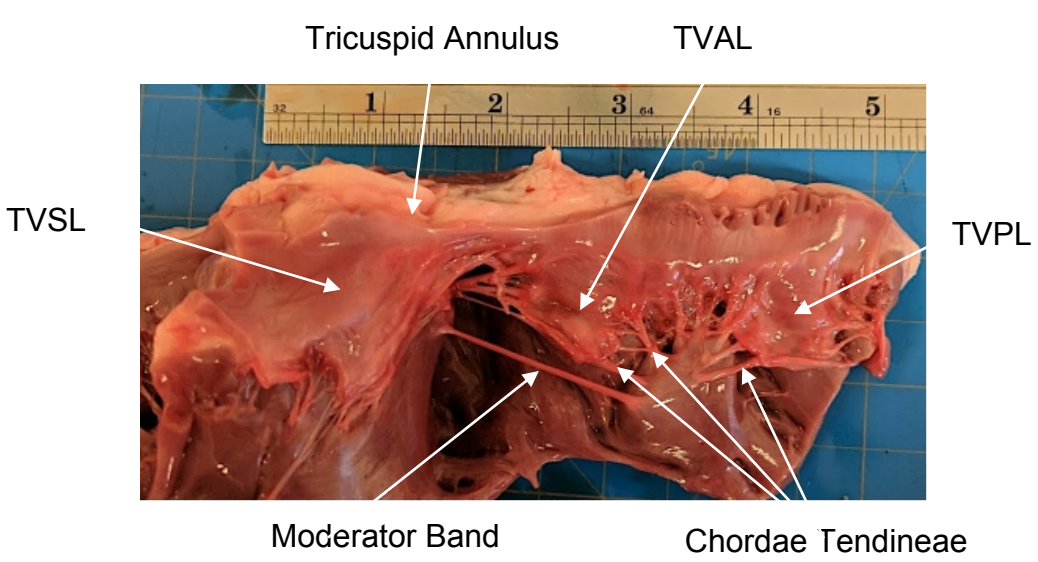
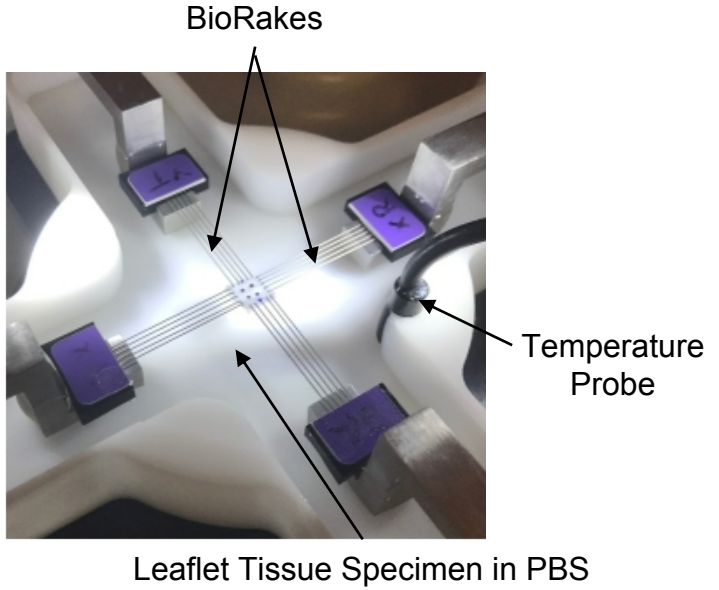
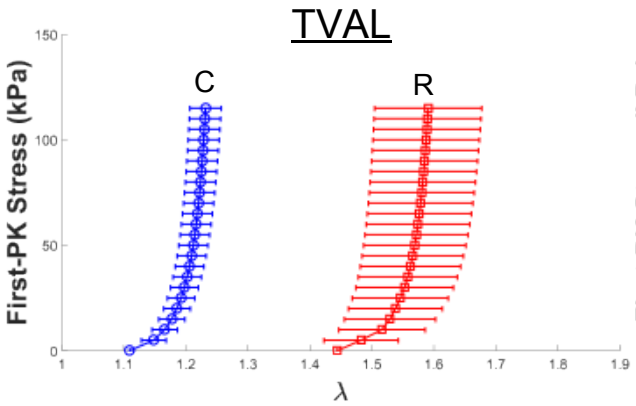
The novelties and research highlights of this work are briefly summarized below:

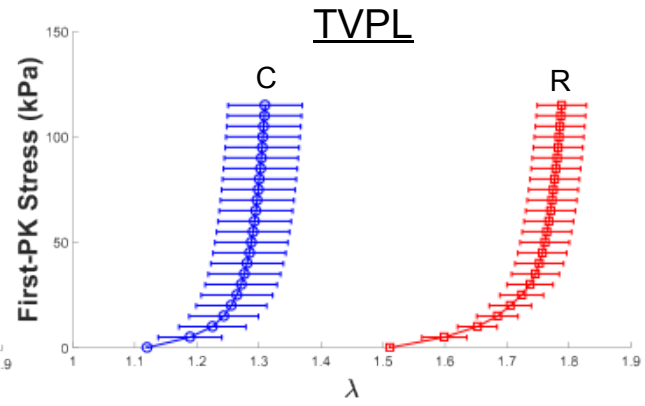
- A comprehensive characterization of the anisotropic mechanical behaviors of porcine atrioventricular heart valves (mitral and tricuspid valves) were thoroughly conducted;
- Loading-rate and temperature effects on the valve tissue mechanics were investigated;
- Histology analyses were performed to quantify four morphologically distinct layers of the valve leaflet tissues and were linked to their mechanical responses;
- Within-species and cross-species commonalities and discrepancies in the tissue mechanical responses were examined and, for the first time, reported; and
- The biaxial mechanical testing data as acquired in this study will be submitted as Data In Brief and shared as open-access resources for future development of heart valve constitutive models.

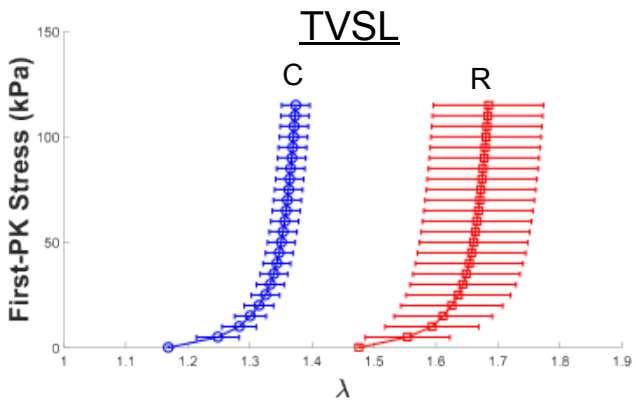
Downloaded from SHAREOK. Persistent Handle URL: https://hdl.handle.net/11244/301842

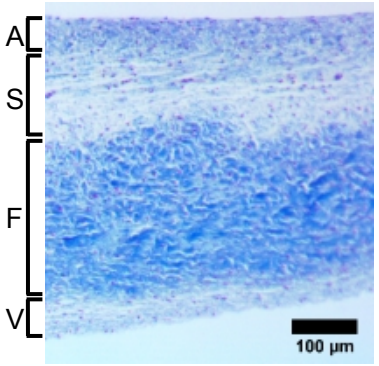

Tricuspid Annulus
TVAL
TVSL
TVPL
Moderator Band
Chordae Tendineae

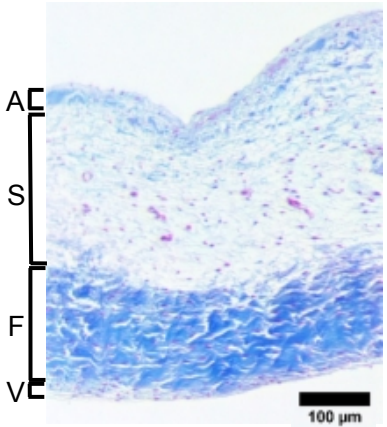

BioRakes
Temperature Probe
Leaflet Tissue Specimen in PBS

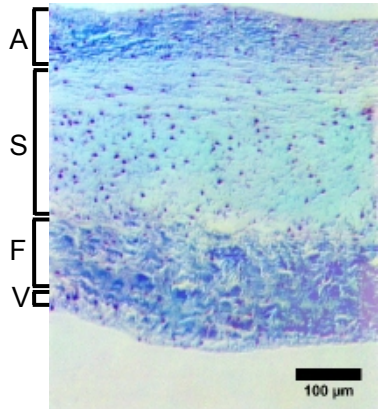

TVAL
C
R
First-PK Stress (kPa)
λ


TVPL
C
R
First-PK Stress (kPa)
λ


TVSL
C
R
First-PK Stress (kPa)
λ


A
S
F
V
100 μm


A
S
F
V
100 μm


A
S
F
V
100 μm

An investigation of the anisotropic mechanical properties and anatomical structure of porcine atrioventricular heart valves

Samuel Jett[1*], Devin Laurence[1*], Robert Kunkel[1], Anju R. Babu[1], Katherine Kramer[1], Ryan Baumwart[2], Rheal Towner[3], Yi Wu[1], and Chung-Hao Lee[1]

[1]School of Aerospace and Mechanical Engineering
The University of Oklahoma
865 Asp Ave., Felgar Hall Rm. 219C
Norman, OK 73019, USA

[2]Center for Veterinary Health Sciences
Oklahoma State University
208 S. McFarland Street
Stillwater, OK 74078, USA

[3]Advanced Magnetic Resonance Center, MS 60
Oklahoma Medical Research Foundation
825 N.E. 13th Street
Oklahoma City, OK 73104, USA

[*]Equal 1st-Authored Contribution

Submitted to the Journal of Mechanical Behaviors of Biomedical Materials

May 5, 2018

For correspondence:
Chung-Hao Lee, Ph.D.
Assistant Professor
School of Aerospace and Mechanical Engineering
Institute for Biomedical Engineering, Science, and Technology
The University of Oklahoma
865 Asp Ave., Felgar Hall Rm. 219C
Norman OK 73019-3609
email: ch.lee@ou.edu
Tel: 405-325-4842

Abstract

Valvular heart diseases are complex disorders, varying in pathophysiological mechanism and affected valve components. Understanding the effects of these diseases on valve functionality requires a thorough characterization of the mechanics and structure of the healthy heart valves. In this study, we performed biaxial mechanical experiments with extensive testing protocols to examine the mechanical behaviors of the mitral valve and tricuspid valve leaflets. We also investigated the effect of loading rate, testing temperatures, species (porcine versus ovine hearts), and age (juvenile vs adult ovine hearts) on the mechanical responses of the leaflet tissues. In addition, we evaluated the structure of chordae tendineae within each valve and performed histological analysis on each atrioventricular leaflet. We found all tissues displayed a characteristic nonlinear anisotropic mechanical response, with radial stretches on average 30.7% higher than circumferential stretches under equibiaxial physiological loading. Tissue mechanical responses showed consistent mechanical stiffening in response to increased loading rate and minor temperature dependence in all five atrioventricular heart valve leaflets. Moreover, our anatomical study revealed similar chordae quantities in the porcine mitral (30.5 ± 1.43 chords) and tricuspid valves (35.3 ± 2.45 chords), but significantly more chordae in the porcine than the ovine valves ($p<0.010$). Our histological analyses quantified the relative thicknesses of the four distinct morphological layers in each leaflet. This study provides a comprehensive database of the mechanics and structure of the atrioventricular valves, which will be beneficial to development of subject-specific atrioventricular valve constitutive models and toward multi-scale biomechanical investigations of heart valve function to improve valvular disease treatments.

Keywords: biaxial mechanical testing; the mitral and tricuspid valves; soft tissue biomechanics and microstructure; morphological analysis

1. Introduction

Efficient heart function relies on the proper mechanical behavior of the mitral and tricuspid valves. Aptly classified as the atrioventricular valves, the mitral and tricuspid valves are located between the atrium and the ventricle of the left heart and the right heart, respectively. These functionally-similar and morphologically-distinct valves are each enclosed by ring-like structures known as the valvular annuli and regulate blood flow between the heart chambers through the motion of membranous tissue known as leaflets. Two anatomically distinct leaflets comprise the mitral valve (MV), namely the anterior leaflet (MVAL) and posterior leaflet (MVPL), whereas the tricuspid valve (TV) is made up of three leaflets: the anterior leaflet (TVAL), posterior leaflet (TVPL), and septal leaflet (TVSL) (**Fig. 1a**). During systole of the cardiac cycle, these leaflets are closed by pressure gradients to enforce the unidirectional blood flow from the ventricles to the arteries. The leaflets are mechanically reinforced by a network of structurally robust tendons known as chordae tendineae. These chordae tendineae connect the valve leaflets to the papillary muscles of the ventricular wall to provide necessary supporting forces for the atrioventricular leaflets during cardiac function [1].

In healthy individuals, blood circulates through the heart with minimal backflow (or regurgitation) via proper opening and closing of the atrioventricular valves. However, valvular heart diseases, such as valve leaflet stenosis, valve prolapse, chordae rupture, leaflet damage and/or congenital defects, can affect the overall function of the valve and eventually lead to significant regurgitation [2-5]. The regurgitation increases the strain on the heart and is a corollary and precedent to the development of other, more threatening, heart conditions. Various treatments have been developed for valve regurgitation, including traditional open-heart surgical repair via an annuloplasty ring, surgical valve replacement with a bio-prosthetic or mechanical valve, transcatheter-based valve replacement, or the novel transcatheter deployable mitral clip device [6-12]. Currently, numerous imaging techniques, such as cardiac magnetic resonance imaging (MRI), cardiac computed tomography (CT) and echocardiographic imaging, are employed to examine the degree of heart valve regurgitation and to select the proper treatment for regurgitated valves [13,14]. However, these clinical imaging modalities may be inadequate to characterize the severity of the valve regurgitation, especially to accurately determine relevant parameters, such as the volume of blood backflow and the orifice area [15]. Recently, significant efforts have been focused on the development of predictive computational modeling tools to provide objective recommendation for cardiologists and cardiac surgeons in the diagnosis of valvular regurgitation and in the improvement of patient-specific treatment. For example, high-fidelity models have been employed toward precision medicine by incorporating patient-specific

valve geometries and essential microstructure information to simulate healthy and diseased heart valves [16-19]. These computational models can then be applied to predictions of various potential patient-specific treatment scenarios to examine the stresses, hemodynamics, and valve functional mechanics for the purpose of determining the optimal treatment solution [20-23].

In order to validate and accurately utilize these computational models in treatment selection, characterization of mechanical and anatomical data of heart valve leaflets is essential [24,25]. In soft tissue biomechanics, planar biaxial mechanical testing employing multiple loading ratios has been utilized to investigate the anisotropic material response of membrane-like connective tissue [26-31]. The observed anisotropic and nonlinear material response of many soft tissues, including heart valve leaflets, stems from microstructural collagen and elastin fiber networks with preferred orientations [32-35]. May-Newman et al. was the first to establish biaxial mechanical testing procedures for heart valve leaflets in her characterization of the material properties of the porcine MVAL and MVPL [36]. Further biaxial testing examined the mechanical dependence of the MVAL on the loading rate [37,38]. A similar study was conducted to examine stress relaxation of the tissue mechanical response, and minor hysteresis was found in the MVAL independent of the loading rate applied [28]. More recently, biaxial mechanical evaluation was used to characterize the leaflet-specific anisotropic nonlinear response of porcine tricuspid valve leaflets [35].

Despite the prior studies on heart valve leaflet biomechanical responses, biaxial testing results can vary significantly based on specific experimental procedures [39]. Therefore, it would be useful to compare leaflet material properties from both mitral and tricuspid valves tested from the same heart under a unified mechanical quantification procedure. Moreover, the relationship between mechanics of the mitral and tricuspid valves is important toward understanding the link between mitral valve surgical intervention and the development of functional tricuspid regurgitation (FTR) which has been observed in clinical studies [40-42].

Hence, the goal of this work is to utilize biaxial testing experiments to compare the unique material response of each mitral and tricuspid valve leaflet. Porcine atrioventricular leaflets were tested at various temperature levels and loading rates to characterize the dependence of tissue response on these selected experimental parameters. In addition, the stretch responses of leaflets from juvenile ovine and adult ovine hearts were characterized and compared to understand the effects of species and animal age on testing results. Lastly, an anatomical study was conducted to examine chordae distributions in porcine and ovine valves, and histological methods were employed to study and examine the layered microstructure of the porcine atrioventricular valve leaflets.

2. Methods

2.1 Tissue acquisition

To characterize the material properties of porcine atrioventricular heart valve leaflets, porcine heart tissues from physically healthy pigs (80-140 kg, 1-1.5 years of age) were acquired from a local USDA approved abattoir (Country Home Meat Company, Edmond, OK). To further compare the leaflet mechanical behaviors between different species and ages, we acquired adult ovine hearts (65-90 kg, 2-5 years of age) and juvenile ovine hearts (35-60 kg, 6-12 months of age) from Chickasha Meat Company (Chickasha, OK). The fresh heart tissues were frozen within 12 hours post-mortem in a standard freezer at -14°C for storage purposes. This tissue storage procedure was based on the previous studies which found that freezing has a minimal impact on the mechanics of collagenous tissues [43-45]. In preparation for testing, the hearts were thawed and leaflets from both the mitral and tricuspid valves were dissected (**Fig. 1a**). The excised leaflets were preserved in phosphate buffered saline (PBS) and refrigerated at 4°C to maintain material properties until testing [46].

2.2 Tissue preparation

For biaxial mechanical testing, a square specimen (8mm x 8mm) was dissected from the central region (called the clear zone) of each valve leaflet. The thickness of the specimen was measured at 3 different locations using digital calipers (WestWard Company, Lake Forest, IL) to determine an average tissue thickness. The tissue specimen was then mounted on a commercial biaxial mechanical testing system (**Fig. 2a**)—BioTester (CellScale, Waterloo, ON, Canada) with a load cell capacity of 1,500 mN, by using 4 BioRakes to create a 6.5mm x 6.5mm effective testing region (**Fig. 2b**). The primary axes of the testing system were aligned with the circumferential and radial directions of the tissue (**Fig.1b**). Four glass beads (with diameters of 300-500 μm) were attached to the specimen in a square configuration (**Fig. 2c**) for measuring the in-plane deformation and strain using a non-contact image-based technique as described in **Section 2.4**. To ensure the tissue remained hydrated, the specimen was submerged in a heated bath of PBS solution for the duration of each biaxial test.

2.3 Biaxial mechanical testing protocols

Comprehensive characterization of the mechanical behaviors of both atrioventricular valve leaflet tissues was then conducted, which included investigations of the material anisotropy, loading rate effect, temperature effect, and the differences within and across species. In brief, we performed force-controlled biaxial mechanical testing on four experimental groups: (i) the baseline testing group (porcine tissues) with a loading rate of 4.42 N/min at room temperature (22°C), (ii)

the loading rate effect group (porcine tissues) with three loading rates of 2.29 N/min, 4.42 N/min, and 7.92 N/m at room temperature (22°C), (iii) the temperature effect group (porcine tissues) with a loading rate of 4.42 N/min at three temperature levels (27°C, 32°C and 37°C), and (iv) the species and age group (porcine, adult ovine, and juvenile ovine tissues) with a loading rate of 4.42 N/min at the physiological temperature (37°C). The sample size for each of these groups was n=6 for each mitral and tricuspid valve leaflet, i.e., MVAL, MVPL, TVAL, TVPL, and TVSL.

For each biaxial mechanical test, the physiological stress levels of the MV and TV leaflets were estimated to be 240 kPa and 115 kPa, respectively, based on the Laplace's law for a spherical surface assuming the mean radius of curvature of the coapted MV and TV to be 2 cm [35] and the transvalvular pressure gradients to be 100 mmHg [47] and 40 mmHg [35], respectively. Then, the applied membrane tensions in both the circumferential and radial directions, i.e., $T_{C,\max}$ and $T_{R,\max}$, were calculated based on the specimen size and the measured thickness of the leaflet tissue.

Each test began with a preconditioning protocol, during which estimated membrane tensions $T_{C,\max}$ and $T_{R,\max}$ were applied for 8 repeated loading-unloading cycles (**Fig. 2d**), considering a preload of 1% of $T_{C,\max}$ and $T_{R,\max}$, to restore the dissected tissue to its respective *in vivo* functional configuration [48,49]. Then, biaxial testing protocols with various loading ratios ($T_{C,\max}$:$T_{R,\max}$ = 1:1, 0.75:1, 1:0.75, 0.5:1, and 1:0.5) were conducted (**Fig. 2d**). In addition to the monitored forces and Biorake separation distance in both directions, a series of 1280x960 images were collected by a high-resolution CCD camera (The Imaging Source LLC, Charlotte, NC) at a rate of 15 Hz. These images were then used in the non-contact in-plane strain calculations as described in the next section.

2.4 Tissue strain and stress calculations

Digital image correlation (DIC) based techniques have been widely utilized in the biomechanics society to track the deformations of a tissue specimen. To avoid the Saint-Venant effects on tissue deformations during our biaxial mechanical testing [50], four fiducial markers (glass beads) were placed in the central delimited region (3mm x 3mm) of the valve leaflet specimen (**Fig. 2c**). A series of images of the tissue specimen were collected by the high-resolution CCD camera and the time-dependent positions of the four fiducial markers were analyzed based on the acquired images using the DIC-based technique in the LabJoy software of the BioTester system:

$$\mathbf{x}_I = \mathbf{X}_I + \mathbf{d}_I\,,\ I = 1 \sim 4 \qquad (2),$$

where $\mathbf{X}_I$'s and $\mathbf{x}_I$'s are the marker positions at the undeformed (reference) configuration (Ω_0) and at the deformed configuration (Ω_t), respectively, and $\mathbf{d}_I$'s are the displacement vectors of the fiducial markers, i.e., $\mathbf{d}_I = [u_I(t), v_I(t)]^T$.

To compute the in-plane strain of the tissue specimen, a four-node bilinear finite element was used based on the 4 markers, and the deformation gradient tensor **F** was determined using an in-house MATLAB program (R2016a, The MathWorks, Natick, MA) based on the strain-calculation technique developed previously [31,33]:

$$\mathbf{F} = \mathbf{F}(\mathbf{X},t) = \frac{\partial \mathbf{x}}{\partial \mathbf{X}} = \begin{bmatrix} \sum_{I=1}^{4} B_{xI} u_I(t) & \sum_{I=1}^{4} B_{yI} u_I(t) \\ \sum_{I=1}^{4} B_{xI} v_I(t) & \sum_{I=1}^{4} B_{yI} v_I(t) \end{bmatrix} \quad (3),$$

where B_{xI}'s and B_{yI}'s are the shape function derivatives associated with node I with respect to the x and y coordinates, respectively. Note the x-y coordinates were aligned with the tissue's circumferential and radial directions, respectively (**Figs. 1b & 2b**). The right Cauchy-Green deformation tensor **C** and the Green strain tensor **E** can then be computed by

$$\mathbf{C} = \mathbf{F}^T\mathbf{F}, \text{ and } \mathbf{E} = \frac{1}{2}(\mathbf{C} - \mathbf{I}) \quad (4),$$

where **I** is the 2nd-order identity tensor. The circumferential and radial stretches, λ_C and λ_R, were determined by the square roots of the principal values of **C**. Next, the first Piola-Kirchhoff (1st-PK) stress tensor **P** was computed from the applied membrane tensions, T_C and T_R, as follows:

$$\mathbf{P} = \frac{1}{L \cdot t}\begin{bmatrix} T_C & 0 \\ 0 & T_R \end{bmatrix} \quad (5).$$

Here, L is the valve leaflet specimen edge length, and t is the tissue thickness.

2.5 Quantification of anatomical structure of valve apparatus

To complement the mechanical response data from biaxial mechanical testing, we further investigated the anatomical and structural features of both the porcine and ovine atrioventricular heart valves (n=6). Briefly, we measured the number of chordae and each chord's length and classified the chordae according to their respective supporting leaflet. Adopting the chordae classification convention by Toma et al. [51], tertiary chordae were not measured, and no distinction was made between primary and secondary chordae in our measurements. The length of the chord was measured from proximal attachment at the papillary muscles to distal attachment at the inferior surface of the leaflets. We also measured each leaflet's thickness in 3 separate locations using digital calipers with a resolution of 0.01 mm.

2.6 Histological analysis

To examine the microstructural organization of the extracellular matrix (ECM) components in the valve leaflets, specimens from all five atrioventricular valve leaflets were fixed in 10% formalin at room temperature. These tissue specimens were then dehydrated in graded solutions of alcohol and embedded in paraffin. Sections of 5-7 μm thickness were sectioned and stained with Masson's trichrome stain. For the morphological characterization, all stained sections were examined with a halogen illumination microscope (AmScope, Irvine, CA) at a magnification of 2X. Images of leaflets were captured using a 10 Mega Pixel camera and analyzed with ImageJ software (National Institute of Health, Bethesda, MD). A single image of each valve leaflet was captured from the stained sections and thickness was measured at three random locations in the collected image. Further, the color deconvolution and image threshold plugins in ImageJ [52] were used to quantify the collagen content in each leaflet specimen.

2.7 Statistical analysis

To focus our statistical analyses of biaxial mechanical testing results on relevant relationships, we chose to analyze only the circumferential and radial stretch values at the peak stress from the equibiaxial tension protocol. These circumferential and radial peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) were analyzed using a one-way Analysis of Variance (ANOVA). This method was also utilized to examine differences between mechanical responses of valve leaflets (MVAL, MVPL, TVAL, TVPL, and TVSL) from our baseline testing group.

Previous studies have demonstrated a difference between the stretch responses due to the preconditioning effect and the applied loading to the tissue [53]. Thus, we decomposed the *peak stretches* (λ_C^{0-peak} and λ_R^{0-peak}) into two components: (i) the *preconditioning stretches* (λ_C^{0-1} and λ_R^{0-1}), defined as the tissue stretch from the mounting configuration to the post-preconditioning state, and (ii) the *mechanical stretches* (λ_C^{1-peak} and λ_R^{1-peak}), defined as the tissue stretch from the post-preconditioning configuration to the peak loading state. We compared these stretch measures using the ANOVA approach described previously to isolate the effects of the loading rate, temperature, and species on the atrioventricular valve leaflet's mechanical response.

In the anatomical study, leaflet thicknesses, chordae lengths, and chordae quantities were all compared using the standard one-way ANOVA method.

All statistical analyses were performed with MATLAB, considering a p-value < 0.05 as statistically significant and a p-value < 0.10 as nearly statistically significant.

3. Results

3.1 Baseline testing

Results for the MV and TV leaflets for loading protocols ($T_{C,\max}$:$T_{R,\max}$ =1:1, 0.5:1, and 1:0.5) are presented in **Figures 3-4**, respectively. **Tables 1-3** provide the mean ± standard error of the mean (SEM) of the stretches for the MV and TV leaflets at specified stress values for loading protocols $T_{C,\max}$:$T_{R,\max}$ = 1:1, 0.5:1, 1:0.5, respectively. Similarly, we report the results associated with the intermediate loading protocols ($T_{C,\max}$:$T_{R,\max}$ = 0.75:1 and 1:0.75) in the **Appendix** (**Figs. A1-A2**). Datasets of each of the atrioventricular heart valve leaflets under all 5 biaxial loading protocols are provided in an accompanying article in the *Data in Brief* Journal [54].

Through the statistical analysis, we found both the MV and TV leaflets exhibited anisotropic material behavior with the peak stretches smaller in the circumferential direction than the radial direction ($p < 0.012$ for all leaflets, **Figs. 3a & 4a**). Our results also revealed statistically significant differences in the mechanical responses of the atrioventricular heart valve leaflets. Specifically, the MVPL and TVAL were stiffer in the circumferential direction than the TVSL (p=0.050), and the TVPL was more compliant in the radial direction than the MVAL (p=0.024).

3.2 Loading rate effect

We present representative results of our loading rate effect group from the equibiaxial loading protocol ($T_{C,\max}$:$T_{R,\max}$ = 1:1) for all mitral and tricuspid valve leaflets in **Fig. 5**. Representative results from the same group under loading protocols $T_{C,\max}$:$T_{R,\max}$ = 0.5:1 and 1:0.5 are reported in the **Appendix** (**Figs. A3**-**A4**). Statistical analysis results of the circumferential and radial stretches of the MVAL are presented in **Figure 6** and **Table 4**. The MVAL was selected as a representative leaflet owing to its similarity to the other leaflets' trends. We have included the statistical analysis results for all other leaflets (MVPL, TVAL, TVPL, and TVSL) in the **Supplementary Material** (**Figs. S1-S4** and **Tables S1-S4**).

Three observations could be drawn from the statistical analysis results: (1) the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}) increase as the loading rate increases in both the circumferential and radial directions (**Fig. 6a**); (2) the mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}) decrease as the loading rate increases in both directions (**Fig. 6b**); (3) as the loading rate increases, the peak stretch in the circumferential direction decreases while the peak stretch (λ_C^{0-peak} and λ_R^{0-peak}) in the radial direction increases (**Fig. 6c**). These observed trends were not statistically significant but were generally consistent across MV and TV leaflets (**Figs. S1-S4** and **Tables S1-S4**).

3.3 Temperature effect

To show the mechanical response of the leaflet to varied temperature levels, we presented representative results under equibiaxial tension for all the mitral and tricuspid valve leaflets in **Figure 7**. Similarly, the representative results associated with other loading protocols ($T_{C,\max}$:$T_{R,\max}$ = 0.5:1 and 1:0.5) are provided in the **Appendix** (**Figs. A5**-**A6**). We found from our statistical analyses that the MVAL is typical of other atrioventricular leaflets (**Fig. 8 and Table 5**), whereas the statistical analysis results of the MVPL, TVAL, TVPL, and TVSL are provided in the **Supplementary Material** (**Figs. S5-S8** and **Tables S5-S8**).

Three observations could be drawn from our statistical analysis results associated with the temperature effect group: (1) an increase in temperature led to the decrease in the circumferential preconditioning stretch (λ_C^{0-1}) but an increase in the radial preconditioning stretch (λ_R^{0-1}) (**Fig. 8a**); (2) the mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}) had no detectable or statistically significant correlation to the increase in temperature for both the circumferential and radial directions (**Table 5**); (3) through comparison of the peak stretches in the MVAL (**Fig. 8c**), we found that an increase in the temperature corresponded to a decrease in the peak circumferential stretch (λ_C^{0-peak}) but an increase in the peak radial stretch (λ_R^{0-peak}). None of these observed trends were found to be statistically significant. Nevertheless, these trends were generally consistent across all the MV and TV leaflets despite the presence of some outliers, specifically for the mechanical responses of the MVPL (**Fig. S5**).

3.4 Comparison of mechanical responses between different species

In the presentation of results from the species effect experimental group, one representative leaflet was examined from each atrioventricular heart valve. Specifically, we chose the MVAL to represent the mitral valve and the TVAL to represent the tricuspid valve. The averaged mechanical responses (n=6) for these two leaflets under equibiaxial loading from porcine, adult ovine, and juvenile ovine hearts are presented in **Figures 9a-c**, respectively. We further present the statistical comparisons of the peak, preconditioning, and mechanical stretches (as defined in **Section 2.7**) between these species in the **Supplementary Material** (**Tables S9-10**).

The cross-species (porcine versus ovine) and within-species (juvenile ovine versus adult ovine) observed behaviors were specific to MVAL or TVAL and did not support conclusions across leaflets. Through interpreting the statistical analysis results, we found three leaflet-specific relations: (1) the preconditioning stretches were lower for the porcine TVAL than the adult ovine TVAL in both circumferential (p=0.028) and radial directions (p=0.057) (**Table S10**); (2) the mechanical stretches were notably lower for the juvenile ovine MVAL than the adult ovine MVAL

in the radial direction (p=0.052) but showed no consistent trends in the circumferential direction (p=0.614) (**Table S9**); (3) In the MVAL, the peak stretch was lower for the porcine tissue than the juvenile ovine tissue in the circumferential direction but higher in the radial direction. These results also showed the more isotropic behavior of the juvenile ovine compared to the porcine tissue (**Table S9**).

3.5 Anatomical quantification of the valve apparatus

The measurements of the anatomical and structural features of porcine and adult ovine atrioventricular valves are presented in **Table 6**. We found the porcine MVAL was significantly thicker than each porcine TV leaflet ($p<0.040$), and the porcine MVPL was thicker than the porcine TVPL and TVSL ($p<0.006$). We also observed the similarity in the length of the chordae anchoring porcine and ovine valve leaflets, as reflected in the MVAL where chordae anchoring porcine leaflets had a length 17.5 ± 1.32 mm while those anchoring the ovine leaflet had a comparable length of 17.9 ± 1.12 mm (**Table 6**). However, we observed the porcine MV and TV had more chordae than the ovine MV and TV ($p<0.001$ and $p=0.002$, respectively). These similarities in chordae lengths and significant differences in chordae quantities are explored further in **Section 4.2.2**.

3.6 Histological analysis

The Masson's trichrome-stained MV and TV leaflets from a representative porcine heart are shown in **Figure 10**. The average thickness obtained from the histology images of each leaflet are presented in **Table 7**. The MV leaflets were distinctly thicker than the TV leaflets to sustain higher transvalvular pressure loading [55]. The characteristic four layers of the leaflets: ventricularis (V), spongiosa (S), fibrosa (F), and atrialis (A) were histologically distinguishable in all the five leaflets. The fibrosa layer is primarily composed of a dense layer of collagen and comprises ~60% of the total thickness of the MV leaflets. Similarly, the histology image suggested the fibrosa is the dominant layer of the TVAL. Interestingly, we found the spongiosa, made of non-fibrous constituents (such as Glycosaminoglycans, or GAGs) and located between the fibrosa and atrialis layers, was the thickest layer in the TVPL and TVSL. The ventricularis layer, facing the ventricle in each atrioventricular leaflet, is twice as thick in the MVAL as the TVAL, whereas it is seven-times thicker in the MVPL than in the TVPL. Histology image-based collagen quantification of the valve leaflets indicated the MVAL and MVPL have 77.7%, and 69.1% collagen fiber contents, respectively, whereas the TVAL and TVPL have 68.5%, 45.2% collagen fiber content, respectively (**Table 7**). Interestingly, we also found the collagen fiber content is lower in the TVSL (~32%) than in other leaflets.

4. Discussion

The anisotropic and nonlinear mechanical responses of mitral and aortic valve leaflet tissues have been characterized in the past two decades [30,31,36,56-58]. However, relatively less attention has been paid to the tricuspid heart valve leaflets [35]. Therefore, this work represents a logical first step to a thorough understanding of the mechanics and structure of the tricuspid valve leaflets, as well as the differences of their mechanical behaviors compared to their relatively well-understood mitral valve counterparts.

4.1 Overall findings

4.1.1 Baseline force-controlled testing

Our baseline testing confirmed the previously quantified anisotropic, nonlinear elastic mechanical response of heart valve leaflets (**Figs. 3-4**), through the presence of a toe portion of the curve (near the low-stress region) and a highly stiff, nearly asymptotic region as the stress approached the maximum physiological stress experienced by the leaflets. All leaflets tested were significantly stiffer in the circumferential direction ($p<0.010$). Specifically, the average stretches in the circumferential and radial directions in our study differed by 0.360 for the MVAL, 0.389 for the MVPL, 0.441 for the TVAL, 0.444 for the TVPL, and 0.284 for the TVSL (**Table 1**). In general, the TV leaflets were more compliant than the MV leaflets at their respective physiological loadings in both circumferential radial directions.

In a prior study, planar biaxial testing was performed on porcine mitral valve tissue and stretches of 1.2 and 1.4 were found for the MVAL in the circumferential and radial directions under an equibiaxial loading protocol [37], whereas stretches of 1.232 ± 0.154 and 1.592 ± 0.136 were found in our study for the MVAL in the same directions (**Table 1**). Recent studies have shown that testing parameters, such as the mounting mechanism, the spacing of attachment points and the specimen size, could have an impact on the biaxial mechanical testing results [39]. Such discrepancy between our results and those of Grashow et al. (2006), particularly in the radial direction, may be attributed to procedural differences such as different tissue mounting mechanisms (rigid BioRake fixture in our study versus suture hooks in the study by Grashow et al. (2006)).

Our observed stretches of the TV leaflets were similar to those reported by Khoiy and Amini (2016) [35]. In their study, the TV posterior leaflet showed the most anisotropy, with the TVAL and TVSL exhibiting similar but less anisotropy than the TVPL. In general, our study's results agreed with these findings for the tricuspid valve leaflets. Khoiy and Amini (2016) also found the TVPL was the most compliant leaflet in the radial direction. Our study affirmed this finding, with our

observed peak radial stretches of 1.651 ± 0.089 for the TVAL, 1.788 ± 0.040 for the TVPL, 1.685 ± 0.089 for the TVSL (**Table 1**). Importantly, both studies found the same highly anisotropic and nonlinear behaviors of the tricuspid valve leaflets.

4.1.2 Effect of various loading rates on the mechanical response

We further conducted mechanical testing (**Fig. 5, A3-A4**) and statistical analysis (**Fig. 6, S1-S4**) on the response of the atrioventricular leaflets considering varied loading rates. An interesting result of this testing group is the decrease in mechanical stretch of the tissue in both tissue directions with an increased loading rate (**Fig. 6b**). Our results support the accepted viscoelastic nature of heart valve leaflets [59] and agrees with the time-dependent stress relaxation as observed in heart valve leaflets [38]. However, our results prove contrary to those of a prior study by Grashow et al. (2006), where a variety of loading rates were examined and no significant dependence of mechanical properties on loading rate was observed [37]. One possible reason for this discrepancy is Grashow et al. (2006) examined the peak stretch of the tissue, while we examined the decomposed components of the peak stretch. As for the MVAL, we observed minor differences in the peak stretch between the loading rates of 2.29 N/min and 7.92 N/min in the circumferential ($p=0.844$) and radial directions ($p=0.614$) (**Fig. 6c** and **Table S1**). Based on these peak stretch comparisons, it would be reasonable to draw a conclusion that there was no significant viscoelastic effect. However, by decomposing the peak stretch and examining only the tissue's mechanical stretch, the statistical comparisons showed the tissue is stiffer under the loading rate of 7.92 N/min than the rate of 2.29 N/min in both the circumferential ($p=0.334$) and radial directions ($p=0.077$) (**Table S1**). These comparisons of mechanical stretches could shed light on the tissue's response to varied loading rates, as decoupled from the preconditioning effect.

4.1.3 Effect of temperature on the mechanical responses

We also performed mechanical testing (**Figs. 7 and A4-A5**) and statistical analysis (**Figs. 8** and **S5-S8**) on each atrioventricular leaflet to examine the temperature dependence of tissue mechanical properties. In examination of these results, we found distinct trends between the circumferential and radial directions (**Figs. 8** and **S5-S8**). Specifically, we found that the preconditioning stretches (**Fig. 8b**) and peak stretches (**Fig. 8c**) decreased in the circumferential direction but increased in the radial direction with increased temperature. We also observed a general lack of temperature dependence in the mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}) of the atrioventricular heart valve leaflet tissues (**Fig. 8c**) which clarified how the temperature dependence observed in the peak stretches was primarily a result of similar temperature effects

in the preconditioning stretches. These findings are potentially useful for informing experimental protocols where maintenance of valve tissue at body temperature would not be feasible. In such experiments, we expect valve tissue's mechanical response would be unimpacted by the non-physiological temperature.

4.2 Comparisons between porcine and ovine atrioventricular valves

Both porcine and ovine hearts are commonly employed in animal studies as analogous to human hearts [60]. A proper comparison between human and ovine or porcine hearts first requires an intimate understanding of the mechanics and anatomy of the non-sapiens heart. To that end, we compared the biaxial mechanical responses of porcine, adult ovine, and juvenile ovine atrioventricular leaflet tissues (**Fig. 9**) and examined specific anatomical features of the atrioventricular valves from both porcine and ovine hearts (**Table 6**).

4.2.1 Mechanical responses

We examined the responses of the MVAL and TVAL of the porcine, adult ovine, and juvenile ovine heart (**Figs. 9a-c**) to show the differences in tissue response between different species and between the juvenile and adult ovine animals.

In this study group, we found that the preconditioning stretches were lower for the porcine heart valve than the ovine heart valves, and that the mechanical stretches of the porcine heart were generally higher than the mechanical stretches of the ovine hearts, particularly in the TVAL (**Figs. 9a-c**). We also observed that the peak stretches for the adult ovine TVAL were higher than the peak stretches for the porcine TVAL in both the circumferential ($p=0.090$) and radial ($p=0.180$) directions. These differences between the porcine and ovine atrioventricular valves imply the necessity to consider the mechanics of the specific analogous heart valve for computational modeling rather than assuming similar properties based on their similar function.

With regards to within-species examinations, we found the juvenile ovine MVAL behaved in a more isotropic manner as compared to the adult ovine heart, whereas the TVAL showed little stretch response differences between the adult and juvenile ovine hearts (**Figs. 9b-c** and **Table S9**). In addition, we also observed some discrepancy between the juvenile and adult ovine MVAL mechanical responses and the similarity between the juvenile and adult ovine TVAL mechanical responses. These findings suggest possible differences in the development rates between the mitral and tricuspid valves, and could motivate further examinations of heart valve growth and remodeling over the pubescent maturation period [61].

458 **4.2.2 Anatomy and structure**

459 Our anatomical comparison allowed a glimpse into the differences and similarities between
460 valvular structures of different species (**Table 6**). We found substantial differences in leaflet
461 thickness between the porcine and ovine valve, with the recorded porcine MVAL thickness of 0.79
462 ± 0.10 mm and the recorded ovine MVAL thickness of 0.40 ± 0.03 mm, but similar average
463 chordae lengths between the porcine and ovine valves (**Table 6**). We interpret these leaflet
464 thicknesses as proportional to the valvular capacity to bear pressure load. As such, it is intriguing
465 that the mitral valves exhibit such similar chordae lengths, despite the apparent difference in load-
466 bearing capacity/necessity [55]. In addition, we found the chordae quantity generally correlates
467 with the leaflet thickness. For example, the porcine TVAL used 11.2 ± 1.30 anchoring chordae to
468 support a leaflet of thickness 0.52 ± 0.06 mm, while the ovine TVAL had only 6.8 ± 0.50 anchoring
469 chordae to support a leaflet of thickness 0.28 ± 0.05 mm. This relationship could provide insight
470 into the growth and development process of chordae tendineae in the atrioventricular valves.
471 To examine the chordae quantities within human heart valves, Lam et al. (1970) and Silver et
472 al. (1971) conducted anatomical studies on the human mitral and tricuspid valves, respectively
473 [62,63]. They found human hearts have, on average, 25 chordae in both mitral and tricuspid
474 valves. Our study revealed average chordae quantities of 30.5 ± 1.7 and 35.30 ± 2.8 chords for
475 porcine mitral and tricuspid valves, respectively, and 14.9 ± 1.3 and 23.7 ± 1.9 chords for the adult
476 ovine mitral and tricuspid valves, respectively. These differences in chordae quantities between
477 species suggest unique valvular density and distribution of chordae tendineae within human,
478 porcine, and ovine atrioventricular valves. The error presented in the chordae quantities also
479 alludes to the inherent differences in physiological structures, and the inability to determine a
480 precise expected value for the number of chordae within a healthy valve.

481 **4.3 Histological results**

482 Valve leaflets are composed of collagen, elastin, GAGs, and proteoglycans (PGs). The
483 corresponding nonlinear anisotropic mechanical response of the valve leaflets is mainly
484 determined by these constituents in the ECM. Studying the microstructural organization of leaflets
485 is crucial to understanding the physiological functions of the leaflets. In this study, we investigated
486 the difference in the thickness of individual leaflet layers as well as the collagen fiber distribution
487 in the MV and TV leaflets based on quantitative histology analyses. The intact layer thickness
488 measured from the histological images was in the same range as the thickness obtained from
489 anatomical measurements (**Section 3.5**). The histological measurements allowed us to precisely
490 quantify the discrepancies in the thickness of intact and individual layers of the MV and TV leaflets
491 (**Table 7)**. The measurements also showed the MV leaflets have thicker collagen-rich fibrosa and

ventricularis layers compared to TV leaflets (**Fig. 10**). The atrialis layer, which is facing the atrium in each leaflet, occupies 6.42 ± 1.36% of the total thickness of the MVPL and 11.32 ± 2.31% of the total thickness of the TVPL (**Table 7**). Although both atrioventricular heart valves are located in between the atrium and the ventricle, their respective leaflets are subjected to distinct hemodynamic loading conditions and transvalvular pressure gradients. Such functional discrepancies lead to the difference in the microstructural organization of the MV and TV leaflets as observed in this study. In addition, the high collagen fiber content and the collagen-rich fibrosa layer in the MV leaflets may serve as a primary load bearing layer to sustain higher pressures during cardiac cycles compared to the TV counterparts.

4.4 Study limitations and future extensions

Biaxial testing methods are notoriously limited in their ability to capture and account for in-plane shear stresses [29]. In our study, planar shear stresses were examined and found to be negligible compared to their direct stress counterparts, and so were not included in the presentation of results. This is standard practice in biaxial testing experiments, and previous studies utilizing biaxial testing methods have found similarly negligible shear stresses [64]. In addition, we encountered difficulty in the determination of an average sample thickness due to the roughness and non-uniformity of heart valve leaflets, particularly in regions with chordae attachments [65]. To address this obstacle, our group averaged thickness data from three measurements taken from different regions of the testing sample. Another potential limitation of our study was in the use of frozen heart valve tissues. Although this is a standard practice in biaxial mechanical testing experiments [37,38], the effect of freezing on the mechanics of heart valve tissues has not been rigorously studied. In addition to these experimental limitations, the natural variance in biological material property and structure led to the substantial calculated errors in biaxial testing and anatomical study results. This challenge persists across many biological domains and leads to larger errors in experimental results compared to other fields.

The essential extension from this study is the development of a structurally-informed and computationally-tractable constitutive model for the atrioventricular valve leaflets, which is currently under investigation by our lab. In the meantime, we make our experimental results associated with all the five loading protocols available in the accompanying *Data in Brief* article [54], for other researcher groups' development and validations of novel computational models for atrioventricular heart valve tissues.

With regards to the anatomical study, we only characterized a few dimensional parameters of the valvular complex. Further research on the annuli, moderator band, papillary muscles, and

ventricular dimensions could allow potential early diagnosis of valvular regurgitation through the establishment of a healthy range of parameter values, and it could also be beneficial to the development of predictive computational models towards increasing the long-term durability of surgically repaired atrioventricular heart valves [25].

Another ongoing research effort from our lab is the investigation of the important relation between the tissue microstructure and its mechanical behavior. For example, the relatively thick and collagen-rich fibrosa layer (**Fig. 10**) accounts for much of the mechanical strength of the heart valves under physiological tensile loading [58,66]. Although the collagen fiber orientation in the fibrosa has been examined, very few studies have comprehensively related the characterized tissue's mechanical responses to the dynamical changes in the collage fiber architecture (such as fiber orientation) under physiological and pathophysiological loading [67-69]. More rigorous studies on examining the interrelationship between the mechanics and collagenous structure of the leaflets will give a better insight into the atrioventricular heart valve function, especially under diseased and/or surgically-intervened conditions.

4.5 Conclusion

In this study, the response of each porcine atrioventricular leaflet has been thoroughly characterized. Our comprehensive results suggest the porcine MVAL and MVPL exhibit similar levels of material anisotropy, and the porcine TVPL was the most compliant in both circumferential and radial directions and the most anisotropic among the three tricuspid leaflets. These findings also suggest the necessity of employing different constitutive model parameters to describe the distinct mechanical behavior of each individual atrioventricular valve leaflet, rather than assuming homogeneous mechanical properties between leaflets. Moreover, we have demonstrated an increased loading rate is associated with a stiffer mechanical response, and increased temperature yields directionally-specific differences in the tissue stretches. We have also provided a novel quantification of the distinct mechanical responses of porcine and ovine leaflets and quantified the difference in leaflet mechanical response between different species and between juvenile and adult ovine animals. This study is also the first of this kind to examine the anatomic chordae distributions within porcine and ovine atrioventricular valves to quantify the structural differences between mitral and tricuspid valves. Future investigation may include the development of high-fidelity, multiscale models of atrioventricular valve function, and incorporation of the material responses acquired in this study into computational models [21,70,71]. These efforts will allow us to predict the response of the leaflet *in vivo* and to examine how the changes in the valve's mechanical properties and structure can lead to suboptimal valvular performance.

Appendix: Biaxial Testing Results of Porcine Valve Leaflet Tissues under Intermediate Loading Protocols

As for the baseline biaxial testing (**Section 3.1**), the mechanical responses of porcine mitral and tricuspid valve leaflets under loading protocols $T_{C,\max}$:$T_{R,\max}$ = 0.75:1 and $T_{C,\max}$:$T_{R,\max}$ = 1:0.75 were presented in **Fig. A1** and **Fig. A2**, respectively. Results under those intermediate loading protocols for the studies on the loading rate effect (**Section 3.2**) and the temperature effect (**Section 3.3**) were presented in **Fig. A3-4** and in **Fig. A5-6**, respectively.

Acknowledgments

Support from the American Heart Association Scientist Development Grant (SDG) Award (16SDG27760143) is gratefully acknowledged. CHL was in part supported by the institutional start-up funds from the School of Aerospace and Mechanical Engineering (AME) and the research funding through the Faculty Investment Program from the Research Council at the University of Oklahoma (OU). SJ, RK, and DL were supported by the Mentored Research Fellowship from the Office of Undergraduate Research at OU. KK and DL were supported by the Undergraduate Research Opportunities Program from the Honors College at OU. We also acknowledge undergraduate researchers Jacob Richardson and Colton Ross for their assistance with the anatomical study, as well as Dr. Kar-Ming Fung from the Department of Pathology at the OU Health Sciences Center for his help in the histology study.

Conflicts of Interest

None of the authors have a conflict of interest with the present work.

Nomenclature

category	abbreviation	definition
anatomy	MVAL	mitral valve anterior leaflet
	MVPL	mitral valve posterior leaflet
	TVAL	tricuspid valve anterior leaflet
	TVPL	tricuspid valve posterior leaflet
	TVSL	tricuspid valve septal leaflet
valve tissue layers	A	atrialis
	S	spongiosa
	F	fibrosa
	V	ventricularis
mechanics	**F**	deformation gradient tensor
	C	right Cauchy-Green deformation tensor = $\mathbf{F}^T\mathbf{F}$
	E	Green strain tensor = 0.5(**C**-**I**)
	I	2nd-order identity tensor
	σ	Cauchy stress tensor
	P	1st Piola-Kirchhoff (PK) stress tensor
	λ	Tissue stretch
	λ_C	Tissue stretch in the circumferential direction
	λ_R	Tissue stretch in the radial direction
	λ_C^{0-peak}	Peak stretch in the circumferential direction
	λ_R^{0-peak}	Peak stretch in the radial direction
	λ_C^{0-1}	Preconditioning stretch in the circumferential direction
	λ_R^{0-1}	Preconditioning stretch in the radial direction
	λ_C^{1-peak}	Mechanical stretch in the circumferential direction
	λ_R^{1-peak}	Mechanical stretch in the radial direction
tissue states	Ω_0	tissue sample at the unstressed reference configuration
	Ω_t	tissue sample at the deformed configuration under loading
collagen fiber directions	C	circumferential direction
	R	radial direction
others	SEM	standard error of the mean
	DIC	digital image correlation
	ECM	extracellular matrix
	GAG	glycosaminoglycan
	PG	proteoglycan

Tables

Table 1 – Circumferential and radial stretches of porcine atrioventricular heart valves (n=6) at selected stress levels (% of maximum 1st-PK stress) under equibiaxial (1:1) tension protocol (**Fig. 3a** and **4a**). Values are presented as mean ± SEM.

% of $P_{11,max}$	λ	MVAL	MVPL	TVAL	TVPL	TVSL
0%	λ_C^{0-peak}	1.061 ± 0.049	1.087 ± 0.032	1.089 ± 0.023	1.120 ± 0.041	1.168 ± 0.035
	λ_R^{0-peak}	1.358 ± 0.042	1.429 ± 0.069	1.434 ± 0.050	1.511 ± 0.035	1.476 ± 0.066
25%	λ_C^{0-peak}	1.207 ± 0.058	1.192 ± 0.031	1.172 ± 0.026	1.270 ± 0.058	1.331 ± 0.023
	λ_R^{0-peak}	1.551 ± 0.054	1.572 ± 0.081	1.569 ± 0.072	1.735 ± 0.037	1.641 ± 0.035
50%	λ_C^{0-peak}	1.219 ± 0.061	1.204 ± 0.032	1.194 ± 0.026	1.292 ± 0.059	1.355 ± 0.023
	λ_R^{0-peak}	1.572 ± 0.055	1.589 ± 0.082	1.624 ± 0.084	1.766 ± 0.040	1.664 ± 0.088
75%	λ_C^{0-peak}	1.225 ± 0.062	1.212 ± 0.034	1.204 ± 0.026	1.303 ± 0.060	1.367 ± 0.023
	λ_R^{0-peak}	1.583 ± 0.055	1.599 ± 0.082	1.640 ± 0.087	1.780 ± 0.040	1.676 ± 0.089
100%	λ_C^{0-peak}	1.231 ± 0.063	1.217 ± 0.034	1.210 ± 0.027	1.310 ± 0.060	1.374 ± 0.023
	λ_R^{0-peak}	1.591 ± 0.055	1.606 ± 0.082	1.651 ± 0.089	1.788 ± 0.040	1.685 ± 0.089

Table 2 – Circumferential and radial stretches of porcine atrioventricular heart valves (n=6) at selected stress levels (% of maximum 1st-PK stress in the radial direction) under 0.5:1 tension protocol (**Figs. 3b** and **4b**). Values are presented as mean ± SEM.

% of $P_{22,max}$	λ	MVAL	MVPL	TVAL	TVPL	TVSL
0%	λ_C^{0-peak}	1.033 ± 0.025	1.045 ± 0.031	1.056 ± 0.029	1.068 ± 0.038	1.114 ± 0.033
	λ_R^{0-peak}	1.407 ± 0.057	1.474 ± 0.071	1.476 ± 0.058	1.567 ± 0.036	1.565 ± 0.075
25%	λ_C^{0-peak}	1.158 ± 0.048	1.134 ± 0.026	1.121 ± 0.029	1.197 ± 0.056	1.229 ± 0.020
	λ_R^{0-peak}	1.583 ± 0.064	1.614 ± 0.083	1.602 ± 0.075	1.783 ± 0.039	1.693 ± 0.087
50%	λ_C^{0-peak}	1.172 ± 0.051	1.145 ± 0.027	1.140 ± 0.029	1.219 ± 0.057	1.251 ± 0.019
	λ_R^{0-peak}	1.606 ± 0.065	1.535 ± 0.084	1.645 ± 0.083	1.824 ± 0.042	1.721 ± 0.091
75%	λ_C^{0-peak}	1.179 ± 0.051	1.152 ± 0.028	1.150 ± 0.029	1.231 ± 0.058	1.264 ± 0.018
	λ_R^{0-peak}	1.618 ± 0.065	1.645 ± 0.085	1.675 ± 0.094	1.840 ± 0.042	1.734 ± 0.092
100%	λ_C^{0-peak}	1.184 ± 0.052	1.158 ± 0.029	1.157 ± 0.029	1.239 ± 0.058	1.273 ± 0.018
	λ_R^{0-peak}	1.627 ± 0.066	1.652 ± 0.085	1.688 ± 0.096	1.850 ± 0.043	1.744 ± 0.093

Table 3 – Circumferential and radial stretches of porcine atrioventricular heart valves (n=6) at selected stress levels (% of maximum 1st-PK stress in the circumferential direction) under 1:0.5 tension protocol (**Figs. 3c** and **4c**). Values are presented as mean ± SEM.

% of $P_{11,max}$	λ	MVAL	MVPL	TVAL	TVPL	TVSL
0%	λ_C^{0-peak}	1.085 ± 0.035	1.099 ± 0.039	1.110 ± 0.023	1.146 ± 0.043	1.216 ± 0.037
	λ_R^{0-peak}	1.313 ± 0.036	1.367 ± 0.075	1.377 ± 0.049	1.435 ± 0.044	1.391 ± 0.070
25%	λ_C^{0-peak}	1.226 ± 0.063	1.224 ± 0.036	1.187 ± 0.029	1.302 ± 0.060	1.382 ± 0.029
	λ_R^{0-peak}	1.502 ± 0.047	1.520 ± 0.085	1.488 ± 0.063	1.643 ± 0.038	1.552 ± 0.078
50%	λ_C^{0-peak}	1.239 ± 0.066	1.237 ± 0.036	1.206 ± 0.031	1.325 ± 0.061	1.407 ± 0.028
	λ_R^{0-peak}	1.523 ± 0.048	1.537 ± 0.085	1.503 ± 0.069	1.679 ± 0.040	1.579 ± 0.084
75%	λ_C^{0-peak}	1.245 ± 0.067	1.245 ± 0.037	1.214 ± 0.032	1.336 ± 0.062	1.419 ± 0.028
	λ_R^{0-peak}	1.533 ± 0.049	1.546 ± 0.085	1.512 ± 0.071	1.697 ± 0.040	1.591 ± 0.085
100%	λ_C^{0-peak}	1.249 ± 0.068	1.251 ± 0.038	1.221 ± 0.033	1.343 ± 0.062	1.426 ± 0.029
	λ_R^{0-peak}	1.542 ± 0.049	1.553 ± 0.086	1.516 ± 0.072	1.708 ± 0.040	1.598 ± 0.086

Table 4 – Statistical analysis of the loading rate effect on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the MVAL (n=6). All quantities are presented as mean ± SEM.

Circ.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_C^{0-1}	1.053 ± 0.029	1.074 ± 0.030	1.081 ± 0.034	0.624	0.879	0.539
λ_C^{1-peak}	1.180 ± 0.038	1.142 ± 0.034	1.129 ± 0.032	0.479	0.785	0.334
λ_C^{0-peak}	1.245 ± 0.069	1.231 ± 0.069	1.225 ± 0.071	0.884	0.956	0.844
Rad.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_R^{0-1}	1.297 ± 0.078	1.366 ± 0.046	1.418 ± 0.053	0.464	0.472	0.227
λ_R^{1-peak}	1.202 ± 0.028	1.164 ± 0.022	1.142 ± 0.012	0.315	0.408	0.077
λ_R^{0-peak}	1.559 ± 0.099	1.590 ± 0.061	1.620 ± 0.065	0.789	0.744	0.614

Table 5 – Statistical analysis of the temperature effect on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the MVAL (n=6). All quantities are presented as mean ± SEM.

Circ.	2.29 N/min	4.42 N/min	7.92 N/min	p value (32-37)	p value (32-37)	p value (27-37)
λ_C^{0-1}	1.046 ± 0.023	1.088 ± 0.035	1.103 ± 0.042	0.349	0.787	0.268
λ_C^{1-peak}	1.165 ± 0.035	1.121 ± 0.019	1.114 ± 0.019	0.295	0.808	0.235
λ_C^{0-peak}	1.218 ± 0.039	1.218 ± 0.037	1.227 ± 0.045	0.999	0.871	0.873
Rad.	2.29 N/min	4.42 N/min	7.92 N/min	p value (32-37)	p value (32-37)	p value (27-37)
λ_R^{0-1}	1.397 ± 0.081	1.424 ± 0.079	1.480 ± 0.084	0.812	0.639	0.492
λ_R^{1-peak}	1.175 ± 0.017	1.129 ± 0.016	1.101 ± 0.007	0.077	0.159	0.0024
λ_R^{0-peak}	1.638 ± 0.089	1.606 ± 0.090	1.630 ± 0.093	0.807	0.859	0.951

Table 6 – Anatomical measurements for chordae and leaflet tissues of both porcine and ovine atrioventricular valves (n=6, each valve leaflet). All quantities are presented as mean ± SEM.

Anatomical Measurements		Mitral Valve		Tricuspid Valve		
		MVAL	MVPL	TVAL	TVPL	TVSL
Porcine	Leaflet Thickness (mm)	0.79 ± 0.10	0.70 ± 0.06	0.52 ± 0.06	0.46 ± 0.04	0.37 ± 0.02
	Number of Chordae	10.0 ± 0.50	20.5 ± 1.20	11.2 ± 1.30	11.8 ± 1.00	12.3 ± 1.80
	Chordae Length (mm)	17.5 ± 1.32	14.1 ± 0.70	10.8 ± 0.80	11.3 ± 1.30	11.3 ± 0.77
Adult Ovine	Leaflet Thickness (mm)	0.40 ± 0.03	0.36 ± 0.02	0.28 ± 0.03	0.26 ± 0.02	0.27 ± 0.03
	Number of Chordae	4.7 ± 0.50	10.2 ± 0.80	6.8 ± 0.50	8.2 ± 0.50	8.7 ± 0.90
	Chordae Length (mm)	17.9 ± 1.12	14.8 ± 0.59	10.5 ± 0.79	10.9 ± 0.82	12.1 ± 0.83

Table 7 – The thickness and collagen fiber content obtained from the histology sections of porcine MV and TV leaflets. Valve leaflet layers include A: atrialis, S: spongiosa, F: fibrosa, and V: ventricularis. All quantities except for collagen content (generated from image analysis) are presented as mean ± SEM with 3 repeated measurements.

		Mitral Valve		Tricuspid Valve		
		MVAL	MVPL	TVAL	TVPL	TVSL
Thickness (μm)	A	43.74 ± 4.22	54.84 ± 7.07	54.11 ± 5.32	50.27 ± 5.16	74.29 ± 4.78
	S	135.21 ± 8.23	99.85 ± 6.93	108.53 ± 5.59	221.26 ± 7.40	260.31 ± 9.26
	F	477.82 ± 6.41	556.35 ± 11.72	250.77 ± 9.69	154.01 ± 10.95	149.21 ± 6.62
	V	121.14 ± 7.64	141.44 ± 15.18	35.93 ± 3.72	20.17 ± 2.78	20.82 ± 2.24
	Intact	777.92 ± 13.46	852.49 ± 7.00	449.33 ± 3.63	445.71 ± 7.51	504.62 ± 9.55
Collagen Content (%)		77.65	69.13	68.51	45.16	32.30

Figure Captions

Figure 1 – (a) A dissected porcine heart showing the mitral valve (top) and tricuspid valve (bottom), with labels describing key anatomical components: valve leaflets, annulus, papillary muscles, and chordae tendineae (ruler shows inches). (b) Schematic of the excised leaflet and the central bulk region (top), and the mounted tissue specimen with preferred collagen fiber orientation on the biaxial mechanical testing system (C: circumferential direct, R: radial direction).

Figure 2 – (a) Image of the biaxial mechanical testing system (BioTester), (b) image of the mounted tissue, with labelled components of the testing system, (c) schematics of the valve leaflet specimen before and after prescribed loading, and (d) illustration of the force-controlled protocols (with a loading rate of 4.42 N/min) employed in both the baseline and temperature-controlled tests (*T*: maximum applied load, black solid line: tension in the circumferential direction, and red dashed line: tension in the radial direction).

Figure 3 – Mean ± SEM of the 1st-PK stress versus stretch results of the porcine MVAL and MVPL tissues (n=6) under various biaxial loading protocols at room temperature (22°C): (a) equibiaxial tension ($T_{C,\max}$:$T_{R,\max}$ = 1:1), (b) $T_{C,\max}$:$T_{R,\max}$ = 0.5:1, and (c) $T_{C,\max}$:$T_{R,\max}$ = 1:0.5.

Figure 4 – Mean ± SEM of the 1st-PK stress versus stretch results of the porcine TVAL, TVPL, and TVSL tissues (n=6) under various biaxial loading protocols at room temperature (22°C): (a) equibiaxial tension ($T_{C,\max}$:$T_{R,\max}$ = 1:1), (b) $T_{C,\max}$:$T_{R,\max}$ = 0.5:1, and (c) $T_{C,\max}$:$T_{R,\max}$ = 1:0.5.

Figure 5 – Representative biaxial mechanical testing results of each porcine atrioventricular leaflet under equibiaxial tension ($T_{C,\max}$:$T_{R,\max}$ = 1:1) at room temperature (22°C), showing the effect of varied loading rates on the quantified 1st-PK stress versus stretch results: (a) MVAL, (b) MVPL, (c) TVAL, (d) TVPL, and (e) TVSL.

Figure 6 – Statistical analyses of the MVAL from the loading rate effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, p>0.10, and **#**: nearly statistically significant difference, p<0.10)

Figure 7 – Representative biaxial mechanical testing results of each porcine atrioventricular leaflet under equibiaxial tension ($T_{C,\max}$:$T_{R,\max}$ = 1:1), showing the effect of temperature on the quantified 1st-PK stress versus stretch results: (a) MVAL, (b) MVPL, (c) TVAL, (d) TVPL, and (e) TVSL.

Figure 8 – Statistical analyses of the MVAL from the temperature effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, p>0.10)

Figure 9 – Comparisons of the biaxial mechanical responses of representative atrioventricular valve leaflet tissues under equibiaxial tension ($T_{C,\max}$:$T_{R,\max}$ = 1:1) at body temperature (37°C) between different species (adult porcine, adult ovine/sheep, and juvenile ovine/lamb).

Figure 10 – Histological sections of the MV and TV leaflets (MVAL, MVPL, TVAL, TVPL, and TVSL) stained with Masson's Trichrome. Four morphologically distinct layers of the leaflets (atrialis: A, spongiosa: S, ventricularis: V, fibrosa: F) were identified.

Figure Captions (Appendix)

Figure A1 – Mean ± SEM of the 1st-PK stress versus stretch results of the porcine MVAL and MVPL tissues (n=6) under intermediate biaxial loading protocols at room temperature (22°C): (a) $T_{C,\max}$:$T_{R,\max}$ = 0.75:1, and (b) $T_{C,\max}$:$T_{R,\max}$ = 1:0.75.

Figure A2 – Mean ± SEM of the 1st-PK stress versus stretch results of the porcine TVAL, TVPL, and TVSL tissues (n=6) under intermediate biaxial loading protocols at room temperature (22°C): (a) $T_{C,\max}$:$T_{R,\max}$ = 0.75:1, and (b) $T_{C,\max}$:$T_{R,\max}$ = 1:0.75.

Figure A3 – Representative biaxial mechanical testing results of each porcine atrioventricular leaflet under biaxial tensions ($T_{C,\max}$:$T_{R,\max}$ = 0.5:1) at room temperature (22°C), showing the effect of varied loading rates on the quantified 1st-PK stress versus stretch results: (a) MVAL, (b) MVPL, (c) TVAL, (d) TVPL, and (e) TVSL.

Figure A4 – Representative biaxial mechanical testing results of each porcine atrioventricular leaflet under biaxial tensions ($T_{C,\max}$:$T_{R,\max}$ = 1:0.5) at room temperature (22°C), showing the effect of varied loading rates on the quantified 1st-PK stress versus stretch results: (a) MVAL, (b) MVPL, (c) TVAL, (d) TVPL, and (e) TVSL.

Figure A5 – Representative biaxial mechanical testing results of each porcine atrioventricular leaflet under biaxial tensions ($T_{C,\max}$:$T_{R,\max}$ = 0.5:1), showing the effect of temperature on the quantified 1st-PK stress versus stretch results: (a) MVAL, (b) MVPL, (c) TVAL, (d) TVPL, and (e) TVSL.

Figure A6 – Representative biaxial mechanical testing results of each porcine atrioventricular leaflet under biaxial tensions ($T_{C,\max}$:$T_{R,\max}$ = 1:0.5), showing the effect of temperature on the quantified 1st-PK stress versus stretch results: (a) MVAL, (b) MVPL, (c) TVAL, (d) TVPL, and (e) TVSL.

References

1. Klabunde R. Cardiovascular physiology concepts: Lippincott Williams & Wilkins; 2011.
2. Shiran A., and Sagie A. (2009) Tricuspid regurgitation in mitral valve disease. **Journal of the American College of Cardiology**, 53(5):401-08.
3. O'rourke R. A., and Crawford M. H. (1984) Mitral valve regurgitation. **Current Problems in Cardiology**, 9(2):1-52.
4. Nkomo V. T., Gardin J. M., Skelton T. N., Gottdiener J. S., Scott C. G., and Enriquez-Sarano M. (2006) Burden of valvular heart diseases: A population-based study. **The Lancet**, 368(9540):1005-11.
5. Waller B. F., Howard J., and Fess S. (1994) Pathology of mitral valve stenosis and pure mitral regurgitation—part i. **Clinical Cardiology**, 17(6):330-36.
6. Taramasso M., Pozzoli A., Guidotti A., Nietlispach F., Inderbitzin D. T., Benussi S., Alfieri O., and Maisano F. (2016) Percutaneous tricuspid valve therapies: The new frontier. **European Heart Journal**, 38(9):639-47.
7. Preston-Maher G. L., Torii R., and Burriesci G. (2015) A technical review of minimally invasive mitral valve replacements. **Cardiovascular Engineering and Technology**, 6(2):174-84.
8. Rodés-Cabau J., Taramasso M., and T O'gara P. (2016) Diagnosis and treatment of tricuspid valve disease: Current and future perspectives. **The Lancet**, 388(10058):2431-42.
9. Schofer J., Bijuklic K., Tiburtius C., Hansen L., Groothuis A., and Hahn R. T. (2015) First-in-human transcatheter tricuspid valve repair in a patient with severely regurgitant tricuspid valve. **Journal of the American College of Cardiology**, 65(12):1190-95.
10. Schoen F. J., and Gotlieb A. I. (2016) Heart valve health, disease, replacement, and repair: A 25-year cardiovascular pathology perspective. **Cardiovascular Pathology**, 25(4):341-52.
11. Franzen O., Baldus S., Rudolph V., Meyer S., Knap M., Koschyk D., Treede H., Barmeyer A., Schofer J., and Costard-Jäckle A. (2010) Acute outcomes of mitraclip therapy for mitral regurgitation in high-surgical-risk patients: Emphasis on adverse valve morphology and severe left ventricular dysfunction. **European Heart Journal**, 31(11):1373-81.
12. Hammerstingl C., Schueler R., Malasa M., Werner N., and Nickenig G. (2016) Transcatheter treatment of severe tricuspid regurgitation with the mitraclip system. **European Heart Journal**, 37(10):849-53.
13. Grigioni F., Enriquez-Sarano M., Zehr K. J., Bailey K. R., and Tajik A. J. (2001) Ischemic mitral regurgitation. **Circulation**, 103(13):1759-64.
14. Lancellotti P., Tribouilloy C., Hagendorff A., Moura L., Popescu B. A., Agricola E., Monin J.-L., Pierard L. A., Badano L., and Zamorano J. L. (2010) European association of echocardiography recommendations for the assessment of valvular regurgitation. Part 1: Aortic and pulmonary regurgitation (native valve disease). **European Journal of Echocardiography**, 11(3):223-44.
15. Shanks M., Siebelink H.-M. J., Delgado V., Van De Veire N. R., Ng A. C., Sieders A., Schuijf J. D., Lamb H. J., Marsan N. A., and Westenberg J. J. (2010) Quantitative assessment of mitral regurgitation. **Circulation: Cardiovascular Imaging**, 3(6):694-700.
16. Lee C.-H., Oomen P. J., Rabbah J. P., Yoganathan A., Gorman R. C., Gorman J. H., Amini R., and Sacks M. S. A high-fidelity and micro-anatomically accurate 3d finite element model for simulations of functional mitral valve. International Conference on Functional Imaging and Modeling of the Heart; 2013: Springer.
17. Aggarwal A., Aguilar V. S., Lee C.-H., Ferrari G., Gorman J. H., Gorman R. C., and Sacks M. S. Patient-specific modeling of heart valves: From image to simulation. International Conference on Functional Imaging and Modeling of the Heart; 2013: Springer.

18. Lee C.-H., Rabbah J.-P., Yoganathan A. P., Gorman R. C., Gorman J. H., and Sacks M. S. (2015) On the effects of leaflet microstructure and constitutive model on the closing behavior of the mitral valve. **Biomechanics and Modeling in Mechanobiology**, 14(6):1281-302.
19. Wang Q., and Sun W. (2013) Finite element modeling of mitral valve dynamic deformation using patient-specific multi-slices computed tomography scans. **Annals of Biomedical Engineering**, 41(1):142-53.
20. Taylor C. A., and Figueroa C. (2009) Patient-specific modeling of cardiovascular mechanics. **Annual Review of Biomedical Engineering**, 11:109-34.
21. Stevanella M., Maffessanti F., Conti C. A., Votta E., Arnoldi A., Lombardi M., Parodi O., Caiani E. G., and Redaelli A. (2011) Mitral valve patient-specific finite element modeling from cardiac mri: Application to an annuloplasty procedure. **Cardiovascular Engineering and Technology**, 2(2):66-76.
22. Rim Y., Laing S. T., Mcpherson D. D., and Kim H. (2014) Mitral valve repair using eptfe sutures for ruptured mitral chordae tendineae: A computational simulation study. **Annals of Biomedical Engineering**, 42(1):139-48.
23. Choi A., Rim Y., Mun J. S., and Kim H. (2014) A novel finite element-based patient-specific mitral valve repair: Virtual ring annuloplasty. **Bio-Medical Materials and Engineering**, 24(1):341-47.
24. Abbasi M., Barakat M. S., Vahidkhah K., and Azadani A. N. (2016) Characterization of three-dimensional anisotropic heart valve tissue mechanical properties using inverse finite element analysis. **Journal of the Mechanical Behavior of Biomedical Materials**, 62:33-44.
25. Lee C.-H., Amini R., Gorman R. C., Gorman J. H., and Sacks M. S. (2014) An inverse modeling approach for stress estimation in mitral valve anterior leaflet valvuloplasty for in-vivo valvular biomaterial assessment. **Journal of Biomechanics**, 47(9):2055-63.
26. Huang H.-Y. S., and Lu J. (2017) Biaxial mechanical properties of bovine jugular venous valve leaflet tissues. **Biomechanics and Modeling in Mechanobiology**:1-13.
27. Kavanagh E. G., Grace P. A., Mcgloughlin T. M., and Doyle B. J. (2014) The biaxial mechanical behaviour of abdominal aortic aneurysm intraluminal thrombus: Classification of morphology and the determination of layer and region specific properties. **Journal of Biomechanics**, 47(6):1430-37.
28. Schmid F., Sommer G., Rappolt M., Regitnig P., Holzapfel G. A., Laggner P., and Amenitsch H. (2006) Bidirectional tensile testing cell for in situ small angle x-ray scattering investigations of soft tissue. **Nuclear Instruments and Methods in Physics Research Section B: Beam Interactions with Materials and Atoms**, 246(1):262-68.
29. Sacks M. (1999) A method for planar biaxial mechanical testing that includes in-plane shear. **Journal of Biomechanical Engineering**, 121(5):551-55.
30. Billiar K. L., and Sacks M. S. (2000) Biaxial mechanical properties of the natural and glutaraldehyde treated aortic valve cusp-part i: Experimental results. **Transactions-American Society of Mechanical Engineers Journal of Biomechanical Engineering**, 122(1):23-30.
31. Sacks M. S. (2000) Biaxial mechanical evaluation of planar biological materials. **Journal of Elasticity**, 61(1):199.
32. Balguid A., Rubbens M. P., Mol A., Bank R. A., Bogers A. J., Van Kats J. P., De Mol B. A., Baaijens F. P., and Bouten C. V. (2007) The role of collagen cross-links in biomechanical behavior of human aortic heart valve leaflets—relevance for tissue engineering. **Tissue Engineering**, 13(7):1501-11.
33. Billiar K., and Sacks M. (1997) A method to quantify the fiber kinematics of planar tissues under biaxial stretch. **Journal of Biomechanics**, 30(7):753-56.

34. De Hart J., Peters G., Schreurs P., and Baaijens F. (2004) Collagen fibers reduce stresses and stabilize motion of aortic valve leaflets during systole. **Journal of Biomechanics**, 37(3):303-11.
35. Khoiy K. A., and Amini R. (2016) On the biaxial mechanical response of porcine tricuspid valve leaflets. **Journal of Biomechanical Engineering**, 138(10):104504.
36. May-Newman K., and Yin F. (1995) Biaxial mechanical behavior of excised porcine mitral valve leaflets. **American Journal of Physiology-Heart and Circulatory Physiology**, 269(4):H1319-H27.
37. Grashow J. S., Yoganathan A. P., and Sacks M. S. (2006) Biaixal stress–stretch behavior of the mitral valve anterior leaflet at physiologic strain rates. **Annals of Biomedical Engineering**, 34(2):315-25.
38. Stella J. A., and Sacks M. S. (2007) On the biaxial mechanical properties of the layers of the aortic valve leaflet. **Journal of Biomechanical Engineering**, 129(5):757-66.
39. Potter S., Graves J., Drach B., Leahy T., Hammel C., Feng Y., Baker A., and Sacks M. S. (2018) A novel small-specimen planar biaxial testing system with full in-plane deformation control. **Journal of Biomechanical Engineering**, 140(5):051001.
40. Dreyfus G. D., Corbi P. J., Chan K. J., and Bahrami T. (2005) Secondary tricuspid regurgitation or dilatation: Which should be the criteria for surgical repair? **The Annals of thoracic surgery**, 79(1):127-32.
41. Anyanwu A. C., and Adams D. H. Functional tricuspid regurgitation in mitral valve disease: Epidemiology and prognostic implications. Seminars in Thoracic and Cardiovascular Surgery; 2010: Elsevier.
42. Navia J. L., Nowicki E. R., Blackstone E. H., Brozzi N. A., Nento D. E., Atik F. A., Rajeswaran J., Gillinov A. M., Svensson L. G., and Lytle B. W. (2010) Surgical management of secondary tricuspid valve regurgitation: Annulus, commissure, or leaflet procedure? **The Journal of Thoracic and Cardiovascular Surgery**, 139(6):1473-82. e5.
43. Woo S. L.-Y., Orlando C. A., Camp J. F., and Akeson W. H. (1986) Effects of postmortem storage by freezing on ligament tensile behavior. **Journal of Biomechanics**, 19(5):399-404.
44. Stemper B. D., Yoganandan N., Stineman M. R., Gennarelli T. A., Baisden J. L., and Pintar F. A. (2007) Mechanics of fresh, refrigerated, and frozen arterial tissue. **Journal of Surgical Research**, 139(2):236-42.
45. Foutz T., Stone E., and Abrams J. C. (1992) Effects of freezing on mechanical properties of rat skin. **American Journal of Veterinary Research**, 53(5):788-92.
46. Liao J., Joyce E. M., and Sacks M. S. (2008) Effects of decellularization on the mechanical and structural properties of the porcine aortic valve leaflet. **Biomaterials**, 29(8):1065-74.
47. Pierlot C. M., Moeller A. D., Lee J. M., and Wells S. M. (2015) Biaxial creep resistance and structural remodeling of the aortic and mitral valves in pregnancy. **Annals of Biomedical Engineering**, 43(8):1772-85.
48. Chuong C.-J., and Fung Y.-C. Residual stress in arteries. Frontiers in biomechanics: Springer; 1986. p. 117-29.
49. Fung Y., Fronek K., and Patitucci P. (1979) Pseudoelasticity of arteries and the choice of its mathematical expression. **American Journal of Physiology-Heart and Circulatory Physiology**, 237(5):H620-H31.
50. Humphrey J. D., Vawter D. L., and Vito R. P. (1987) Quantification of strains in biaxially tested soft tissues. **Journal of Biomechanics**, 20(1):59-65.
51. Toma M., Jensen M. Ø., Einstein D. R., Yoganathan A. P., Cochran R. P., and Kunzelman K. S. (2016) Fluid–structure interaction analysis of papillary muscle forces using a comprehensive mitral valve model with 3d chordal structure. **Annals of Biomedical Engineering**, 44(4):942-53.

52. Ruifrok A. C., and Johnston D. A. (2001) Quantification of histochemical staining by color deconvolution. **Analytical and Quantitative Cytology and Histology**, 23(4):291-99.
53. Hosseini S. M., Wilson W., Ito K., and Van Donkelaar C. C. (2014) How preconditioning affects the measurement of poro-viscoelastic mechanical properties in biological tissues. **Biomechanics and Modeling in Mechanobiology**, 13(3):503-13.
54. Jett S., Laurence D., Kunkel R., Babu A. R., Kramer K., Baumwart R., Towner R., Wu Y., and Lee C.-H. (submitted) Biaxial mechanical data of porcine atrioventricular valve leaflets. **Data in Brief**.
55. Bernacca G. M., O'connor B., Williams D. F., and Wheatley D. J. (2002) Hydrodynamic function of polyurethane prosthetic heart valves: Influences of young's modulus and leaflet thickness. **Biomaterials**, 23(1):45-50.
56. Christie G. W., and Barratt-Boyes B. G. (1995) Age-dependent changes in the radial stretch of human aortic valve leaflets determined by biaxial testing. **The Annals of thoracic surgery**, 60:S156-S59.
57. Christie G. W., and Barratt-Boyes B. G. (1995) Mechanical properties of porcine pulmonary valve leaflets: How do they differ from aortic leaflets? **The Annals of thoracic surgery**, 60:S195-S99.
58. Huang H.-Y. S., Balhouse B. N., and Huang S. (2012) Application of simple biomechanical and biochemical tests to heart valve leaflets: Implications for heart valve characterization and tissue engineering. **Proceedings of the Institution of Mechanical Engineers, Part H: Journal of Engineering in Medicine**, 226(11):868-76.
59. Wang Z., Golob M. J., and Chesler N. C. Viscoelastic properties of cardiovascular tissues. Viscoelastic and viscoplastic materials: InTech; 2016.
60. Lelovas P. P., Kostomitsopoulos N. G., and Xanthos T. T. (2014) A comparative anatomic and physiologic overview of the porcine heart. **Journal of the American Association for Laboratory Animal Science**, 53(5):432-38.
61. Hinton R. B., and Yutzey K. E. (2011) Heart valve structure and function in development and disease. **Annual review of physiology**, 73:29-46.
62. Lam J., Ranganathan N., Wigle E., and Silver M. (1970) Morphology of the human mitral valve. **Circulation**, 41(3):449-58.
63. Silver M., Lam J., Ranganathan N., and Wigle E. (1971) Morphology of the human tricuspid valve. **Circulation**, 43(3):333-48.
64. Labrosse M. R., Jafar R., Ngu J., and Boodhwani M. (2016) Planar biaxial testing of heart valve cusp replacement biomaterials: Experiments, theory and material constants. **Acta Biomaterialia**, 45:303-20.
65. Chen L., Yin F. C., and May-Newman K. (2004) The structure and mechanical properties of the mitral valve leaflet-strut chordae transition zone. **Journal of Biomechanical Engineering**, 126(2):244-51.
66. Sacks M. S., and Yoganathan A. P. (2007) Heart valve function: A biomechanical perspective. **Philosophical Transactions of the Royal Society of London B: Biological Sciences**, 362(1484):1369-91.
67. Hadian M., Corcoran B. M., Han R. I., Grossmann J. G., and Bradshaw J. P. (2007) Collagen organization in canine myxomatous mitral valve disease: An x-ray diffraction study. **Biophysical Journal**, 93(7):2472-76.
68. Cole W., Chan D., Hickey A., and Wilcken D. (1984) Collagen composition of normal and myxomatous human mitral heart valves. **Biochemical Journal**, 219(2):451-60.
69. Alavi S. H., Sinha A., Steward E., Milliken J. C., and Kheradvar A. (2015) Load-dependent extracellular matrix organization in atrioventricular heart valves: Differences and similarities. **American Journal of Physiology-Heart and Circulatory Physiology**, 309(2):H276-H84.

70. Lee C.-H., Zhang W., Feaver K., Gorman R. C., Gorman J. H., and Sacks M. S. (2017) On the in vivo function of the mitral heart valve leaflet: Insights into tissue–interstitial cell biomechanical coupling. **Biomechanics and Modeling in Mechanobiology**, 16(5):1613-32.
71. Stevanella M., Votta E., Lemma M., Antona C., and Redaelli A. (2010) Finite element modelling of the tricuspid valve: A preliminary study. **Medical Engineering and Physics**, 32(10):1213-23.

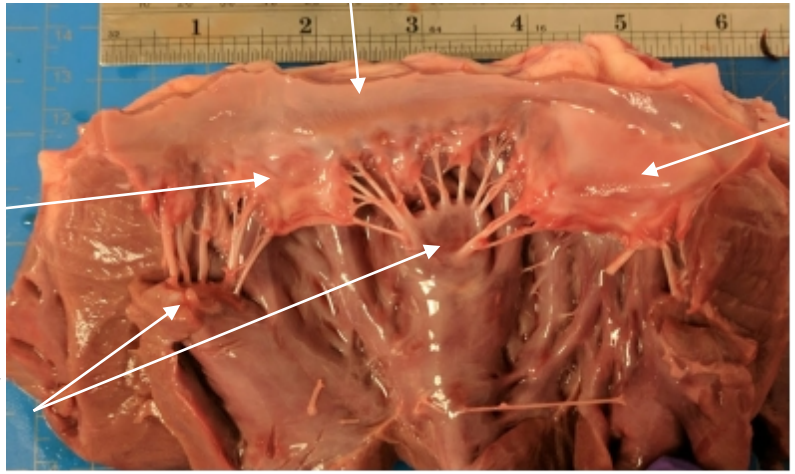
(a)

Mitral Annulus

MVAL

MVPL

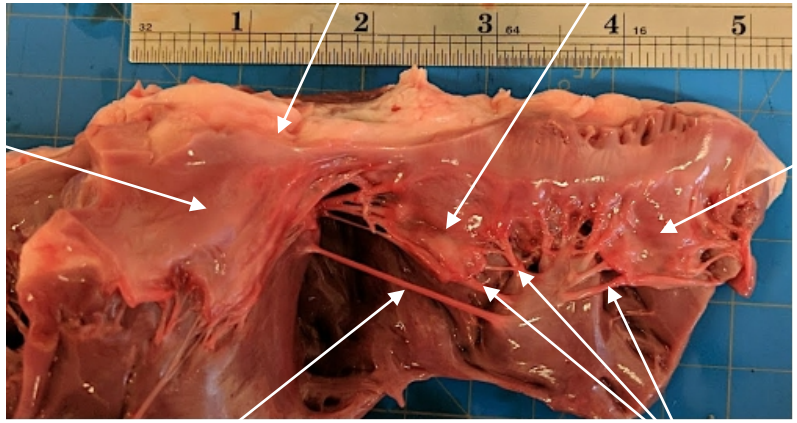
Papillary Muscles



Tricuspid Annulus TVAL

TVSL

TVPL



Moderator Band

Chordae Tendineae

(b)

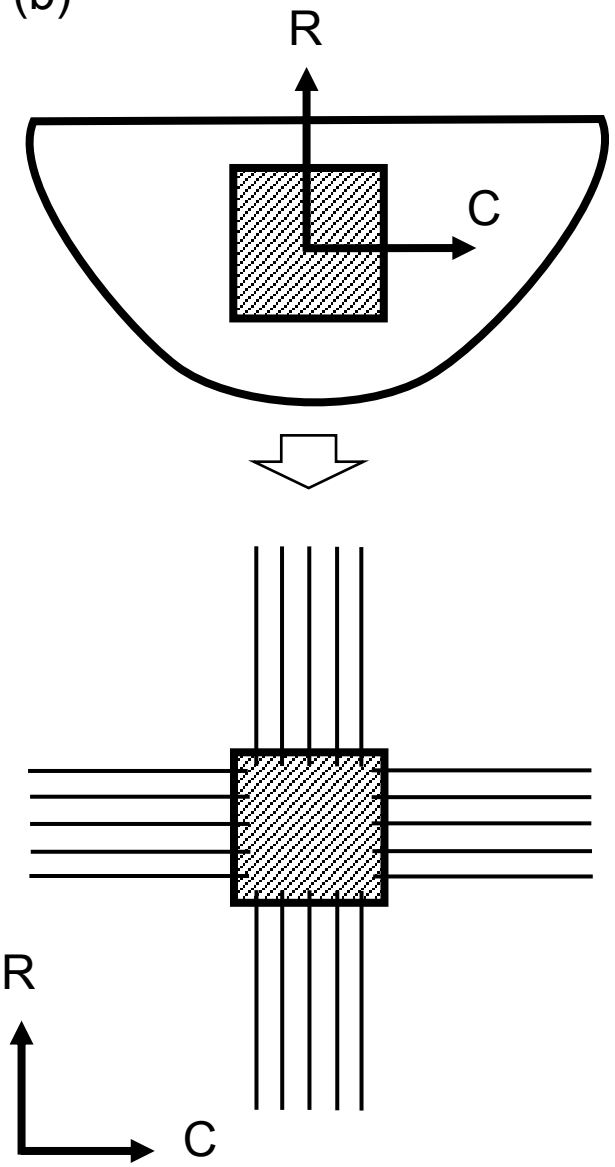
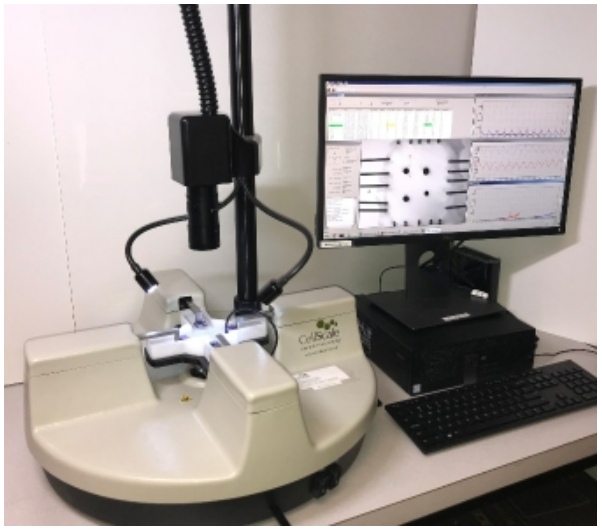
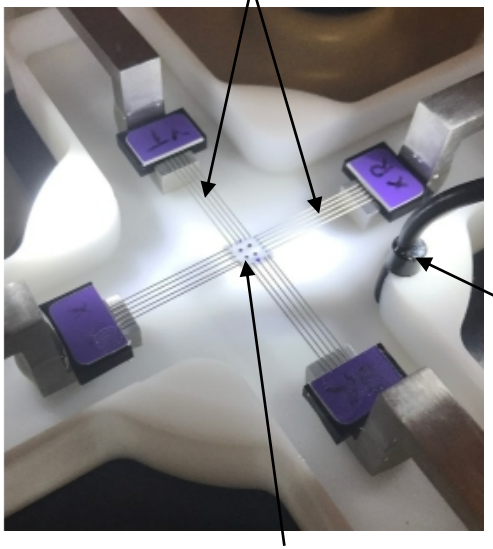


Figure 2

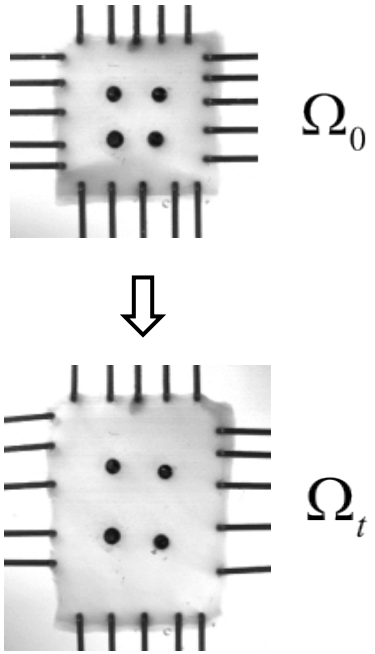
(a)



(b)



(c)



(d)

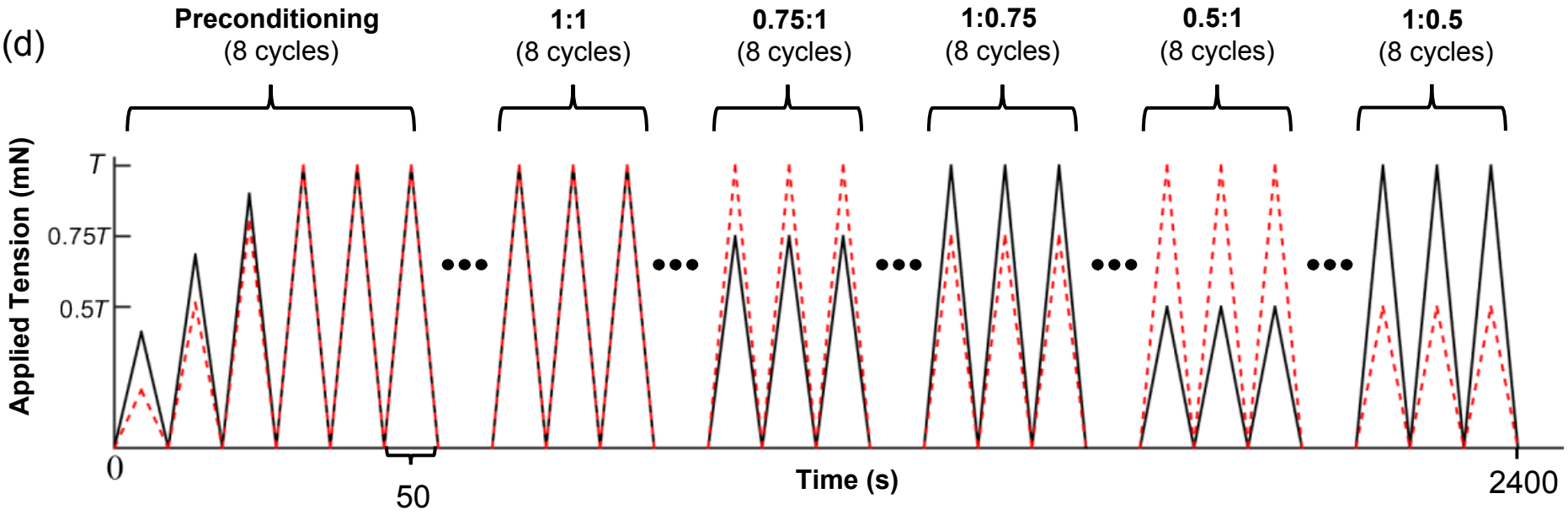


Figure 3

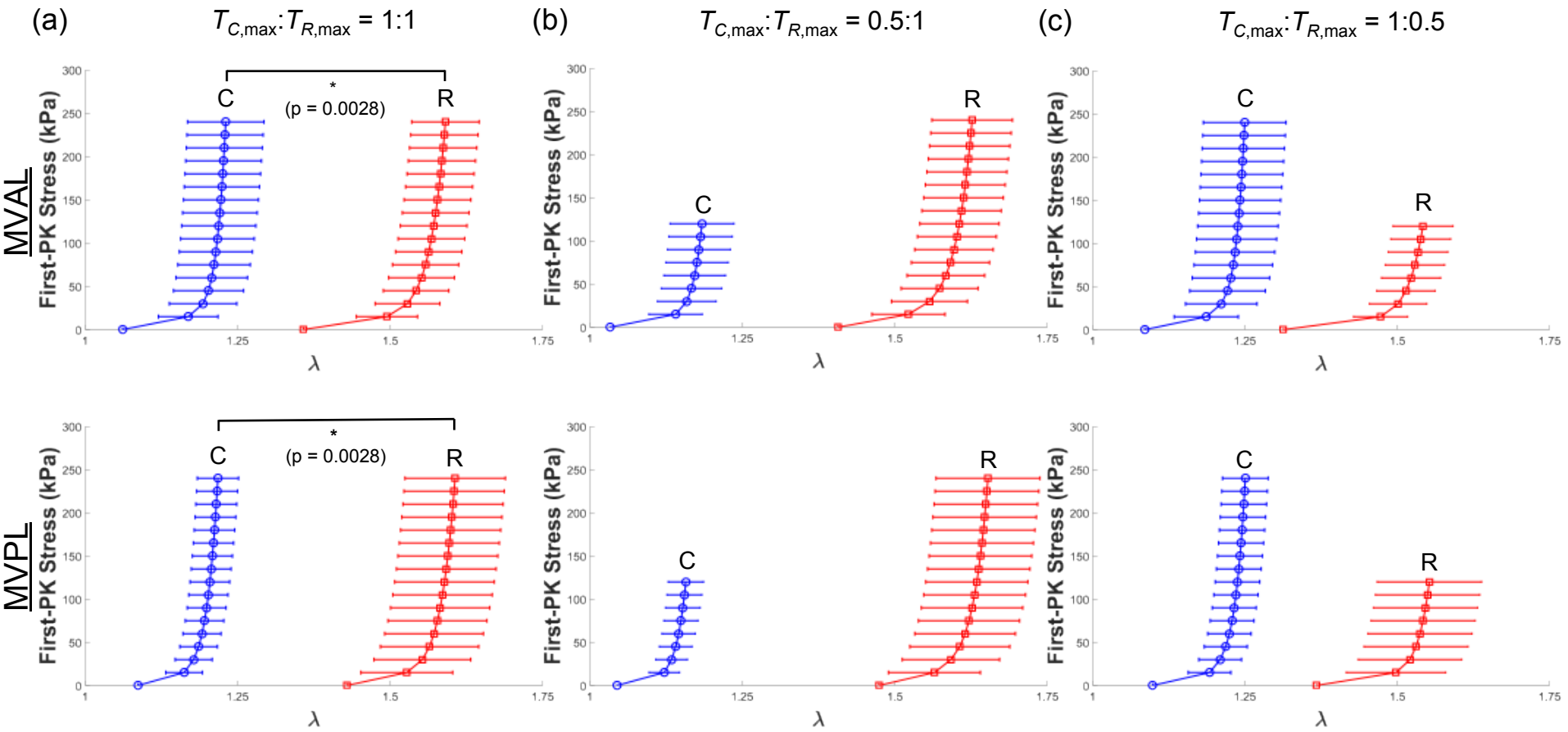

(a)
$T_{C,max}$:$T_{R,max}$ = 1:1
(b)
$T_{C,max}$:$T_{R,max}$ = 0.5:1
(c)
$T_{C,max}$:$T_{R,max}$ = 1:0.5
MVAL
MVPL
First-PK Stress (kPa)
λ
C
R
*
(p = 0.0028)

Figure 4

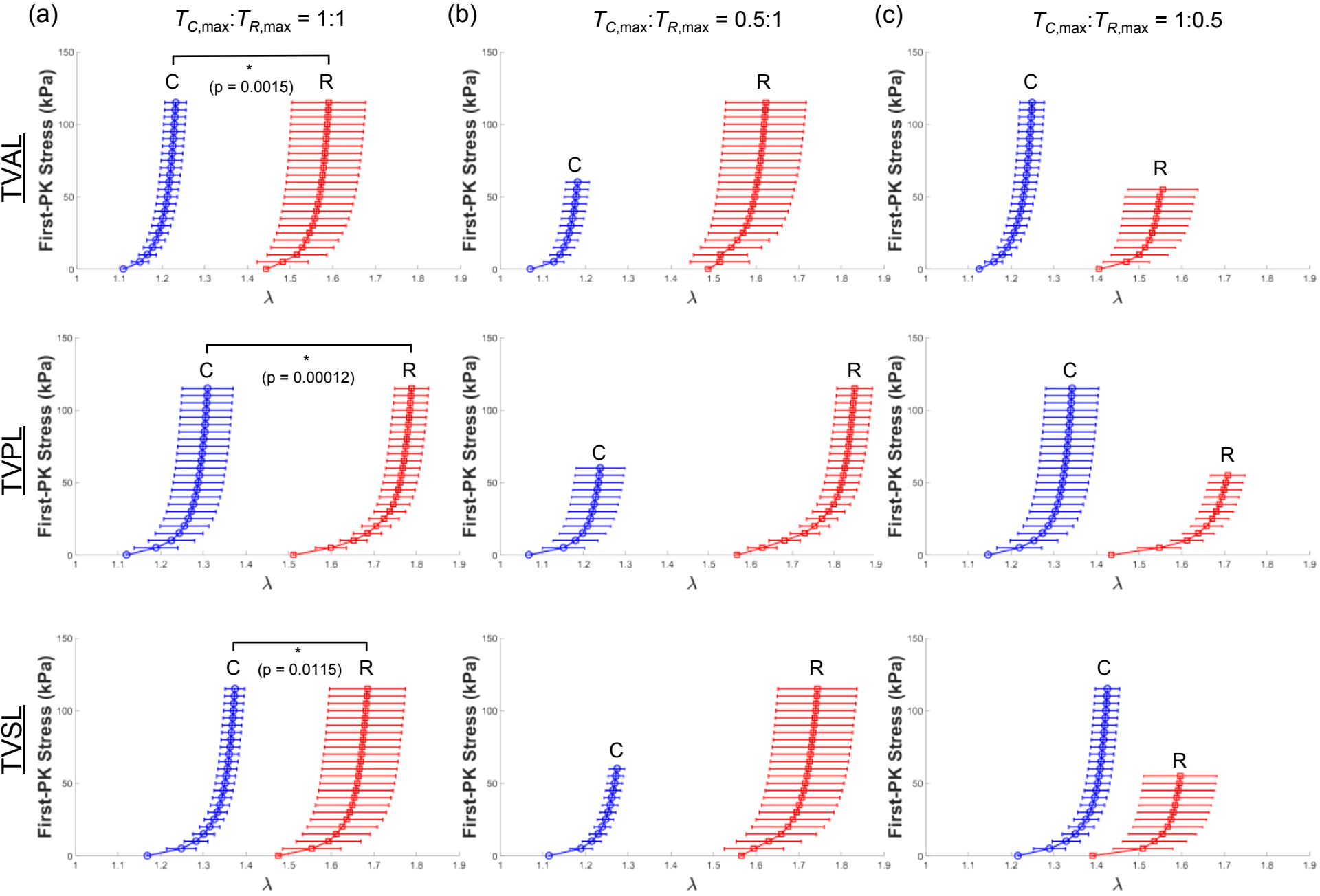

(a)
$T_{C,max}$:$T_{R,max}$ = 1:1
(b)
$T_{C,max}$:$T_{R,max}$ = 0.5:1
(c)
$T_{C,max}$:$T_{R,max}$ = 1:0.5
TVAL
TVPL
TVSL
First-PK Stress (kPa)
λ
C
R
*
(p = 0.0015)
(p = 0.00012)
(p = 0.0115)

Figure 5

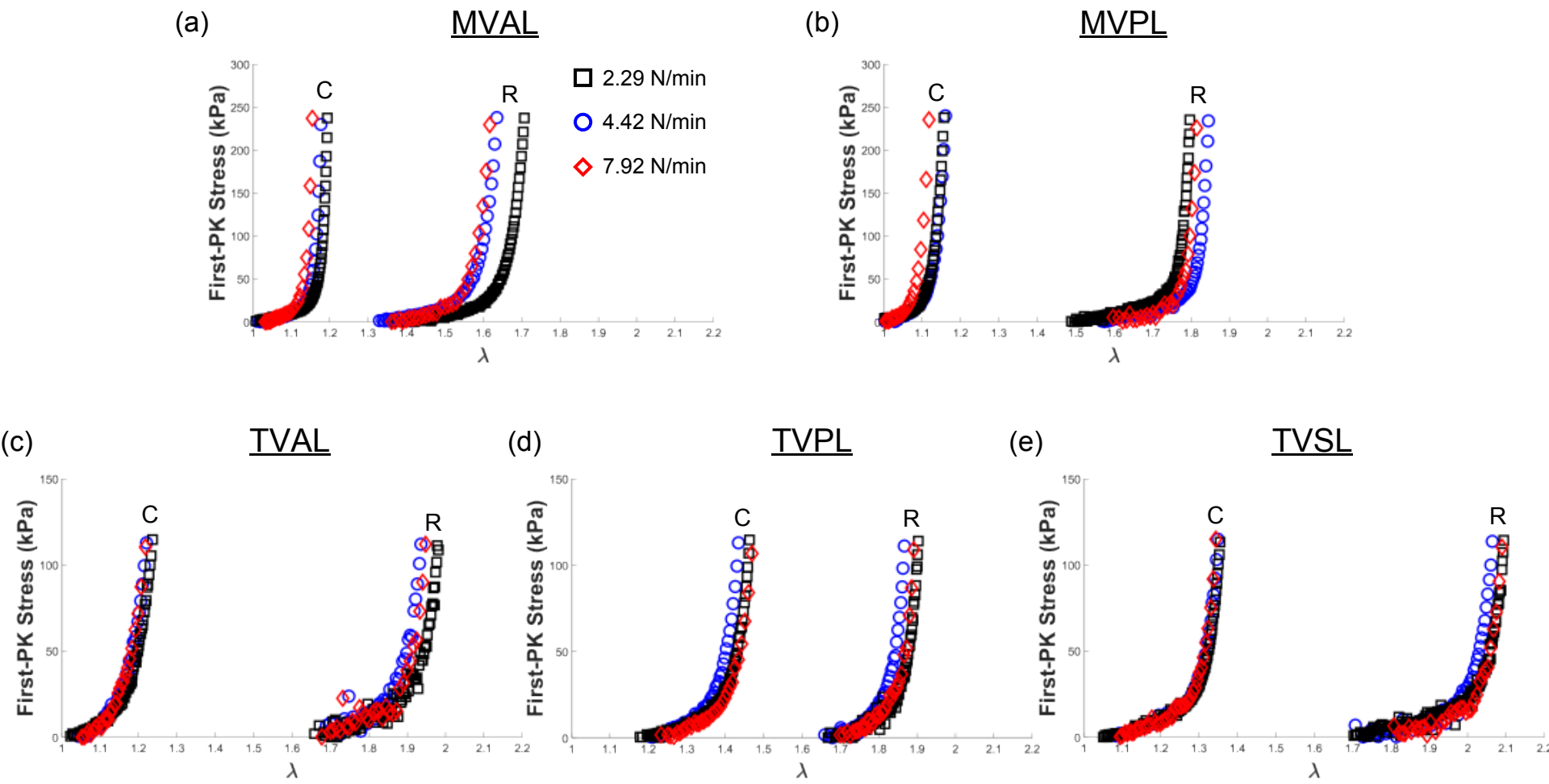
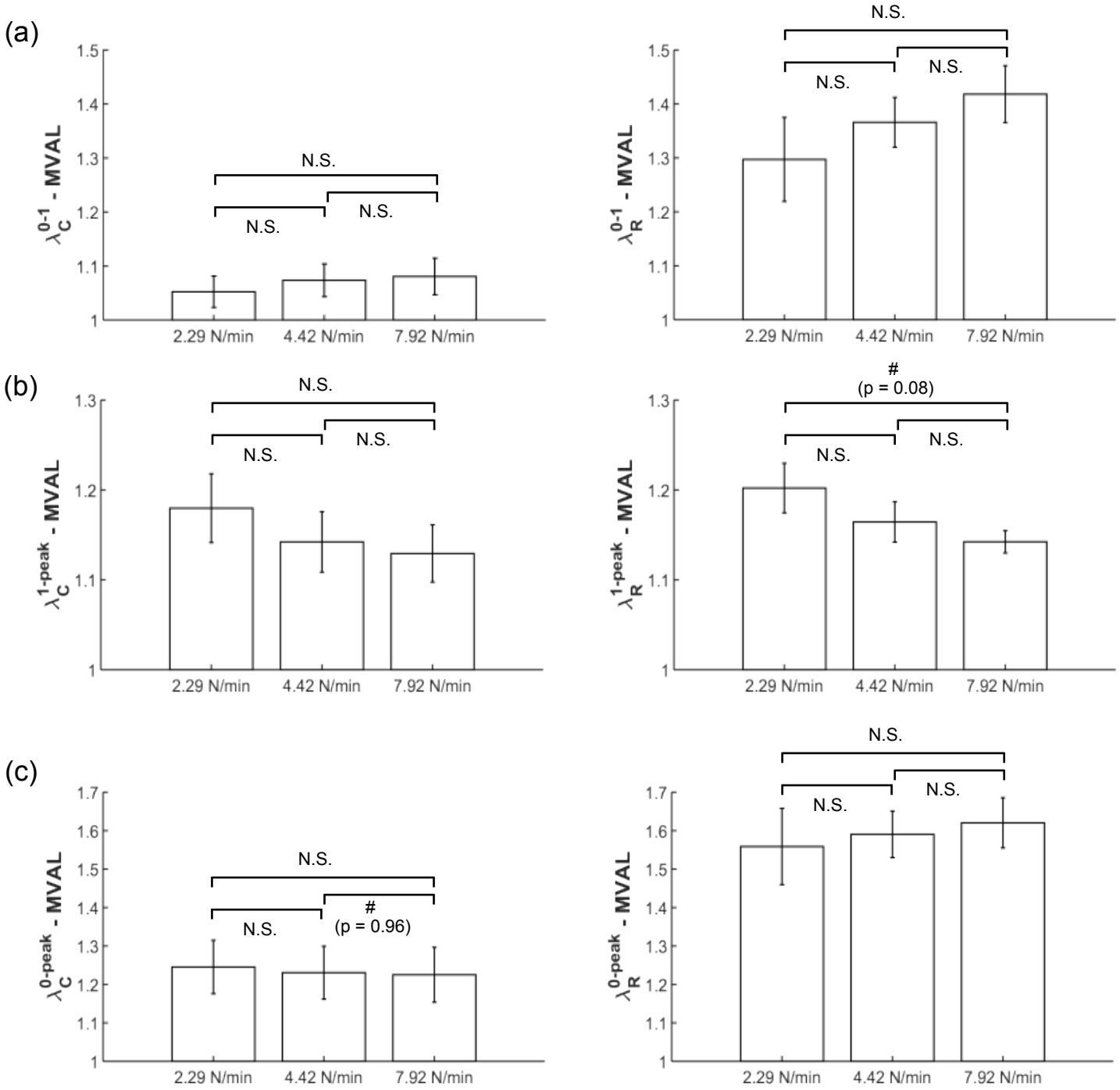

(a)
MVAL
(b)
MVPL
(c)
TVAL
(d)
TVPL
(e)
TVSL
C
R
2.29 N/min
4.42 N/min
7.92 N/min
First-PK Stress (kPa)
λ

Figure 6


(a)
$\lambda_C^{0\text{-}1}$ - MVAL
$\lambda_R^{0\text{-}1}$ - MVAL
N.S.
N.S.
N.S.
(b)
$\lambda_C^{1\text{-peak}}$ - MVAL
$\lambda_R^{1\text{-peak}}$ - MVAL
#
(p = 0.08)
(c)
$\lambda_C^{0\text{-peak}}$ - MVAL
$\lambda_R^{0\text{-peak}}$ - MVAL
#
(p = 0.96)
2.29 N/min
4.42 N/min
7.92 N/min

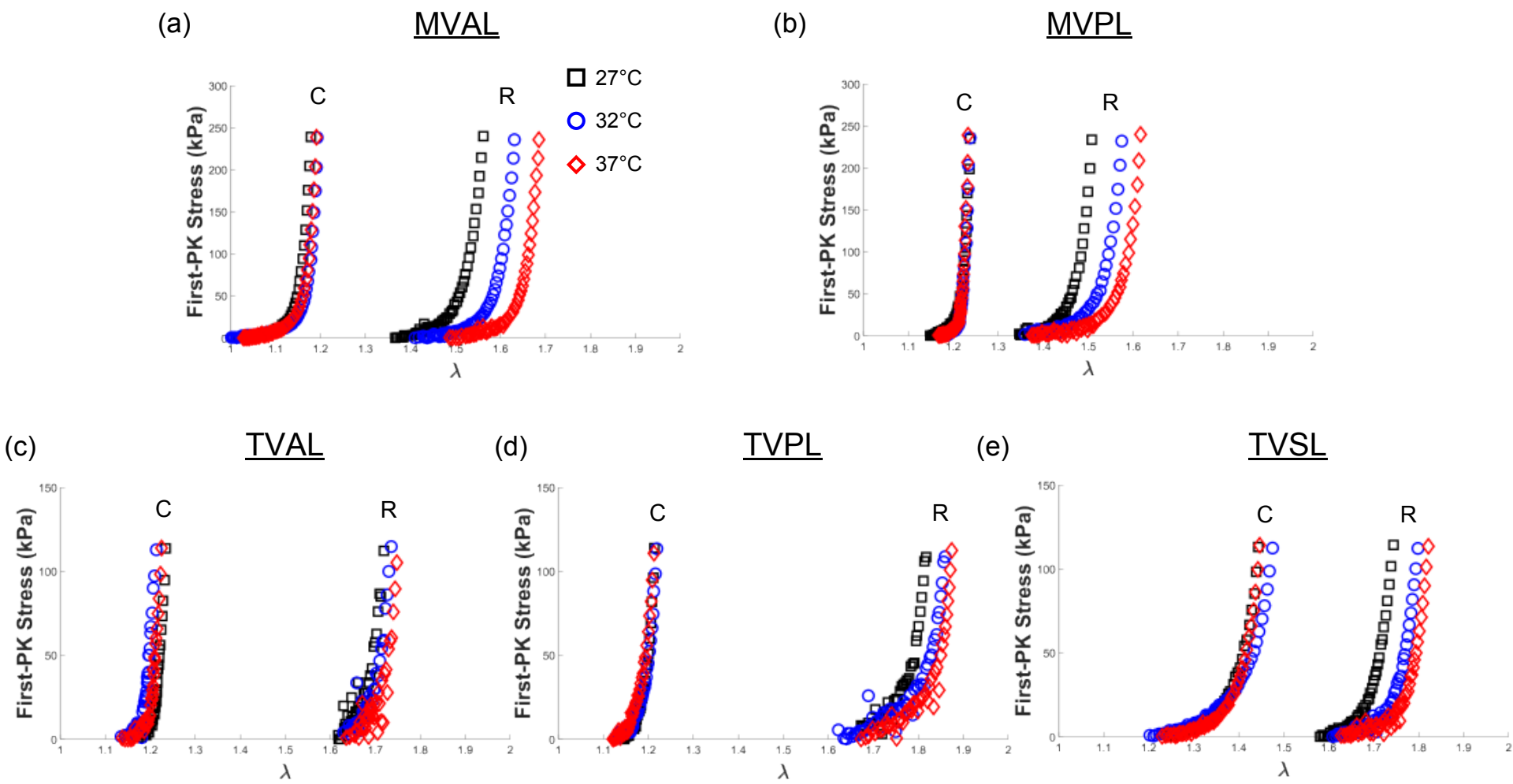

(a)
MVAL
(b)
MVPL
(c)
TVAL
(d)
TVPL
(e)
TVSL
27°C
32°C
37°C
C
R
First-PK Stress (kPa)
λ

Figure 8

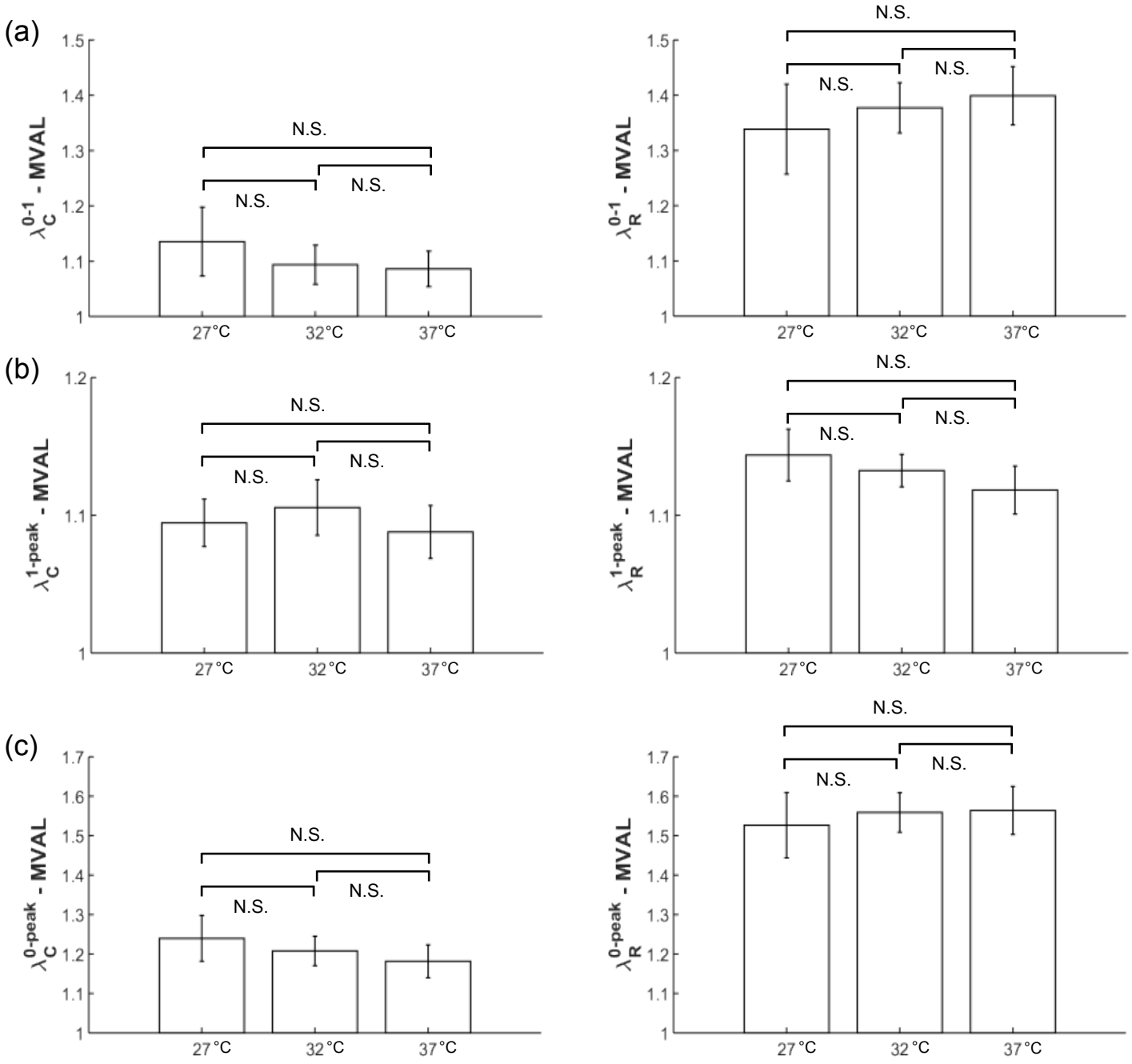

(a)
λ_C^{0-1} - MVAL
λ_R^{0-1} - MVAL
(b)
$\lambda_C^{1\text{-peak}}$ - MVAL
$\lambda_R^{1\text{-peak}}$ - MVAL
(c)
$\lambda_C^{0\text{-peak}}$ - MVAL
$\lambda_R^{0\text{-peak}}$ - MVAL
N.S.
27°C
32°C
37°C

Figure 9

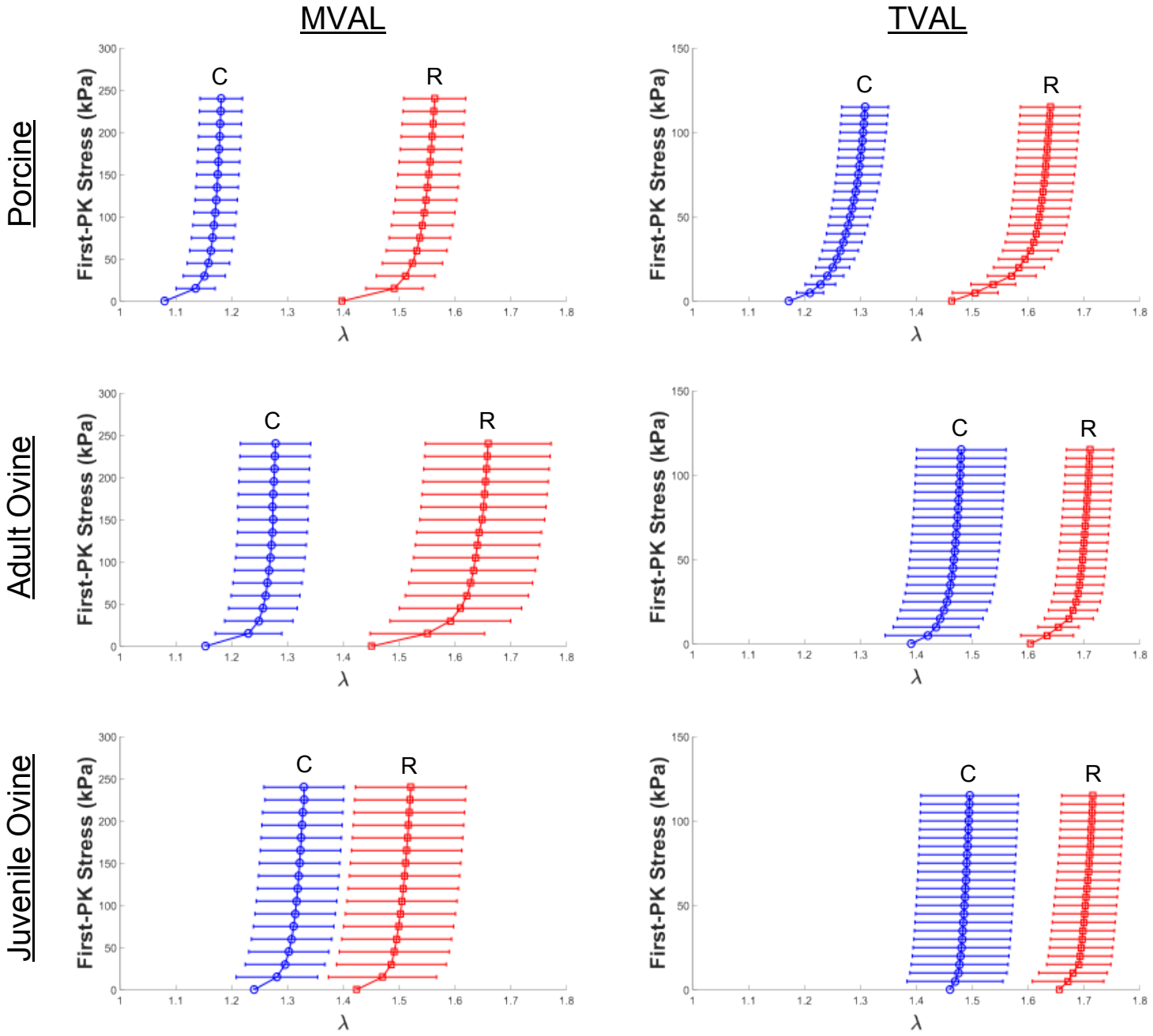
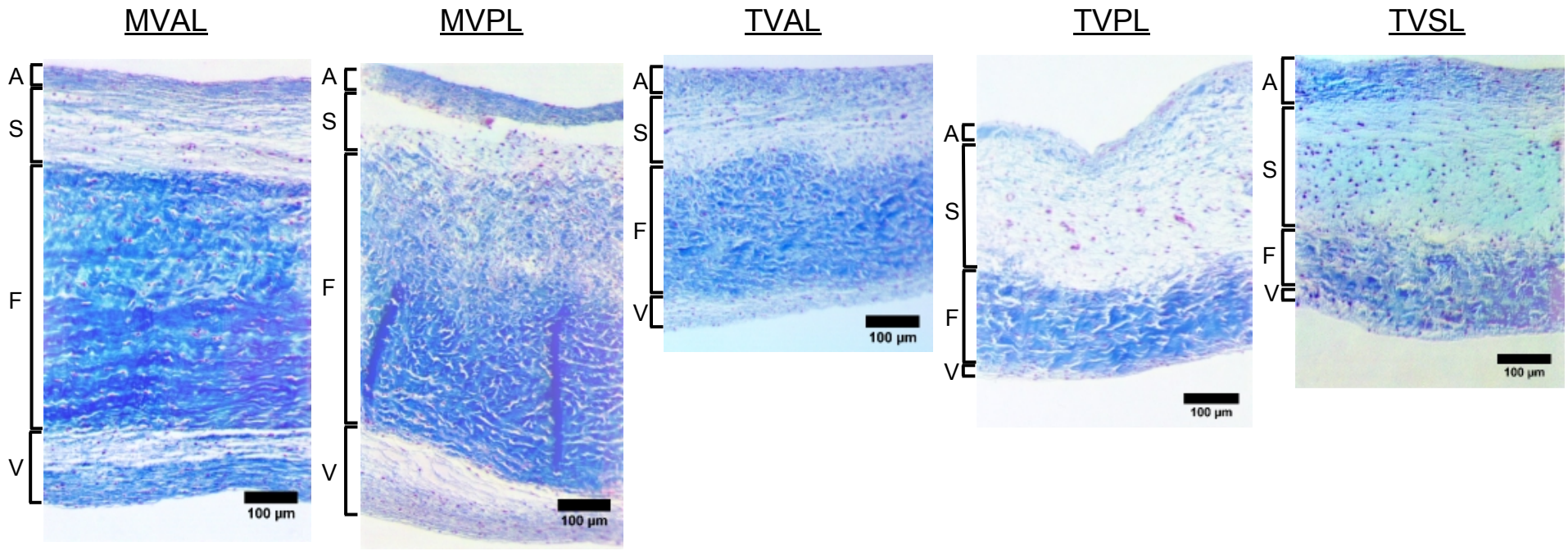
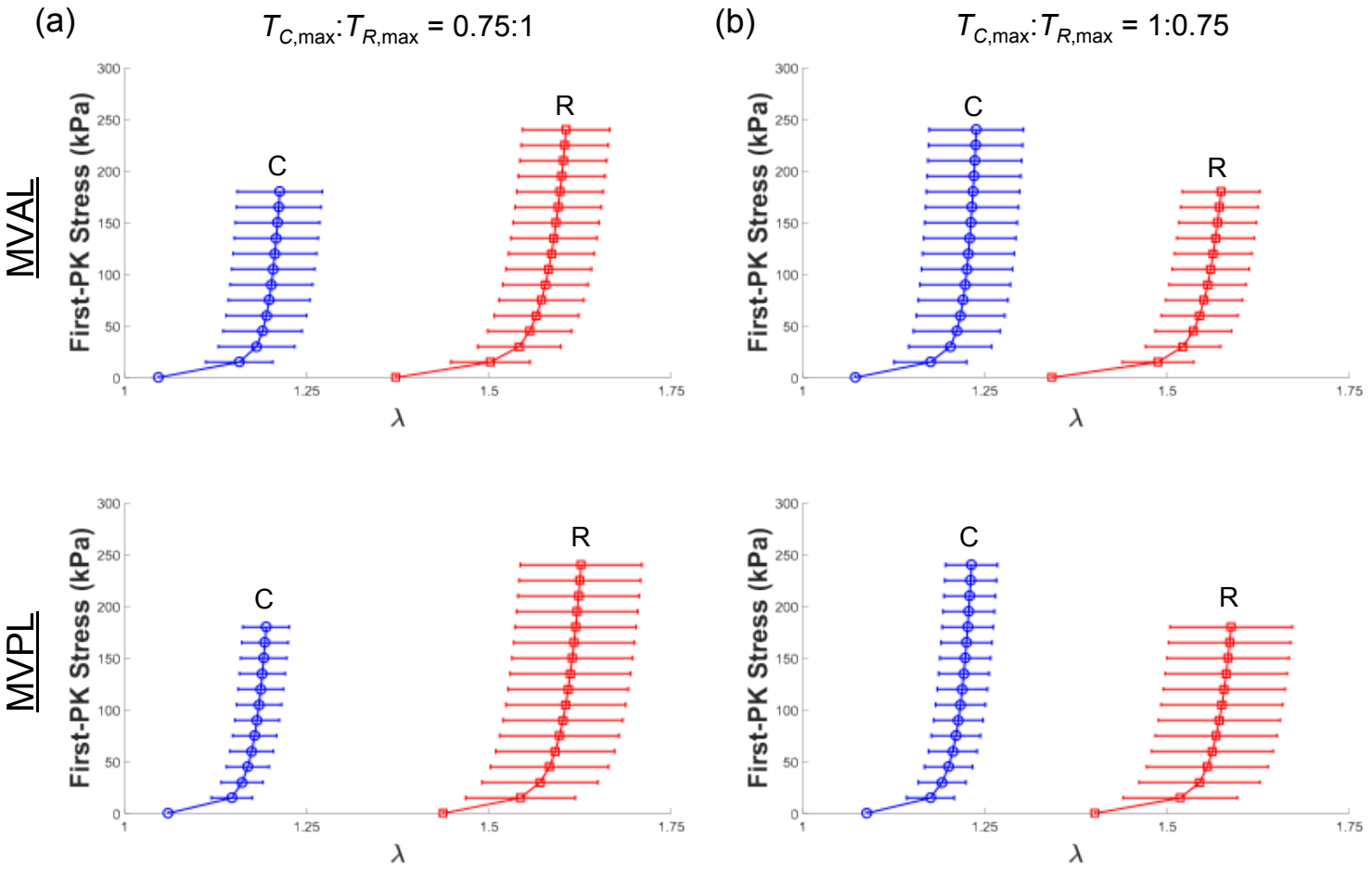

MVAL
TVAL
Porcine
Adult Ovine
Juvenile Ovine
First-PK Stress (kPa)
λ
C
R

Figure 10




(a)
$T_{C,max}$:$T_{R,max}$ = 0.75:1
(b)
$T_{C,max}$:$T_{R,max}$ = 1:0.75
MVAL
MVPL
First-PK Stress (kPa)
λ
C
R
0
50
100
150
200
250
300
1
1.25
1.5
1.75

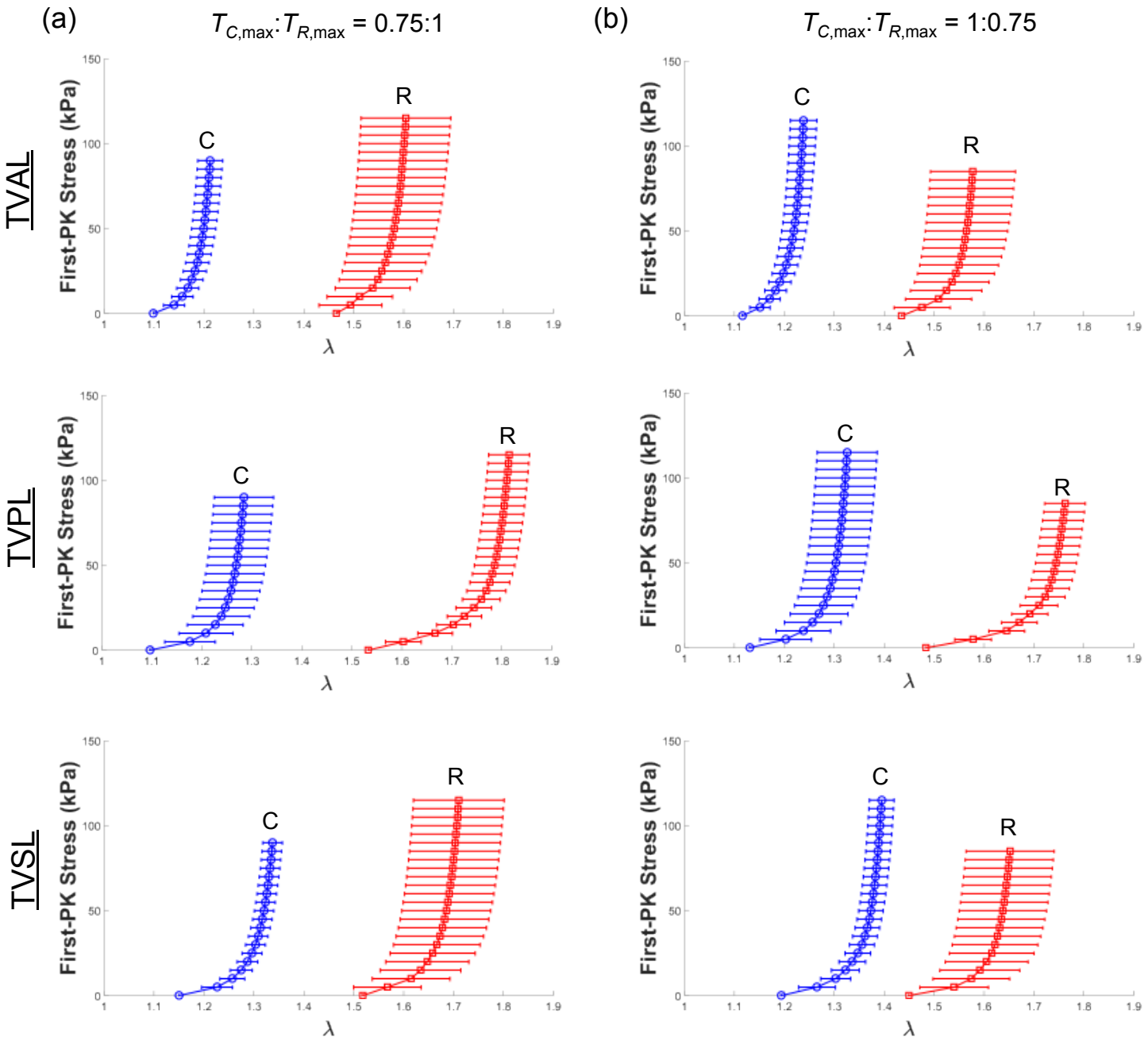
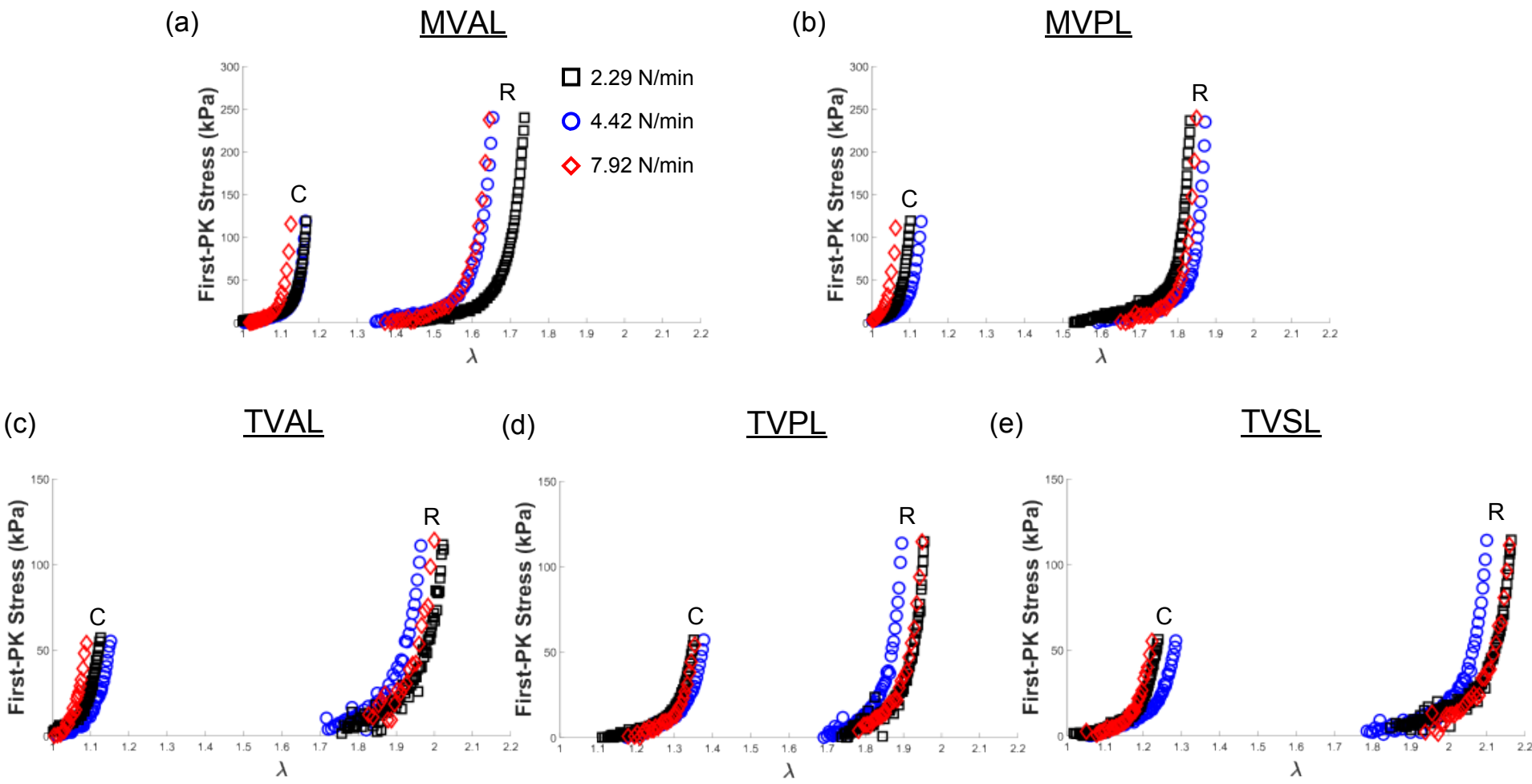
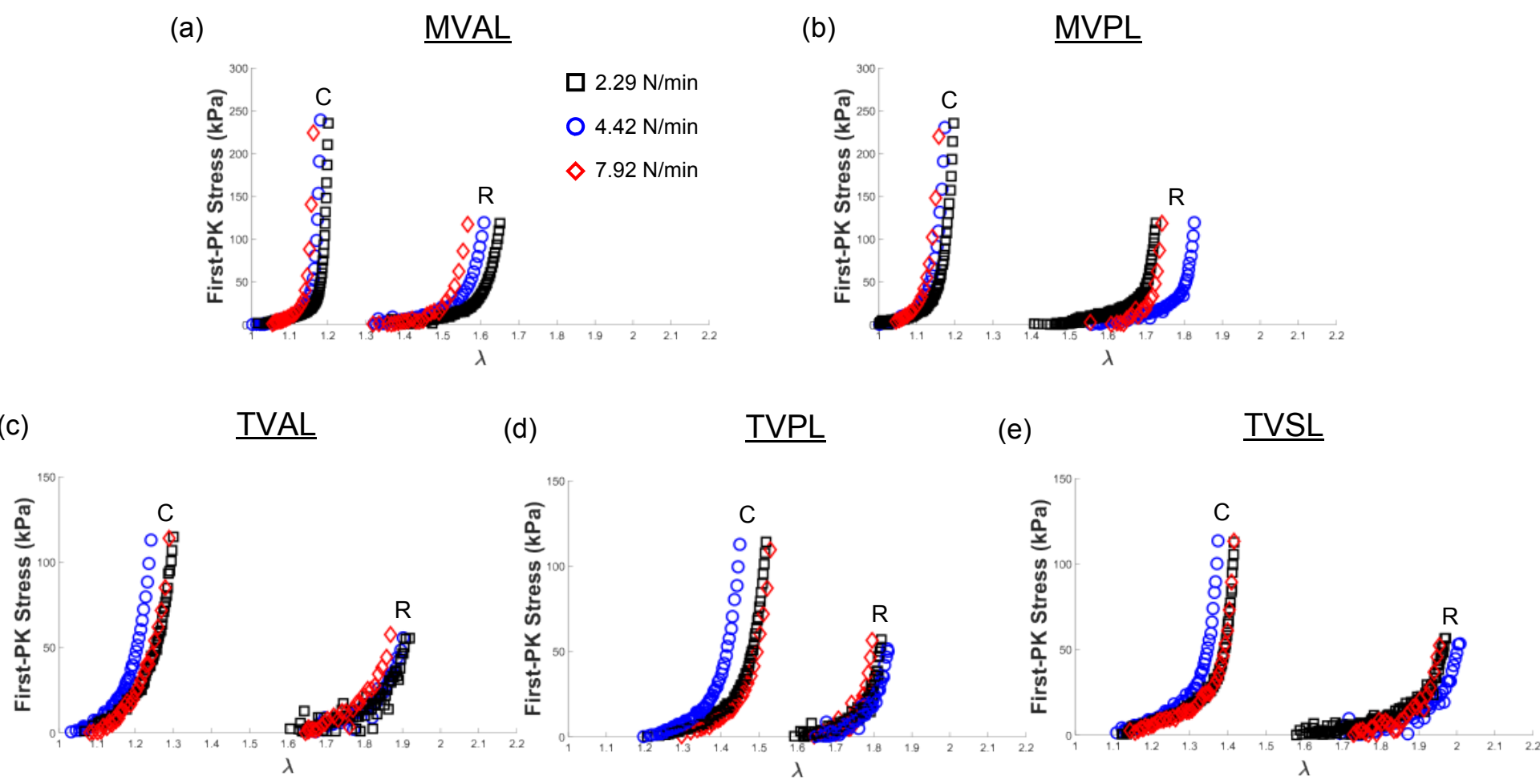
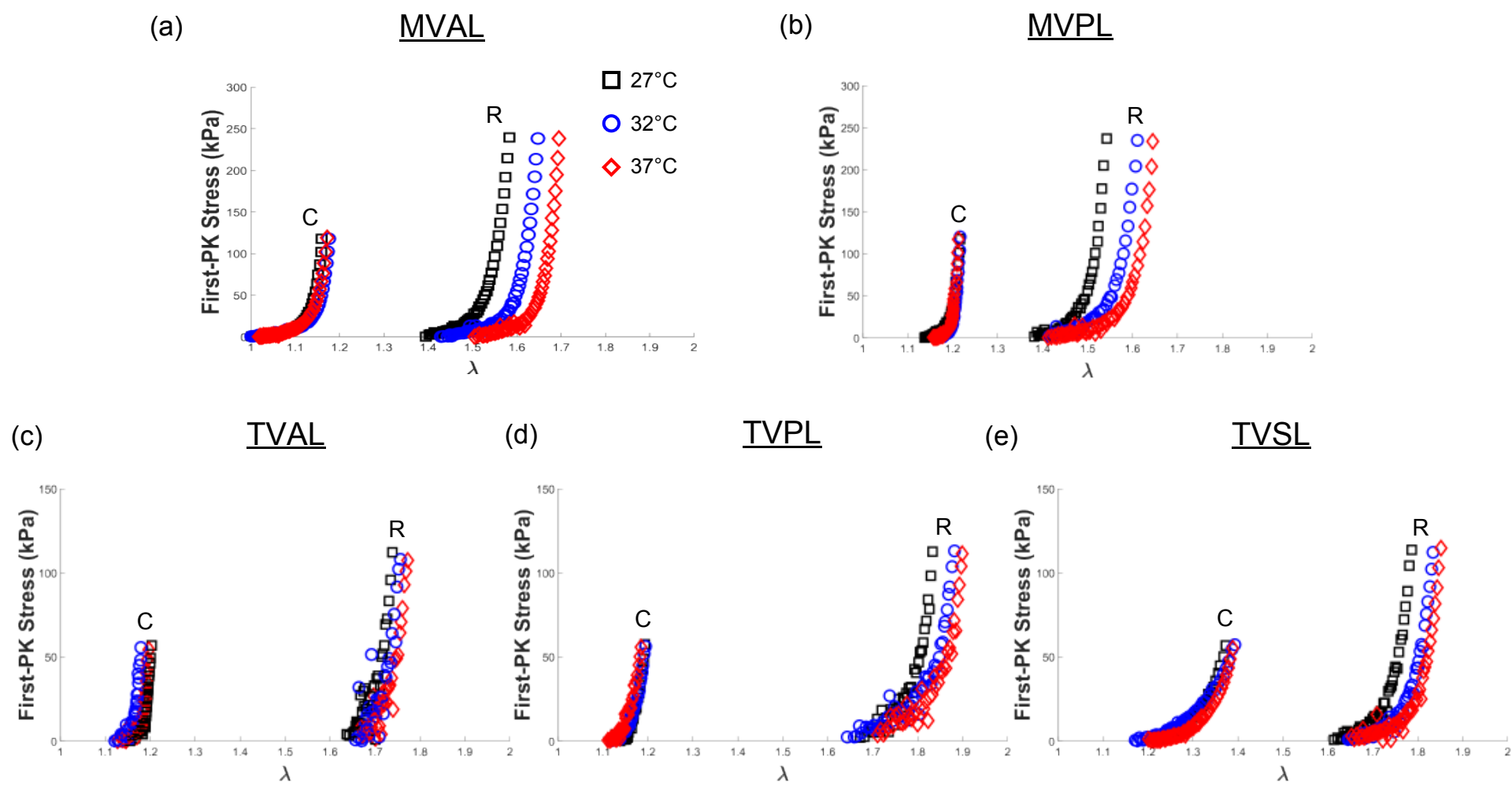
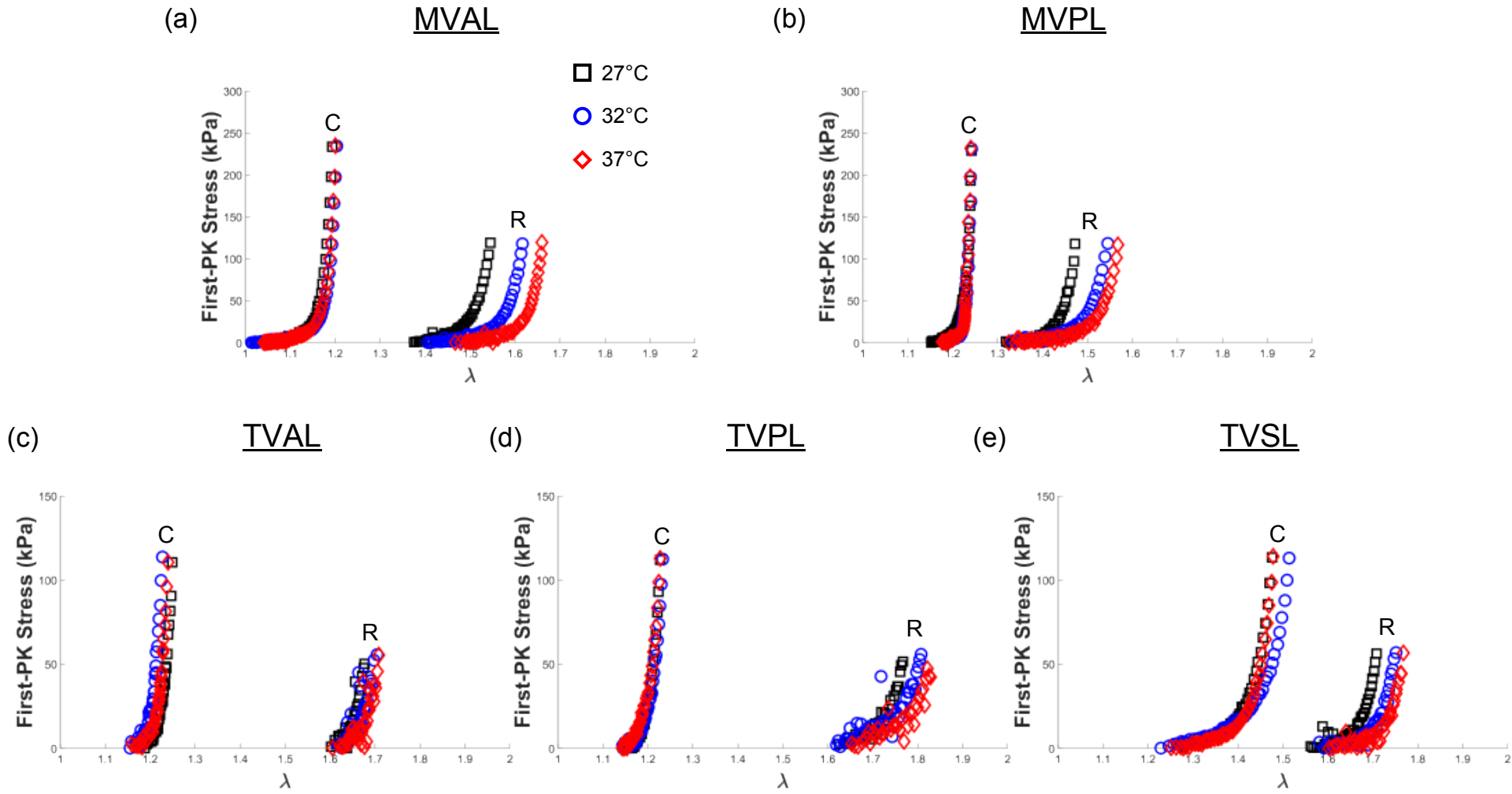

(a)
$T_{C,max}$:$T_{R,max}$ = 0.75:1
(b)
$T_{C,max}$:$T_{R,max}$ = 1:0.75
TVAL
TVPL
TVSL
First-PK Stress (kPa)
λ
C
R

Figure A3


(a)
MVAL
(b)
MVPL
(c)
TVAL
(d)
TVPL
(e)
TVSL
2.29 N/min
4.42 N/min
7.92 N/min
First-PK Stress (kPa)
λ
C
R


(a)
MVAL
(b)
MVPL
(c)
TVAL
(d)
TVPL
(e)
TVSL
2.29 N/min
4.42 N/min
7.92 N/min
First-PK Stress (kPa)
λ
C
R


(a)
MVAL
27°C
32°C
37°C
First-PK Stress (kPa)
C
R
λ
(b)
MVPL
(c)
TVAL
(d)
TVPL
(e)
TVSL


(a)
MVAL
27°C
32°C
37°C
First-PK Stress (kPa)
C
R
λ
(b)
MVPL
First-PK Stress (kPa)
C
R
λ
(c)
TVAL
First-PK Stress (kPa)
C
R
λ
(d)
TVPL
First-PK Stress (kPa)
C
R
λ
(e)
TVSL
First-PK Stress (kPa)
C
R
λ

Supplementary Material

This Supplementary Material section contains the statistical analysis results for the biaxial mechanical testing data presented in the manuscript: "*An investigation of the anisotropic mechanical properties and anatomical structure of porcine atrioventricular heart valves*".

We present statistical analyses for the equibiaxial protocol ($T_{C,\max}$:$T_{R,\max}$ = 1:1) of 3 different experimental groups, as described in **Section 2.3** of the manuscript: (1) the loading rate effect study, (2) the temperature effect group, and (3) the species effect group. We consider tissue responses in the circumferential and radial directions separately in the statistical analysis of each of these studies. For each of these experimental groups, three different stretch measures are examined, as described in detail in **Section 2.7** of the manuscript.

For each leaflet within the loading rate effect group, the statistical analysis of stretch in each direction for each valve leaflet of MVPL, TVAL, TVPL, and TVSL is presented in its respective table and figure (**Tables S1-S4**, **Figs. S1-S4**). Similarly, our statistical analysis results associated with the temperature effect group are presented in **Tables S5-S8** and **Figures 5-8**. As for the species group, the statistical analysis results for the MVAL and TVAL tissues are presented in **Tables S9-S10**, respectively.

Table S1 – Statistical analysis results of the loading rate effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the MVPL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_C^{0-1}	1.046 ± 0.023	1.088 ± 0.035	1.103 ± 0.042	0.349	0.787	0.268
λ_C^{1-peak}	1.165 ± 0.035	1.121 ± 0.019	1.114 ± 0.019	0.295	0.808	0.235
λ_C^{0-peak}	1.218 ± 0.039	1.218 ± 0.037	1.227 ± 0.045	0.999	0.871	0.873
Rad.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_R^{0-1}	1.397 ± 0.081	1.424 ± 0.079	1.480 ± 0.084	0.812	0.639	0.492
λ_R^{1-peak}	1.175 ± 0.017	1.129 ± 0.016	1.101 ± 0.007	0.077	0.159	0.0024
λ_R^{0-peak}	1.638 ± 0.089	1.606 ± 0.090	1.630 ± 0.093	0.807	0.859	0.951

Table S2 – Statistical analysis results of the loading rate effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVAL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_C^{0-1}	1.090 ± 0.038	1.093 ± 0.026	1.091 ± 0.027	0.954	0.966	0.982
λ_C^{1-peak}	1.127 ± 0.031	1.108 ± 0.024	1.105 ± 0.022	0.647	0.929	0.587
λ_C^{0-peak}	1.226 ± 0.041	1.210 ± 0.029	1.206 ± 0.037	0.758	0.936	0.728
Rad.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_R^{0-1}	1.393 ± 0.061	1.435 ± 0.054	1.459 ± 0.052	0.618	0.747	0.424
λ_R^{1-peak}	1.171 ± 0.039	1.147 ± 0.039	1.140 ± 0.038	0.678	0.888	0.576
λ_R^{0-peak}	1.635 ± 0.104	1.650 ± 0.098	1.668 ± 0.101	0.919	0.899	0.823

Table S3 – Statistical analysis results of the loading rate effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVPL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_C^{0-1}	1.150 ± 0.026	1.161 ± 0.030	1.178 ± 0.031	0.780	0.708	0.506
λ_C^{1-peak}	1.165 ± 0.048	1.129 ± 0.048	1.114 ± 0.045	0.610	0.829	0.461
λ_C^{0-peak}	1.337 ± 0.054	1.310 ± 0.066	1.312 ± 0.064	0.763	0.983	0.776
Rad.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_R^{0-1}	1.449 ± 0.050	1.474 ± 0.043	1.523 ± 0.040	0.712	0.433	0.281
λ_R^{1-peak}	1.232 ± 0.027	1.215 ± 0.032	1.190 ± 0.028	0.700	0.567	0.306
λ_R^{0-peak}	1.783 ± 0.057	1.788 ± 0.044	1.810 ± 0.052	0.947	0.746	0.729

Table S4 – Statistical analysis results of the loading rate effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVSL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_C^{0-1}	1.144 ± 0.041	1.166 ± 0.041	1.184 ± 0.044	0.712	0.773	0.521
λ_C^{1-peak}	1.241 ± 0.026	1.195 ± 0.029	1.181 ± 0.023	0.264	0.701	0.115
λ_C^{0-peak}	1.414 ± 0.023	1.388 ± 0.020	1.393 ± 0.027	0.404	0.888	0.549
Rad.	2.29 N/min	4.42 N/min	7.92 N/min	p value (2.29-4.42)	p value (4.42-7.92)	p value (2.29-7.92)
λ_R^{0-1}	1.443 ± 0.076	1.487 ± 0.071	1.525 ± 0.078	0.682	0.725	0.469
λ_R^{1-peak}	1.209 ± 0.029	1.155 ± 0.026	1.146 ± 0.024	0.186	0.796	0.117
λ_R^{0-peak}	1.739 ± 0.081	1.716 ± 0.088	1.745 ± 0.087	0.853	0.823	0.9613

Table S5 – Statistical analysis results of the temperature effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the MVPL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_C^{0-1}	1.168 ± 0.035	1.270 ± 0.052	1.170 ± 0.045	0.134	0.178	0.973
λ_C^{1-peak}	1.089 ± 0.008	1.105 ± 0.016	1.117 ± 0.031	0.370	0.749	0.410
λ_C^{0-peak}	1.273 ± 0.044	1.397 ± 0.068	1.301 ± 0.074	0.154	0.358	0.749
Rad.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_R^{0-1}	1.389 ± 0.072	1.412 ± 0.059	1.375 ± 0.087	0.816	0.733	0.899
λ_R^{1-peak}	1.099 ± 0.072	1.090 ± 0.020	1.127 ± 0.016	0.747	0.175	0.385
λ_R^{0-peak}	1.524 ± 0.068	1.545 ± 0.059	1.558 ± 0.099	0.820	0.908	0.779

Table S6 – Statistical analysis results of the temperature effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVAL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_C^{0-1}	1.192 ± 0.021	1.166 ± 0.023	1.151 ± 0.029	0.421	0.695	0.275
λ_C^{1-peak}	1.107 ± 0.038	1.102 ± 0.018	1.123 ± 0.019	0.739	0.427	0.716
λ_C^{0-peak}	1.321 ± 0.055	1.285 ± 0.039	1.293 ± 0.044	0.610	0.895	0.706
Rad.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_R^{0-1}	1.472 ± 0.084	1.513 ± 0.073	1.417 ± 0.068	0.719	0.360	0.623
λ_R^{1-peak}	1.119 ± 0.084	1.513 ± 0.073	1.417 ± 0.068	0.719	0.360	0.623
λ_R^{0-peak}	1.640 ± 0.080	1.709 ± 0.112	1.603 ± 0.065	0.627	0.435	0.733

Table S7 – Statistical analysis results of the temperature effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVPL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_C^{0-1}	1.141 ± 0.028	1.108 ± 0.027	1.148 ± 0.021	0.421	0.268	0.832
λ_C^{1-peak}	1.068 ± 0.020	1.082 ± 0.022	1.076 ± 0.020	0.636	0.831	0.786
λ_C^{0-peak}	1.219 ± 0.043	1.199 ± 0.033	1.235 ± 0.025	0.711	0.403	0.760
Rad.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_R^{0-1}	1.456 ± 0.065	1.433 ± 0.098	1.515 ± 0.056	0.847	0.474	0.497
λ_R^{1-peak}	1.108 ± 0.013	1.130 ± 0.019	1.123 ± 0.017	0.352	0.791	0.487
λ_R^{0-peak}	1.611 ± 0.067	1.618 ± 0.107	1.701 ± 0.057	0.960	0.507	0.330

Table S8 – Statistical analysis results of the temperature effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVSL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_C^{0-1}	1.227 ± 0.046	1.201 ± 0.039	1.804 ± 0.049	0.670	0.753	0.505
λ_C^{1-peak}	1.079 ± 0.028	1.104 ± 0.043	1.105 ± 0.035	0.630	0.987	0.569
λ_C^{0-peak}	1.326 ± 0.066	1.323 ± 0.054	1.305 ± 0.065	0.979	0.829	0.825
Rad.	27 °C	32 °C	37 °C	p value (27-32)	p value (32-37)	p value (27-37)
λ_R^{0-1}	1.462 ± 0.075	1.475 ± 0.080	1.486 ± 0.078	0.907	0.924	0.829
λ_R^{1-peak}	1.087 ± 0.017	1.101 ± 0.016	1.093 ± 0.021	0.550	0.770	0.820
λ_R^{0-peak}	1.589 ± 0.087	1.625 ± 0.096	1.629 ± 0.107	0.787	0.978	0.777

Table S9 – Statistical analysis results of the species effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the MVAL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	Porcine (P)	Adult Ovine (AO)	Juvenile Ovine (JO)	p value (P-AO)	p value (AO-JO)	p value (P-JO)
λ_C^{0-1}	1.086 ± 0.032	1.149 ± 0.057	1.196 ± 0.087	0.355	0.662	0.262
λ_C^{1-peak}	1.088 ± 0.019	1.113 ± 0.014	1.073 ± 0.075	0.314	0.614	0.854
λ_C^{0-peak}	1.181 ± 0.042	1.280 ± 0.069	1.270 ± 0.094	0.248	0.929	0.409
Rad.	Porcine (P)	Adult Ovine (AO)	Juvenile Ovine (JO)	p value (P-AO)	p value (AO-JO)	p value (P-JO)
λ_R^{0-1}	1.399 ± 0.053	1.443 ± 0.105	1.449 ± 0.102	0.717	0.967	0.372
λ_R^{1-peak}	1.118 ± 0.017	1.146 ± 0.031	1.074 ± 0.012	0.446	0.052	0.059
λ_R^{0-peak}	1.564 ± 0.061	1.654 ± 0.123	1.552 ± 0.104	0.527	0.542	0.925

Table S10 – Statistical analysis results of the species effect group on the preconditioning stretches (λ_C^{0-1} and λ_R^{0-1}), mechanical stretches (λ_C^{1-peak} and λ_R^{1-peak}), and peak stretches (λ_C^{0-peak} and λ_R^{0-peak}) of the TVAL tissue (n=6). All quantities are presented as mean ± SEM.

Circ.	Porcine (P)	Adult Ovine (AO)	Juvenile Ovine (JO)	p value (P-AO)	p value (AO-JO)	p value (P-JO)
λ_C^{0-1}	1.151 ± 0.029	1.379 ± 0.084	1.463 ± 0.095	0.028	0.522	0.010
λ_C^{1-peak}	1.123 ± 0.019	1.071 ± 0.016	1.023 ± 0.007	0.065	0.021	0.0005
λ_C^{0-peak}	1.293 ± 0.044	1.476 ± 0.044	1.495 ± 0.096	0.090	0.880	0.084
Rad.	Porcine (P)	Adult Ovine (AO)	Juvenile Ovine (JO)	p value (P-AO)	p value (AO-JO)	p value (P-JO)
λ_R^{0-1}	1.417 ± 0.068	1.601 ± 0.052	1.657 ± 0.076	0.057	0.555	0.040
λ_R^{1-peak}	1.137 ± 0.029	1.073 ± 0.014	1.038 ± 0.018	0.077	0.147	0.016
λ_R^{0-peak}	1.606 ± 0.064	1.716 ± 0.046	1.716 ± 0.061	0.181	0.999	0.239

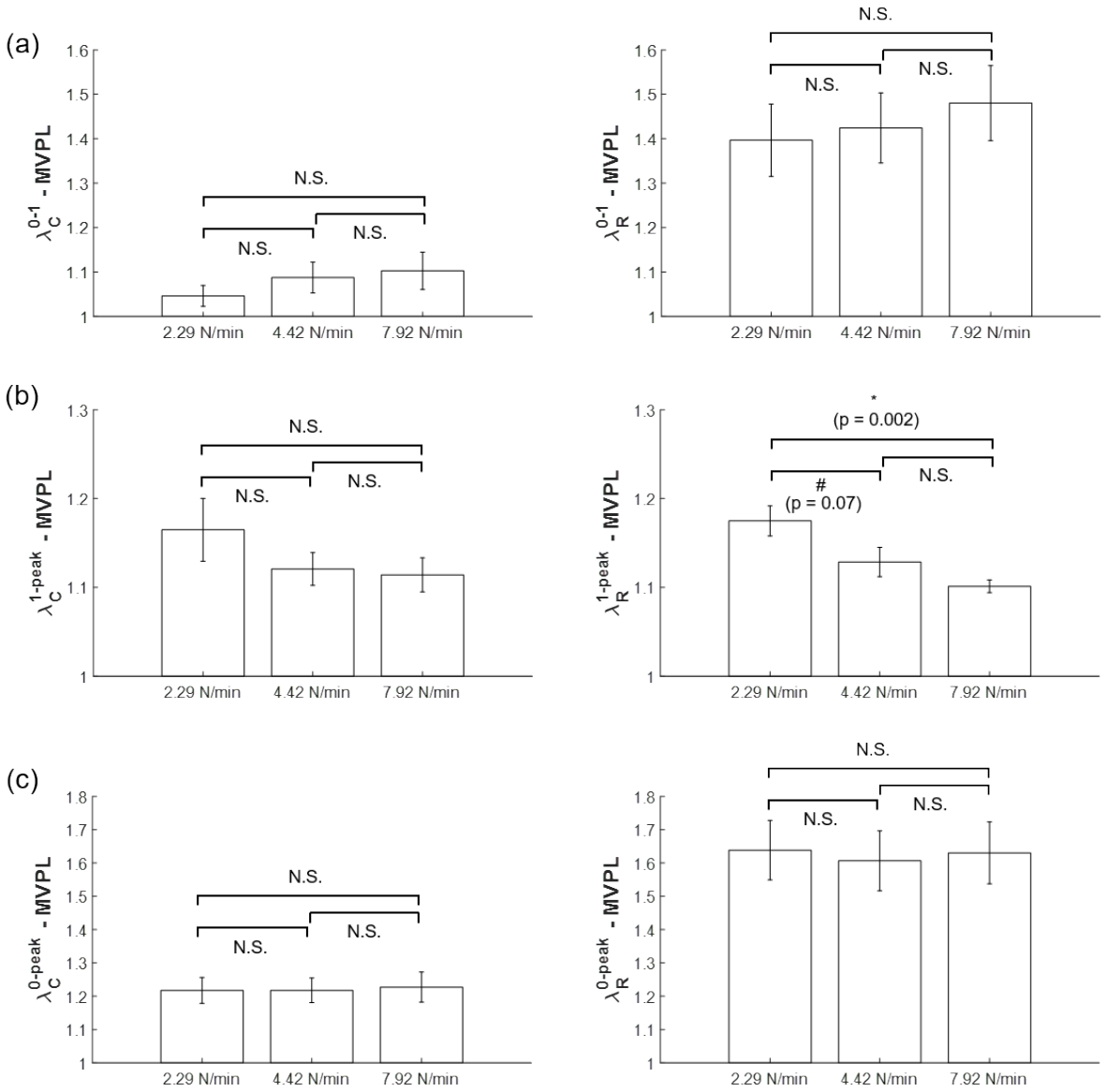


Figure S1 – Statistical analyses of the MVPL from the loading rate effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, $p>0.10$, **#**: nearly statistically significant difference, $p<0.10$, and *****: statistically significant difference, $p<0.05$)

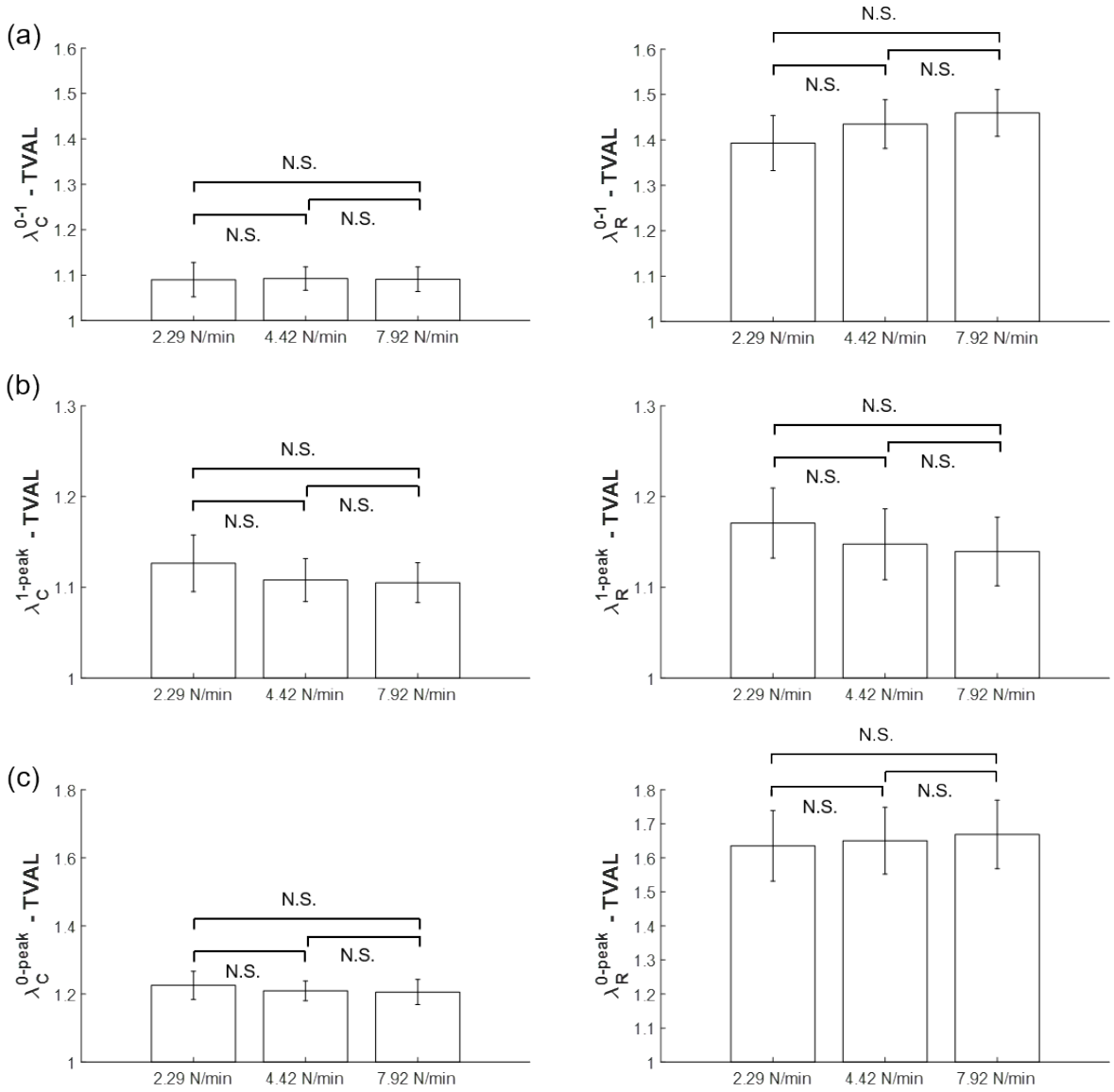


Figure S2 – Statistical analyses of the TVAL from the loading rate effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, $p>0.10$)

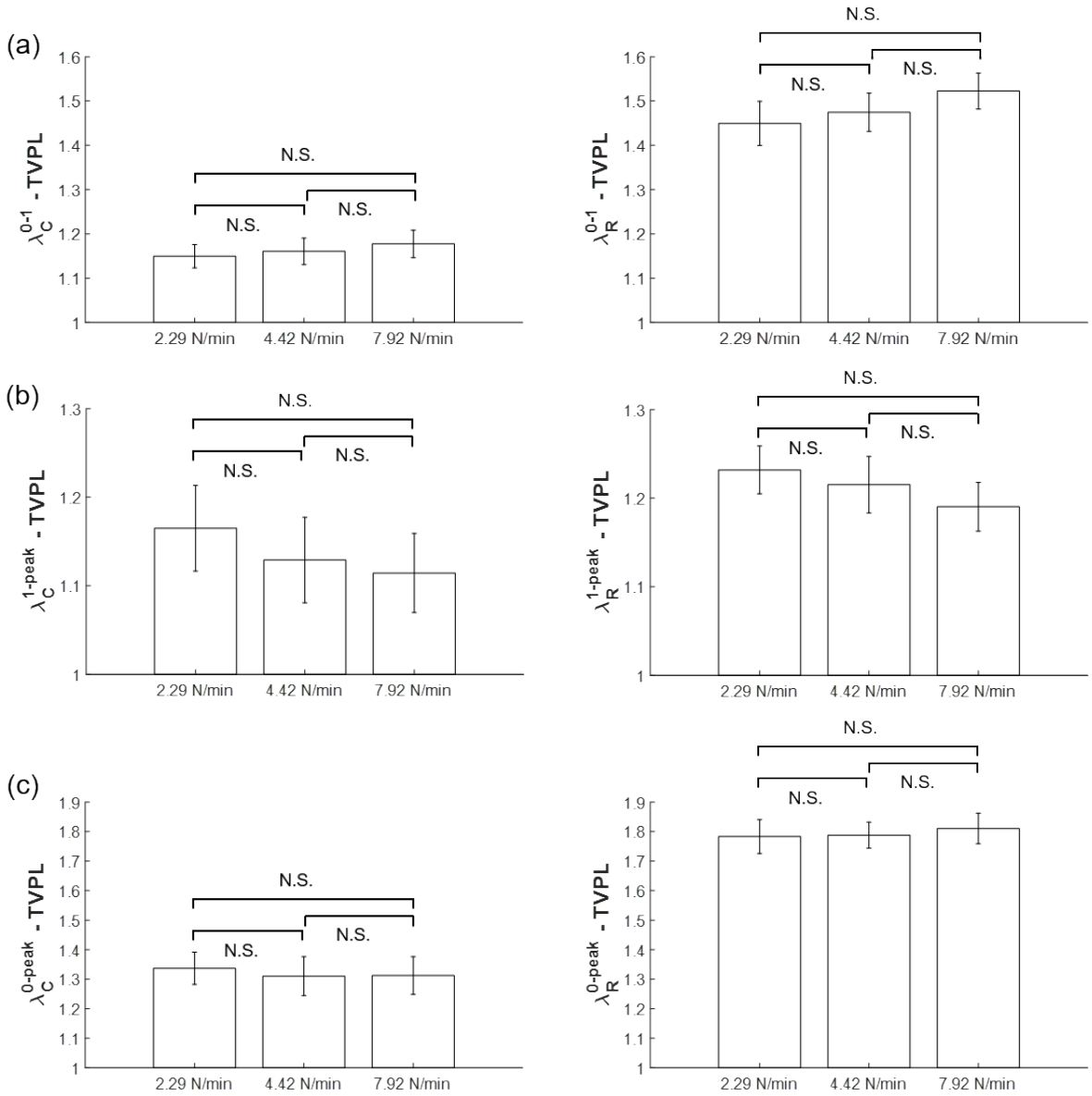


Figure S3 – Statistical analyses of the TVPL from the loading rate effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, p>0.10)

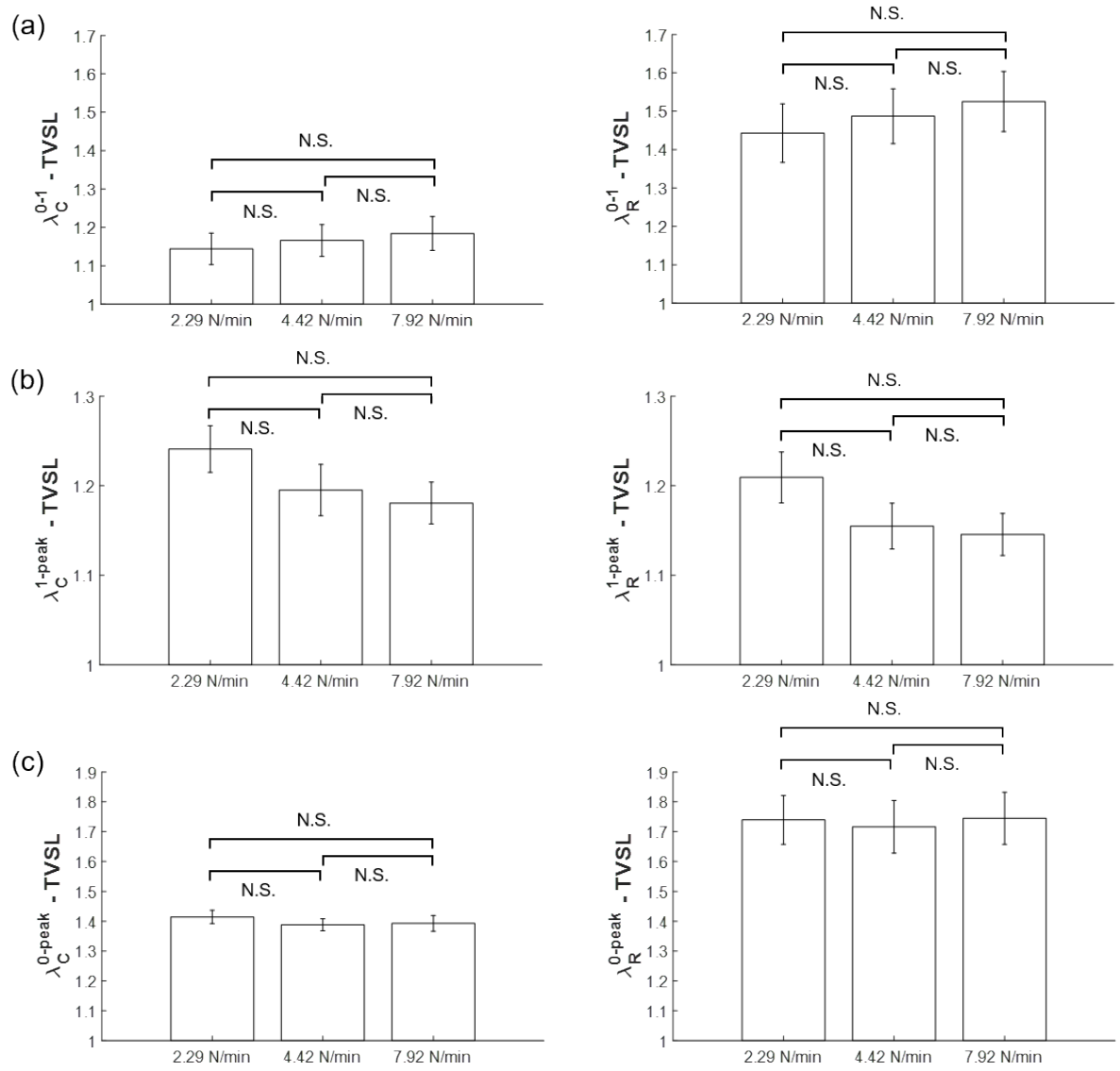


Figure S4 – Statistical analyses of the TVSL from the loading rate effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, $p>0.10$)

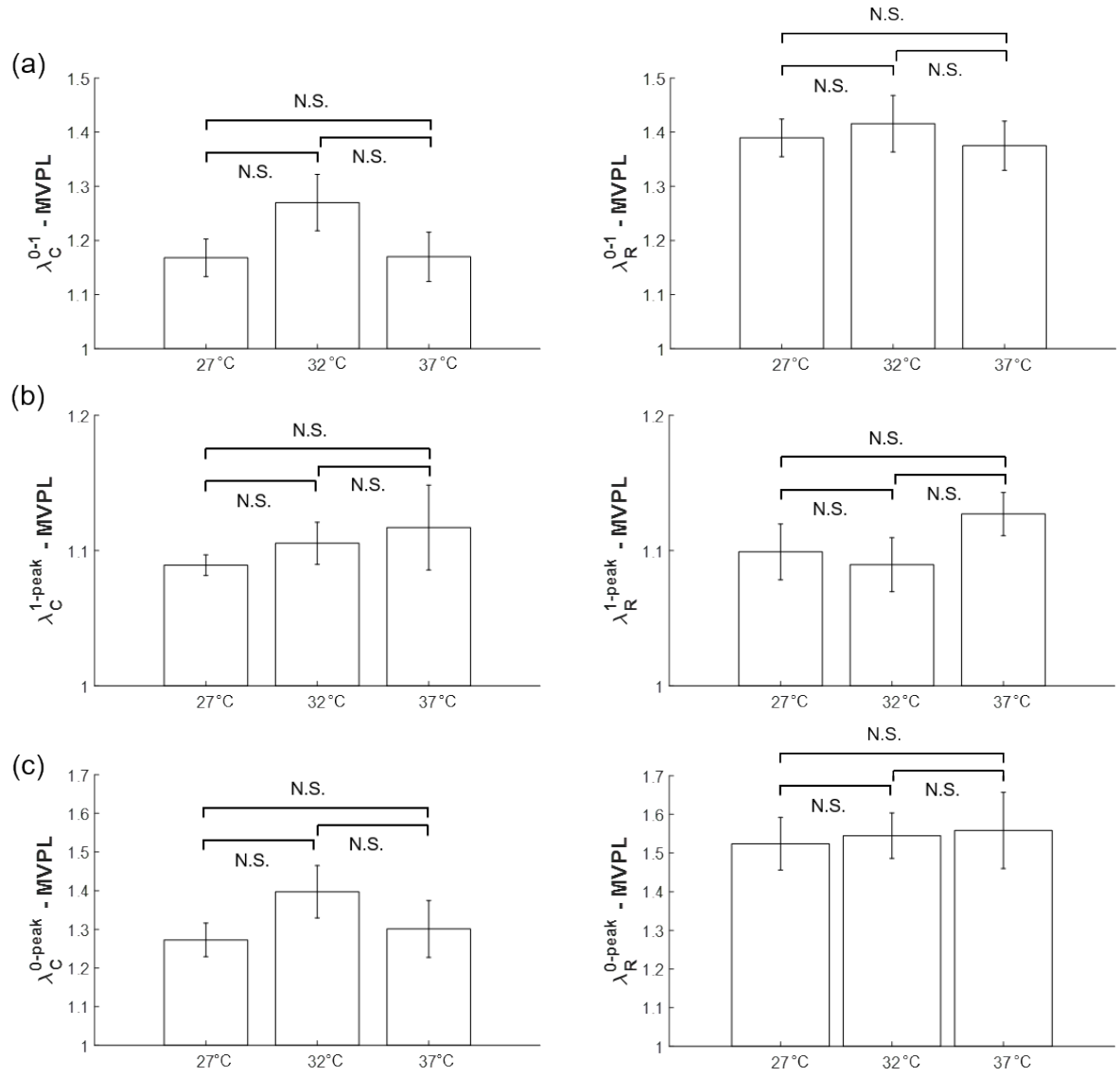


Figure S5 – Statistical analyses of the MVPL from the temperature effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, $p>0.10$)

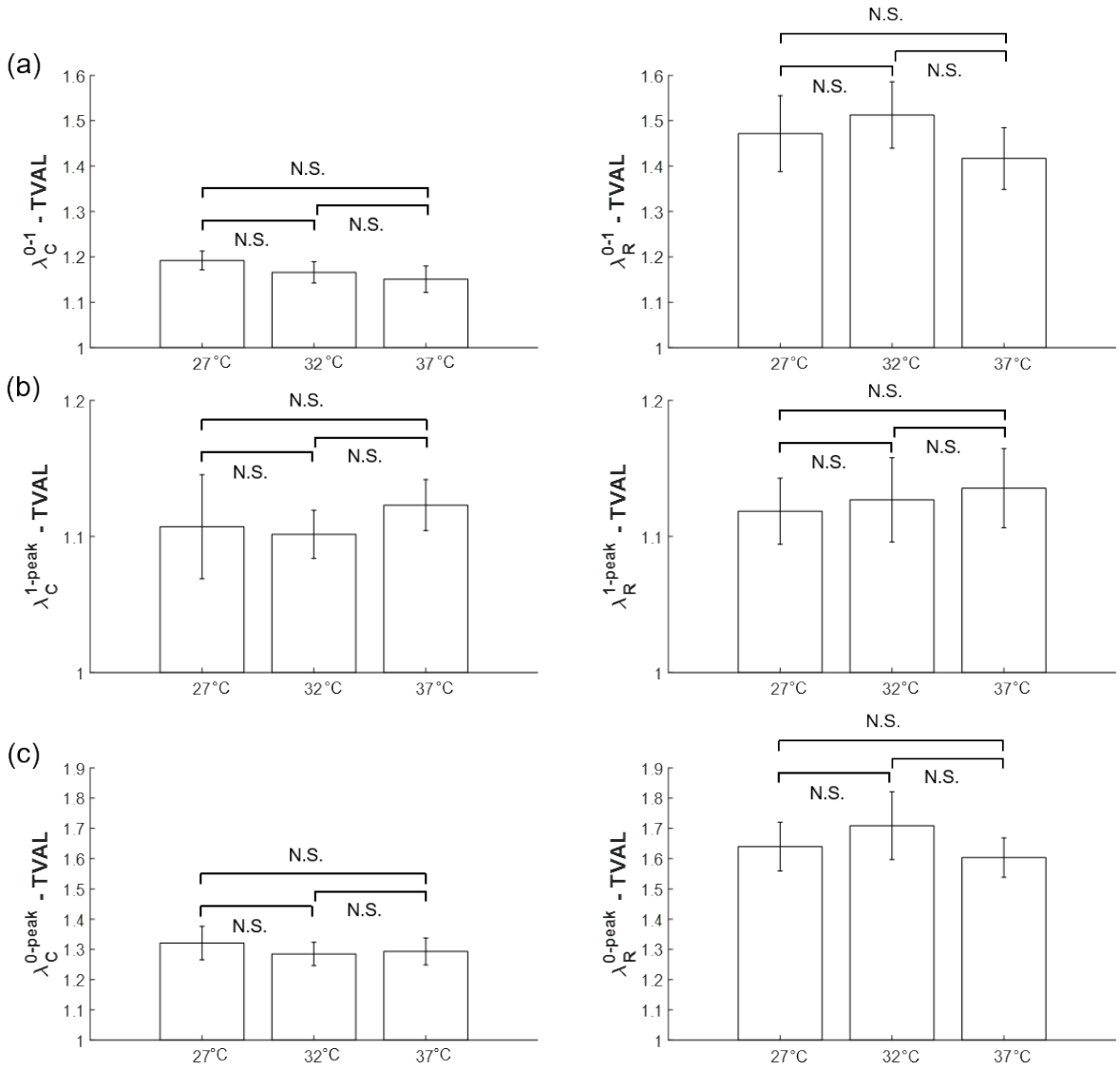


Figure S6 – Statistical analyses of the TVAL from the temperature effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, p>0.10)

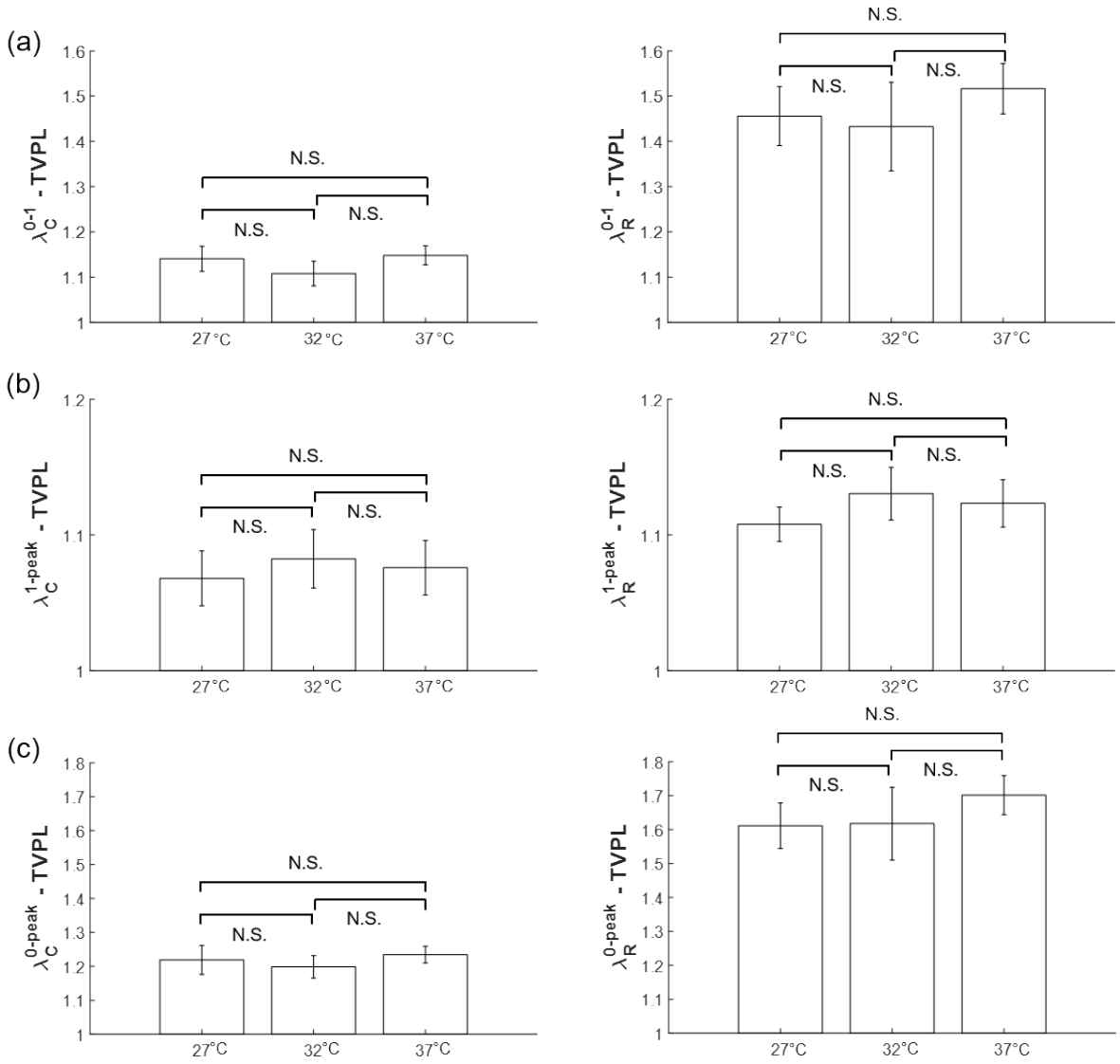


Figure S7 – Statistical analyses of the TVPL from the temperature effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, $p>0.10$)

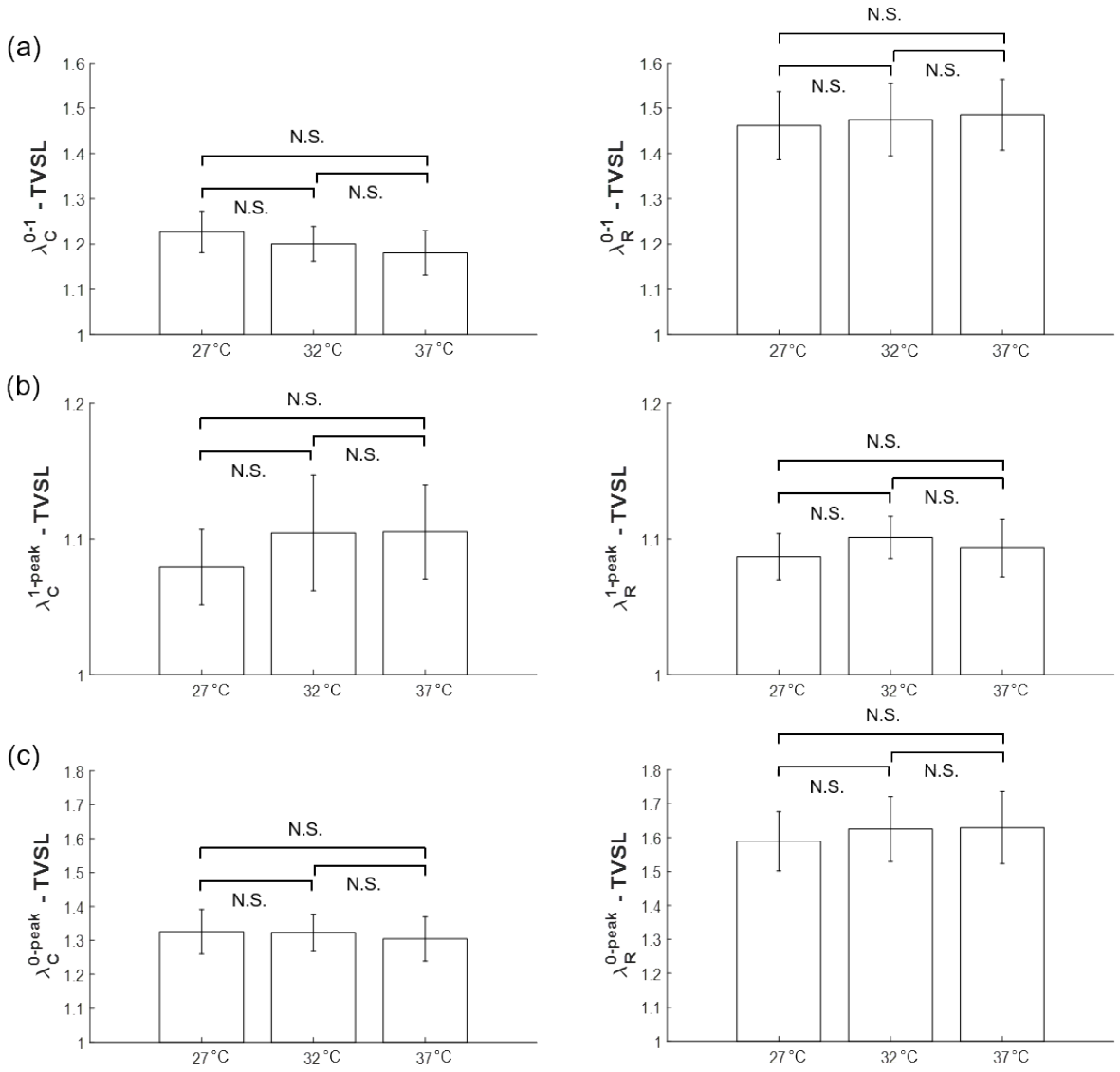


Figure S8 – Statistical analyses of the TVSL from the temperature effect group (n=6), with plots showing trends in (a) preconditioning stretches, (b) mechanical stretches, and (c) peak stretches. All bars show mean ± SEM. (N.S.: no statistically significant difference, p>0.10)