

A Study on the Spatial Variance in the Mechanics and Microstructure of Heart Valves

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Valvular Heart Disease Background

• Valvular heart diseases (VHD), including regurgitation and stenosis, impair proper valve function and affect 2.5% of adults^[1] (Fig. 1b). VHD can accompany alterations to the *mechanics* and *structure* of the heart valve.



Research Objectives

• While previous studies examined the center of the *mitral and tricuspid* valve (MV and TV) leaflets (Fig. 2a), our objective is to quantify the spatial variance in mechanics and microstructure of the MV and TV leaflets to inform *computational models* and *material design* for replacement valves.



Courtesy of Gorman Research Group (Upenn)

Figure 1. (a) Schematic of a heart with chambers, valves, and major arteries labelled. (b) Echocardiogram showing mitral valve *regurgitation*, i.e., backflow from atrium to ventricle.

Figure 2. (a) Schematic of a TV leaflet, showing principal directions. (b) Biaxial testing of the central region for quantification of (c) *anisotropic* and *nonlinear* mechanics of the TV leaflets.

Biaxial Mechanical Testing to Quantify Regional Variance in Leaflet Mechanics



Figure 3. (a) Dissected and labelled TV. (b) Schematic of a leaflet showing regional specimen locations. (c) Specimen mounted on the *biaxial tester* with markers for optical strain tracking. (d) Representative mechanical responses for the TVAL.

- In each region of the MV and TV anterior leaflets (MVAL and TVAL), the tissues were more compliant in the Rad than the Circ direction (**Fig. 4**).
- the TVAL, we observed similar In mechanics between regions **B**, **D**, and **E**, and between regions **A** and **C**, but unique mechanics in **F** (**Fig. 4a**).

In the MVAL, we found increased Rad compliance and Circ rigidity in **B** and **E**



compared to A, C, D, and F (Fig. 4b).

applied *membrane tensions* in Rad and Circ directions for all regions of the (a) TVAL and (b) MVAL.

Analysis of Heart Valve Microstructure

Our group developed a polarized Spatial Frequency Domain Imaging (pSFDI) system, (Fig. 5a) based on structural birefringence (Fig. 5b), to quantify *local collagen fiber orientations* (Fig. 5c) in heart valve tissue.



Study Applications

- Our mechanical analysis will allow development of *spatially-varied* leaflet constitutive models (Fig. 6a) for use in computational models (Fig. 6b).
- Our results could eventually improve understanding of the *complex relationship* between tissue microstructure and mechanical behavior.



Figure 6. (a) The development of a spatially-varied material model to describe leaflet behavior for use in computational models of the MV and (b) **TV closure** and valvular function.

Future Research

The microstructural quantification from this study could inform design of *bioreactor protocols* for *tissue engineering* biomimetic valves (Fig. 7).

Figure 5. (a) Schematic of the PSFDI system showing major components. (b) Schematic showing the polarized *light scattering* off *structurally birefringent* collagen fibers in valve tissue. (c) Original and analyzed images of **system validation testing** on bovine tendon.

• **Goal** – Improve system *resolution* to provide accurate analysis of local fiber

orientations in tissues with *spatially-heterogeneous* fiber networks.



Figure 7. (a) Spatially-varied collagen fiber networks in heart valve leaflets. (b) Local fiber orientation informs loading protocols to engineer (c) *high-fidelity replacement* heart valves.



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