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Ambroise Duprey, Olfa Trabelsi, Marco Vola, Jean-Pierre Favre, Stéphane Avril. Biaxial rupture properties of ascending thoracic aortic aneurysms. Acta Biomaterialia, 2016. hal-01380216

HAL Id: hal-01380216 https://hal.science/hal-01380216v1

Submitted on 12 Oct 2016

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Accepted Manuscript

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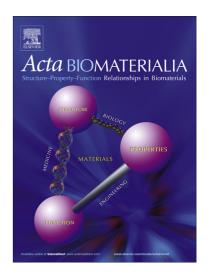
PII: S1742-7061(16)30305-1

DOI: http://dx.doi.org/10.1016/j.actbio.2016.06.028

Reference: ACTBIO 4300

To appear in: Acta Biomaterialia

Received Date: 25 February 2016 Revised Date: 11 May 2016 Accepted Date: 22 June 2016



Please cite this article as: Duprey, A., Trabelsi, O., Vola, M., Favre, J-P., Avril, S., Biaxial rupture properties of ascending thoracic aortic aneurysms, *Acta Biomaterialia* (2016), doi: http://dx.doi.org/10.1016/j.actbio. 2016.06.028

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Biaxial rupture properties of ascending thoracic aortic aneurysms

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ABSTRACT

Although hundreds of samples obtained from ascending thoracic aortic aneurysms (ATAA) of patients undergoing elective surgical repair have already been characterized biomechanically, their rupture properties were always derived from uniaxial tensile tests. Due to their bulge shape, ATAAs are stretched biaxially in vivo. In order to understand the biaxial rupture of ATAAs, our group developed a novel methodology based on bulge inflation and full-field optical measurements. The objective of the current paper is threefold. Firstly, we will review the failure properties (maximum stress, maximum stretch) obtained by bulge inflation testing on a cohort of 31 patients and compare them with failure properties obtained by uniaxial tension in a previously published study. Secondly, we will investigate the relationship between the failure properties and the age of patients, showing that patients below 55 years of age display significantly higher strength. Thirdly, we will define a rupture risk based on the extensibility of the tissue and we will show that this rupture risk is strongly correlated with the physiological elastic modulus of the tissue independently of the age, ATAA diameter or the aortic valve phenotype of the patient.

KEYWORDS: aneurysm; human thoracic aorta; bulge inflation test; rupture properties; extensibility.

1. INTRODUCTION

Ascending thoracic aortic aneurysms (ATAA), described at tissue level by medial degeneration and biomechanical weakening of the aneurysmal wall [1,2], are a life-threatening pathology causing a permanent dilation associated with a high risk of aortic rupture or dissection and death of the patient. ATAA affects approximately 10 out of 100 000 persons per year [3] and is the 17th most common cause of death [4].

ATAAs are treated by replacing them with synthetic grafts when the aortic diameter exceeds 5.5 cm [5]. However the rupture or the dissection of ATAAs remains rather unpredictable on a patient-specific basis. Even if the yearly risk of dissection or rupture rises from 3% to 7% with aneurysms >6 cm, rupture of ATAAs has been documented to occur at diameters less than 4.5 cm [6]. Factors other than the aneurysm diameter that may affect the predisposition to rupture are age, hypertension, aortic valve phenotype (bicuspid or tricuspid) or the presence of genetic disorders (Marfan syndrome, Ehler-Danlos syndrome).

Rupture of an ATAA occurs when the stress applied to the aortic wall locally exceeds its capacity to sustain stress. Finite-element analyses can be used to estimate the local distribution of the stress applied by the blood pressure onto the aortic wall [7–9]. An open question is to estimate the patient-specific strength, which can vary from a few tenths of MPa to a few units of MPa from one individual to another [10,11]. Another open question is that mean physiological wall stresses acting on pathologic aortas were found to be far from rupture, with factors of safety (defined as the ratio of tensile strength to the mean wall stress) larger than six [12].

Another equivalent definition of rupture can be stated when the stretch applied to the tissue exceeds its extensibility. For instance, Martin et al. [13] defined a diameter risk, which is the ratio between the current diameter of the aneurysm and the rupture diameter. The authors show that the diameter risk increases significantly with the physiological elastic modulus of the artery. Indeed, if the aortic wall is stiff, a rather large increase of pressure can be induced by a small increase of volume.

In vitro mechanical characterization of ATAA rupture on specimens, obtained from ascending aorta tissue of patients undergoing elective surgical repair, has been instrumental to improve the understanding of the aortic wall properties and the potential risk for rupture of ATAAs [14]. Several authors have focused on elastic properties of ATAAs [9,15–17]. They showed that ATAAs have altered collagen fiber architecture in the medial plane and have significantly less elastin than healthy tissue. The tissue is generally stiffer in the circumferential direction than in the longitudinal direction and the elastic properties vary regionally [16]. Stress-strain analyses also aided in determining material properties [18–20]. Kim and Baek showed significant spatial variations of mechanical properties in a porcine thoracic aorta using optical full-field measurements and inflation-extension tests [21].

Despite the interest of these studies on elastic properties, we will focus on the rupture properties of ATAAs. At least 11 papers reported rupture properties of ascending thoracic aortas and ATAAs in uniaxial tension (Table 1). In these studies, the strength was always determined in uniaxial tension, except for a few samples that were reported in the second paper of Mohan and Melvin where they performed bulge inflation tests [22].

Due to their bulge shape, aneurysm tissues are stretched biaxially in vivo [23], and there is no guarantee that results obtained from a uniaxial test can be applied to a biaxial state. The lack of rupture analysis in biaxial tension can be explained by the fact that biaxial testing is less suitable for strength assessment, due to attachment techniques and the square shape of the specimen, which may prevent failure of the specimen in the gauge area.

Bulge inflation test is another technique for investigation of biaxial mechanical behavior, which has been applied in the past to the study of human and pig aortas [22,24] as well as other soft tissues [25]. The basic testing protocol consists in obtaining square specimens from an excised cylindrical aortic tissue laid flat. These are clamped in the inflation device forming a hermetically sealed cavity in which a fluid (water) is injected at a controllable rate while simultaneously measuring pressure.

Our group recently conducted bulge inflation tests on human ATAA tissues [26–28], to test media/adventitia samples as well as complete layers. Stereo-digital image correlation (SDIC) was

used to obtain the strain field of the entire inflated membrane. Video-based tracking techniques can allow simultaneous evaluation of material parameters for constitutive modeling purposes and of localized stress in the area that eventually ruptures, providing a powerful experimental tool to characterize the mechanical properties of vascular tissues [17,26,29–32].

The objective of the current paper is threefold. Firstly, we will review the failure properties (maximum stress, maximum stretch) obtained by bulge inflation testing on a cohort of 31 patients undergoing elective surgical repair and compare them with failure properties obtained by uniaxial tension in a previously published study. Secondly, we will investigate the relationship between the failure properties and the age of patients. Thirdly, we will define a rupture risk based on the brittleness of the tissue and investigate the correlation between this rupture risk and the physiological elastic modulus of the tissue.

2. METHODS

2.1. Materials for bulge inflation tests

ATAA samples for bulge inflation tests were obtained from patients undergoing elective surgery to replace the pathological segment of aorta with a synthetic graft. The collection of the aortic tissues was carried out in accordance with the guidelines of the Institutional Review Board of the University Hospital Center of Saint-Etienne and patients signed informed consent before surgery. Specimens were kept at +4°C in a 0.9% physiological saline solution and tests were performed within 24 hours [33]. The demographic information for all the aortic specimens used in this study is presented in Table 2.

The aortic samples (Fig. 1a) were cut in the axial direction (in the direction of blood flow) at the level of the inner curvature (Fig. 1b). Then a flat square piece was cut consisting of the tunica intima, media, adventitia and loose connective tissue (Fig. 1c). For all the specimens the loose connective tissue was carefully removed to prevent sliding in the experimental device using dissecting blunt-point scissors. The thickness of each specimen, denoted h₀, was measured using a digital caliper

while holding the specimen between two plates. All the aortic tissues were tested in bulge inflation. The entire process was recorded using a SDIC system to reconstruct the 3D shape of the inflated tissues. The experimental set-up and the bulge inflation tests performed by our group are explained in details in previous publications [27,28,34]. The aim of the present study is to review the failure properties (maximum stress, maximum stretch) obtained by the bulge inflation test and compare them with failure properties obtained by uniaxial testing.

2.2. Materials for uniaxial tensile tests

We carried out tensile tests on 103 ATAA samples collected from 12 patients at the University of Michigan in 2008. Elastic properties were reported in a previous publication [16] but the rupture properties were not reported. They will be reported herein and compared to the biaxial rupture properties.

All procedures for the uniaxial tensile tests were carried out in accordance with the guidelines of the Institutional Review Board of the University of Michigan and patients signed informed consent. A segment of aortic wall was excised from surgical specimens obtained from patients undergoing elective surgical repair of their ATAA. These segments were cut with custom-designed tissue cutters and multiple test specimens were available when the surgical aortic specimen was large enough. About half of the specimens were cut along the circumferential direction of the aorta and the other half along the axial direction, for the sake of comparison.

Experiments were carried out using a tensile testing machine (Instron model 5542, Norwood, MA, USA). To avoid damaging the tissue from excessive compression, fine grit adhesive sand paper was placed on the surface of the pneumatic grips to prevent slippage during the test. The tissue was kept wet by a spray of phosphate buffering solution. The specimen was stretched at a rate of 10 mm.min⁻¹ until failure. Load and cross-head displacement were continuously recorded by the data acquisition software Merlin, provided by Instron.

2.3. Analysis method in bulge inflation tests

The 3D shape of the inflated ATAA layer captured in vitro, was numerically discretized in order to create a mesh across the 3D shape [27]. Using a finite difference method, the membrane deformation gradient was calculated based on the deformed and initial tangent vectors at each element of the mesh, and the Cauchy-Green stretch tensor was derived from the deformation gradient [27].

Assuming that the tested samples behave as a pure membrane, it is also possible to derive the Cauchy stress at each element of the mesh [27,34,35].

Let $(\mathbf{e}_1, \ \mathbf{e}_2)$ denote the cartesian basis of the circumferential and axial directions of the tissue, respectively. In the current paper, where we analyze experiments for which the rupture occurred in the central region of the specimen, we derived the following data for each sample:

- 1. λ_1 and λ_2 are the circumferential and axial components of the stretch at the center of the specimen, respectively.
- 2. σ_1 and σ_2 are the circumferential and axial components of the Cauchy stress tensor at the center of the specimen, respectively.

For each pressure stage P(t), $\lambda_1(t)$, $\lambda_2(t)$, $\sigma_1(t)$ and $\sigma_2(t)$ were derived as the average value of the respective stretch or stress fields across a region of 2 cm radius around the center. It was checked that the in plane shear stresses and strains are negligible in the central region.

In order to determine the stage t_{physio} corresponding to the average physiological state of the tissue in vivo, we approximated the circumferential stress in the bulging region of the aneurysm with the Laplace law:

$$\sigma_1 = (P_{in} D)/(4h) \tag{1}$$

where σ_1 is an estimate of the average circumferential stress in vivo, P_{in} is the average physiological pressure (mean between diastolic and systolic pressure measured by sphygmomanometry for each patient), D is the diameter of the aneurysm measured from the CT scan, and h is the current thickness of the wall in vivo, defined as:

$$h = h_0 / [\lambda_1(t_{\text{physio}}) \lambda_2(t_{\text{physio}})]$$
 (2)

The physiological tangent modulus (E) of the specimen is defined as:

$$E(t_{physio}) = d\sigma_1(t_{physio})/d\varepsilon_1(t_{physio})$$
 (3)

where $d\epsilon_1$ denotes the infinitesimal circumferential strain that occurs in vivo when the circumferential Cauchy stress varies due to blood pressure variations. The infinitesimal circumferential strain is defined as the ratio between the change of circumference and the average circumference in vivo. An equivalent ratio in the bulge test is $d\epsilon_1 = d\lambda_1/\lambda_1$. Accordingly, we define the physiological tangent modulus as:

$$E(t_{physio}) = \lambda_1(t_{physio}) d\sigma_1(t_{physio})/d\lambda_1(t_{physio})$$
(4)

where $d\sigma_1(t_{physio})/d\lambda_1(t_{physio})$ is the slope of the $\sigma_1(t)$ versus $\lambda_1(t)$ curve (Figure 3a). We consider the elastic modulus along the circumferential direction (σ_1 versus λ_1) instead of elastic modulus along the axial direction (σ_2 versus λ_2) because the changes of blood pressure during the cardiac cycle produce effects which are more significant circumferentially than axially.

For all the bulge inflation tests reported here, the artery burst with a straight crack splitting the media or the entire artery into two halves (Fig 1d and Fig 2). For each test where the crack made a straight line, the vector normal to this line was denoted \mathbf{e}_{θ} , θ being the angle between \mathbf{e}_{θ} and \mathbf{e}_{2} (values reported in Tab. 2). For 8 samples, the crack made an undetermined pattern or covered a large part of the sample and we could not estimate an angle. In these cases, an arbitrary value of θ =45° was assigned.

Using σ and C which are the Cauchy stress tensor and the Right Cauchy Green stretch tensor, respectively, the stress and the stretch components along e_{θ} can be derived with the following formulas:

$$\sigma = \mathbf{e}_{\theta} \cdot \mathbf{\sigma} \cdot \mathbf{e}_{\theta} \tag{5}$$

$$\lambda^2 = \mathbf{e}_{\theta} \cdot \mathbf{C} \cdot \mathbf{e}_{\theta} \tag{6}$$

The stage at which the rupture occurred was denoted t_{rup} . Applying equations 5 and 6 at t_{rup} and neglecting shear (implying that σ and C are diagonal), it can be written:

$$\sigma_{\text{rup}} = \sigma_1(t_{\text{rup}}) \cos^2 \theta + \sigma_2(t_{\text{rup}}) \sin^2 \theta \tag{7}$$

$$\lambda_{\text{rup}} = \{ [\lambda_1(t_{\text{rup}})]^2 \cos^2\theta + [\lambda_2(t_{\text{rup}})]^2 \sin^2\theta \}^{1/2}$$
 (8)

Eventually we derived two rupture criteria to evaluate the proximity between the physiological state and the ultimate state where rupture occurred:

1. the first criterion is based on the stress values:

$$\gamma_{\text{stress}} = [\sigma_1(t_{\text{physio}})/\sigma_{\text{rup}}] \cos^2\theta + [\sigma_2(t_{\text{physio}})/\sigma_{\text{rup}}] \sin^2\theta$$
 (9)

2. the second criterion is based on the stretch values:

$$\gamma_{\text{stretch}} = \{ \left[\lambda_1(t_{\text{physio}}) / \lambda_{\text{rup}} \right]^2 \cos^2 \theta + \left[\lambda_2(t_{\text{physio}}) / \lambda_{\text{rup}} \right]^2 \sin^2 \theta \}^{1/2}$$
 (10)

These criteria take values between 0 and 1. The closer they are to 1, the larger is the risk of rupture. In structural mechanics, $1/\gamma$ represents a safety factor.

2.4. Analysis method in uniaxial tensile tests

In the uniaxial tensile tests, the tensile Cauchy stress and the tensile stretch were derived from the recorded data F (axial load measured with a load cell) and L (current length of the specimen equal to the initial length L_0 plus the motion of the cross-head) according to: $\lambda = L/L_0$, $\sigma = F\lambda / S_0$, where S_0 is the initial cross section area.

 σ_{rup} and λ_{rup} were estimated as the stress and stretch values reached at first maximum just before dropping abruptly. This indicated a partial rupture (most of the time in the media) that was followed by tearing of the remaining tissue.

2.5. Data classification

We investigated the differences of rupture properties according to 3 different parameters: age, type of test and physiological modulus. Accordingly, patients were classified:

- 1. in 2 groups of different age: 8 patients ≤ 55 years old and 23 patients > 55 years old;
- in 3 groups of different test type (only for patients > 55): bulge inflation (23 patients), uniaxial tension along the circumferential direction (9 patients) and uniaxial tension along the axial direction (8 patients);

- 3. in 2 groups of different physiological modulus: 7 patients with E_{physio} < 1 and 24 patients with $E_{physio} \geq 1$.
- in 2 groups of different type of aortic valve: 8 patients with a bicuspid aortic valve (BAV) and
 23 patients with a tricuspid aortic valve (TAV).

2.6. Statistics

Relationships among different relevant parameters (age, diameter, E_{physio} , $\gamma_{stretch}$, γ_{stress} , λ_{rup} , σ_{rup}) was assessed using Pearson correlation analysis. A normality test was performed using Anderson-Darling test for the parametric inference, to determine whether the data in each group of patients followed a normal distribution. One-Way ANOVA with Bonferroni post hoc test was performed to define the statistical differences between groups of patients. The results are presented as mean \pm typical error. The statistical significance was set at p<0.05. The statistical methods applied were calculated using Minitab®.

3. RESULTS

3.1. Stress-stretch responses in bulge inflation

A total of 31 different samples were successfully tested with the bulge inflation test. They were collected from the bulging region of ATAAs of 31 patients ranging from 24 to 80 years old. Of interest is the mode of rupture that was observed in these bulge inflation tests. Although some samples burst completely and suddenly at the maximum pressure, in 55% of them (17 out of 31), the intima/media layer failed initially which was characterized by a pressure drop. Afterwards the pressure could be increased again as the adventitia still contained the inflating liquid and resisted rupture. This failure of the intima/media layer occurred with a sudden crack crossing at least half of the sample width. Following this crack, the intima/media layer did not bear any load and recoiled elastically. During this elastic recoil the media slipped over the adventitia, indicating that there had been a partial disruption between the two layers. This mode of failure has many similarities with an aortic

dissection (failure of the media and delamination between the media and the adventitia). We show a few pictures of ruptured samples in Fig. 2 for the sake of illustrating this typical dissection-like mode of rupture.

The stretch/stress curves of all the tests are shown in Fig 3, with the circumferential direction shown in Fig. 3a and the axial direction shown in Fig. 3b. For 32% of the samples (10 out of 31), the axial stretch tended to be larger than their circumferential counterparts for equivalent stress. For 29% of the samples (9 out of 31), the tendency was the opposite, whereas for the rest (39% - 12 out of 31) similar stretch was obtained in both directions.

3.2. Statistical analysis of rupture properties determined in bulge inflation tests

For the samples which showed the dissection-like mode of rupture, the maximum stress and stretch were measured when the media ruptured, even if the aorta could still bear load due to its preserved adventitia. It was observed that the ultimate stress values and the ultimate stretch values show no correlation with the aneurysm diameter (respectively p=0.5 and p=0.35). However, a correlation with age was noted (both p<0.01).

We show a graphical representation of the ultimate stress and stretch values as a function of age in Fig 4. The biaxial strength measured from the bulge inflation test shows a significant decrease between 20 and 55 years old. After 55 years of age, the values are dispersed around an average of about 0.9 MPa (Fig. 4a). The statistical analysis determined significant differences for the rupture stress (σ_{rup}) and stretch (λ_{rup}), and for the physiological modulus (E_{physio}) between samples from patients > 55 years old and samples from patients \leq 55 years old (p<0.05) (Fig. 5 and Table A1). Normal distributions were satisfied in both groups of patients for σ_{rup} , λ_{rup} and E_{physio} .

We also compared statistically the rupture stress (σ_{rup}) and stretch (λ_{rup}) between BAV and TAV patients, but no correlation was found (respectively p=0.48 and p=0.12).

3.3. Stress-stretch responses in uniaxial tension

A total of 103 different samples were successfully tested in uniaxial tension. They were collected from ATAAs of 12 patients ranging from 39 to 76 years old. For each patient, the average ultimate stress and stretch in both directions (circumferential and axial) are reported in Table 3. The values of ultimate stress are superimposed with the biaxial results in Fig 4a. It can be observed that there is a significant difference between the circumferential strength and the axial strength, the latter being on average the half of the former (p<0.01). The variation with age appears marginal for the axial strength whereas a net age-related decrease of the circumferential strength is visible (p=0.53 and p<0.05 respectively). For patients over 55 years old, the biaxial strength is similar to the axial strength and significantly lower than the circumferential strength (Fig. 6a), whereas for younger patients the biaxial strength tends to be similar to the circumferential strength and higher than the axial strength (confirmation of this trend by statistical analysis would require more patients in this group).

We show a graphical representation of the ultimate stretch values as a function of age in Fig. 4b. All the values show a global decrease with age, without reaching significance (p=0.06 and p=0.3 for circumferential and axial direction respectively).

The data from the three different mechanical tests (bulge and uniaxial in the circumferential and axial direction) were statistically compared for patients aged >55 (Fig 6 and Table A2), confirming that:

- 1- σ_{rup} is significantly different between bulge and uniaxial tests in the circumferential direction, and between both uniaxial tests (circumferential and axial).
- 2- λ_{rup} is significantly different between bulge and both uniaxial tests.
- 3.4. Correlation between rupture criteria, age and physiological modulus for the bulge inflation tests

There was a strong correlation between age and the two rupture criteria (γ_{stress} and γ_{stretch}) as seen in Fig. 7a (p<0.01). The stress based rupture criterion varies from 0.04 to 0.45, with an average value

of 0.15, showing that the stress in vivo could be multiplied, on average, by a factor higher than 6 to reach the rupture stress. The stretch based rupture criterion is much closer to 1, with an average value of 0.9. This means that in the aneurysm, on average, an 11% increase (γ_{stretch} =0.9) of the circumference/diameter due to blood flow variations increases the risk of rupture. For the weakest aneurysm, this maximum increase reduces to 3% (γ_{stretch} =0.97) and for the strongest, to 25% (γ_{stretch} =0.8).

Interestingly, a strong correlation is observed between $\gamma_{stretch}$ and E_{physio} (p<0.01). In Fig 7b, it is shown that most of the patients for whom $\gamma_{stretch}$ <0.9 have moduli below 1 MPa whereas patients for whom $\gamma_{stretch}$ >0.9 have moduli ranging from 1 MPa to 7 MPa. Accordingly, two groups were defined with respect to their physiological modulus ($E_{physio} \ge 1$ and $E_{physio} < 1$). The statistical analysis revealed that all variables (σ_{rup} , λ_{rup} , γ_{stress} , $\gamma_{stretch}$ and E_{physio}) had a normal distribution for a pre-determined level of significance of 95% (p≥0.05). It also showed a non-significant difference between both groups of samples for γ_{stress} , which means that the γ_{stress} criterion is independent of the physiological modulus (Fig. 8a and Table A3). Conversely $\gamma_{stretch}$ presented a significant difference between both groups of samples (Fig. 8b and Table A3). This means that brittle aneurysms (having a $\gamma_{stretch}$ very close to 1) are also stiffer. For instance, for an aneurysm having a diameter of 55 mm, a thickness of 2 mm, an elastic modulus of 1 MPa, and for a diastolic pressure of 80 mmHg and a systolic pressure of 130 mmHg, we can expect a variation of 9% of the circumference between diastole and systole by the simple application of the Laplace law.

4. DISCUSSION

4.1. Comparison between uniaxial and biaxial rupture properties of ATAA

In the present study we have reported for the first time the rupture properties under biaxial loading of 31 human ATAA samples. Due to their bulge shape, ATAA tissue is stretched biaxially in vivo [23]. However, all the rupture properties of ATAAs reported in the literature were characterized

utilizing uniaxial tension and there was no guarantee that results obtained from a uniaxial test can be applied to a biaxial state.

An interesting aspect of the bulge inflation test is the dissection-like rupture that we observed in most of the samples [26,27], which has never been reported for uniaxial tensile tests but is a common mode of rupture of ATAAs in vivo [4]. Only Iliopoulos et al. showed that the inner layer ruptures first in uniaxial tensile tests [36]. Even if the stresses are probably not perfectly equibiaxial in vivo, the bulge inflation test still represents a more relevant mode of loading than the uniaxial tension for characterizing rupture properties of ATAAs.

Uniaxial tensile tests showed that the axial direction of the aortic tissue is always weaker than the circumferential direction [12,36,37]. This was observed in most of the published studies which characterized both directions of ATAA samples independently. In the bulge inflation tests, we observed that approximately half of the samples ruptured with a crack perpendicular to the axial direction (θ =90°). We also observed that the ultimate stress in bulge inflation testing is similar to the axial ultimate stress in uniaxial tension tests for patients >55 years old. This confirms that the axial direction is weaker in biaxial loading. A possible explanation for this lower strength in the axial direction is that collagen in the media tend to be aligned in the circumferential direction [38]. This is also consistent with the observed anisotropy in the stress/strain curves, the circumferential direction being generally stiffer. This is a common result with the passive mechanical behavior of the aorta [18,39,40].

Although the circumferential strength is almost double that of the axial direction in uniaxial tests, we observed that the ultimate stretch is similar in both directions, which corroborates other studies [41]. This property of the tissue to reach equivalent ultimate stretch in both direction, despite the anisotropy, brings some complexity to the biaxial behavior which may explain why the rupture does not occur at θ ~90° in all the bulge inflation tests.

When stress analyses are performed on ATAAs using finite element analyses [28], different combinations of axial and circumferential stress can be found. In order to determine if rupture may

occur, failure envelopes taking into account all these different possible combinations are required. Such failure envelopes were previously proposed for abdominal aortic aneurysms [42]. On the one hand, the maximum stretches reported in our study, both in uniaxial tension and equibiaxial tension, are in fairly good agreement with the stretch-based failure envelopes of Volokh [42]. On the other hand, our maximum stresses do not match with the stress-based failure envelopes reported in the Volokh study [42]. The disagreement for the stress may be explained by the assumption of isotropy used to derive the failure envelopes in [42]. Isotropy is not satisfied for ATAA tissue, the strength in the axial direction being half the strength in the circumferential direction. An appropriate failure envelope for ATAAs should be in agreement with both our uniaxial and biaxial data, which requires that it considers anisotropy.

4.2. Correlation of rupture properties with age and valve phenotype

Interestingly we also observed that the biaxial strength for patients ≤55 years old tends to be larger than the axial strength, whereas both are similar for patients >55 years old. The trends seems to be confirmed for instance by the data of García-Herrera et al. who characterized the rupture properties of 23 subjects between 16 and 57 years old without ATAA and obtained an average axial strength of about 1.2 MPa, which is consistent with our axial strength values at this range of age [12].

These observations lead us to the conclusion that there is an important correlation between the rupture properties of ATAAs and age. This was already established using uniaxial data by Okamoto et al. [10]. In our data, the Pearson test yields a significant correlation between age and both the ultimate stress and ultimate stretch.

Several authors have compared the uniaxial rupture properties of ATAAs between BAV and TAV patients [12,38,43,44]. TAV specimens were found significantly weaker than BAV specimens in some studies [38,43] whereas others found no correlation between rupture properties and valve phenotypes [12,44]. No correlation was found between the valve phenotype and rupture properties in our biaxial data.

4.3. Rupture risk based on the extensibility of the tissue

The strength of the aortic tissue is generally defined as the maximum stress that the tissue can withstand before failing. However, when we derived the γ_{stress} ratio between the stress applied to the tissue in vivo and its strength, we noted that most of the collected ATAA samples were far from rupture. Another equivalent definition of rupture can be when the stretch applied to the tissue exceeds its extensibility. This way of defining rupture can even be more physiologically meaningful as it is reported that aneurysm rupture or dissections often occur at a time of severe emotional stress or physical exertion [45]. Emotionally or physically stressful situations can induce significant changes of blood volumes in the aorta and more compliant aneurysms are less prone to rupture as they can sustain such changes of volume. Based on this analysis, Martin et al. also defined a similar criterion, named the diameter risk, which is the ratio between the current diameter of the aneurysm and the rupture diameter [13]. They showed that the diameter risk increases significantly with the physiological elastic modulus of the artery. Indeed, if the aortic wall is stiff, a rather large increase of pressure can be induced by a small increase of volume. We showed similar trends in Fig. 7b with our biaxial data.

A likely explanation of this phenomenon is the mechanism of collagen recruitment described by Hill et al. [46]. As collagen recruitment increases with the load, the tangent stiffness of the tissue increases too. We can expect that collagen recruitment is delayed in the tissue when it still contains a significant fraction of non-fragmented elastin. At physiological pressures, when elastin is not fragmented, only a small fraction of collagen contributes to the stiffness, which is relatively small (below 1MPa). Conversely, when elastic lamella and elastic fibers are highly disrupted, collagen will tend to be recruited earlier and will already contribute significantly to the stiffness at physiological pressures, making the stiffness relatively high (larger than 1 MPa).

This description of higher recruitment of collagen at physiological pressures corroborates with Iliopoulos et al. [41] who showed that wall elastin, but not collagen content, decreased in ATAA

specimens, displaying lower wall thickness and failure strain, higher peak elastic modulus, and equal failure stress than control specimens. Moreover, it is commonly observed that the elastin fraction decreases with age, which may explain the trends observed in Fig. 4.

The level of collagen recruitment may explain why γ_{stretch} can be very close to 1, but not the relatively high values of γ_{stress} that occur for some patients. Indeed, some specimens showed a relatively large extensibility but ruptured at a relatively small stress. When collagen is recruited, it can withstand stresses until damage initiates with ruptures at the fiber level as demonstrated by Weisbecker et al. [47,48]. A dense and crosslinked network of collagen will reach larger stresses before the initiation of damage and ultimate stresses.

4.4. Relationships between rupture properties and tissue histology

From the above discussion, we submit that two factors play a role in the risk of rupture of ATAAs:

- content and crosslinking of the network of elastic fibers in the media which delays the collagen recruitment and ensures a higher extensibility. The degradation/fragmentation of the elastin network renders ATAAs more brittle (γ_{stretch} >0.9).
- content and crosslinking of the network of collagen fibers in the media which delays the initiation of damage leading to rupture. The degradation of the collagen network renders ATAAs weaker (σ_{rup} <1 MPa)

Several authors have highlighted the relationship between the quality of protein networks in the extracellular matrix of ATAAs and the strength and extensibility of the tissue [43,44,49]. The degradation of protein networks in the extracellular matrix of ATAAs is explained by the fact that protein production is overwhelmed by protein destruction by the matrix metalloproteases (MMPs), notably MMP-9 which is responsible for elastin degradation. Once the aneurysm process begins, the changes that occur in the thoracic aortic wall lead to increased wall stress, which leads to greater proteolytic activity, which leads to further remodeling, perpetuating a vicious cycle. Once a critical state is reached, the aorta has lost its elastic properties. In the 31 ATAA samples that we tested, the

state of degradation varied significantly between individuals but it appeared that it was less advanced for patients ≤55 years old. This may mean that protein destruction is slower in young individuals even though they are affected by an ATAA.

4.5. Limitations and future work

Due to some current limitations, the following improvements are planned for our future experimental characterizations of ATAA:

- 1. In this study we used the Laplace law to estimate in vivo physiological conditions (in vivo stress, in vivo tangent modulus) in the bulge region of the aneurysms. Using dynamic gated CT scan and finite-element analyses, this estimation could be refined with a simulation of in vivo physiological conditions as we started to do in preliminary studies [50].
- 2. The initial or zero-stress thickness was measured using a digital caliper assuming a homogeneous thickness. This should be replaced by a more precise and local measurement where the natural thickness variation of the tissue is taken into account, for example with the use of thickness gauges [51].
- 3. It would be interesting to distinguish the anterior, posterior and lateral regions of ATAAs as in Iliopoulos et al. and Kim and Baek [21,36], in order to assess possible regional variations of the rupture properties. Here we tested only the outer curvature of the ATAA where the rupture usually occurs.
- 4. We could apply our approach to healthy ascending aortas as done by Azadani et al. [52], in order to better understand the differences caused in the aortic wall by a pathological state such as aortic insufficiency.
- 5. Refined histological or immune-histological analyses could be undertaken in order to quantify for instance fiber disruption, mucoid accumulation, loss or hyperplasia of SMCs which are typical of ATAAs.
 - 6. A constitutive model accounting for damage [53–59] could be identified from the stress/strain

curves.

7. It was assumed that the samples were in a stress-free state at the beginning of each inflation test but residual stresses are actually present in arteries in vivo [60] and it may be interesting to determine how they could affect the ultimate strength and stretch. Similarly the specimens were not perfectly flat at the beginning of the test and this led to some small initial deformation.

5. CONCLUSIONS

In this study we characterized the rupture properties of 31 human ATAA samples using bulge inflation testing. To our knowledge this is the first time that in vitro rupture properties of ATAAs are reported and analyzed using biaxial loading conditions. The results have highlighted the relationship between failure properties and the age of patients, showing that patients below 55 years of age display significantly higher strength. We have also shown the correlation between the extensibility of the tissue (which indicates its brittleness) and the physiological elastic modulus, independently of the age, ATAA diameter or the aortic valve phenotype of the patient. The results indicate the combined importance of the quality of the collagen and elastin networks to prevent rupture – elastin quality delaying collagen fiber recruitment and bringing extensibility, and collagen quality delaying fiber damage and bringing strength. Further studies should be done to characterize the underlying mechanobiological mechanisms yielding to the degeneration of these protein networks during the growth of an aneutysm.

CONFLICT OF INTEREST

None.

ACKNOWLEDGEMENT

The authors are grateful to the European Research Council for grant ERC-2014-CoG BIOLOCHANICS.

The authors would like to thank Prof Ramon Berguer for sharing rupture data and for the inspiring

discussions.



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Table 1. Published studies reporting uniaxial rupture properties of ascending thoracic aortas and ATAAs.

Reference	Tests	Main conclusions
Mohan and Melvin (1982)	34 aortas in uniaxial tension	Failure properties of human aortic tissue at quasi- static and dynamic strain rates
Okamoto et al (2002)	25 ATAAs tested up to failure in uniaxial tension	Age clearly influences the measured elastic and strength properties in ATAA
Vorp et al. (2003)	36 ATAAs and 10 non aneurysmal ascending thoracic aortas	ATAA samples are stiffer and approximately 30% weaker than healthy samples
Sokolis et al. (2012)	Uniaxial tensile tests on intima, media and adventitia	Similar extensibility for all the layers but larger strength for the adventitia
Iliopoulos et al. (2009a)	Tensile tests on ATAAs sample of 12 patients	Inverse correlations of failure stress with thickness and direct correlations with peak elastic modulus
Iliopoulos et al. (2009b)	Uniaxial tensile tests on 26 ATAAs and 15 non aneurysmal ascending thoracic aortas	ATAA display smaller failure strain, smaller elastin content and higher peak elastic modulus. Circumferential specimens show higher failure stress and peak elastic modulus but equal failure strain
Forsell et al. (2014)	Uniaxial tensile tests on 24 ATAAs, 14 patients with bicuspid aortic valve (BAV) and 10 with tricuspid aortic valve (TAV)	The strength was almost two times higher in BAV samples. Collagen related stiffness was significantly increased in BAV samples whereas elastin related stiffness was similar in both groups
Pichamuthu et al. (2013)	Uniaxial tensile tests on 38 ATAAs, 23 from BAV patients and 15 from TAV patients	Circumferential and longitudinal tensile strengths were higher for BAV ATAAs than TAV ATAAs. Uniform collagen content between groups
García-Herrera et al. (2012)	Uniaxial tensile tests on 26 ATAAs (12 BAV and 14 TAV) and 23 non aneurysmal ascending thoracic aortas	Age causes a major reduction in the mechanical parameters of healthy ascending aortic tissue, no significant differences between the mechanical strength of aneurysmal TAV or BAV aortic specimens and the corresponding age-matched control group
Pham et al. (2013)	Uniaxial tensile tests on 35 ATAAs (20 BAV and 15 TAV) and 20 non aneurysmal ascending thoracic aortas	No difference of failure properties between the groups. Moderate medial degeneration characterized by elastin fragmentation, cell loss, mucoid accumulation and fibrosis. The ultimate tensile strength decreases with age
Sokolis and Iliopoulos (2014)	Uniaxial tensile tests on 35 ATAAs (20/15 male/female ratio)	Male ATAAs are stronger but equally extensible in the circumferential axis compared to female ones. Longitudinally, instead, gender differences at each region are insignificant

Table 2. Demographic information and results of the bulge inflation tests.

Id	Sex/ Age/Valve	Thickness (mm)	Diameter (mm)	E _{physio} (MPa)	σ _{rup} (MPa)	λ_{rup}	γ_{stress}	$\gamma_{stretch}$	θ (°) [†]
1	F/76/TAV	1.82	55	3.69	1.31	1.19	0.09	0.94	80
2	M/64/TAV	1.99	55	1.01	1.28	1.40	0.10	0.86	х
3	M/55/BAV	2.10	65	0.96	0.79	1.48	0.22	0.88	80
4	M/76/TAV	1.98	51	1.69	1.12	1.34	0.11	0.92	х
5	M/68/TAV	1.79	55	2.05	1.17	1.51	0.16	0.94	х
6	M/58/BAV	2.74	52	0.61	1.11	1.53	0.10	0.88	х
7	M/66/TAV	1.86	50	0.72	0.50	1.48	0.28	0.84	90
8	M/75/BAV	1.70	55	2.22	1.07	1.33	0.14	0.92	80
9	M/68/BAV	2.50	51	2.59	0.90	1.34	0.12	0.95	90
10	M/78/TAV	2.72	50	0.62	0.65	1.39	0.13	0.84	90
11	M/68/TAV	2.20	50	0.78	0.78	1.56	0.20	0.91	100
12	M/70/TAV	1.79	55	2.85	1.24	1.32	0.12	0.94	Х
13	M/24/BAV	1.77	50	0.61	4.84	1.70	0.04	0.81	75
14	M/64/TAV	1.79	55	1.31	1.29	1.64	0.17	0.92	90
15	M/66/TAV	1.94	55	1.85	0.45	1.28	0.27	0.92	80
16*	M/36/TAV	0.51	50	0.69	3.60	1.52	0.04	0.83	0
17*	M/55/TAV	1.09	53	2.20	0.82	1.33	0.18	0.95	0
18*	M/49/BAV	0.95	52	1.48	1.51	1.45	0.10	0.88	0
19	M/64/TAV	1.65	52	2.58	0.86	1.40	0.21	0.94	х
20	M/79/TAV	1.79	50	4.51	0.86	1.22	0.14	0.96	100
21	F/65/TAV	1.82	49	5.37	0.85	1.18	0.13	0.96	100
22	F/80/TAV	1.68	52	5.34	0.40	1.12	0.30	0.98	100
23	F/80/TAV	1.68	52	3.77	0.77	1.25	0.20	0.96	30
24	M/40/TAV	1.59	55	1.20	2.30	1.55	0.09	0.88	х
25	M/79/TAV	1.76	52	4.41	0.95	1.21	0.15	0.96	0
26	M/76/BAV	1.82	58	3.28	0.33	1.25	0.51	0.98	100
27	M/72/TAV	1.90	51	2.37	0.76	1.18	0.16	0.97	100
28	F/76/TAV	2.41	65	2.53	1.31	1.28	0.10	0.95	70
29	M/79/TAV	1.89	54	6.52	1.67	1.20	0.07	0.94	80
30	M/55/BAV	2.35	55	1.00	0.98	1.52	0.16	0.89	100
31	M/54/TAV	2.11	55	1.36	2.60	1.45	0.05	0.87	х
Mean	65 (age)	1.97	53.52	2.33	1.26	1.37	0.16	0.92	_
STD	13.8 (age)	0.31	3.80	1.60	0.94	0.15	0.09	0.05	

^{*} for 3 patients, the media and the adventitia were separated as explained in (Romo et al, 2014). The thickness value reported in the table is the thickness of the layer tested in the bulge inflation experiment and it is not taken into account to derive the average and standard deviation of the thickness of the whole population.

 $[\]dot{\theta}$ an angle $\dot{\theta}$ of x means that the crack pattern was undetermined and that it was not possible to estimate an angle.

Table 3. Demographic information and results of the uniaxial tensile tests.

8 M/76/TAV 1.52 46 0.67 0.39 1.48	2 F/72/BAV 1.48 53 1.17 - 1.36 3 M/61/TAV 2.02 51 1.54 0.51 1.77 4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	2 F/72/BAV 1.48 53 1.17 - 1.36 3 M/61/TAV 2.02 51 1.54 0.51 1.77 4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36	Id	Sex/ Age/Valve	Thickness (mm)	Diameter (mm)	Circ σ _{rup} (MPa)	Axial σ _{rup} (MPa)	$\frac{Circ}{\lambda_{rup}}$
3 M/61/TAV 2.02 51 1.54 0.51 1.77 4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	3 M/61/TAV 2.02 51 1.54 0.51 1.77 4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	3 M/61/TAV 2.02 51 1.54 0.51 1.77 4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20						-	1.50
4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	4 M/57/BAV 2.31 55 1.31 0.45 1.69 5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20							
5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	5 M/39/BAV 1.80 52 2.81 0.76 1.95 6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	3	M/61/TAV	2.02	51	1.54	0.51	1.77
6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/75/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	6 M/47/BAV 1.70 50 - 0.80 - 7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/75/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20				55		0.45	1.69
7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	7 M/75/TAV 1.65 48 2.07 0.64 1.77 8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	5	M/39/BAV	1.80	52	2.81	0.76	1.95
8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	8 M/76/TAV 1.52 46 0.67 0.39 1.48 9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20		M/47/BAV	1.70	50	-	0.80	-
9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	9 M/75/TAV 2.01 53 1.00 0.82 1.60 10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	7	M/75/TAV	1.65	48	2.07	0.64	1.77
10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	10 M/68/TAV 2.02 67 2.19 1.03 2.00 11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	8	M/76/TAV	1.52	46	0.67	0.39	1.48
11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	11 M/61/TAV 1.50 45 1.30 0.80 1.70 12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	9	M/75/TAV	2.01	53	1.00	0.82	1.60
12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	12 M/55/TAV 1.79 50 1.87 1.65 1.81 Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	10	M/68/TAV	2.02	67	2.19	1.03	2.00
Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	Mean 63 (age) 1.79 51.08 1.57 0.79 1.69 STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	11	M/61/TAV	1.50	45	1.30	0.80	1.70
STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	STD 11.8 (age) 0.25 6.16 0.61 0.36 0.20	12	M/55/TAV	1.79	50	1.87	1.65	1.81
			Mean	63 (age)	1.79	51.08	1.57	0.79	1.69
			STD	11.8 (age)	0.25	6.16	0.61	0.36	0.20
A =						O			

Legends of figures

- **Figure 1.** (a) Picture of an ATAA sample just after collection during aneurysm surgical repair; (b) Picture of an ATAA sample after cutting the inner curvature and opening; (c) Picture of an ATAA sample after cutting the square sample that will be tested in the bulge inflation bench; (d) Picture of an ATAA sample after rupture in the bulge inflation test.
- **Figure 2.** Pictures taken by the digital camera of the DIC system after rupture of ATAA samples in the bulge inflation test. (a) patient 10; (b) patient 13; (c) patient 14; (d) patient 11. Pictures are taken from above the sample. More details about the experimental arrangements may be found in [27].
- **Figure 3.** Stress/stretch curves obtained on the 31 samples tested with the bulge inflation test. (a) Circumferential Cauchy stress versus circumferential stretch; (b) Axial Cauchy stress versus axial stretch.
- **Figure 4.** (a) Age related variations of the ultimate strength σ_{rup} in uniaxial tensile tests and in bulge inflation tests (p<0.01); (b) Age related variations of the ultimate stretch λ_{rup} in uniaxial tensile tests and in bulge inflation tests (p<0.01)
- **Figure 5.** Significant differences between both populations (age \leq 55 and age > 55) of (a) σ_{rup} , (b) λ_{rup} and (c) E_{physio} .
- **Figure 6.** Statistical comparison between the 3 different types of tests (bulge inflation, uniaxial tension along the circumferential and uniaxial direction) of the distributions of (a) σ_{rup} and (b) λ_{rup} .
- **Figure 7.** (a) Age related variations of both rupture criteria γ_{stress} and $\gamma_{stretch}$ determined from bulge inflation tests (p<0.01); (b) Variations of $\gamma_{stretch}$ with respect to the physiological elastic modulus (p<0.01).
- **Figure 8.** Statistical comparison between both populations ($E_{physio} < 1$ and $E_{physio} \ge 1$) of the distributions of (a) γ_{stress} and (b) $\gamma_{stretch}$.

Appendix

Table A1. Statistical distributions (mean and typical error) for both populations (age \leq 55 and age > 55) for σ_{rup} , λ_{rup} and E_{physio}

Bulge test	σ _{rup} (MPa)		λ	rup	E _{physio} (MPa)		
	≤55	>55	≤55	>55	≤55	>55	
Mean ± TE	2.18 ± 0.52	0.94 ± 0.07	1.5 ± 0.04	1.33 ± 0.03	1.19 ± 0.18	2.72 ± 0.35	
N	8	23	8	23	8	23	

Table A2. Statistical distributions (mean and typical error) for each testing type (bulge inflation, uniaxial tension along the circumferential and axial direction) for σ_{rup} and λ_{rup} .

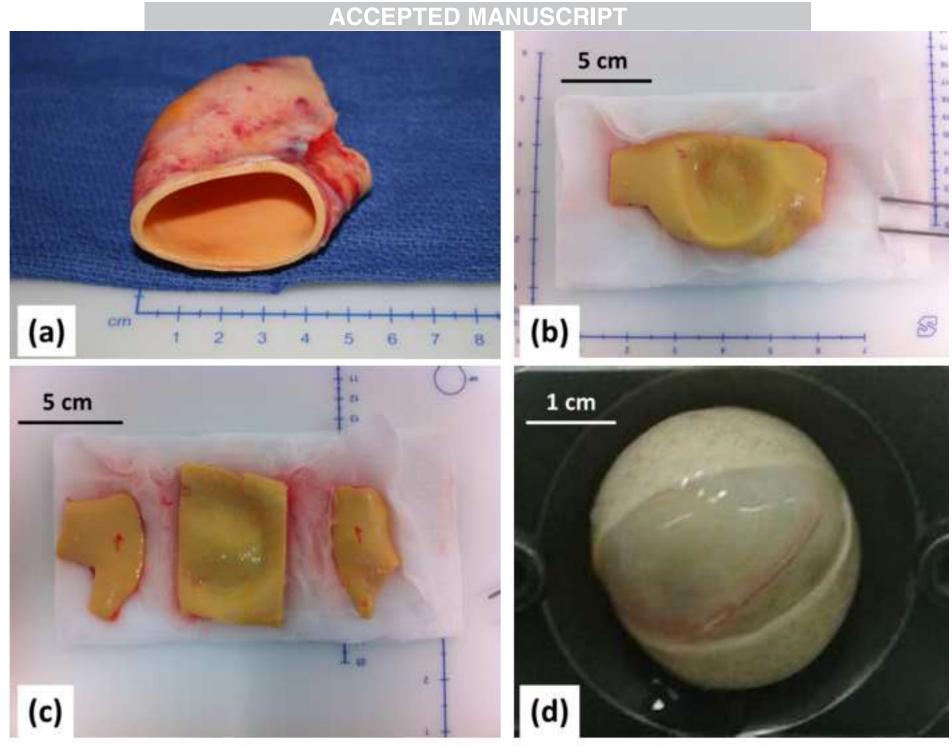
Patients > 55	Uniaxial test (c	ircumferential)	Uniaxial	test (axial)
years old	$\sigma_{rup}(MPa)$	λ_{rup}	σ _{rup} (MPa)	λ_{rup}
Mean ± TE	1.4 ± 0.16	1.65 ± 0.06	0.66 ± 0.09	1.58 ± 0.07
N	9	9	7	7

Table A3. Statistical distributions (mean and typical error) for both populations ($E_{physio} < 1$ and $E_{physio} \ge 1$) for σ_{rup} , λ_{rup} γ_{stress} , $\gamma_{stretch}$ and E_{physio} .

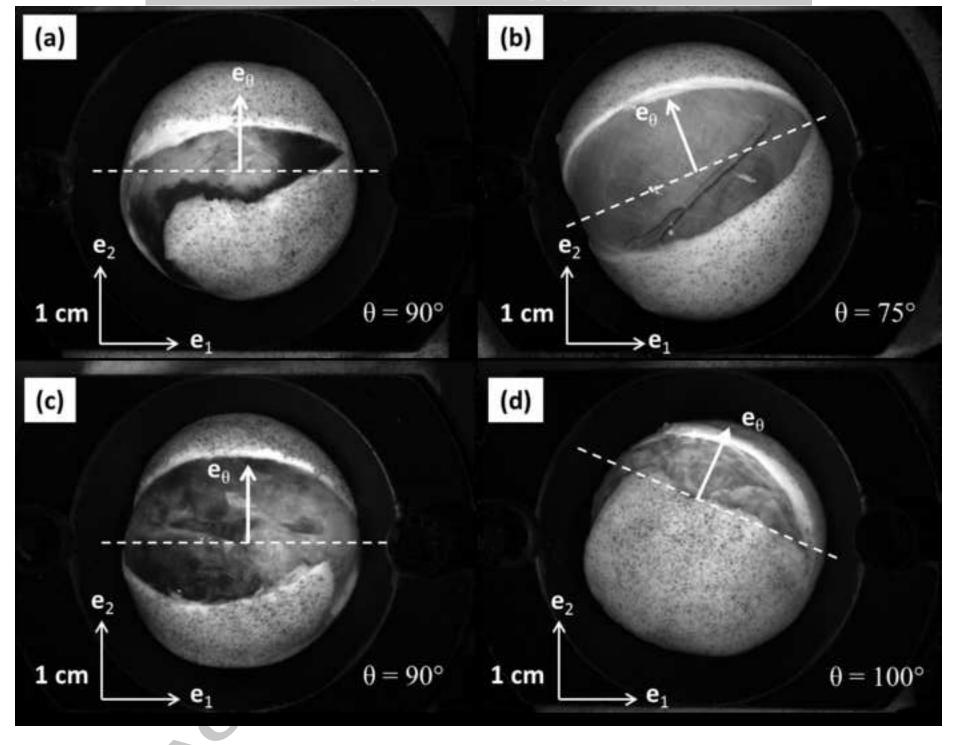
Bulge test	σ_{rup}	(MPa)	λ	rup	γ _{st}	ress	γ _{st}	retch	E _{physio}	(MPa)
	E _{physio} < 1	$E_{physio} \ge 1$	$E_{physio} < 1$	$E_{physio} \ge 1$	$E_{physio} < 1$	$E_{physio} \ge 1$	$E_{physio} < 1$	$E_{physio} \ge 1$	E _{physio} < 1	$E_{physio} \ge 1$
Mean ± TE	0.77 ± 0.1	1.09 ± 0.09	1.49 ± 0.03	1.33 ± 0.03	0.19 ± 0.03	0.15 ± 0.01	0.87 ± 0.01	0.93 ± 0.01	0.74 ± 0.06	2.84 ± 0.33
N	5	22	5	22	5	22	5	22	5	22

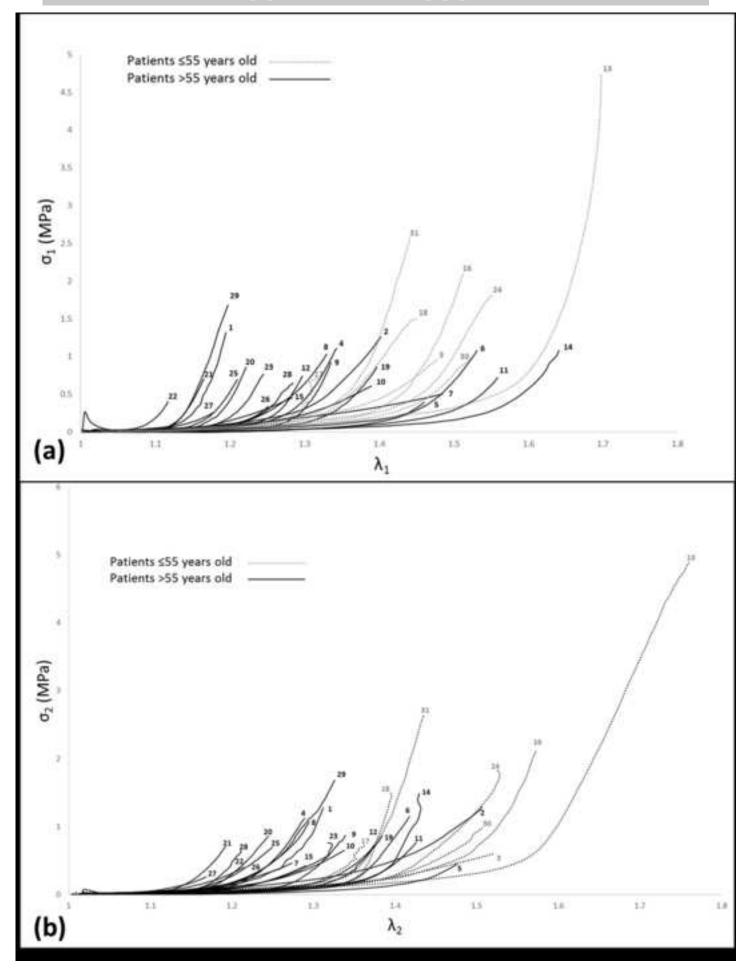
Despite their medical importance, rupture properties of ascending thoracic aortic aneurysms (ATAA) subjected to biaxial tension were inexistent in the literature. In order to address this lack, our group developed a novel methodology based on bulge inflation and full-field optical measurements. Here we report rupture properties obtained with this methodology on 31 patients. It is shown for the first time that rupture occurs when the stretch applied to ATAAs reaches the maximum extensibility of the tissue and that this maximum extensibility correlates strongly with the elastic properties. The outcome is a better detection of at-risk individuals for elective surgical repair.

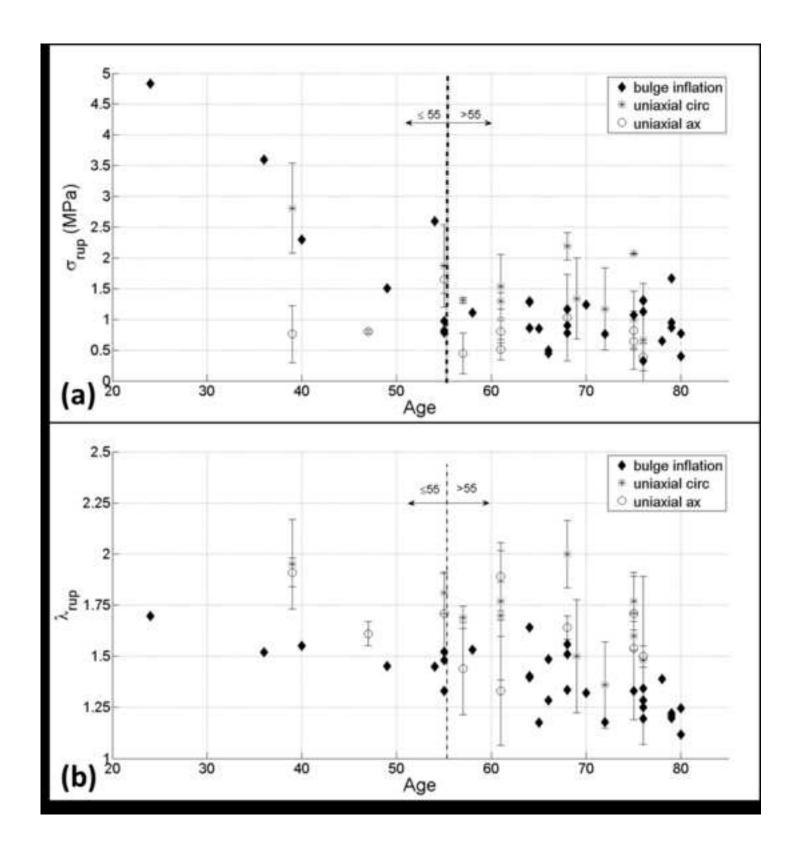


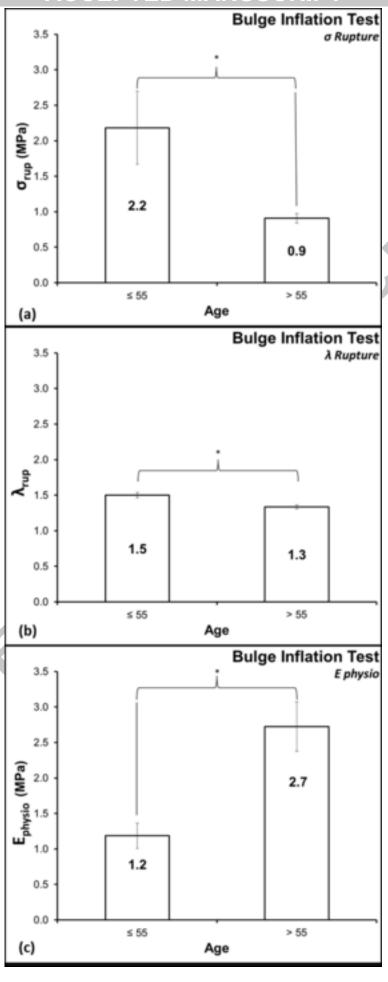




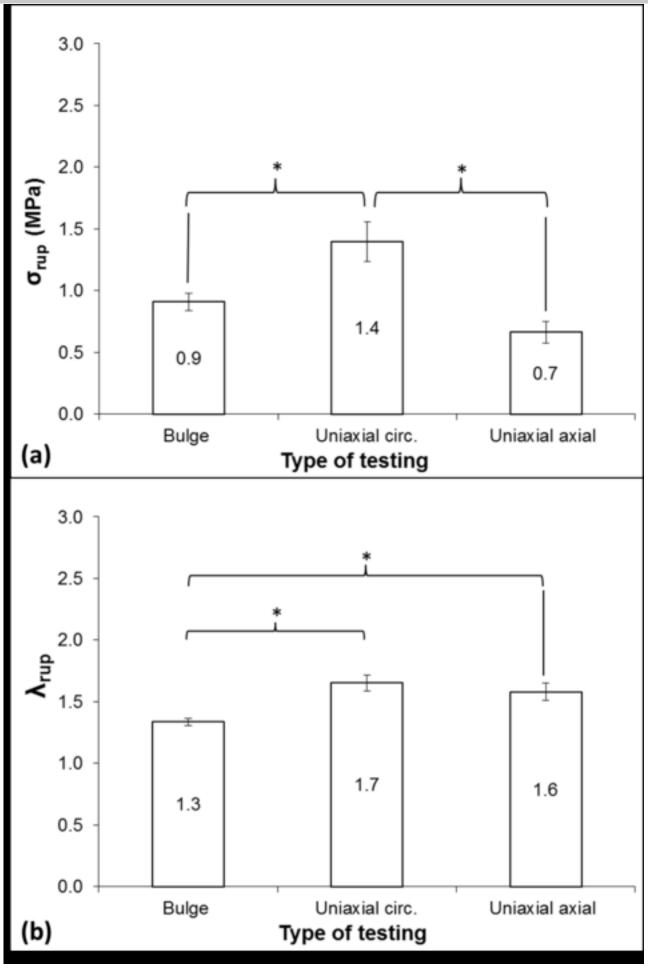


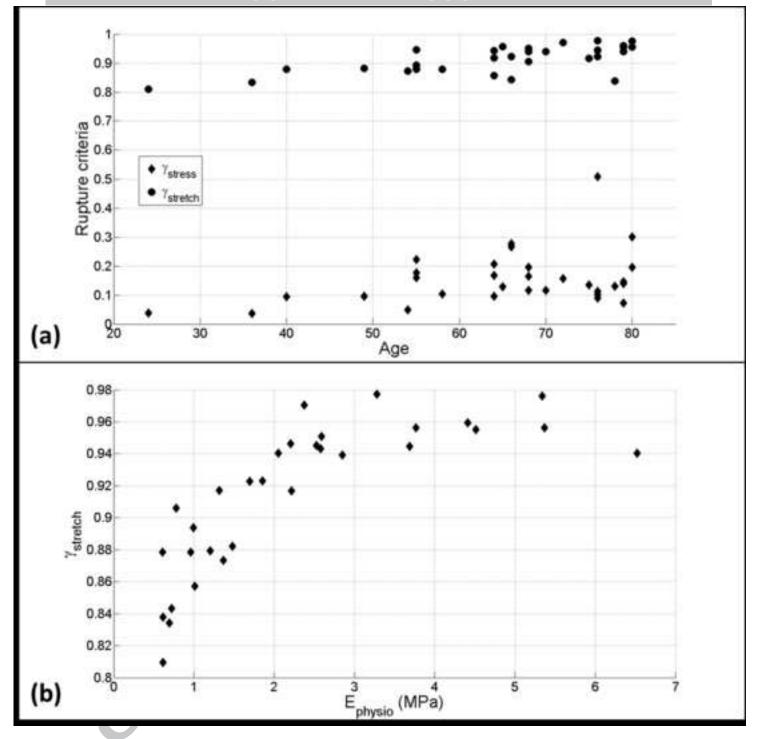












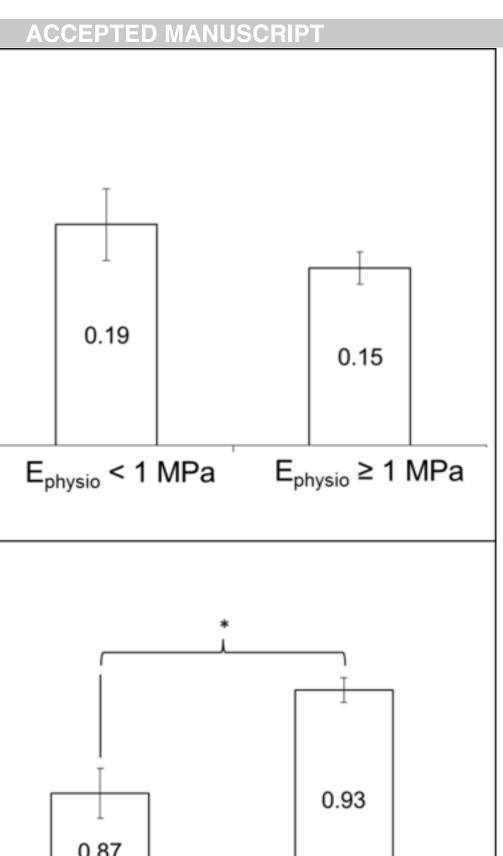
0.3 -

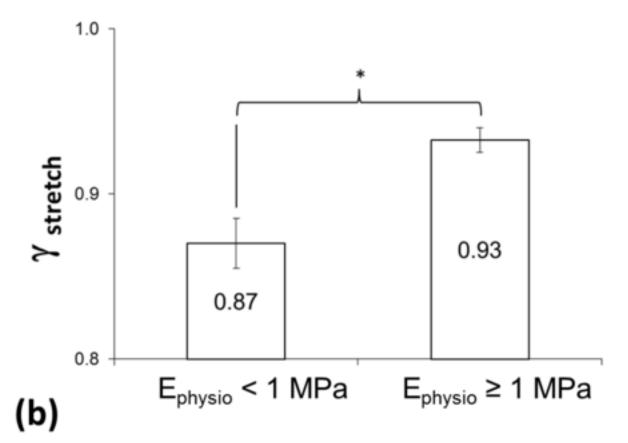
0.2

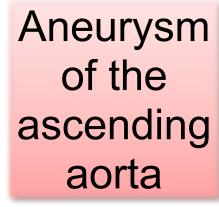
γ stress

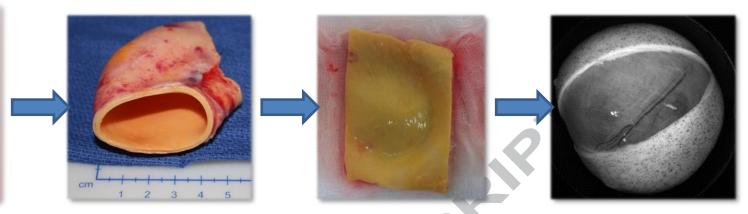
0.0 -

(a)









Bulge inflation rupture test 31 human specimens

