



Centre for Health Engineering
Centre Ingénierie et Santé
LCG CNRS UMR 5146

Pierre Badel + colleagues



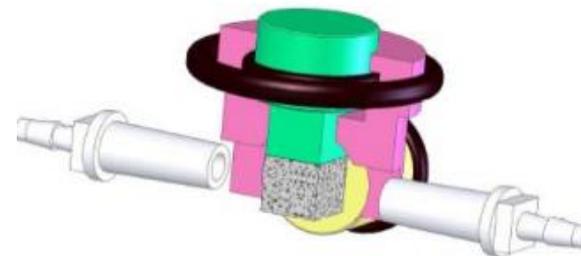
Identification of vascular soft tissue
mechanical properties

➔ Center for Biomedical and Healthcare Engineering

Multi-disciplinary

➔ **Translational research**

**Close collaboration with
clinical and industrial partners**



➔ Soft Tissue Biomechanics group

Diagnosis



Healthcare & MD optimization



Translational research



Soft tissue / structure relationship

Multi-modality imaging
+
Mechanical modeling

Pathological (large) arteries

Osteo-articular system

Skin and sub-cutaneous tissue

Understanding the mechanical behavior of these tissues

Study of the mechanical action of medical devices

➔ Introduction

➔ A few words on inverse identification methods applied to soft tissue biomechanics

➔ Ascending thoracic aortic aneurysm

- Hyper-elastic model identification
- Rupture characterization

➔ Perspectives

Inverse identification

➔ Introduction

➔ **A few words on inverse identification methods applied to soft tissue biomechanics**

➔ **Ascending thoracic aortic aneurysm**

- Hyper-elastic model identification
- Rupture characterization

➔ Perspectives

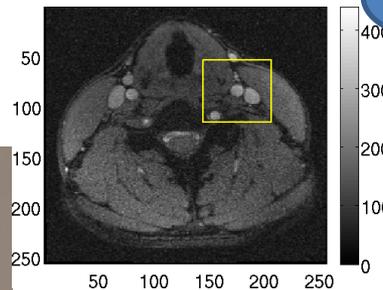
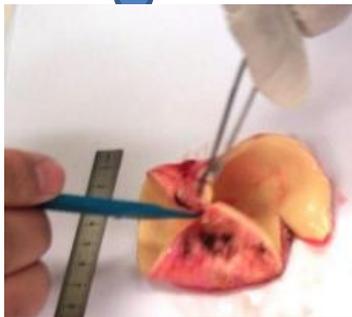
➔ Why inverse identification in soft tissue biomechanics?

What kind of specimen?

What kind of test?

In vitro?
Access to tissue ?
Destructive?
Representative of what we are looking for?

In vivo?
What is the loading?
What is the geometry?



[sigvaris.fr]

➔ Why inverse identification in soft tissue biomechanics?

What kind of specimen?

What kind of test?

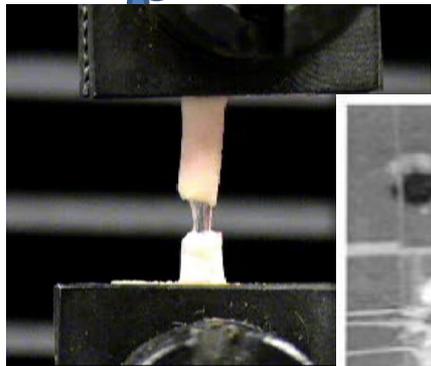
In vitro?

Access to tissue ?

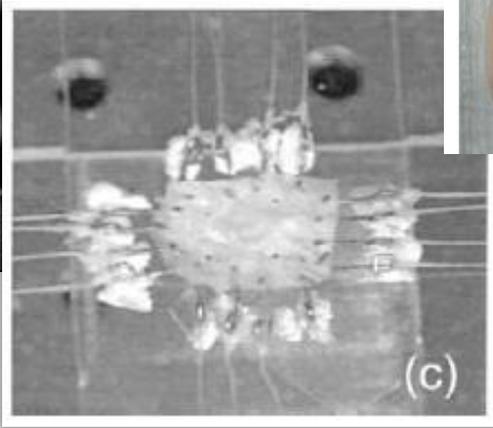
Destructive?

Representative of what we are looking for?

In vivo?



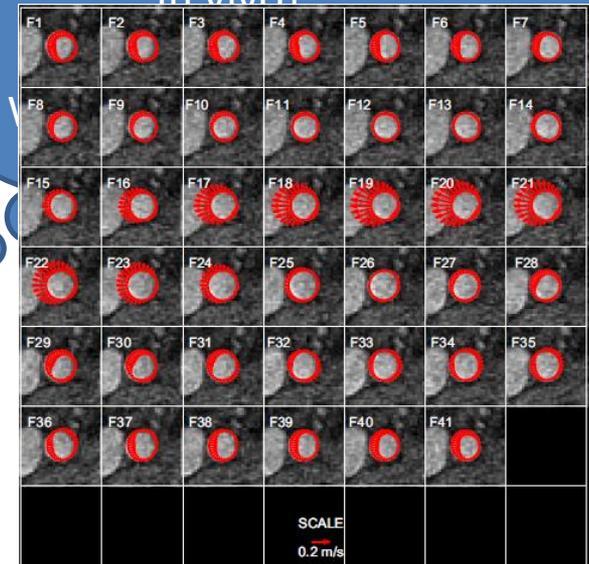
[Duprey, 2010]



[Sacks, 2000]



[Genovese, 2009]



[Avril, 2011]

⇒ Why inverse identification in soft tissue biomechanics?

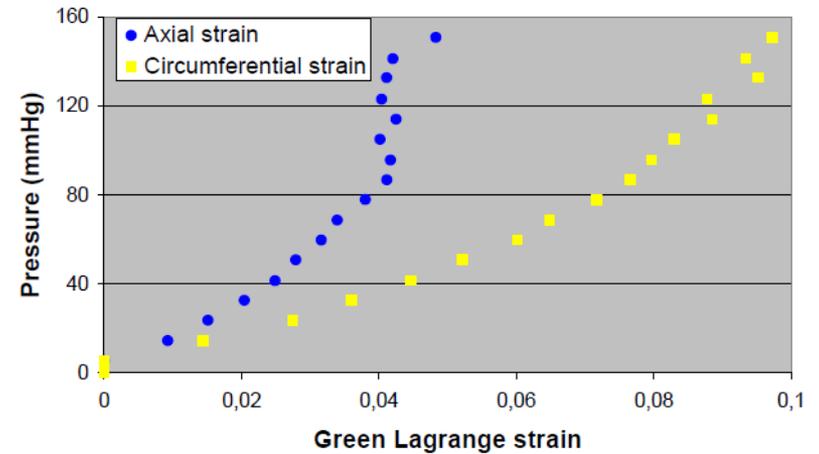
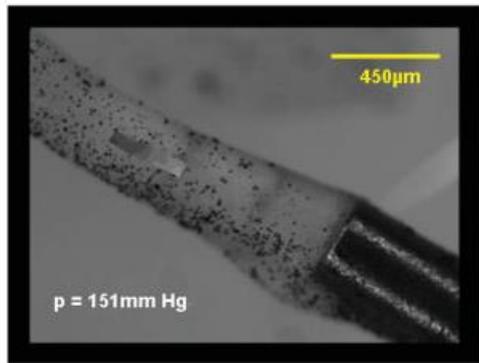
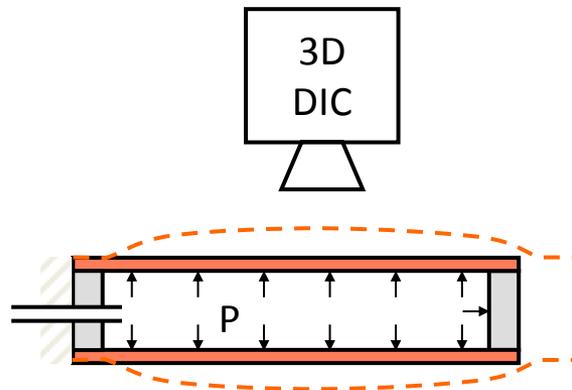
⇒ **Relevant experimental data is complex (geometry, boundary conditions...)**

⇒ **Complex tissues (non-linearity, heterogeneity...)**

⇒ **Models are often complex**

➔ In vitro ex.: Inflation/extension, mouse carotid artery

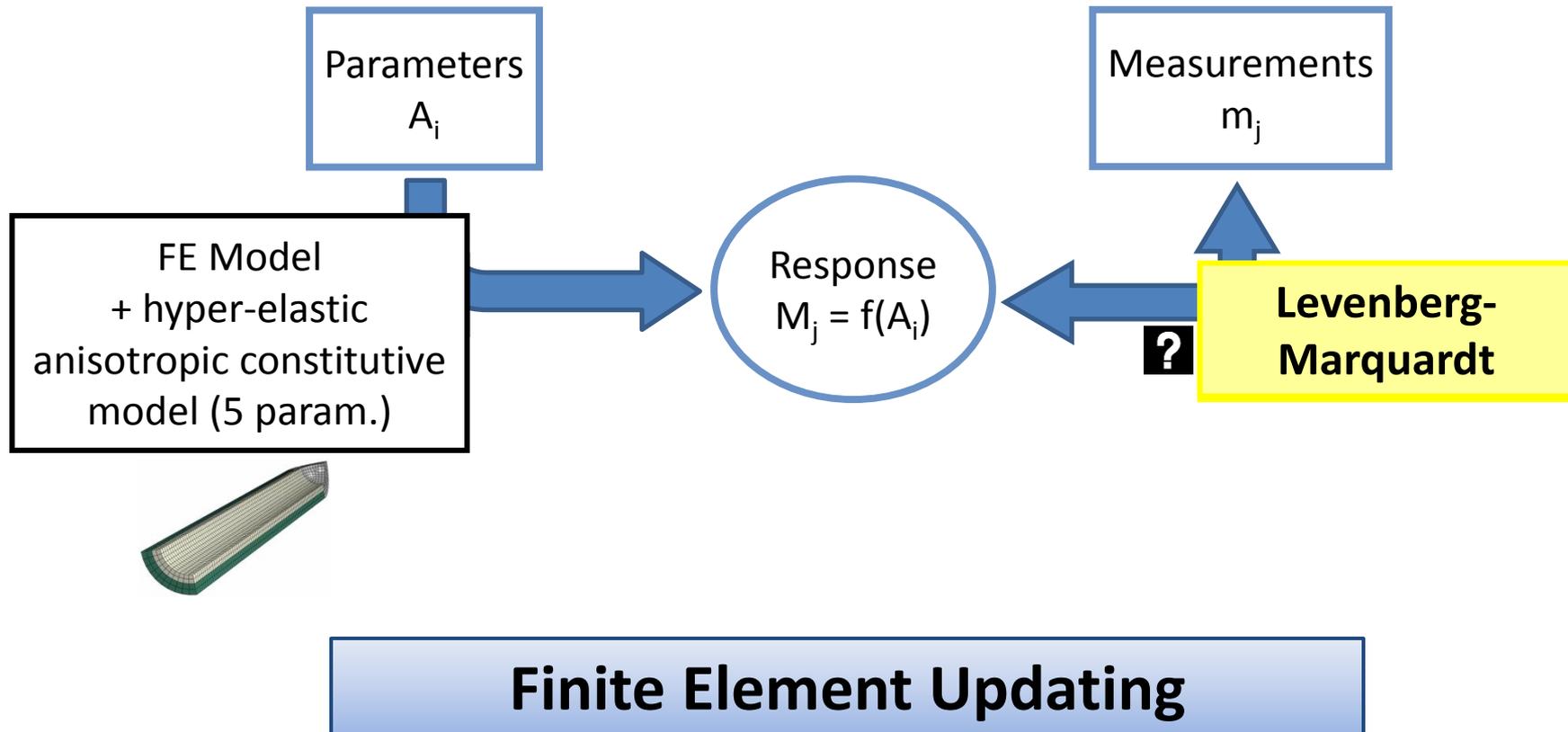
(coll. M. Sutton, U South Carolina, USA)



Mechanical properties of the artery?

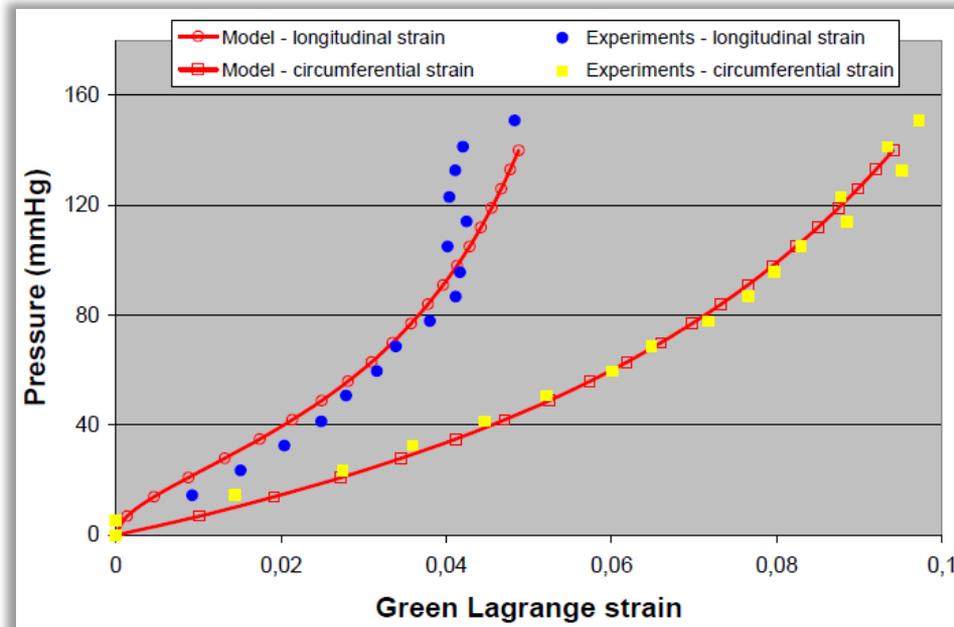
➔ In vitro ex.: Inflation/extension, mouse carotid artery

(coll. M. Sutton, U South Carolina, USA)



➔ In vitro ex.: Inflation/extension, mouse carotid artery

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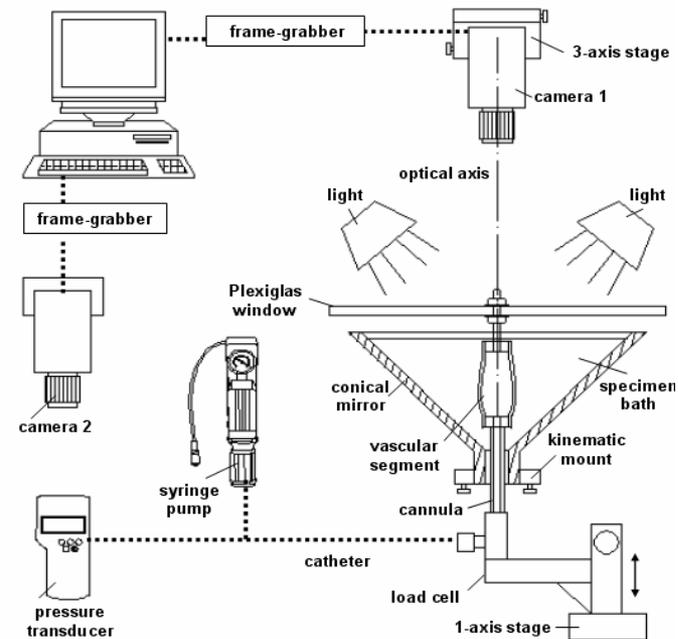


[Badel, 2012]

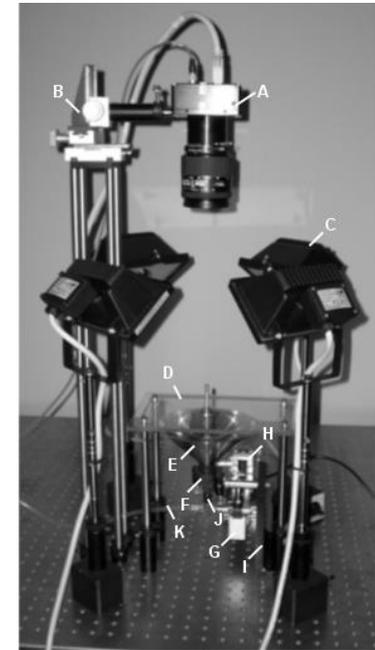
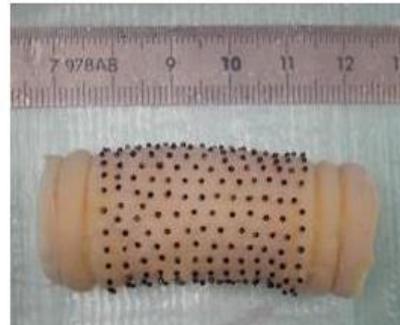
Finite Element Updating

➔ In vitro ex.: Inflation of a pig aorta

(coll. K. Genovese, U della Basilicata, Italy)



[Genovese, 2009]

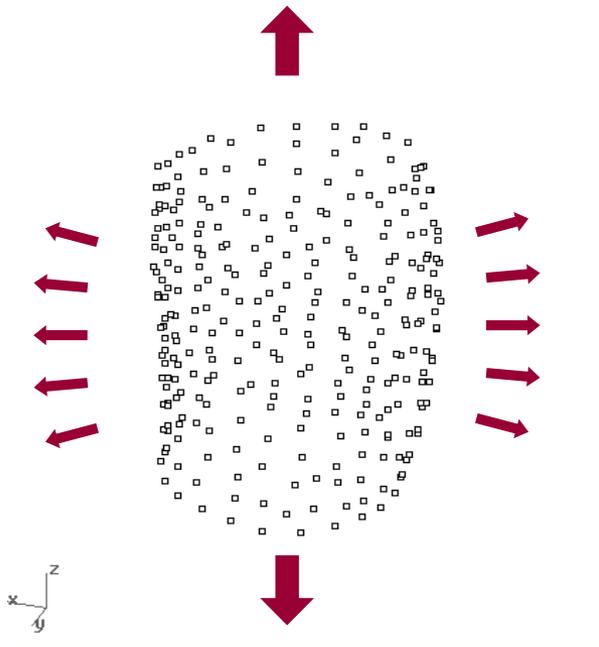


initial length
initial outer radius
initial thickness

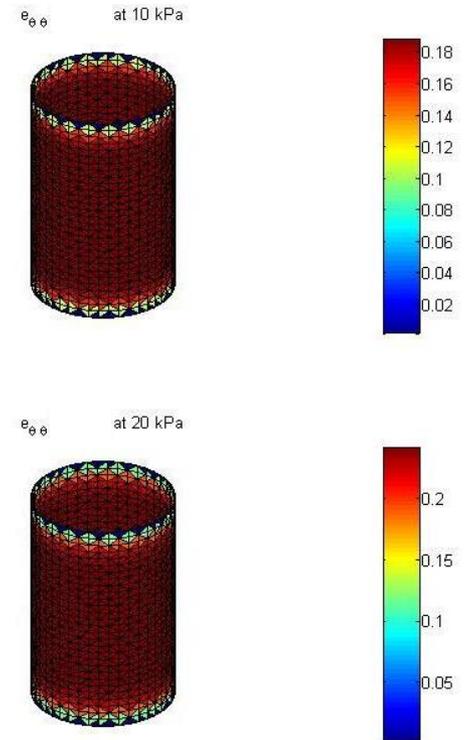
$L_0 \approx 35 \text{ mm}$
 $r_0 \approx 10 \text{ mm}$
 $e_0 \approx 1.3 \text{ mm}$

➔ In vitro ex.: Inflation of a pig aorta

(coll. K. Genovese, U della Basilicata, Italy)



Green Lagrange strain reconstruction



➔ In vitro ex.: Inflation of a pig aorta

Principle of virtual work
with given test functions

$$-\int_V \sigma_{ij} : \varepsilon_{ij}^* dV + \int_{\partial V} T_i u_i^* dS = 0$$

Constitutive model
(unknown parameters)

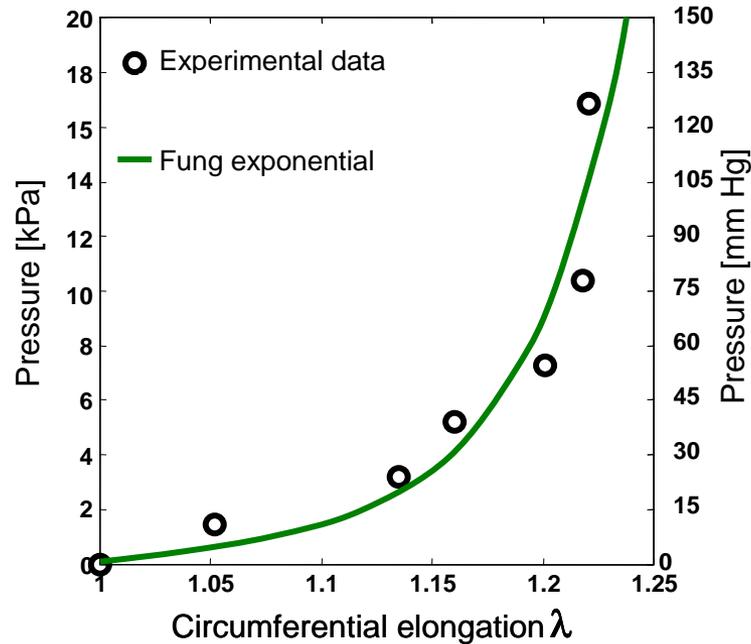
$$-\int_V \sigma_{ij}(\underline{\underline{\mathbf{E}}}, A) : \varepsilon_{ij}^* dV + \int_{\partial V} T_i u_i^* dS \overset{?}{\rightleftharpoons} 0$$

Equilibrium \Leftrightarrow Actual properties

Virtual Fields Method

➔ In vitro ex.: Inflation of a pig aorta

(coll. K. Genovese, U della Basilicata, Italy)



[Avril S, Badel P, Duprey A, 2010]

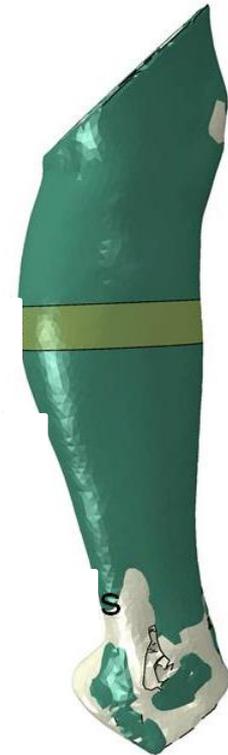
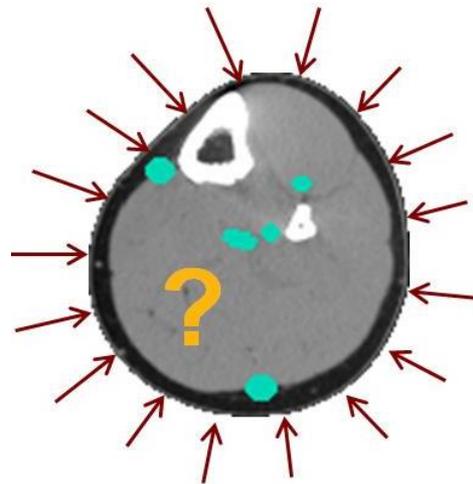
Virtual Fields Method

➔ In vivo ex.: Soft tissues of the leg

Pressure transmission mechanisms?? ... FE modeling of the action of compression socks



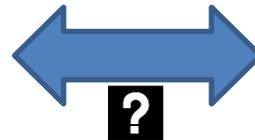
<http://www.Sigvaris.fr>



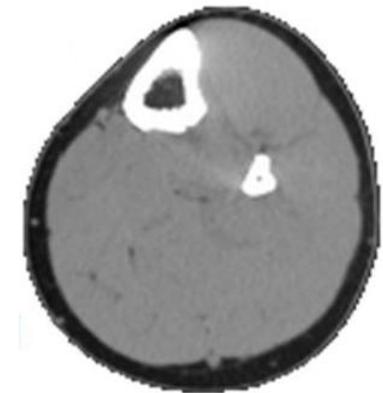
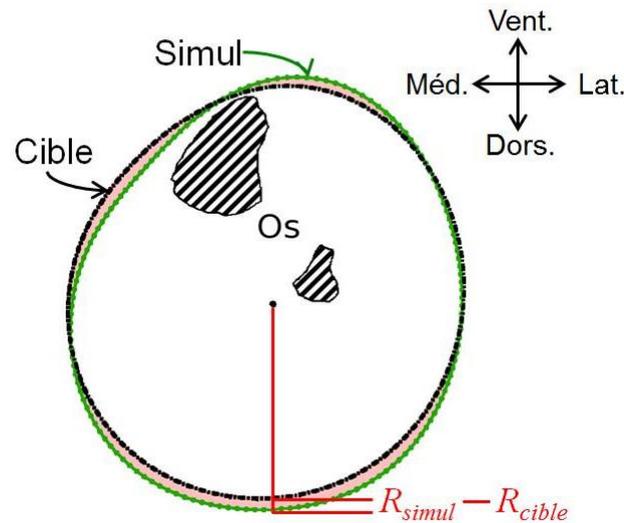
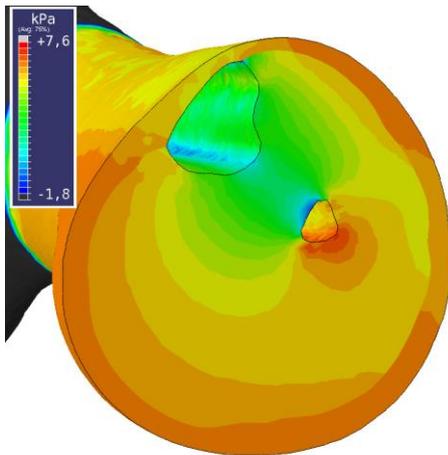
Properties of the leg's soft tissues?

➔ In vivo ex.: Soft tissues of the leg

FE model
Compressed leg



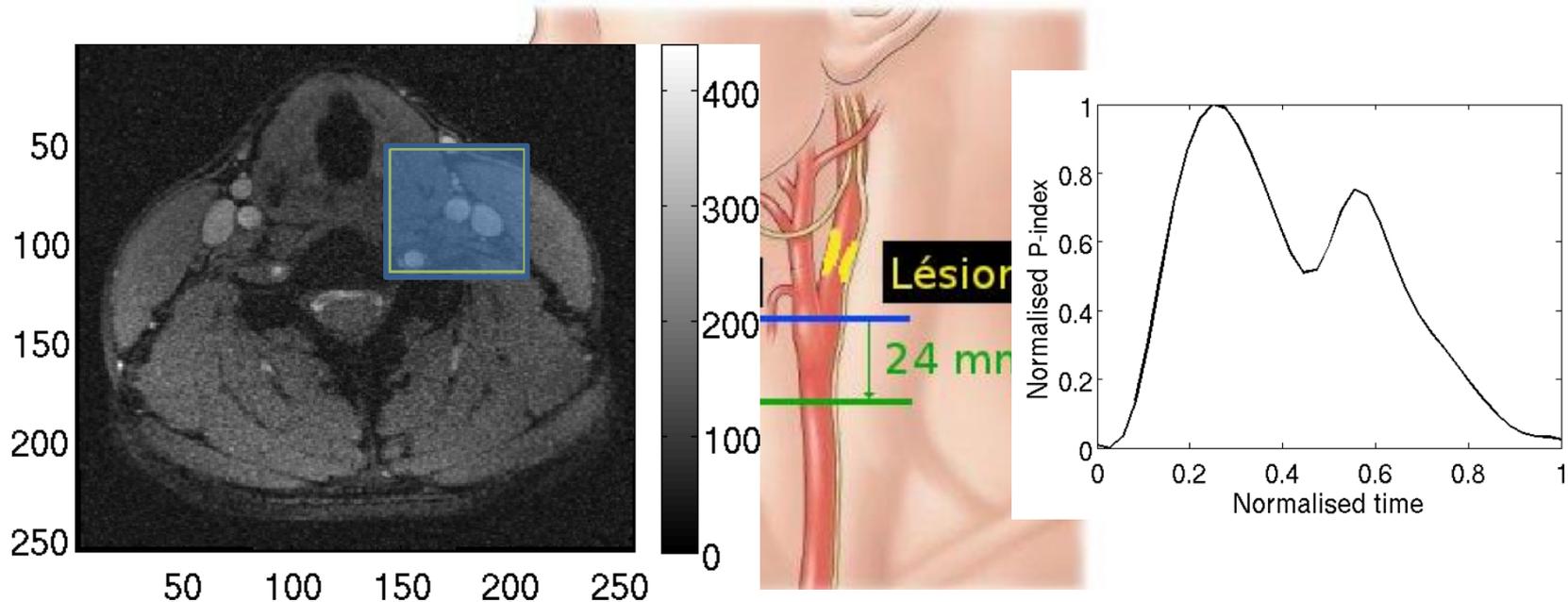
CT-scan image
leg + compression sock



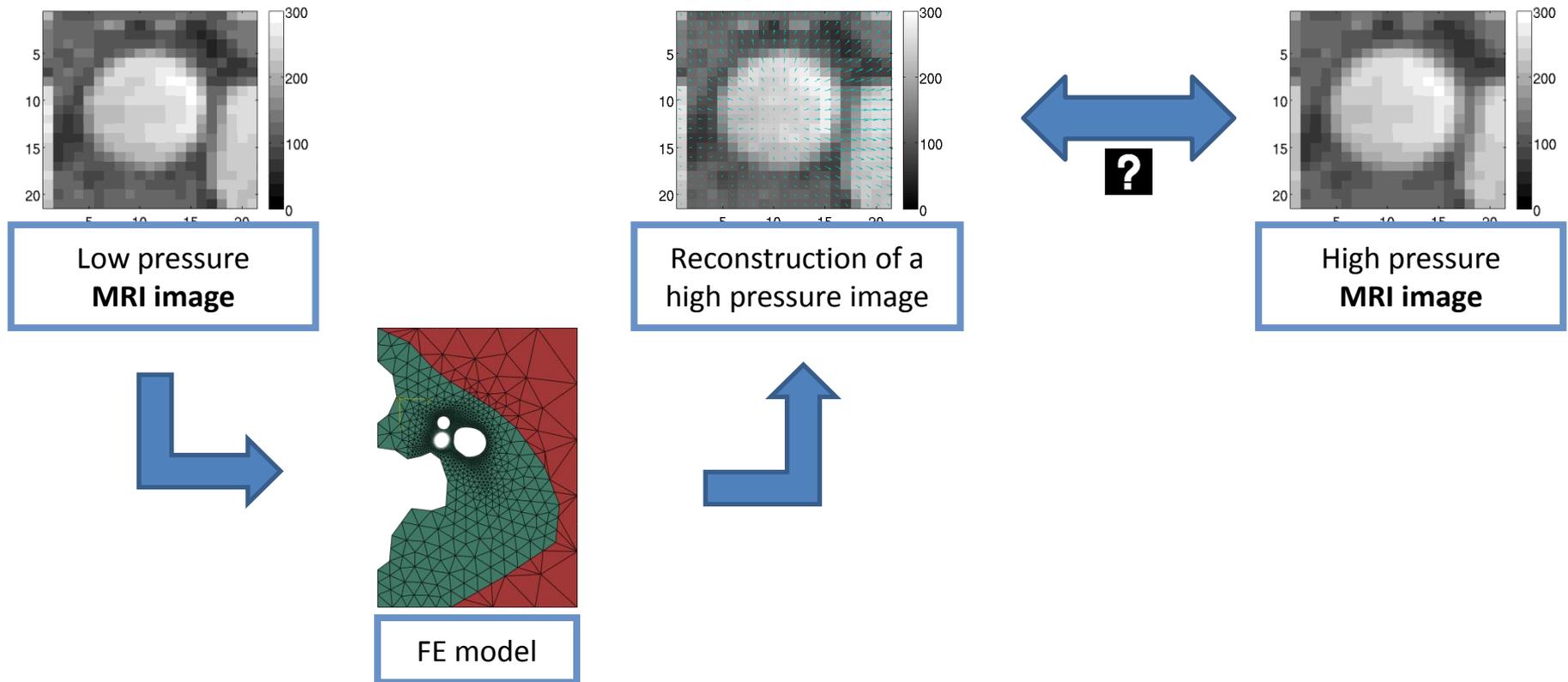
FE Updating + specific cost function

➔ In vivo ex.: elastic properties of human carotid arteries

The data we can access



➔ In vivo ex.: elastic properties of human carotid arteries



FE Updating + specific cost function

Application to ATAA

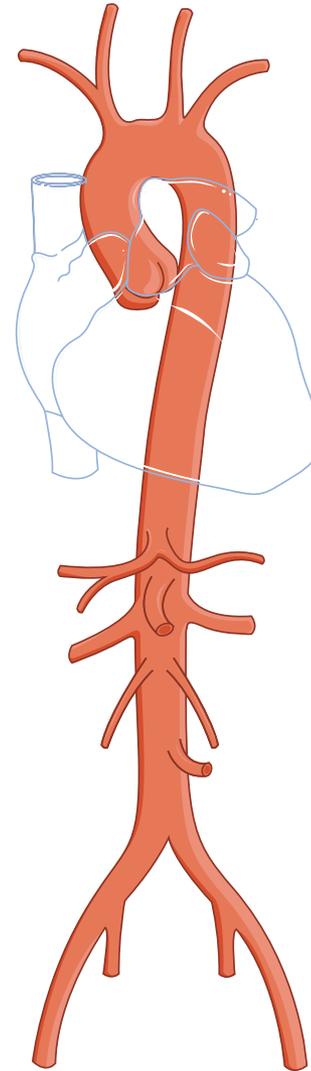
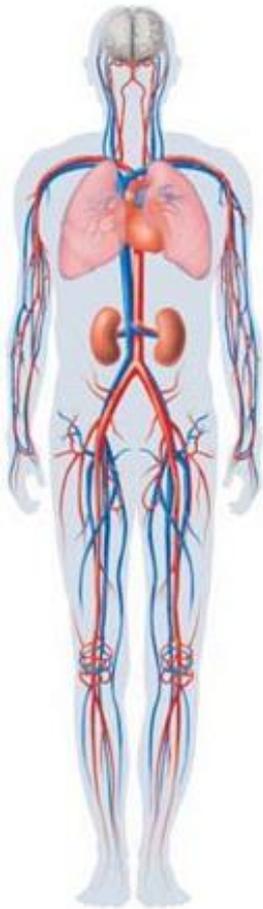
➔ Introduction

➔ A few words on identification methods applied to soft tissue biomechanics

➔ **Ascending thoracic aortic aneurysm**
(PhD A. Romo)

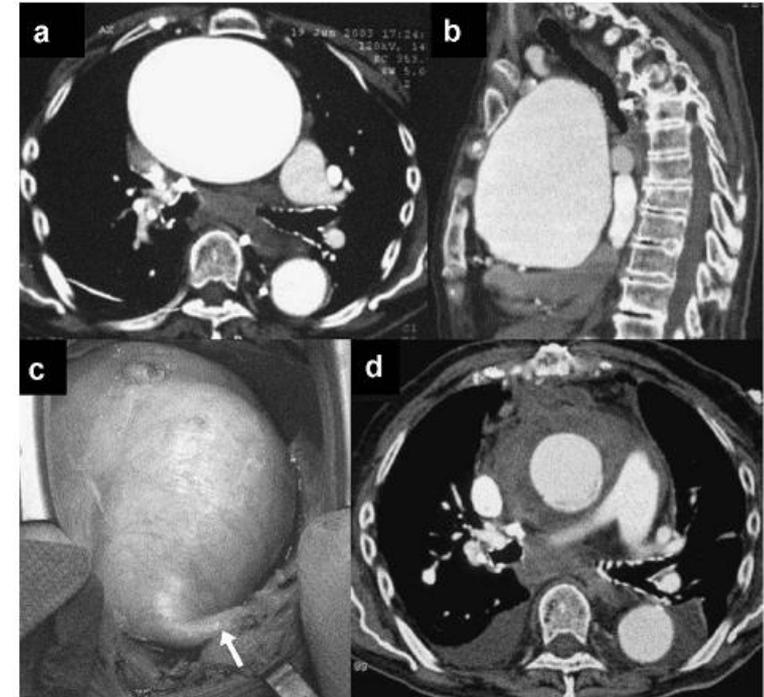
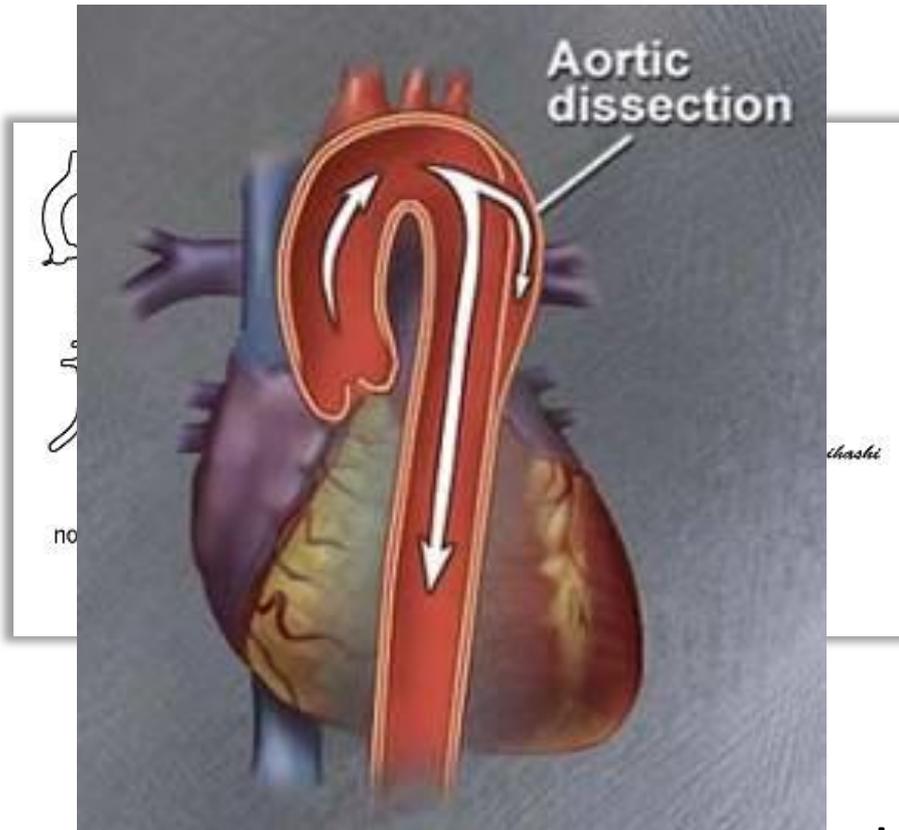
- Hyper-elastic model identification
- Rupture characterization

➔ Perspectives



Motivation

➔ ... examples of diseases...

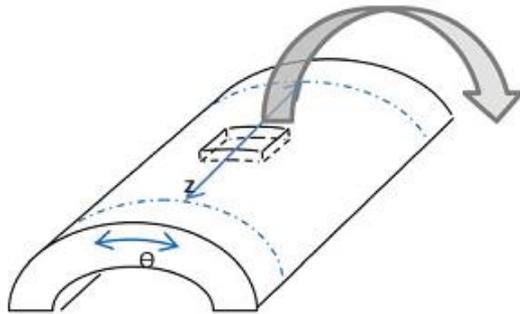


[Chavanon, 2006]

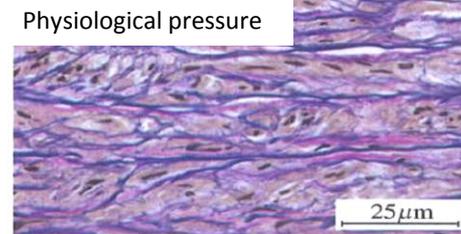
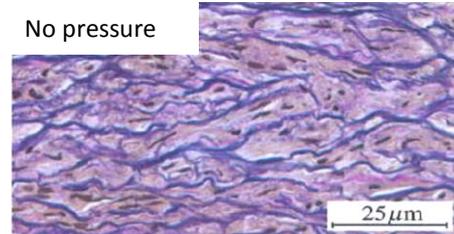
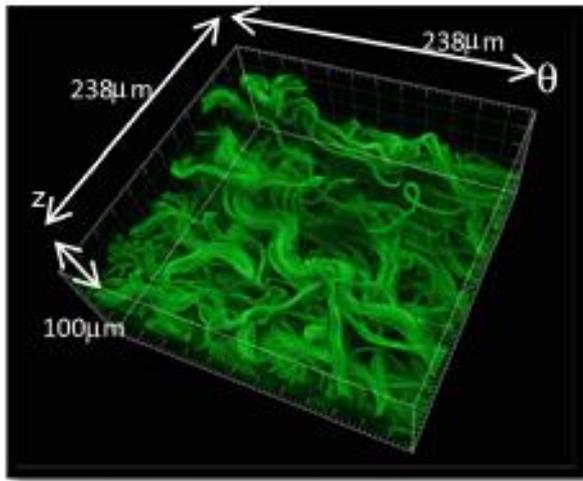
Ascending aorta

... which motivate research!

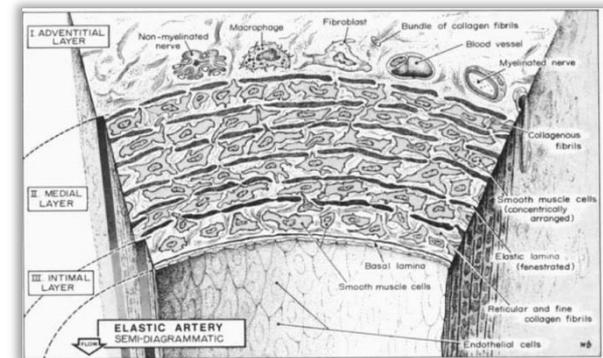
Structure



[Rezakhaniha et. al., 2011]



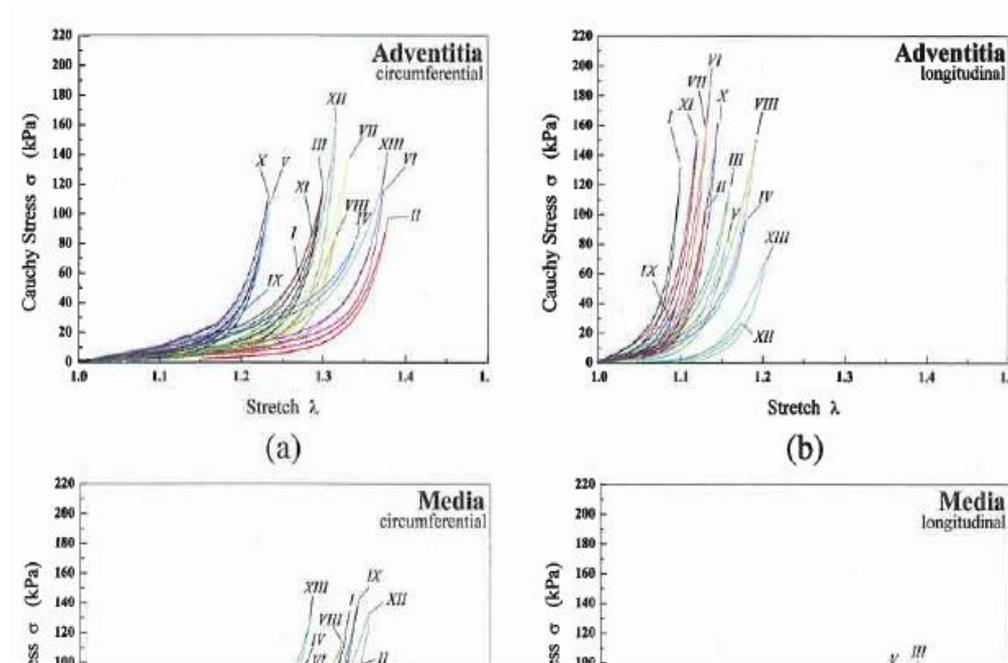
[Sommer et. al. 08]



[Rhodin, 1979]

Motivation

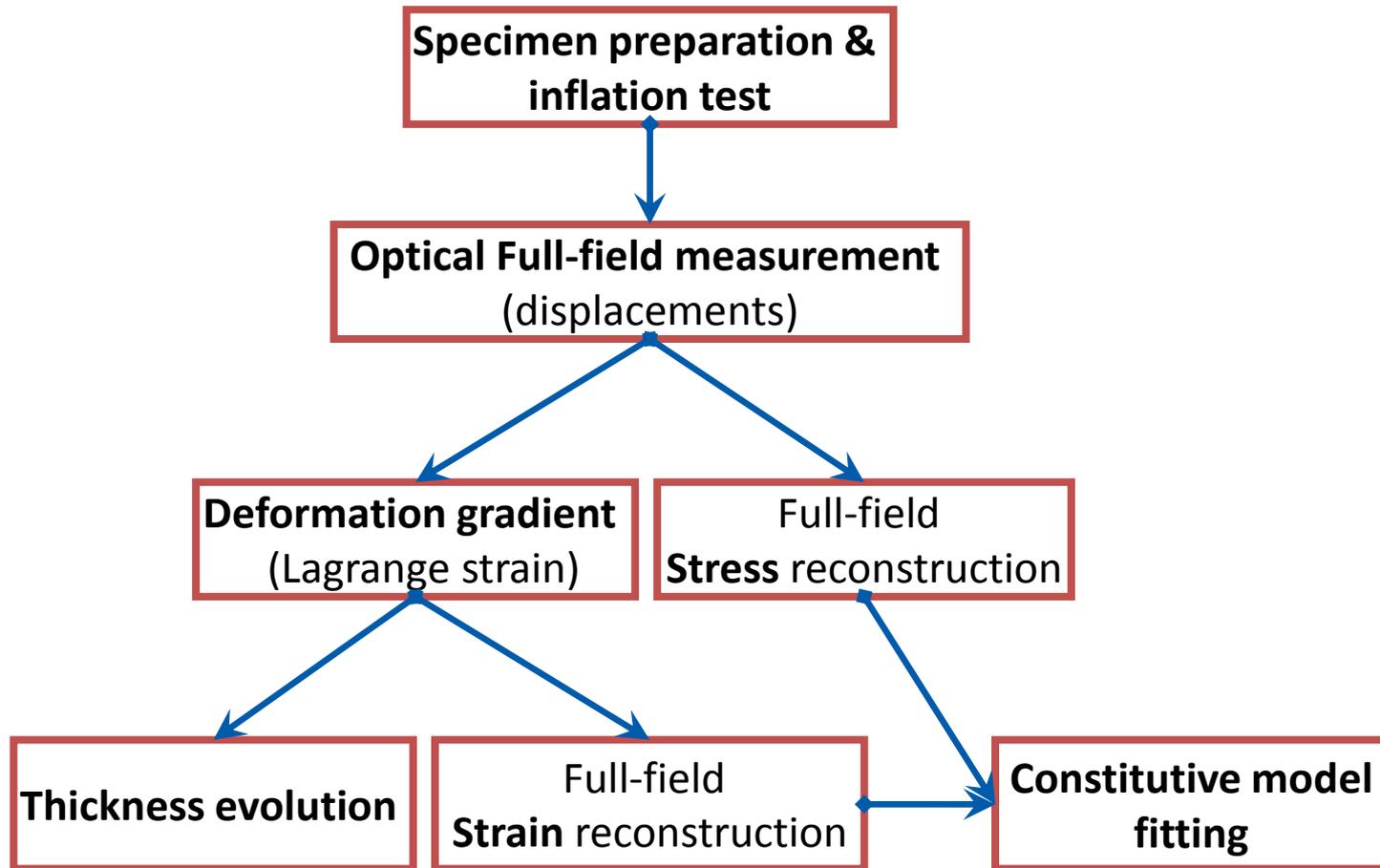
➔ Mechanical behavior well known, qualitatively



- ➔ Patient-specific properties?
- ➔ Aneurysm rupture strength/characterization?
- ➔ Damage/rupture mechanisms?

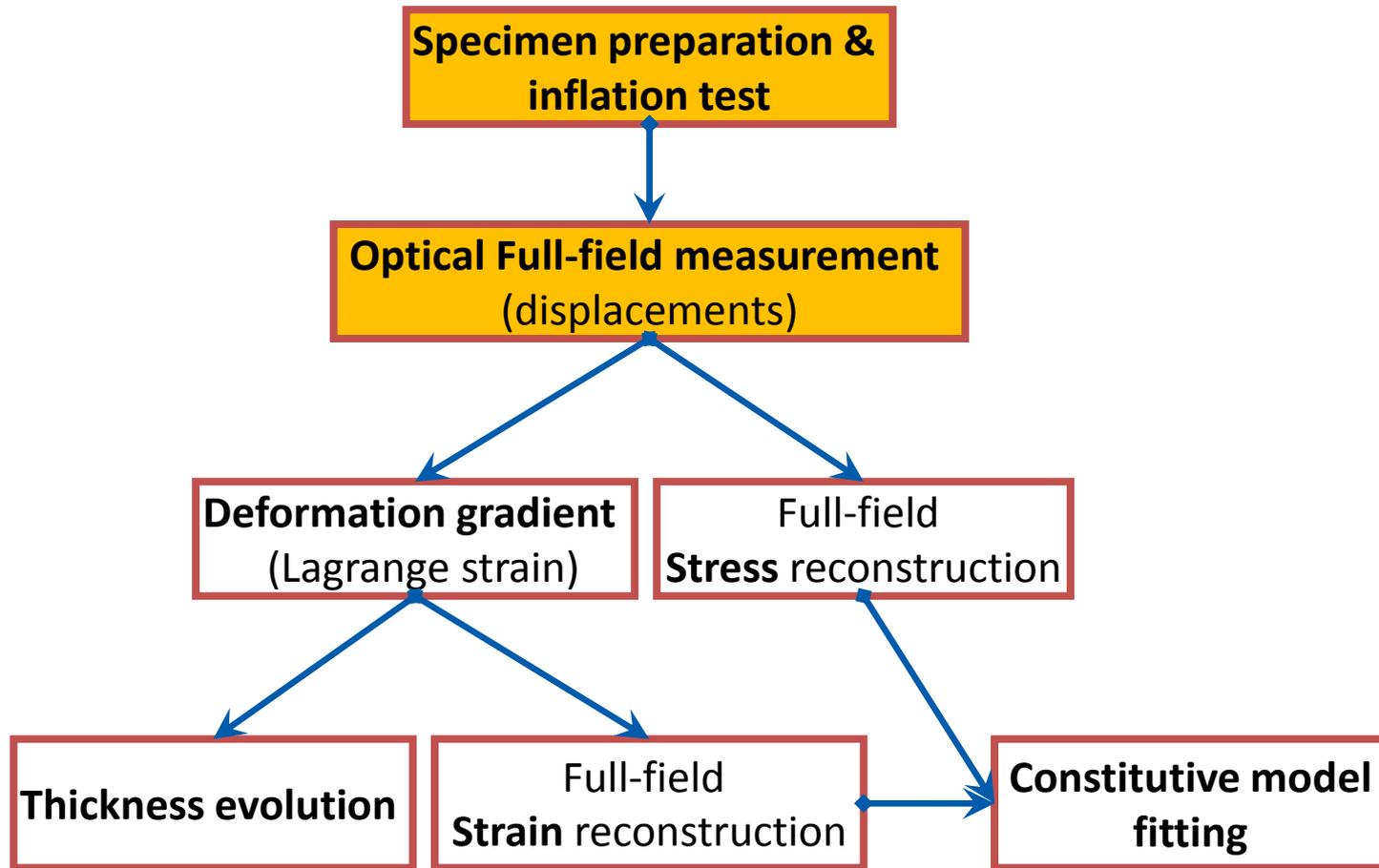
➔ Our methodology

[Romo A, Badel P, Duprey A, Favre J-P, Avril S. In vitro Analysis of Localized Aneurysm Rupture. *J Biomech.* 2014]

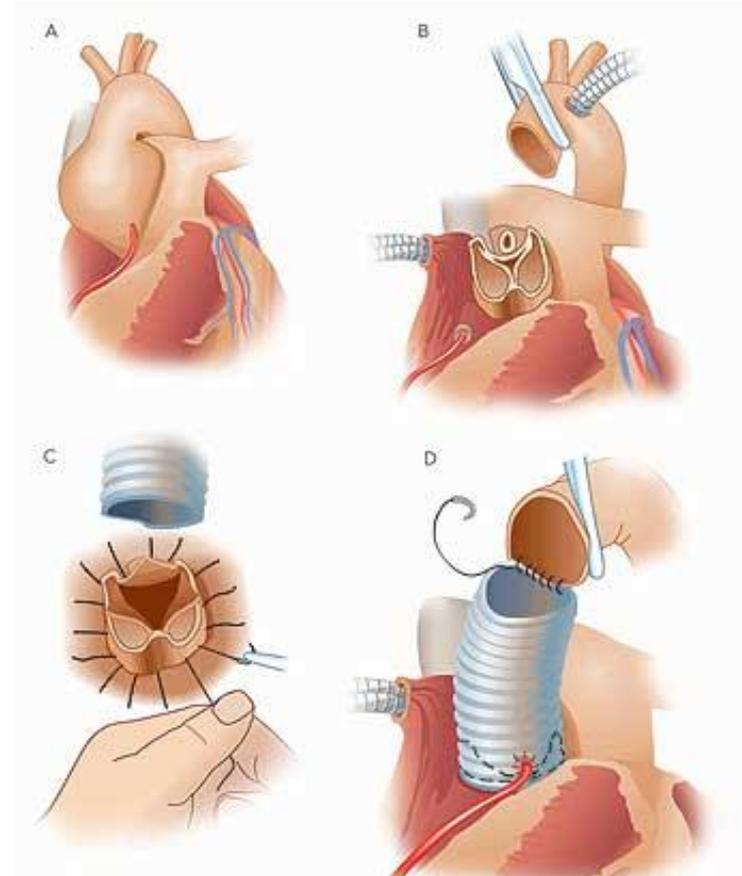
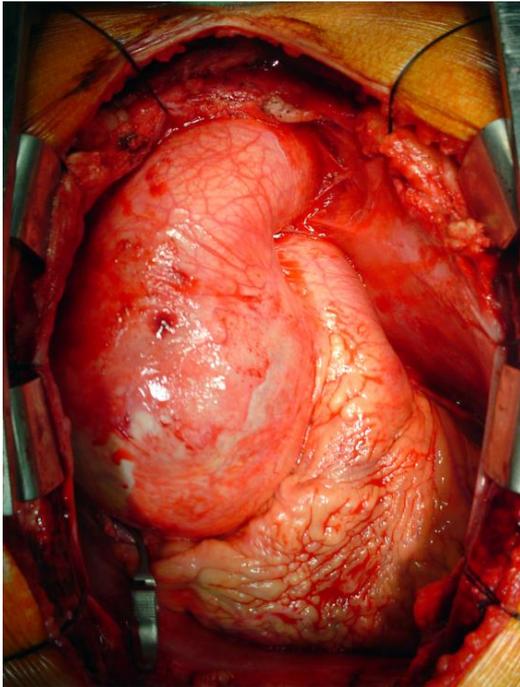


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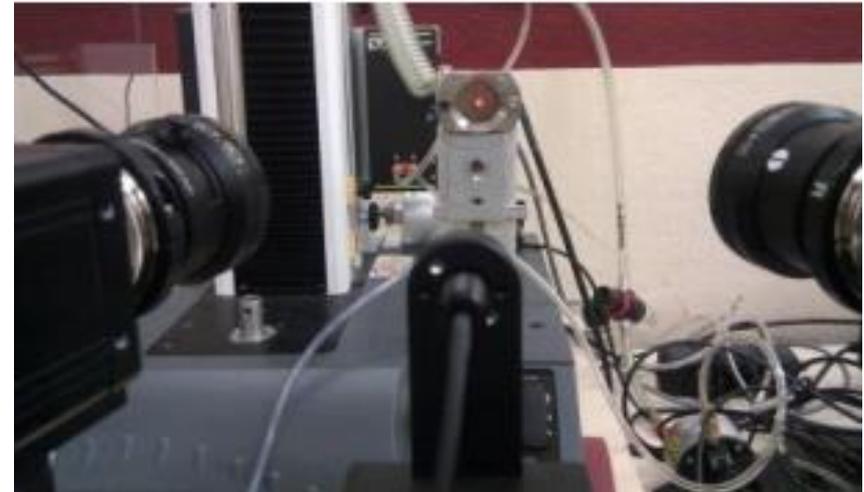
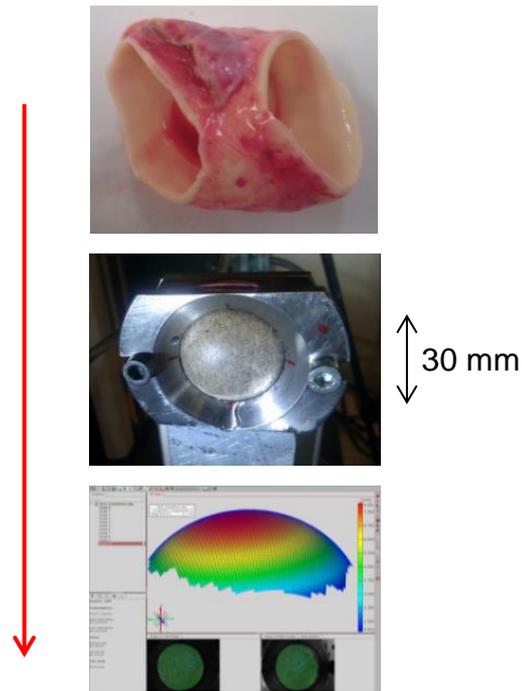


➔ Ascending Thoracic Aortic Aneurysm: Surgery



Experiment

➔ Exp: bulging test + full-field measurements

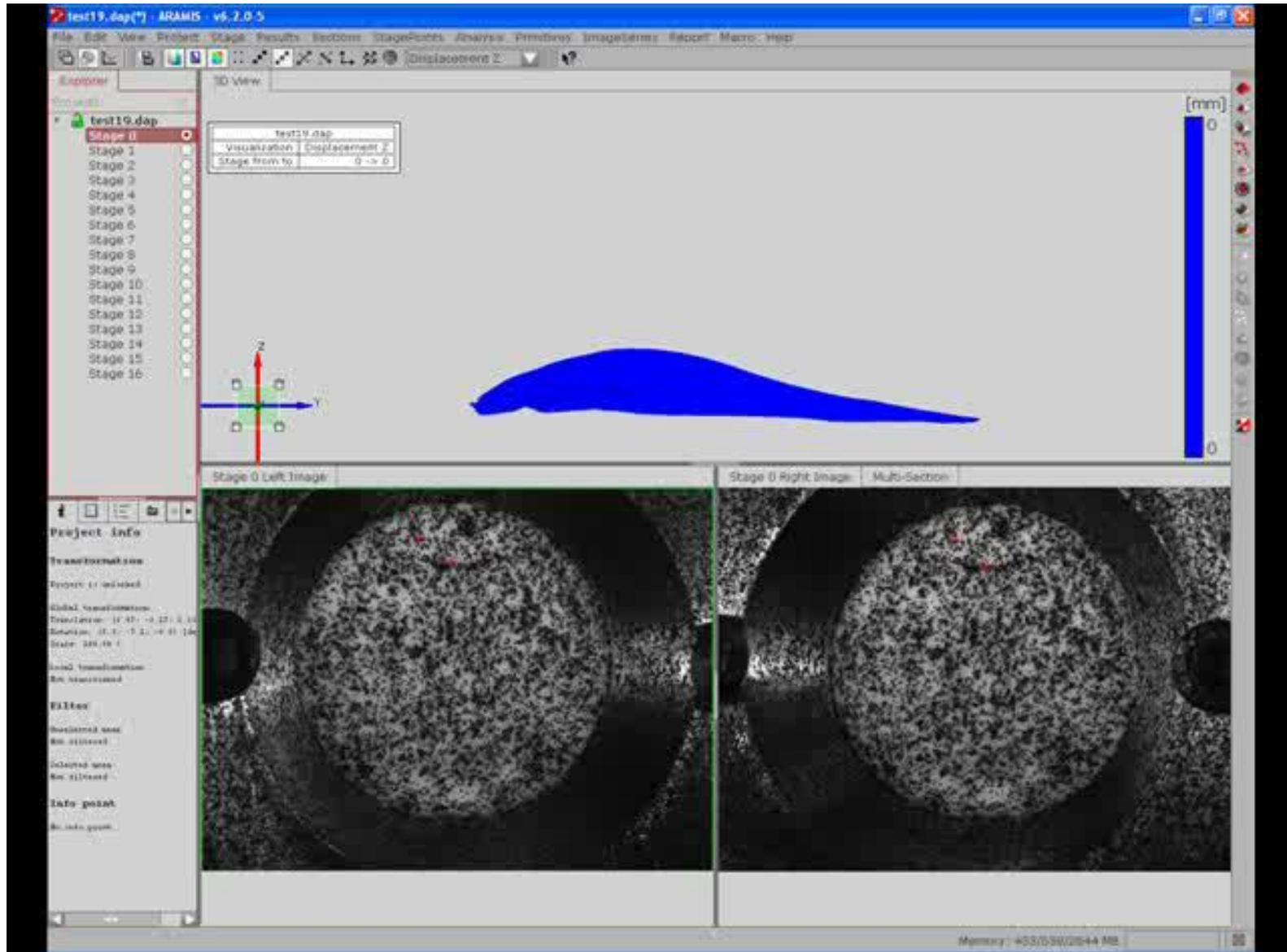


Reconstruction of a **3D surface**

+

Rupture stress calculation

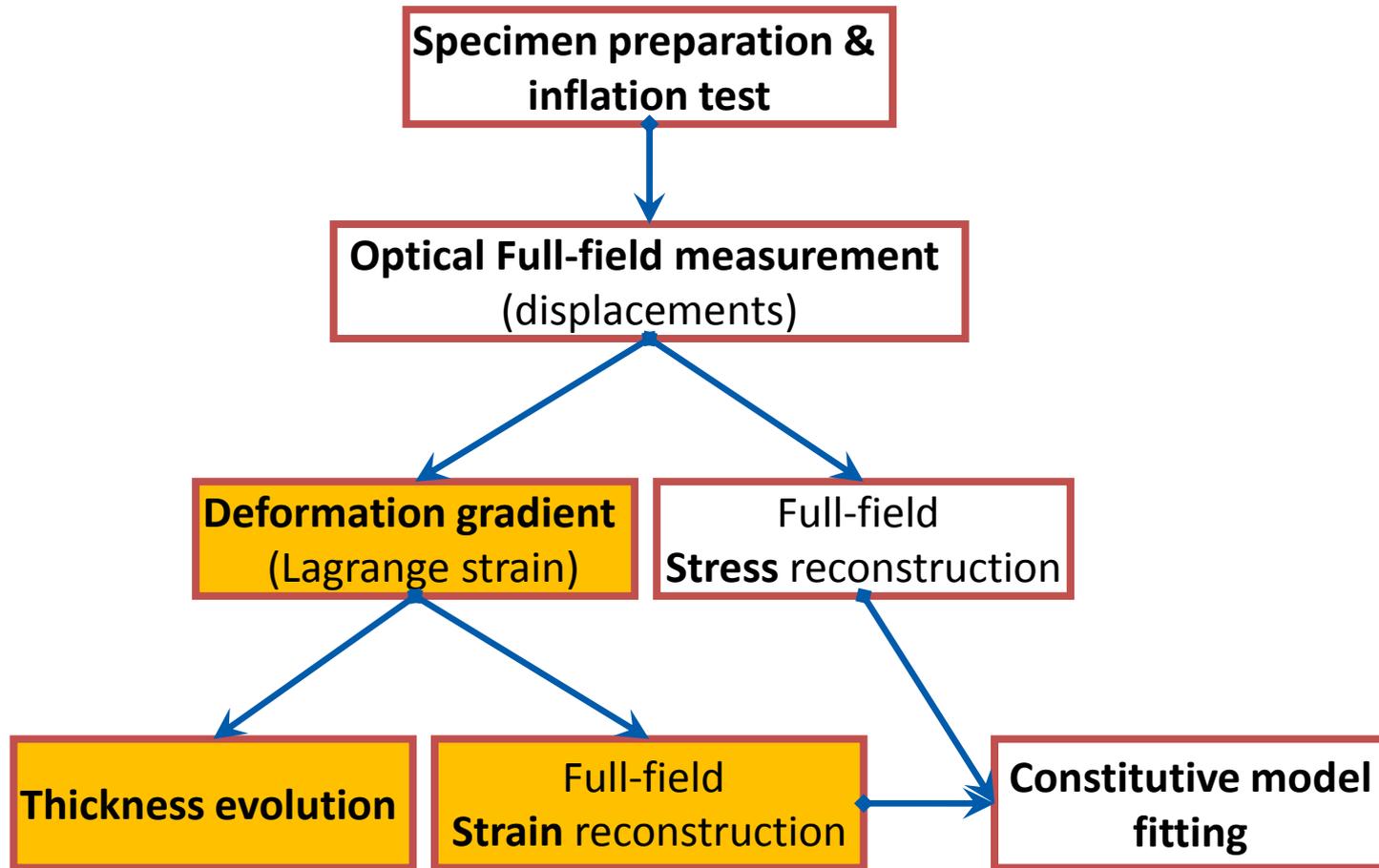
Experiment



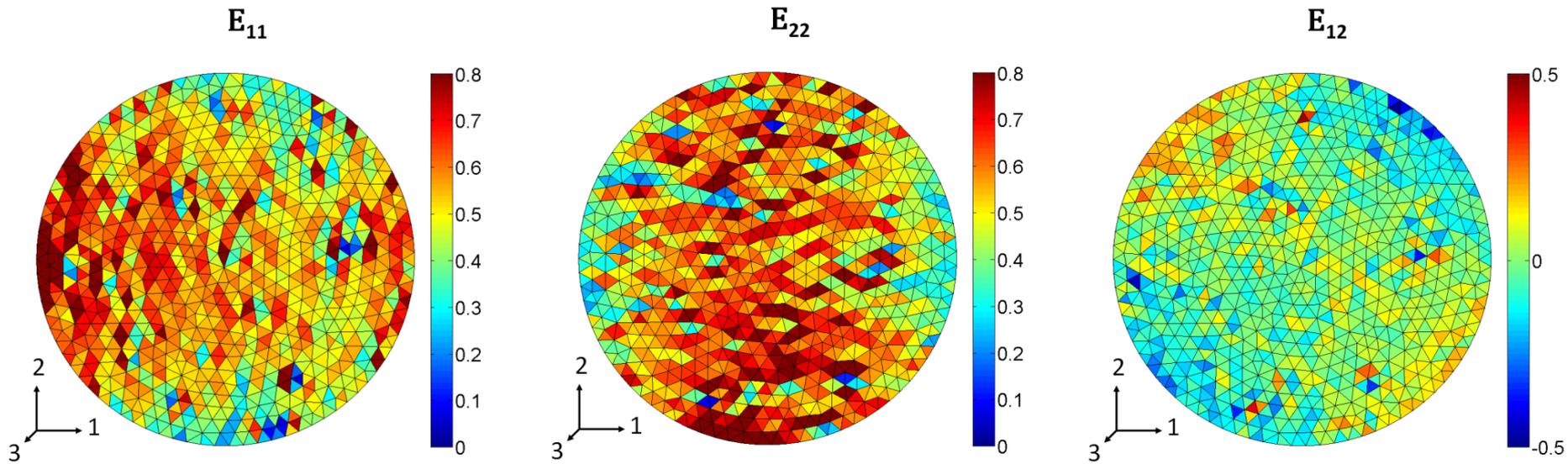
Methodology

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➔ Local strain field

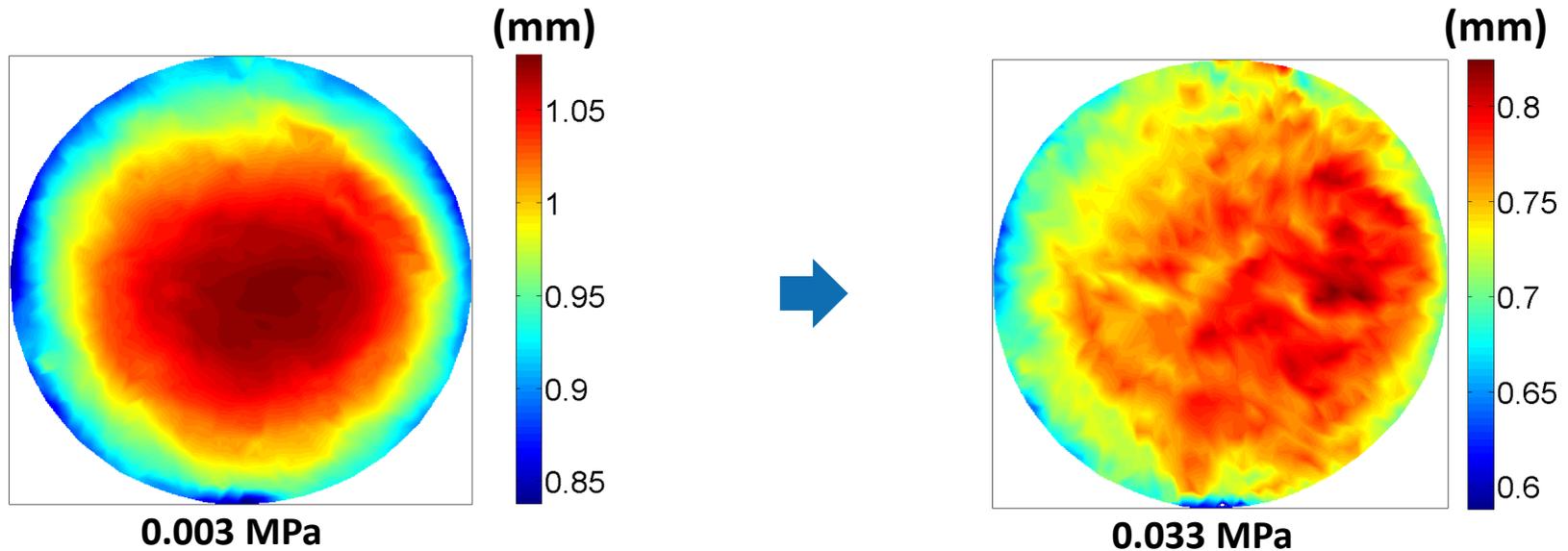


Green-Lagrange strain tensor

$$E = \frac{1}{2} (F^T F - I) = \begin{bmatrix} E_{11} & E_{12} \\ E_{21} & E_{22} \end{bmatrix}$$

➔ Thickness evolution

h_0 is the **homogeneous initial thickness**
+ assumption: **incompressibility**

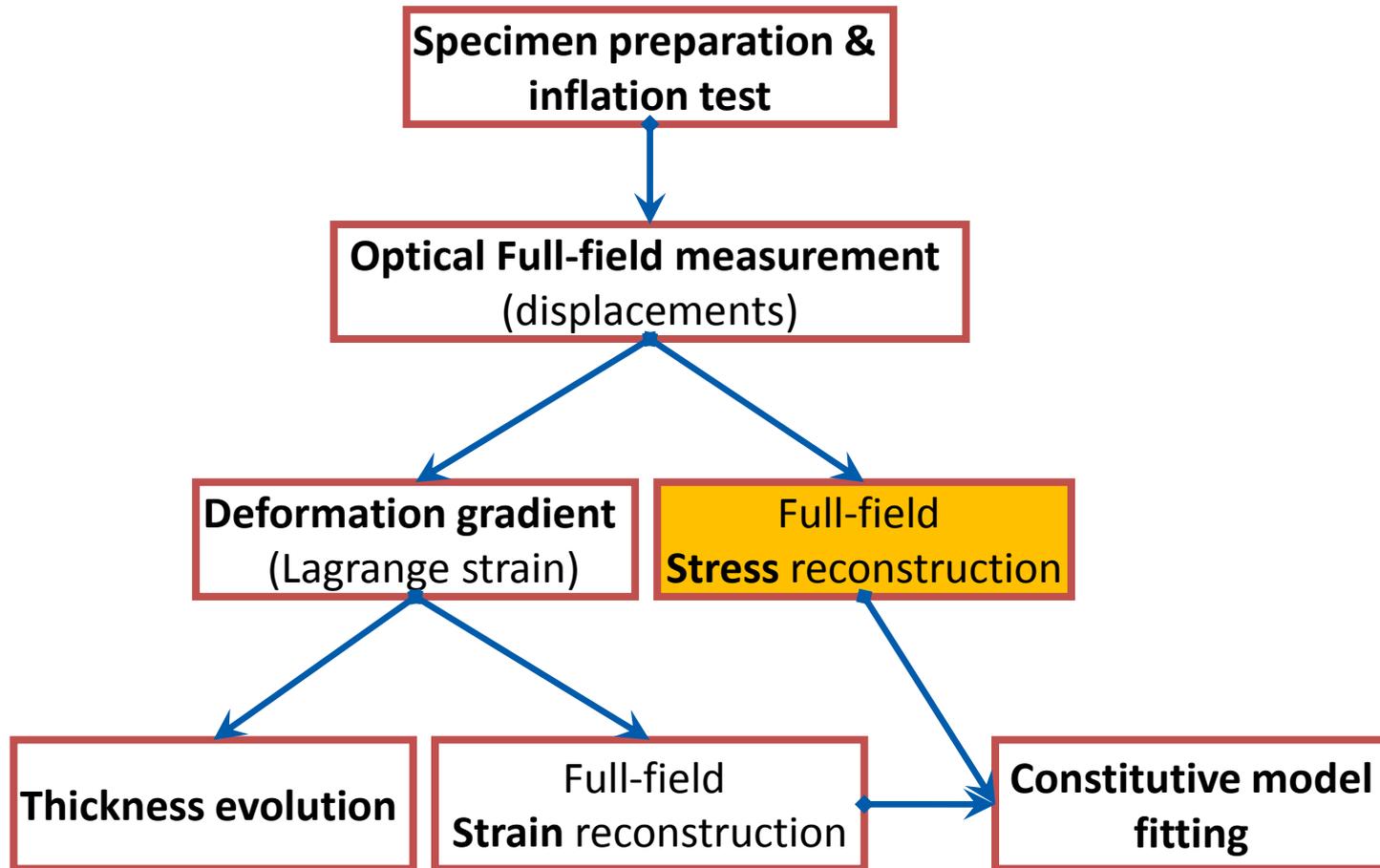


$$h = \frac{h_0}{F_{11}F_{22} - F_{21}F_{12}}$$

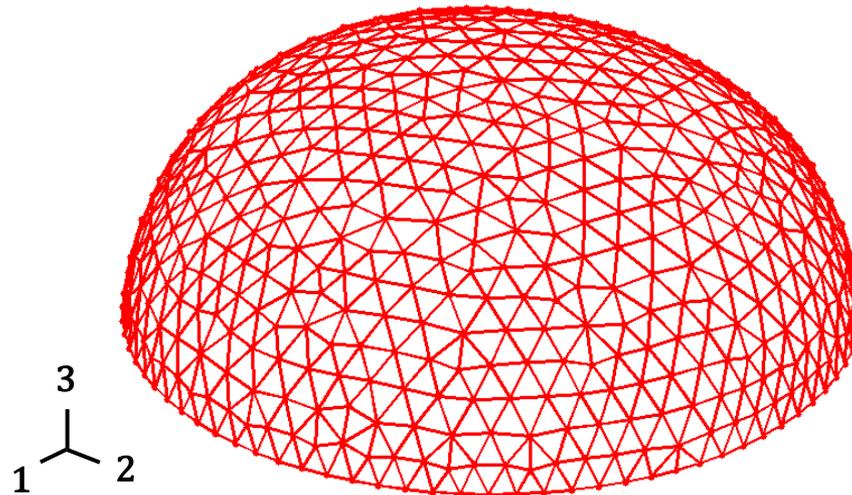
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➔ Membrane elastostatics

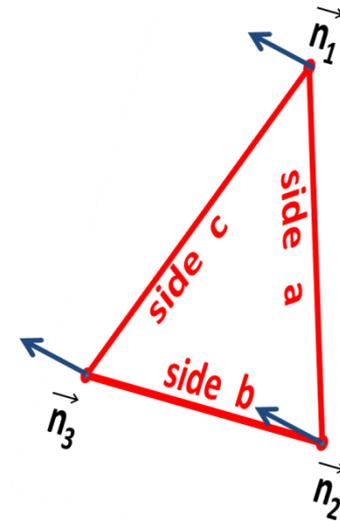
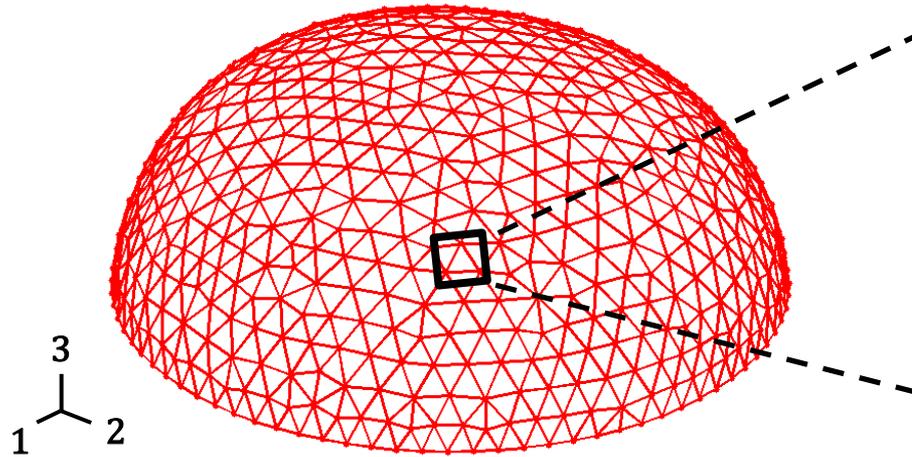


$$\text{div}(\boldsymbol{\sigma}) + f = 0$$

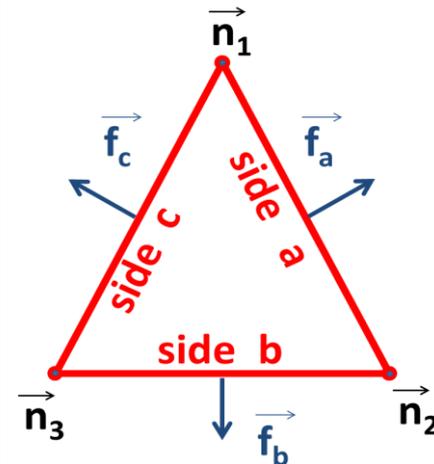


$$[A] \cdot [\boldsymbol{\sigma}] = [B]$$

➔ Membrane elastostatics



Define **normal** (at nodes) and **facet vectors**



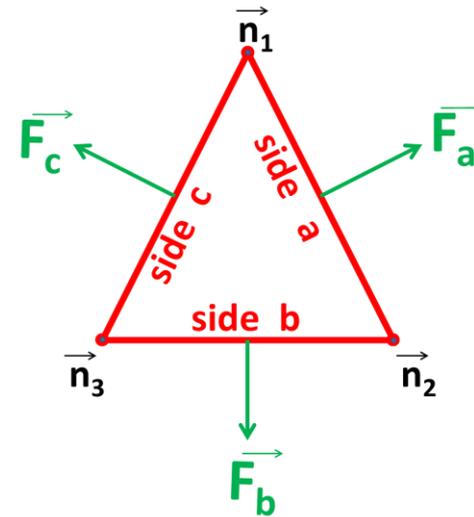
Membrane elastostatics

➔ Membrane elastostatics

Compute facet forces

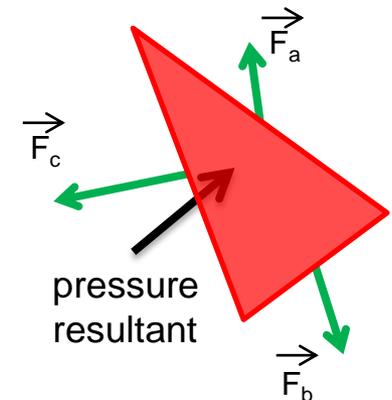
$$\vec{F}_b = \frac{\sigma_{n_2} + \sigma_{n_3}}{2} \cdot \vec{f}_b \cdot l_b \cdot h$$

$$\sigma = \begin{bmatrix} \sigma_{11} & \sigma_{12} \\ \sigma_{12} & \sigma_{22} \end{bmatrix}$$



LOCAL EQUILIBRIUM

$$\vec{F}_a + \vec{F}_b + \vec{F}_c = p \cdot s \cdot \frac{\vec{n}_1 + \vec{n}_2 + \vec{n}_3}{3}$$



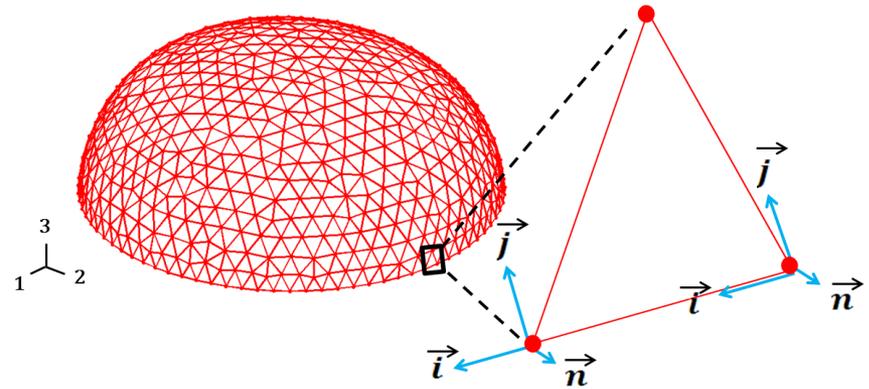
Membrane elastostatics

➔ Membrane elastostatics

Add boundary conditions

$$(\boldsymbol{\sigma} \cdot \vec{j}) \cdot \vec{n} = 0 \rightarrow \text{In-plane traction}$$

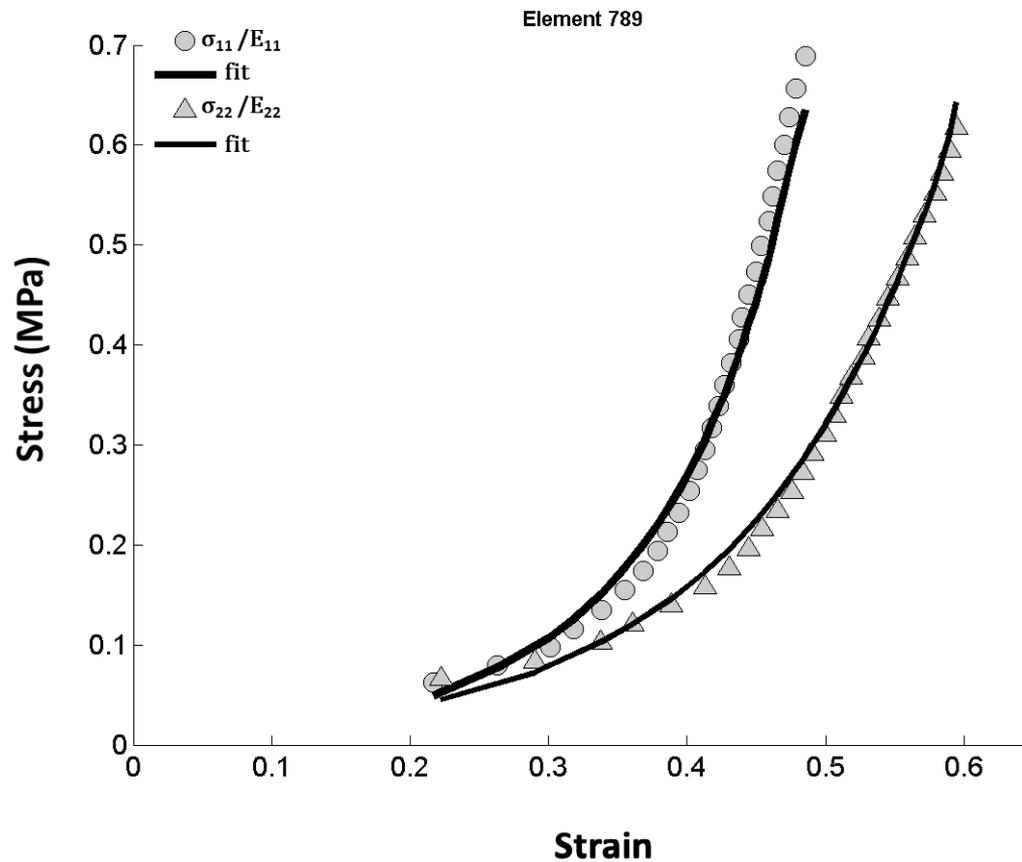
$$(\boldsymbol{\sigma} \cdot \vec{j}) \cdot \vec{i} = 0 \rightarrow \text{No shear}$$



FINAL SYSTEM TO BE SOLVED

$$[A] \cdot [\boldsymbol{\sigma}] = [B]$$

	$\langle 3609 \times 1932 \rangle$	$\langle 1932 \times 1 \rangle$		$\langle 3609 \times 1 \rangle$	
	↓	↓	↓	↓	
3xElements	3xNodes	Unknowns		3xElements	



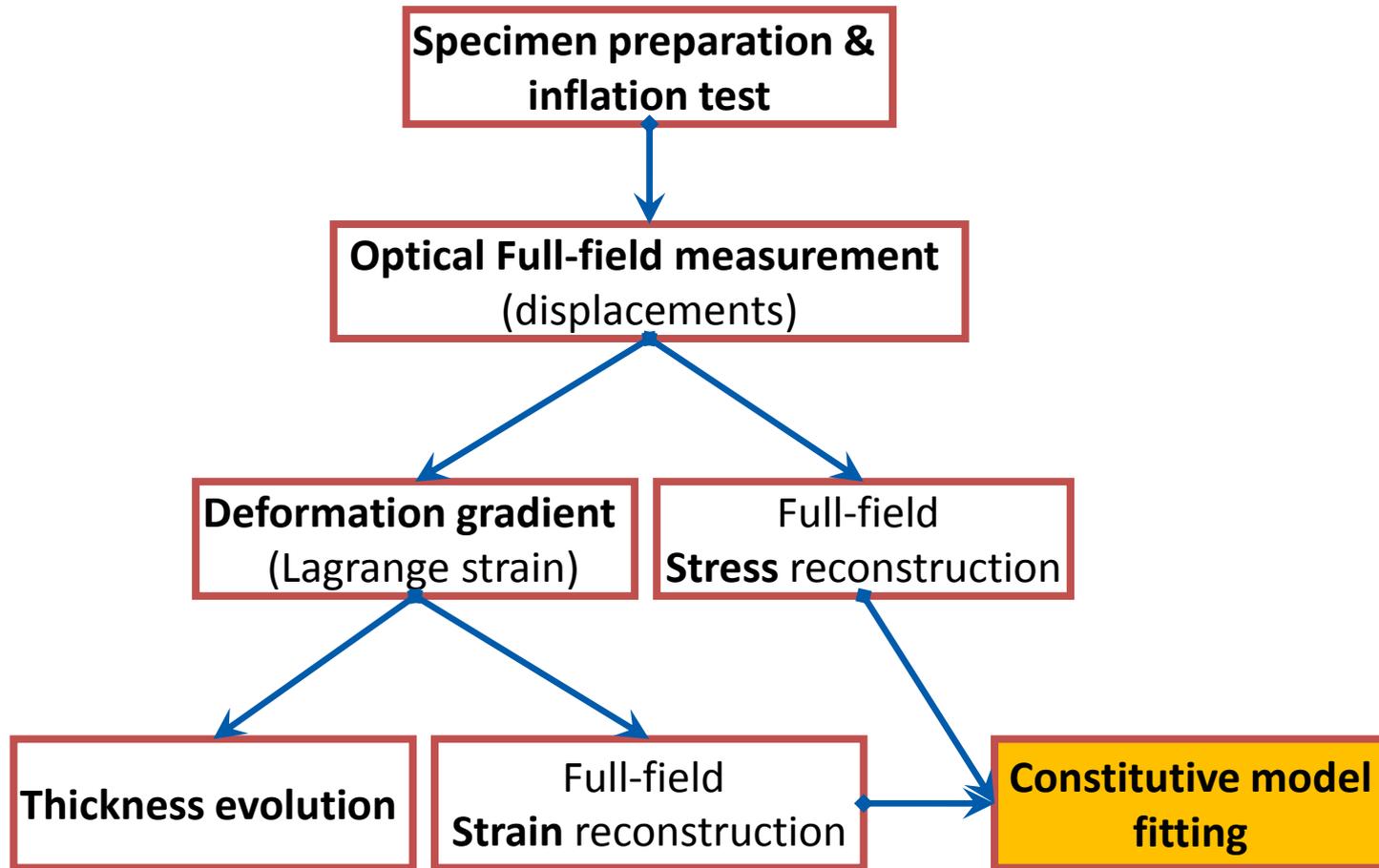
Stress field reconstructed without a constitutive model

!!

Methodology

➔ Our methodology

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Hyper-elastic model identification

Strain energy function

$$\Psi = \Psi_g + \Psi_f$$

Ground substance + elastin
Isotropic

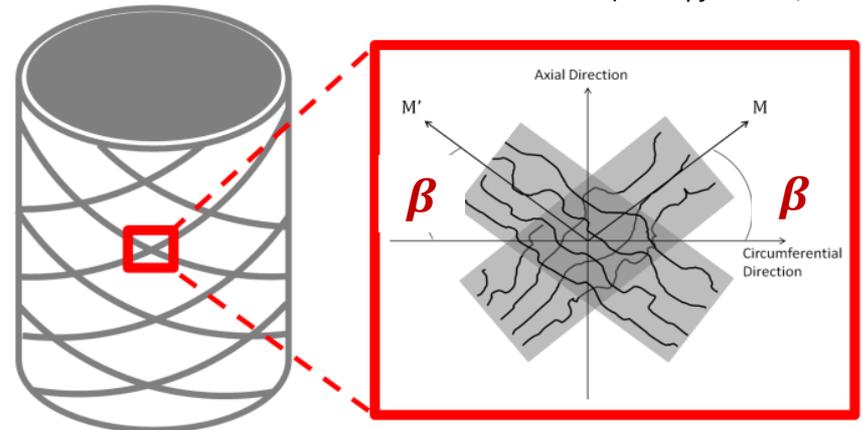
Collagen fibers
Anisotropic

$$\Psi_g = C_{10}(\bar{I}_1 - 3)$$

C_{10} parameter fixed at **5 kPa**
(Weisbecker et al. 2012)

$$\Psi_f(I_4, I_6) = \frac{k_1}{2k_2} \sum_{i=M, M'} \left\{ \exp \left[k_2 (\lambda_i^2 - 1)^2 \right] - 1 \right\}$$

(Holzapfel et al., 2000)



+ incompressibility

➔ Hyper-elastic model identification

Strain energy function

$$\Psi = \Psi_g + \Psi_f$$



2nd Piola-Kirchhoff stress tensor:

$$S = S_g + S_f = \frac{\partial \Psi_g}{\partial E} + \frac{\partial \Psi_f}{\partial E}$$



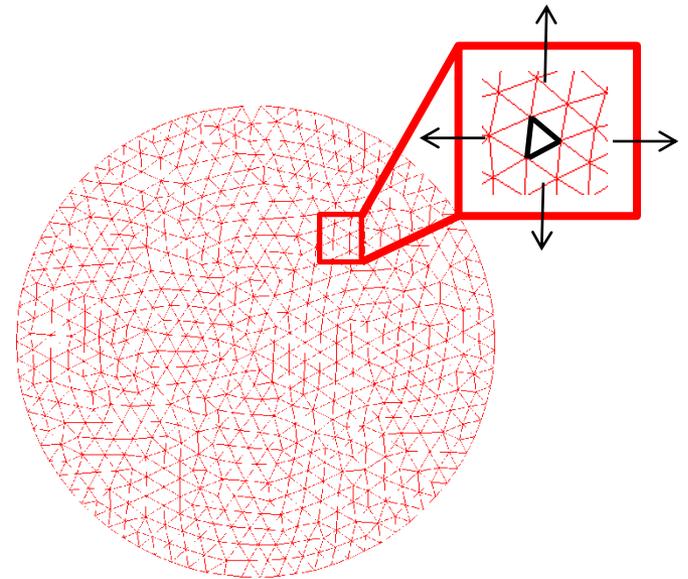
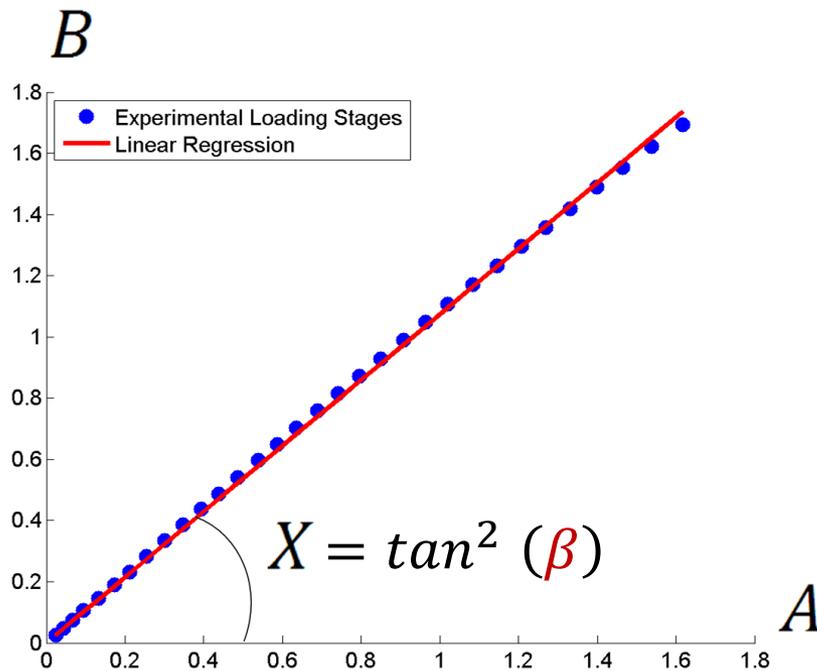
Cauchy stress tensor:

$$\sigma = F \cdot S_g \cdot F^T + \underbrace{F \cdot S_f \cdot F^T}_{\text{circled}} + cI$$

Local identification method

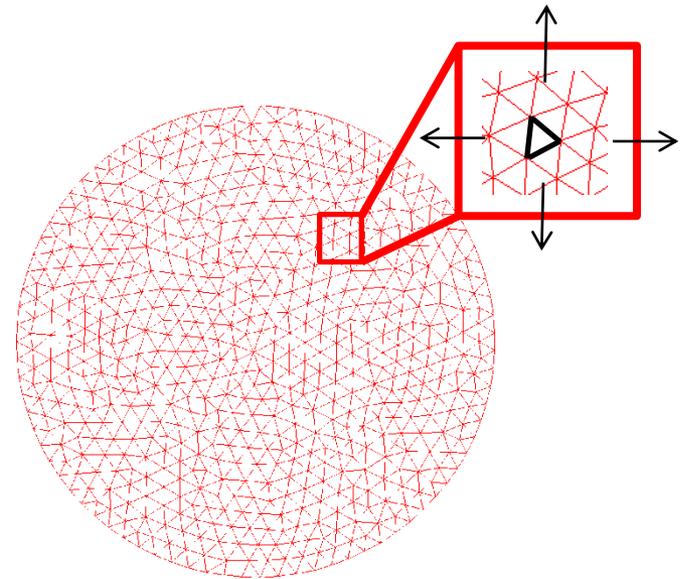
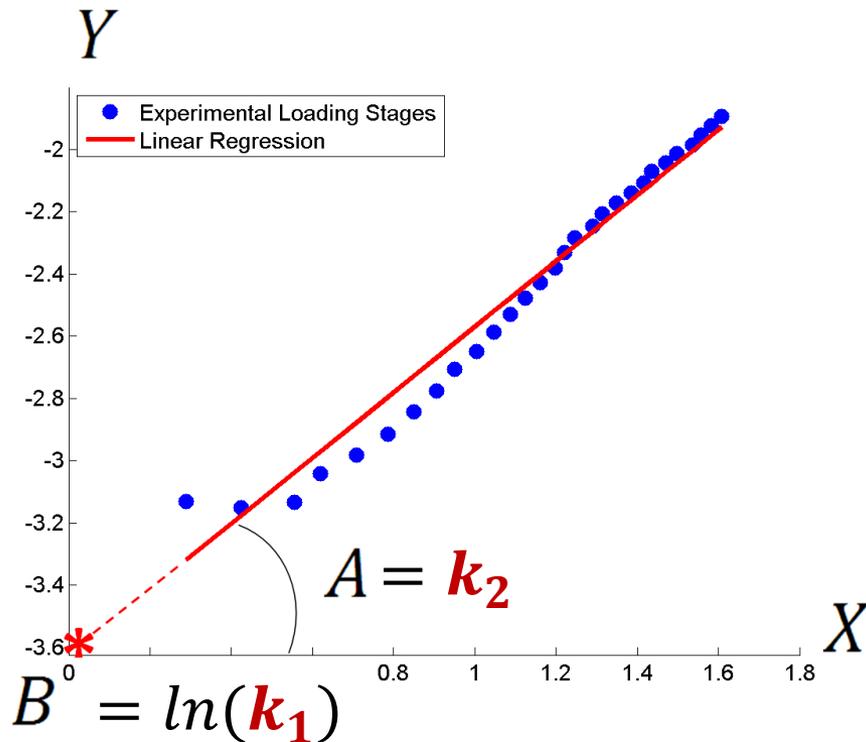
➔ Hyper-elastic model identification

$$A(\sigma', F) \tan^2 \beta + B(\sigma', F) = 0$$



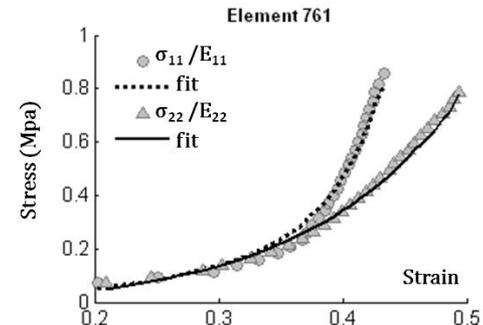
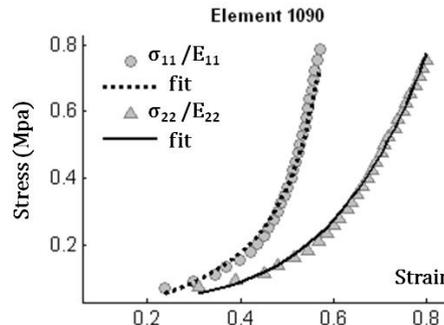
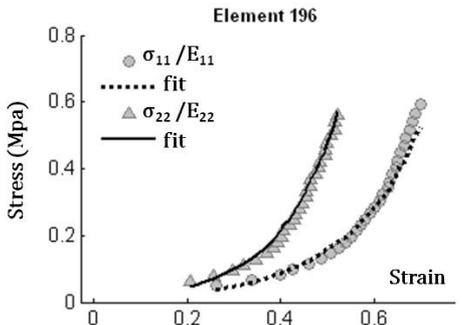
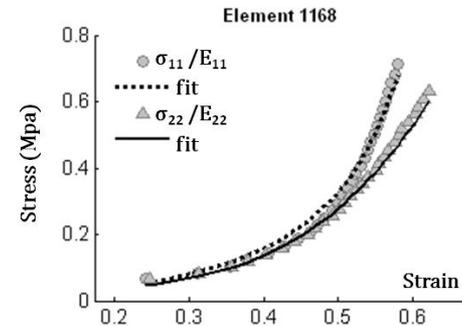
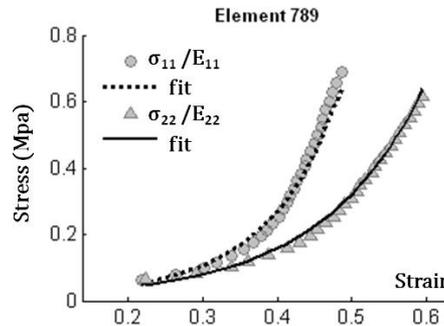
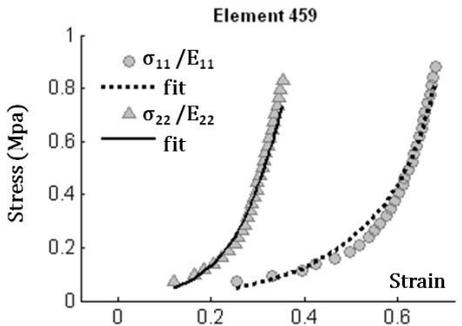
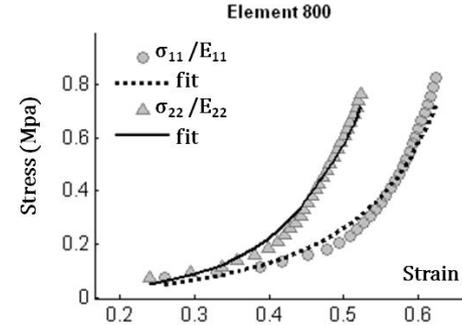
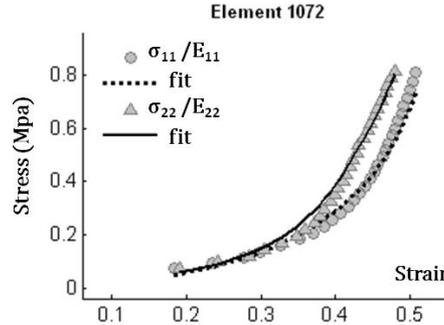
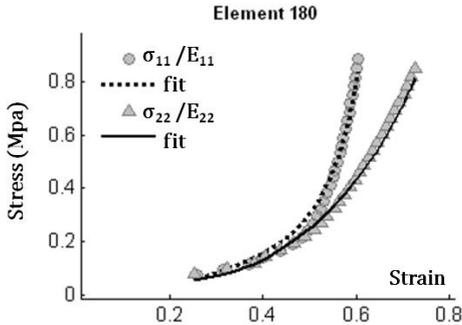
Hyper-elastic model identification

$$\sigma'_{11} = 4k_1 \lambda_1^2 \cos^2(\beta) (\lambda_f^2 - 1) e^{k_2 (\lambda_f^2 - 1)^2}$$



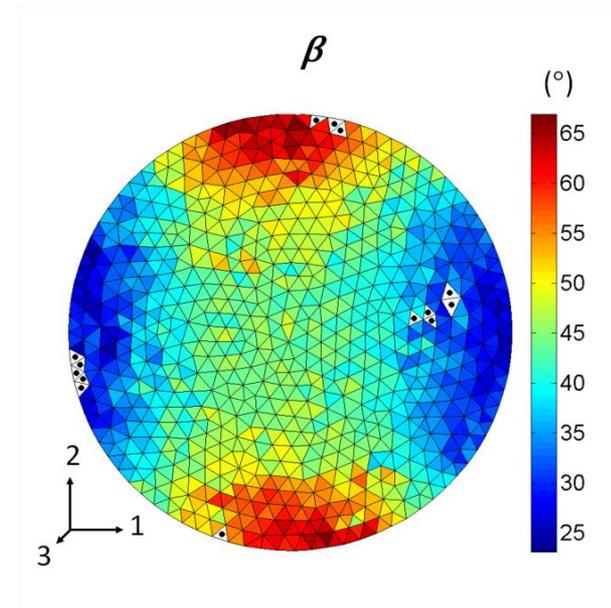
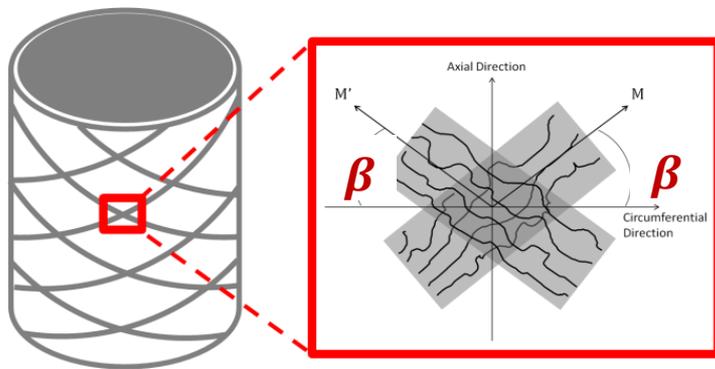
Results

➔ Results: bi-directional experimental fitting



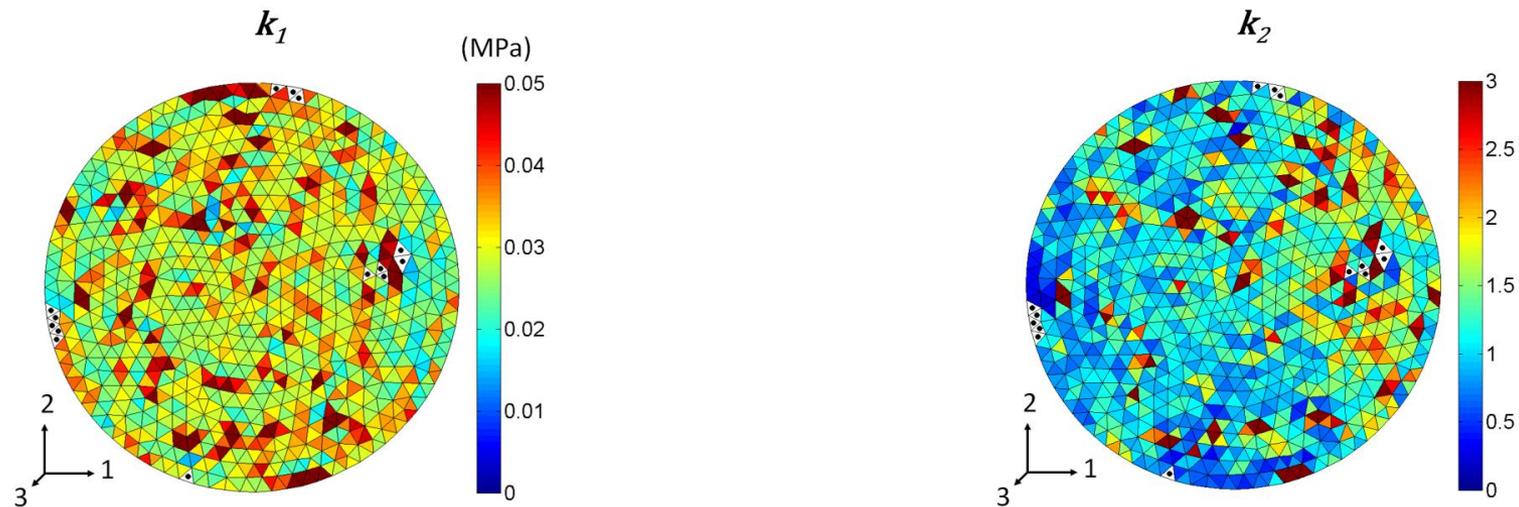
Results

➔ Results: β identification (typical result)



Results

➔ Results: k_1 and k_2 identification (typical result)



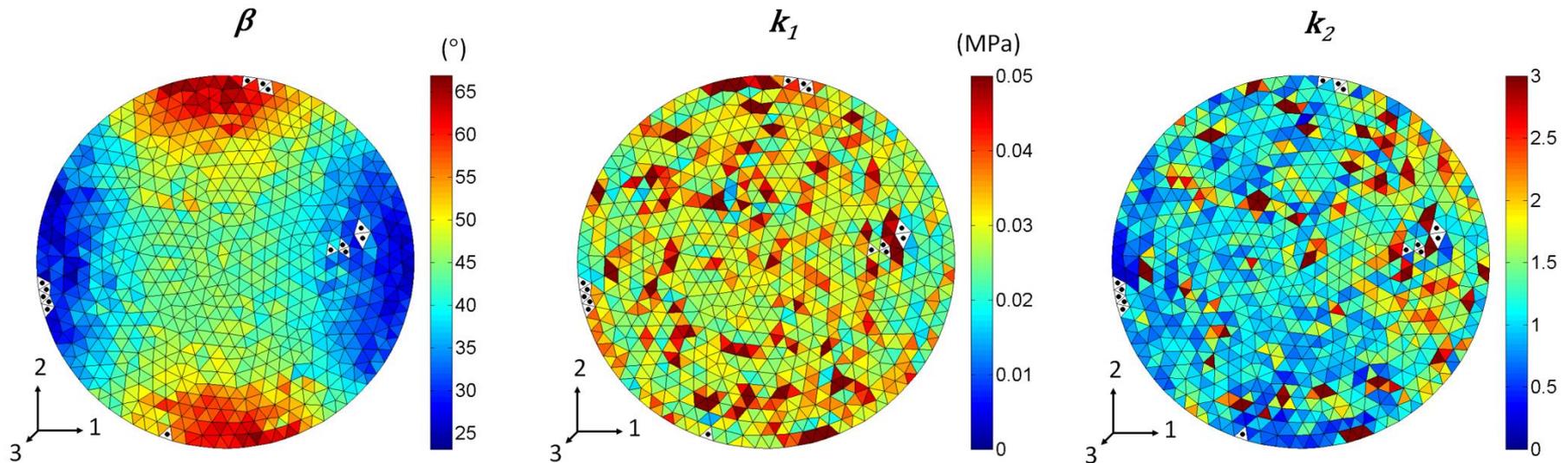
**Very heterogeneous,
... choice of the model?**

Results

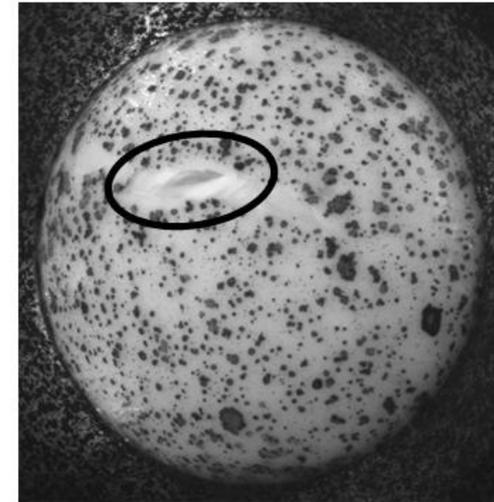
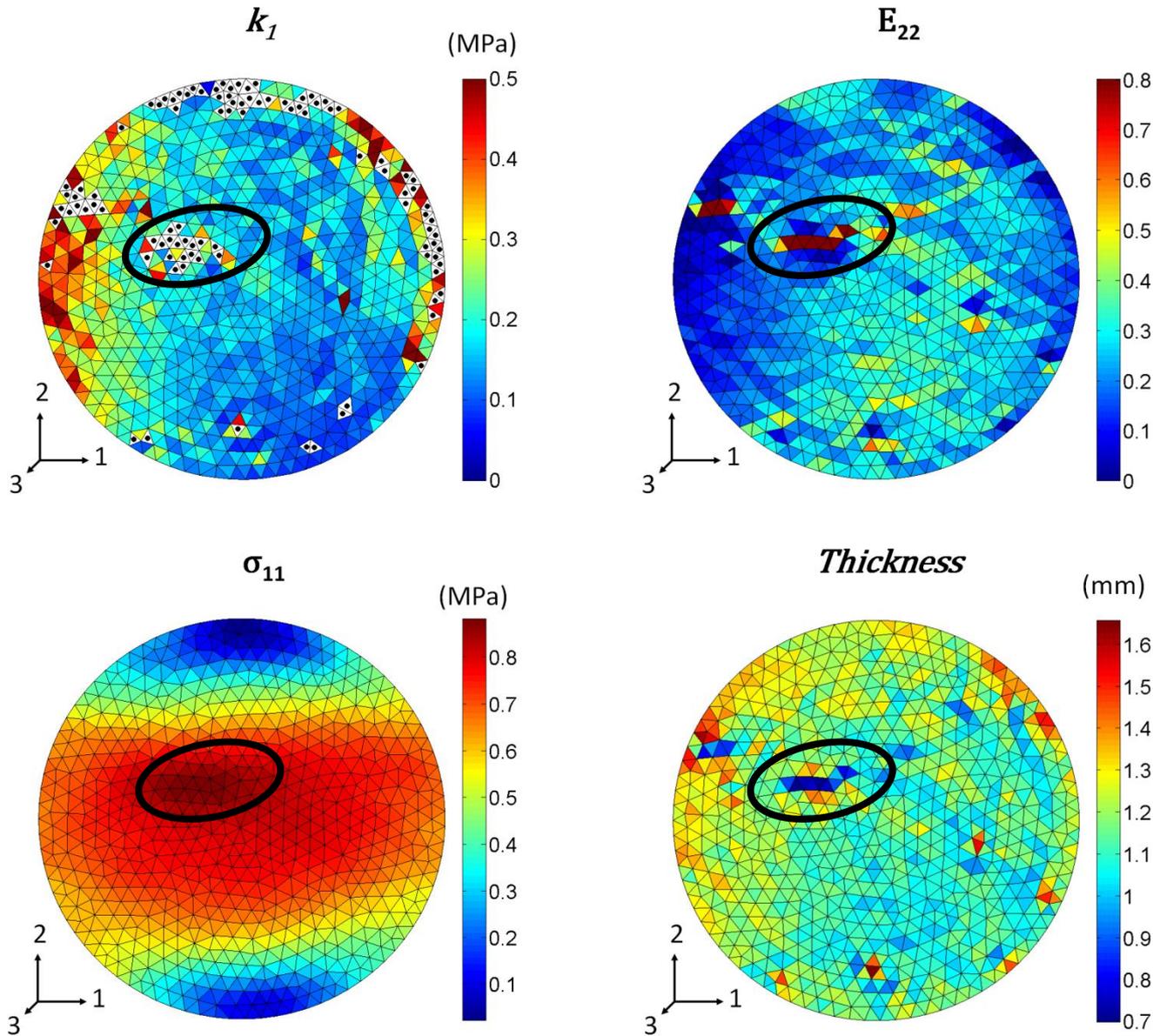
➔ Results: local identification: discarded elements

Quality of fit (classical criterion)

$$R^2 = 1 - \frac{SS_r}{SS_t}$$



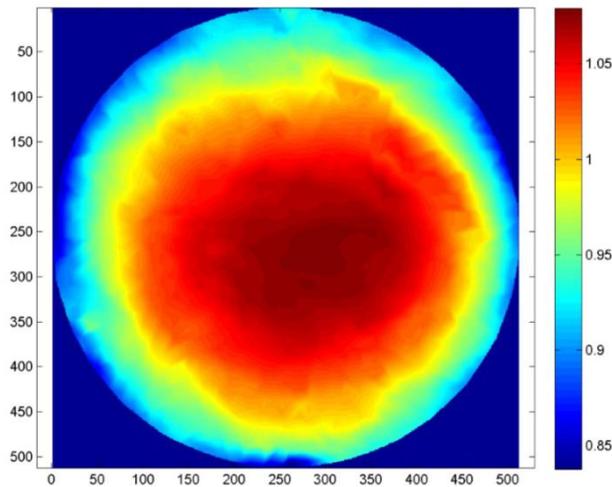
Results



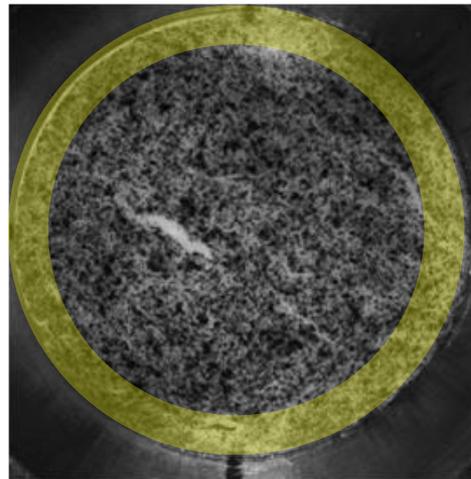
Results

➔ Results: thickness evolution

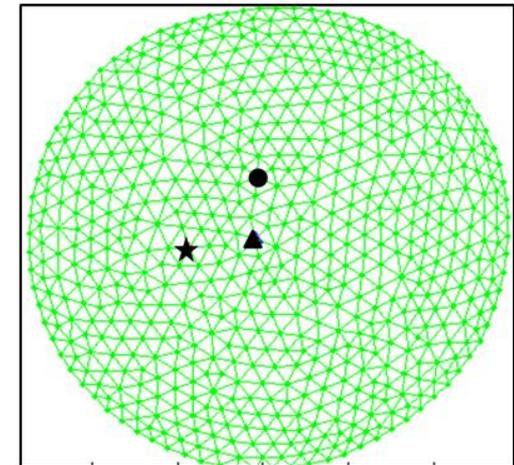
Local thickness evolution (mm)



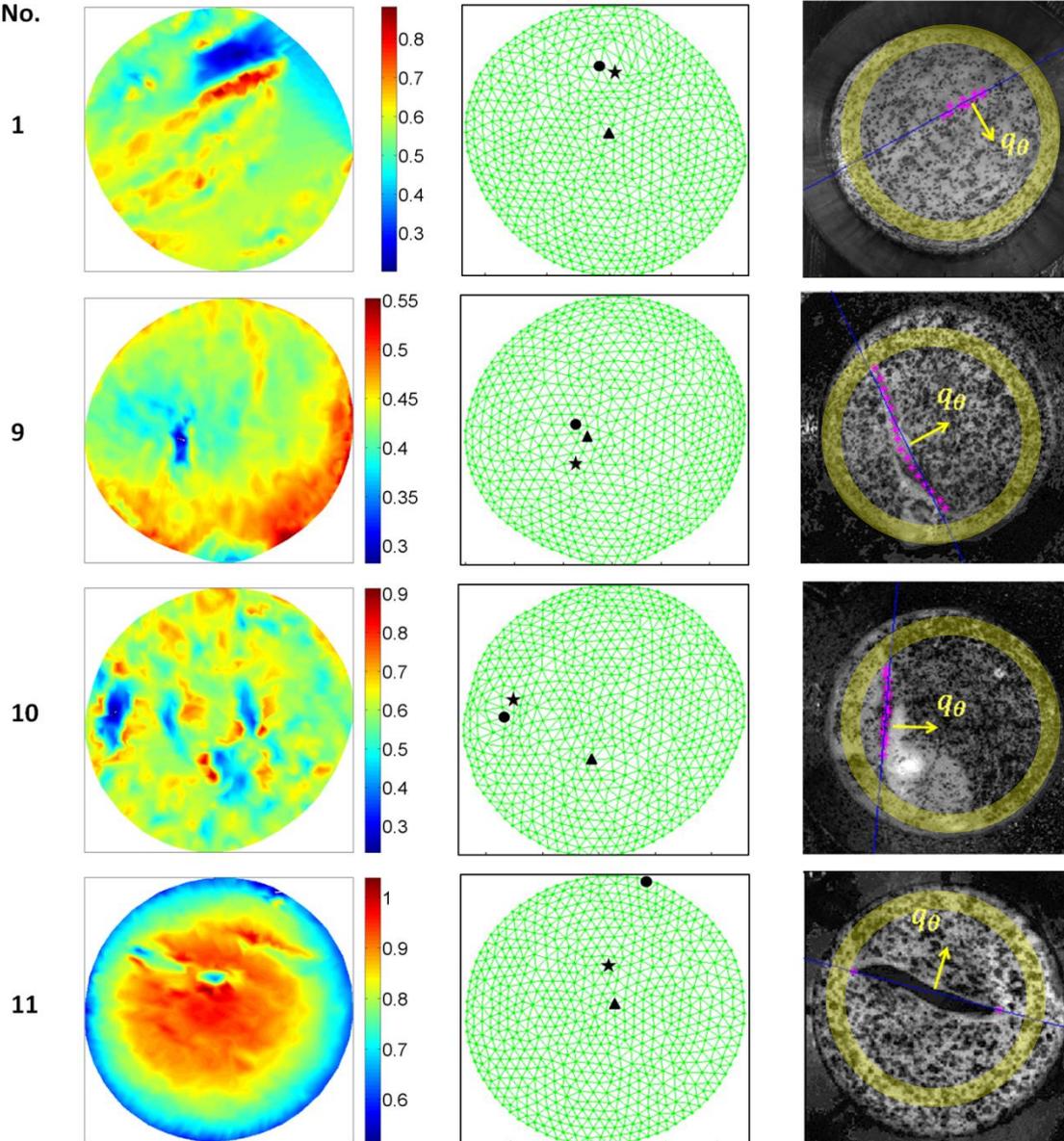
Rupture picture and area of interest (AOI)



Mesh



- = NodeMAX
- ▲ = NodeTOP
- ★ = NodeRUP

Te
No.

Four tests showing, a) the color map of the thickness measurement, b) the deformed mesh (● = NodeMAX, ▲ = NodeTOP, ★ = NodeRUP) and c) the rupture picture and the area of interest (yellow circle).

Conclusion/perspectives

⇒ Able to

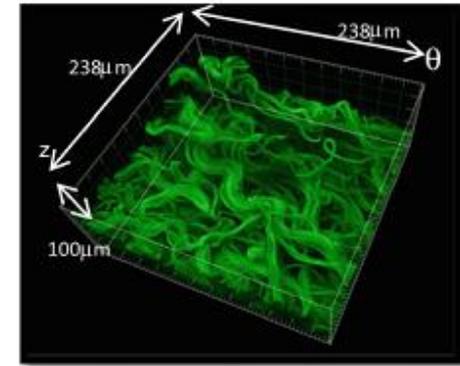
- detect weak zones prior to rupture. This questions the hypothesis of maximal stress at rupture!
- identify local elastic properties and rupture strength

⇒ Issues

- Choice of the model (sensitivity to induced anisotropy, heterogeneities)
- Choice of the hypotheses?

➔ Perspectives

- Local rupture criterion.
- Microstructural investigations: what are the **determinants of rupture???**
- Macroscopic modeling from such evidence to address **in vivo rupture risk assessment**
- This would require in vivo stress reconstruction... FSI?



Still much to do!

MERCI

➔ Special thanks

Stéphane Avril, Jérôme Molimard (colleagues)

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➔ Thank you!