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Dual in vitro-in silico modeling of cell aggregates: from tumor growth to tissue engineering

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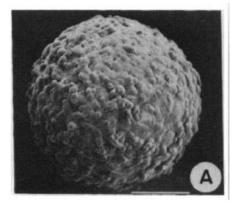




In vitro 3D model of a tumor

The Multicell Spheroid Model

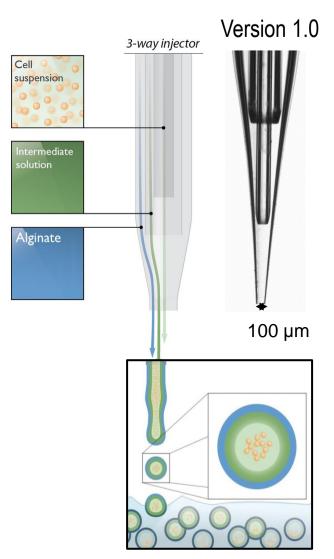
ROBERT M. SUTHERLAND 8 APRIL 1988

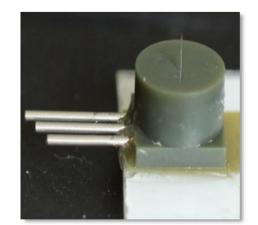


more relevant than 2D cultures



A microfluidic technology





Version 2.0



Version 18.0



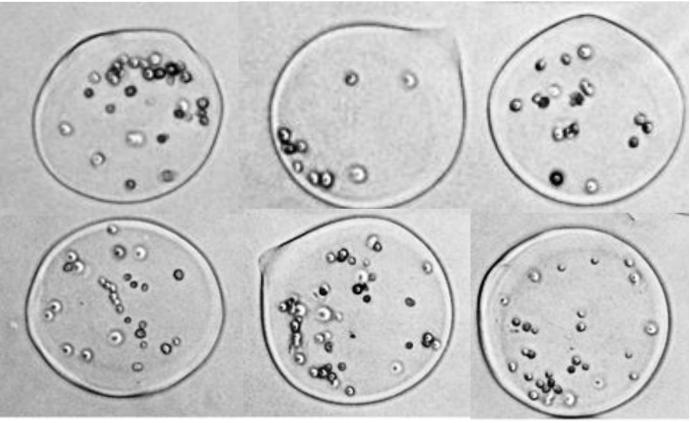
Dripping

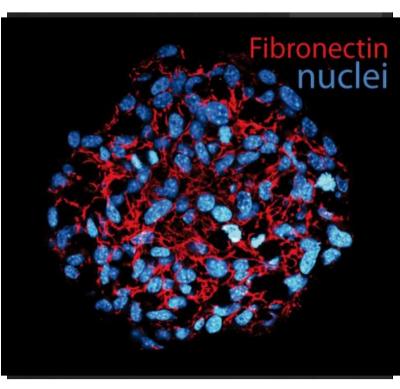




Cellular capsules







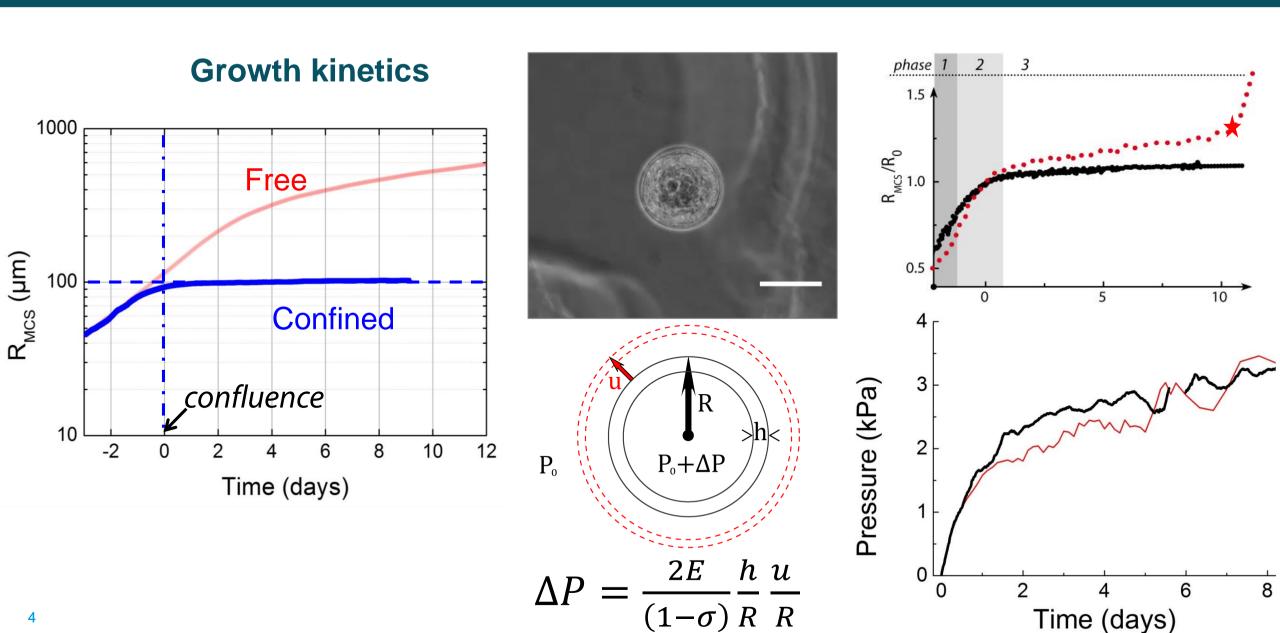
Fall

After encapsulation

Confined growth



Role of confinement-induced mechanical stress?

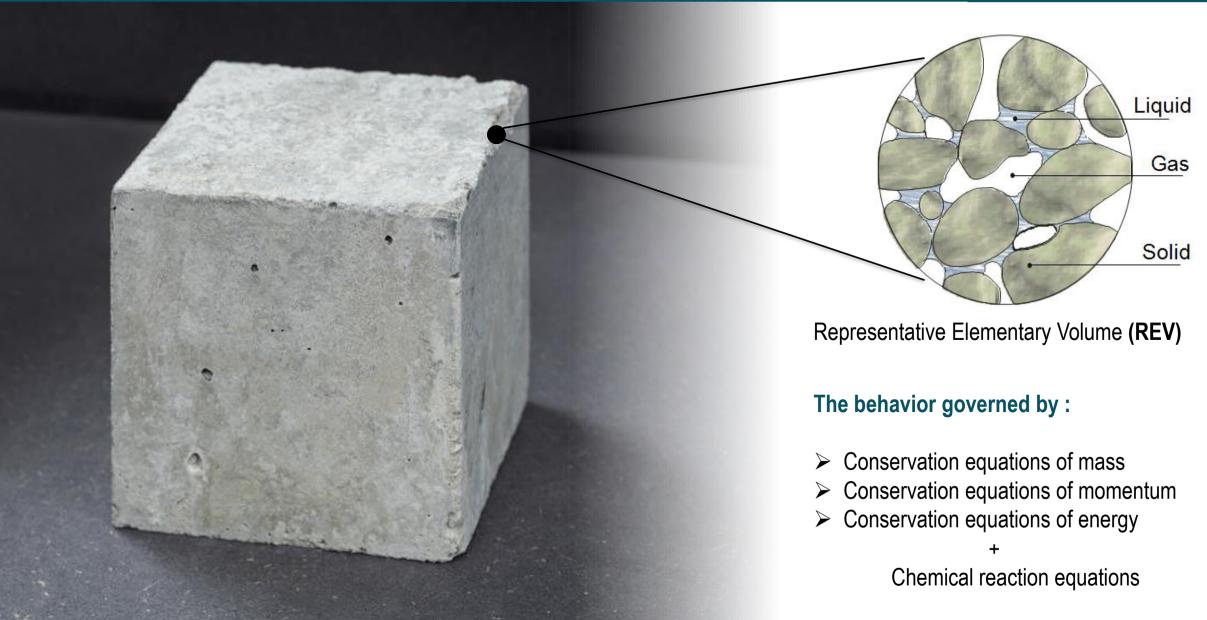


Can we provide a comprehensive mathematical model for the CCT experiment mimicking a tumor surrounded by host tissues?

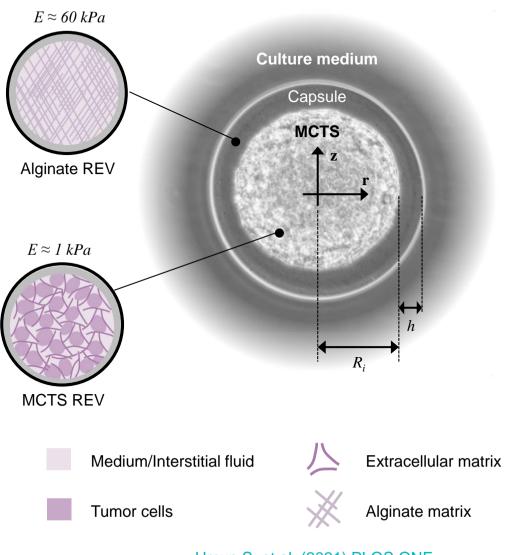
Key issues:

- Link between parameters of the *in silico* and *in vitro* models
- Reduction of the complexity of the in silico model
- Assessment of parameters' sensitivity
- Predictive model (extrapolation and not interpolation)

Porous medium analogy (reactive and deformable)







Volume fraction constraint: $\varepsilon^s + \varepsilon^t + \varepsilon^l = 1$

Mass conservation eqs of S, TC, IF:

$$\frac{\mathbf{D}^{s}}{\mathbf{D}\mathbf{t}} (\rho^{s} \varepsilon^{s}) + \rho^{s} \varepsilon^{s} \nabla \cdot \mathbf{v}^{\bar{s}} = 0$$

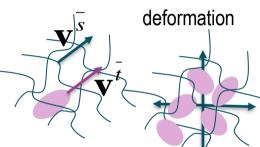
$$\frac{\mathbf{D}^{s}}{\mathbf{D}\mathbf{t}} (\rho^{l} \varepsilon S^{l}) + \nabla \cdot (\rho^{l} \varepsilon S^{l} \mathbf{v}^{\overline{l}s}) + \rho^{l} \varepsilon S^{l} \nabla \cdot \mathbf{v}^{\overline{s}} = - \underbrace{M}_{growth}^{l \to t}$$

$$\frac{\mathbf{D}^{s}}{\mathbf{D}t} (\rho^{t} \varepsilon S^{t}) + \nabla \cdot (\rho^{t} \varepsilon S^{t} \mathbf{v}^{\bar{t}s}) + \rho^{t} \varepsilon S^{t} \nabla \cdot \mathbf{v}^{\bar{s}} = M_{growth}$$

Accumulation rate

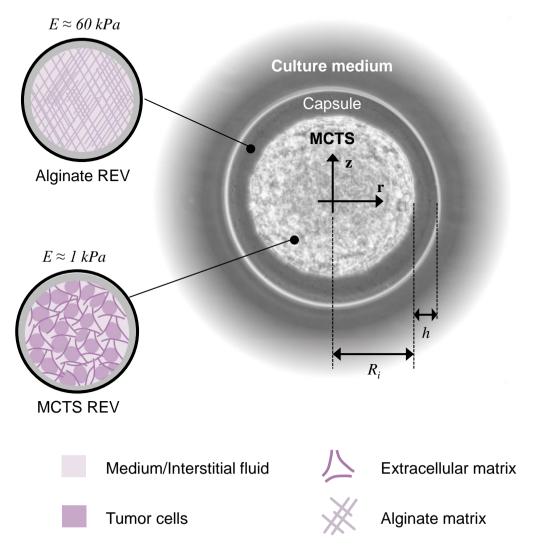
Infiltration

ECM



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Volume fraction constraint: $\varepsilon^s + \varepsilon^t + \varepsilon^l = 1$

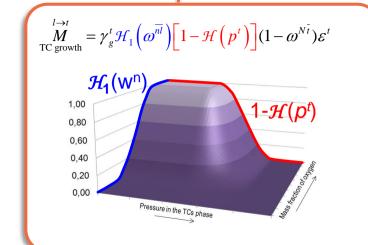
Mass conservation eqs of S, TC, IF:

$$\frac{\mathbf{D}^{s}}{\mathbf{D}\mathbf{t}} (\rho^{s} \varepsilon^{s}) + \rho^{s} \varepsilon^{s} \nabla \cdot \mathbf{v}^{\bar{s}} = 0$$

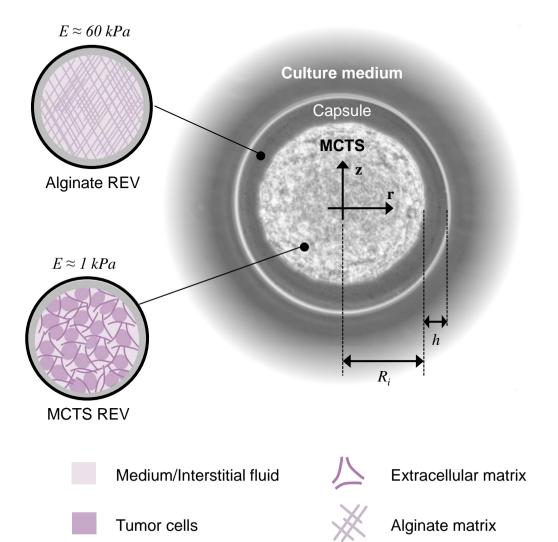
$$\frac{\mathbf{D}^{s}}{\mathbf{D}t} (\rho^{l} \varepsilon S^{l}) + \nabla \cdot (\rho^{l} \varepsilon S^{l} \mathbf{v}^{\bar{l}s}) + \rho^{l} \varepsilon S^{l} \nabla \cdot \mathbf{v}^{\bar{s}} = - \underset{growth}{\stackrel{l \to t}{M}}$$

$$\frac{\mathbf{D}^{s}}{\mathbf{Dt}} \left(\rho^{t} \varepsilon S^{t} \right) + \nabla \cdot \left(\rho^{t} \varepsilon S^{t} \mathbf{v}^{\bar{ts}} \right) + \rho^{t} \varepsilon S^{t} \nabla \cdot \mathbf{v}^{\bar{s}} = M_{growth}^{l \to t}$$

Cell division



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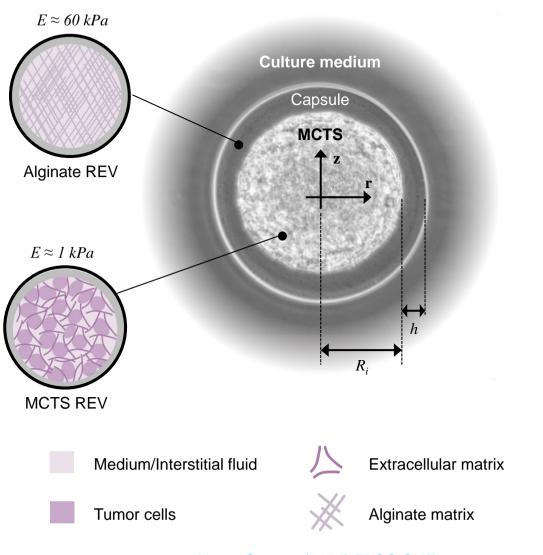


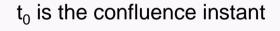
Constitutive equations:

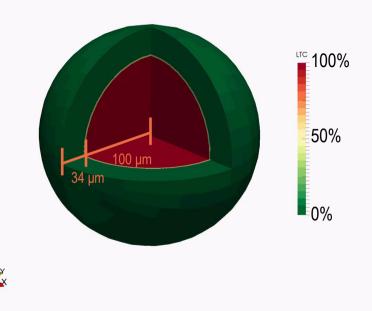
- Generalized Darcy's law for cell and interstitial fluid velocities;
- Capillary pressure-saturation relationship;
- Biot's effective stress with a poroelastic behavior of the ECM scaffold;
- Fick's law for oxygen diffusion.

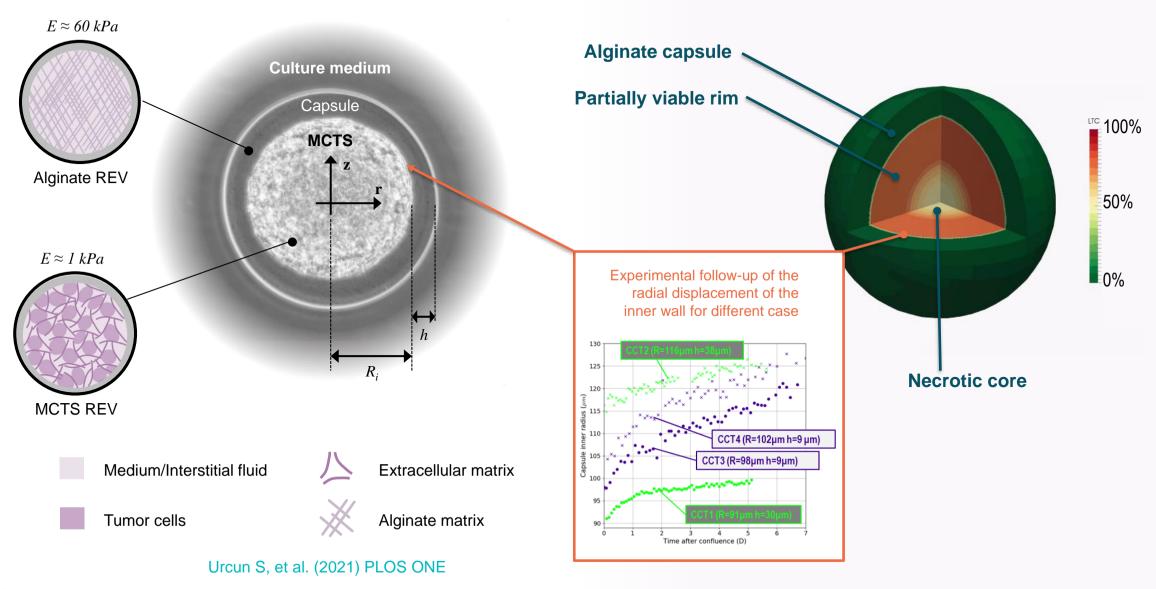


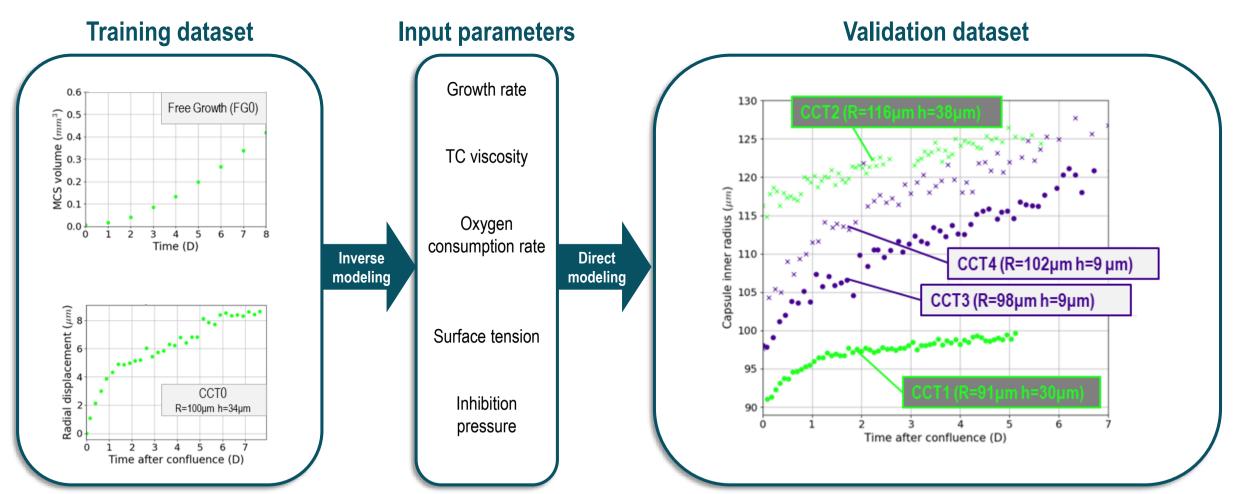




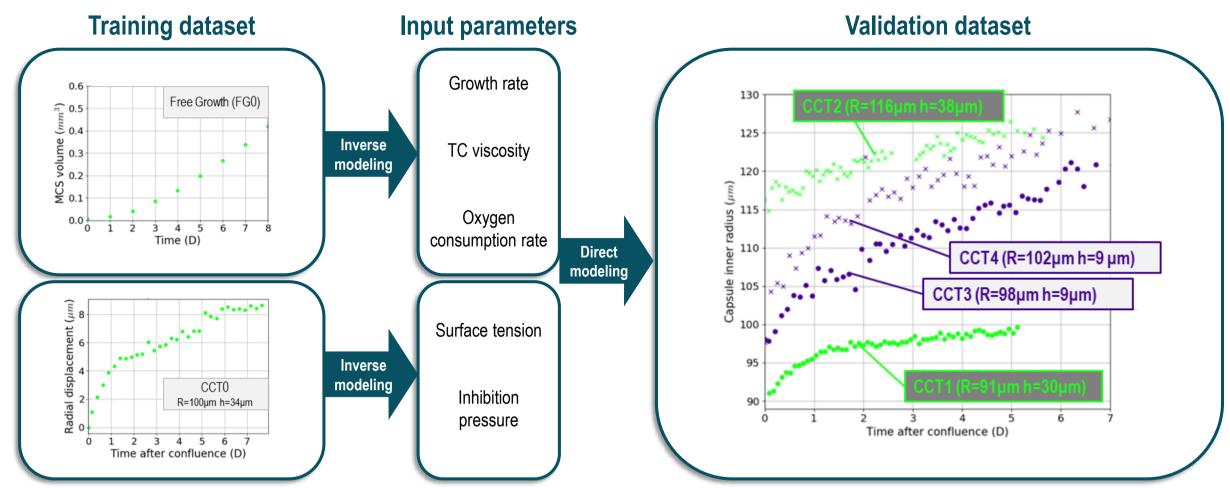




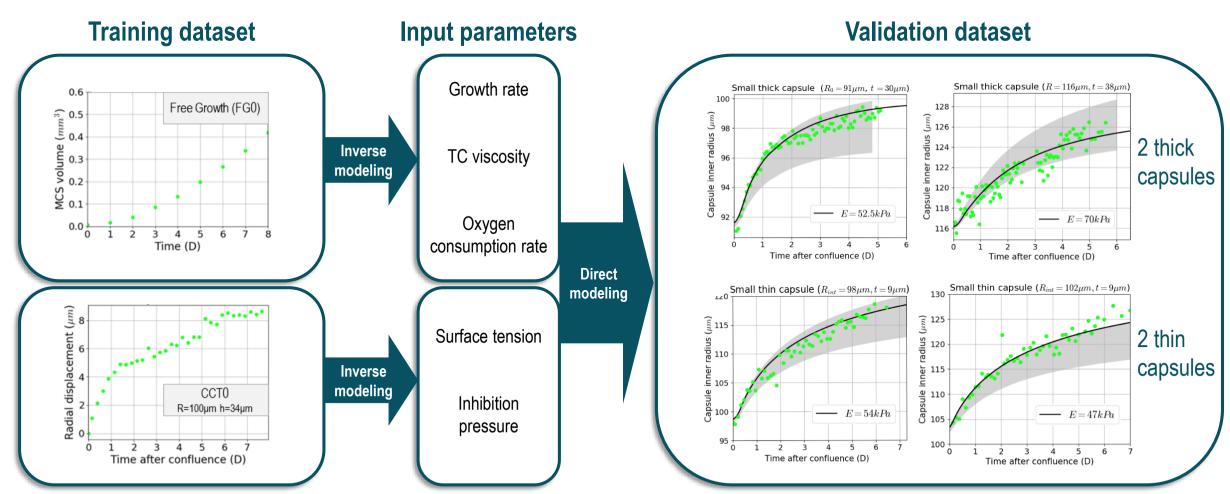




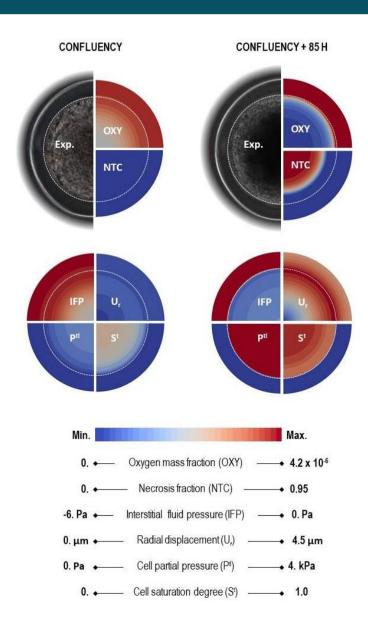




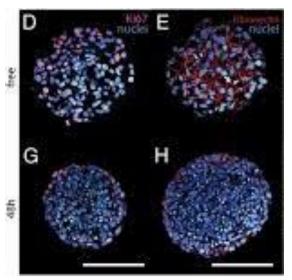




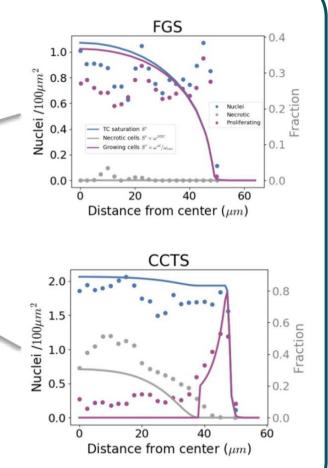




Necrotic core and living rim



Confocal images of free (D and E) and confined (G and H) spheroids after cryosection and immunolabeling for DAPI (blue), KI67 (magenta), and fibronectin (red).

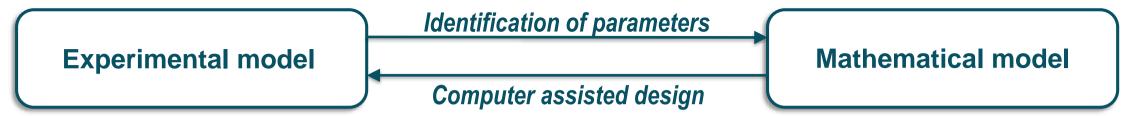




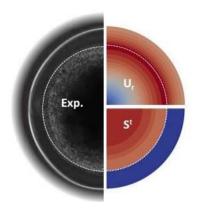
Digital-twinning of the CTT

Utility of the achieved in vitro-in silico synergy

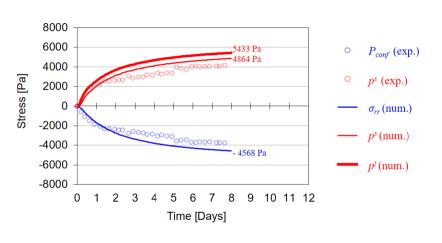
Virtual testing



- Sensibility analyses to identify dominant phenomena in a specific experiment
- Better interpretation of experimental results



The cells deform the ECM and the capsule so the stress retrieved from the capsule deformation is not a direct representation of the TC pressure





Can we provide comprehensive mathematical model for the CCT experiment accounting for alginate deformability?

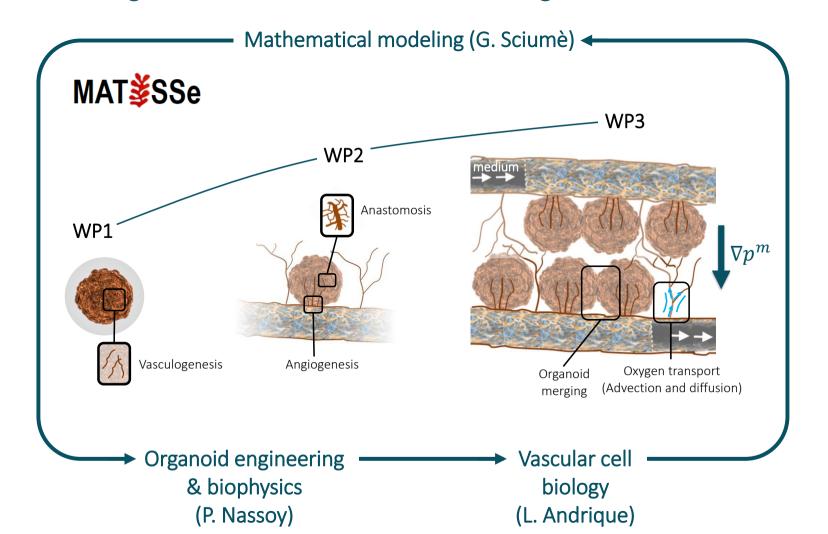
The answer is YES

Opportunities:

- Obtain a digital twin of cellular capsules that can be used reliably for further experimentfree testing of drugs.
- Extend this digital twinning approach to healthy tissues (or assemblies of organoids) for engineered design of tissues (vascularized) at the centimeter scale (ANR MATISSE)

Project: Engineered design of vascularized tissues (MATISSe)

MAthematical TISSues: an *in vitro-in silico* approach for engineering design and production of a new generation of vascularized organoid-based tissues





Thank you for your attention!

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BiOf lab (https://biof-lab.org/)