

# How can we improve the understanding of arterial endofibrosis? J. Jansen, X. Escriva, F. Godeferd and P. Feugier

### Context

High level endurance sports may lead to a vascular pathology called endofibrosis. It induces an **arterial wall thickening** and **reduction of the artery lumen caliber**. Cyclists are impacted mostly, but other athletes may be vulnerable like rowers, triathletes, or rugby players [1,2]. Contrary to all other vascular pathologies, endofibrosis affects **young athletes**, from 19 years old [2]. In collaboration with Lyon's Civil Hospices, we develop a numerical modeling of arterial growth under endofibrosis.

### **Preliminary results**

We experimented our model in a testcase by following the arterial wall dynamic after an **initial damage** as in experimental animal models [4].





### Arterial cut of an endofibrotic lesion [2]: (1) wall thickening (2) normal wall (3) reduced lumen

## Numerical modeling

We propose a phenomenological and multiscale modeling of arterial growth. We loosely couple **hemodynamics** with models of **endofibrotic tissue growth** and **biochemical kinetics**. Blood flow is modeled by Navier-Stokes equations. Based on a previous numerical study [3], we develop a set of ordinary and delay differential equations for biochemical and tissue growth models.

#### weeks

Model prediction of **luminal normalized radius narrowing**. The zoom in (- -) shows different phases at short time during the pathology development.

The pathology development is a superposition of **complex biological multiscale processes** which ends by a **steady state regime**. The behavior of our model at short and long timescale is qualitatively consistent with data from the literature [4].

### References

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