

Competitive interplay of independent remodeling mechanisms in saccular aneurysms

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Outline

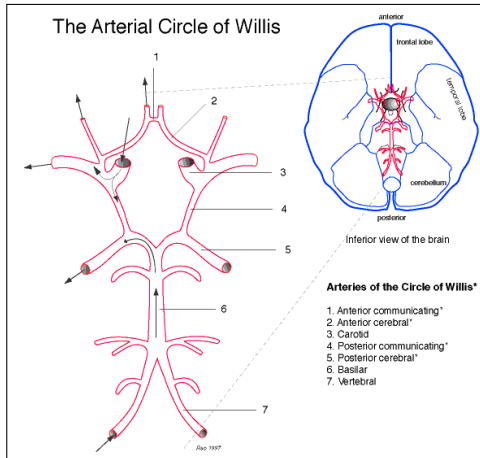
- 1 Histology of saccular aneurysms
- 2 A mechanical model of saccular aneurysms
 - Geometry & kinematics
 - Working & balance
 - Constitutive issues
 - Multiple remodeling mechanisms
- 3 Numerical simulations
 - Natural histories
 - Passive slipping, recovery, null tissue apposition
 - Passive slipping, slow recovery, tissue apposition

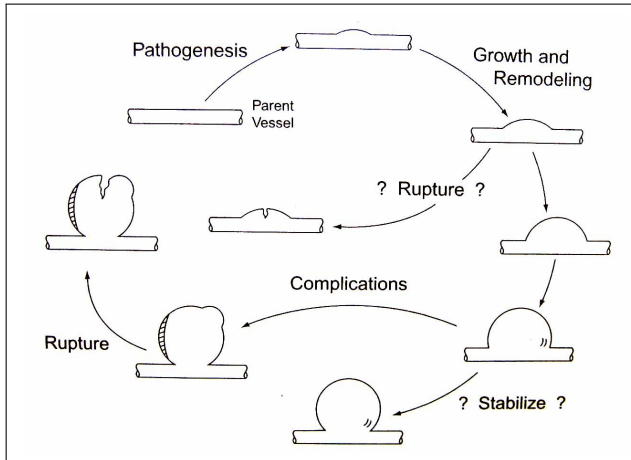
Part I

Aneurysms

1 Histology of saccular aneurysms

Intracranial saccular aneurysms are dilatations of the arterial wall.





[J.D. Humphrey, *Cardiovascular Solid Mechanics*, 2001]

	Time scale of development				Clinical presentation			
	Days–Months	Years	Decades	Multiple	Single	With SAH	Total	
				(148 aneurysms)	(232 aneurysms)	(30 aneurysms)	(380 aneurysms)	
				Mean follow up (months)				
				13.3	11.8	14.2	13.8	
Type 1								
Type 2				3 (2.0%)	1 (0.4%)		4 (1.0%)	
Type 3				9 (6.1%)	9 (3.9%)	4 (13.3%)	18 (4.7%)	
Type 4				136 (91.9%)	222 (95.7%)	26 (86.7%)	358 (94.2%)	

[M. Yonekura, *Neurologia medico-chirurgica*, 2004]

Part II

A mechanical model

- 2 A mechanical model of saccular aneurysms
 - Geometry & kinematics
 - Working & balance
 - Constitutive issues
 - Multiple remodeling mechanisms

Growth mechanics

Growth as change in the zero-stress reference state.

\mathfrak{p} : *gross placement*

$\nabla \mathfrak{p}$: *gradient of the gross placement*

\mathbb{P} : *prototype*

\mathbb{F} : *warp (Kröner-Lee decomposition)*



refined motion

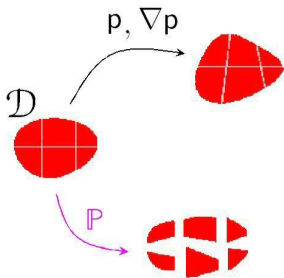
$$(\mathfrak{p}, \mathbb{P}) : \mathcal{D} \times \mathcal{T} \rightarrow \mathcal{E} \times (\mathbb{V}\mathcal{E} \otimes \mathbb{V}\mathcal{E})$$

$$(x, \tau) \mapsto (\mathfrak{p}(x, \tau), \mathbb{P}(x, \tau))$$

(\mathcal{D} : reference shape, \mathcal{T} : time line)

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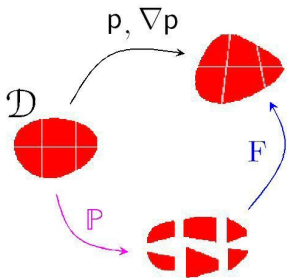
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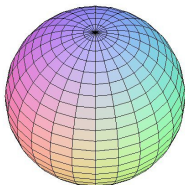
$$(x, \tau) \mapsto (p(x, \tau), \mathbb{P}(x, \tau))$$

(\mathcal{D} : reference shape, \mathcal{T} : time line)

Saccular aneurysms

paragon shape \mathcal{D} of the vessel

$$\mathcal{B}(x_o, \xi_+) - \bar{\mathcal{B}}(x_o, \xi_-)$$



spherical coordinates

$$\hat{\xi}(x), \hat{\vartheta}(x), \hat{\varphi}(x)$$

spherically symmetric vector fields

$$\mathbf{v}(x) = v(\xi) \mathbf{e}_r(\vartheta, \varphi)$$

spherically symmetric tensor fields

$$\mathbf{L}(x) = \mathbf{L}_r(\xi) \mathbf{P}_r(\vartheta, \varphi) + \mathbf{L}_h(\xi) \mathbf{P}_h(\vartheta, \varphi)$$

orthogonal projectors

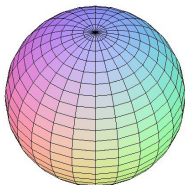
$$\mathbf{P}_r := \mathbf{e}_r \otimes \mathbf{e}_r$$

$$\mathbf{P}_h := \mathbf{I} - \mathbf{P}_r$$

Saccular aneurysms

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orthogonal projectors

$$\mathbf{P}_r := \mathbf{e}_r \otimes \mathbf{e}_r$$

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Geometry & kinematics

gross placement

$$\mathbf{p} = \mathbf{x}_o + \rho \mathbf{e}_r$$

gradient of the gross placement

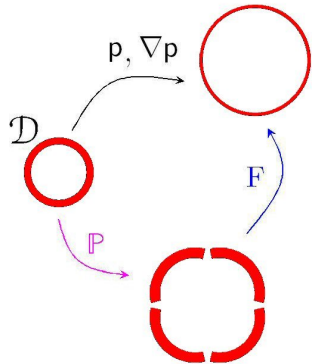
$$\nabla \mathbf{p} = \rho' P_r + \frac{\rho}{\xi} P_h$$

prototype

$$\mathbb{P} = \alpha_r P_r + \alpha_h P_h$$

warp

$$\mathbf{F} := (\nabla \mathbf{p}) \mathbb{P}^{-1} = \lambda_r P_r + \lambda_h P_h$$



Refined motion

Refined motion: (\mathbf{p}, \mathbb{P})

Refined velocity: $(\dot{\mathbf{p}}, \dot{\mathbb{P}} \mathbb{P}^{-1})$

$$\dot{\mathbf{p}} = \dot{\rho} \mathbf{e}_r$$

$$\dot{\mathbb{P}} \mathbb{P}^{-1} = \frac{\dot{\alpha}_r}{\alpha_r} \mathbb{P}_r + \frac{\dot{\alpha}_h}{\alpha_h} \mathbb{P}_h$$

Test velocity: (\mathbf{v}, \mathbb{V})

$$\mathbf{v} = v \mathbf{e}_r$$

$$\mathbb{V} = V_r \mathbb{P}_r + V_h \mathbb{P}_h$$

(gross velocity and growth velocity)

Working

The basic balance structure of a mechanical theory is encoded in the way in which forces expend *working* on a general test velocity.

$$\int_{\mathcal{D}} \left(\mathbb{A}^i \cdot \mathbb{V} - \mathbb{S} \cdot \nabla \mathbb{V} \right) + \int_{\mathcal{D}} \mathbb{A}^0 \cdot \mathbb{V} + \int_{\partial \mathcal{D}} \mathbf{t}_{\partial \mathcal{D}} \cdot \mathbf{v}$$

Balance laws

$$\left. \begin{aligned} 2(S_r(\xi) - S_h(\xi)) + \xi S_r'(\xi) &= 0 \\ A_r^i(\xi) - A_r^o(\xi) &= 0 \\ A_h^i(\xi) - A_h^o(\xi) &= 0 \end{aligned} \right\} (\xi_- < \xi < \xi_+)$$

$$\mp S_r(\xi_{\mp}) = t_{\mp}$$

Energetics

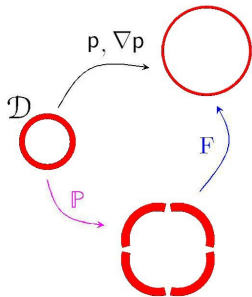
$$\Psi(\mathcal{P}) = \int_{\mathcal{P}} J \psi, \quad J := \det(\mathbb{P}) = \alpha_r \alpha_h^2 > 0$$

ψ free energy per unit *prototypal* volume

$J\psi$ free energy per unit *paragon* volume

(H1): the value of the free energy $\psi(x)$ depends solely on the value of the warp $\mathbf{F}(x)$

$$\psi(x) = \phi(\lambda_r(\xi), \lambda_h(\xi); \xi)$$



Characterizing the passive mechanical response

(H2): incompressible elasticity

$$\det F = \lambda_r \lambda_h^2 = 1 \quad \iff \quad \lambda_r = 1/\lambda_h^2.$$

$$\tilde{\phi} : \lambda \mapsto \phi(1/\lambda^2, \lambda)$$

Fung strain energy density

$$\tilde{\phi}(\lambda) = (c/\delta) \exp((\Gamma/2)(\lambda^2 - 1)^2)$$

[J.D.Humphrey, *Cardiovascular Solid Mechanics*, 2001]

Dissipation principle

$$\mathbf{S} \cdot \nabla \dot{\mathbf{p}} - \mathbf{A}^i \cdot \dot{\mathbf{P}} \mathbf{P}^{-1} - (\mathbf{J} \psi)' \geq 0$$

$$\mathbf{S}_r = \mathbf{J} \phi_{,r} / \alpha_r + \overset{+}{\mathbf{S}}_r$$

$$\mathbf{S}_h = \mathbf{J} \phi_{,h} / (2\alpha_h) + \overset{+}{\mathbf{S}}_h$$

$$\mathbf{A}_r^i = \mathbf{J} [\mathbf{S}_r \alpha_r \lambda_r / \mathbf{J} - \phi] + \overset{+}{\mathbf{A}}_r^i$$

$$\mathbf{A}_h^i = \mathbf{J} [\mathbf{S}_h \alpha_h \lambda_h / \mathbf{J} - \phi] + \overset{+}{\mathbf{A}}_h^i$$

reduced dissipation inequality

$$\overset{+}{\mathbf{S}} \mathbf{P}^T \cdot \dot{\mathbf{F}} - \overset{+}{\mathbf{A}}^i \cdot \dot{\mathbf{P}} \mathbf{P}^{-1} \geq 0$$

$$\overset{+}{\mathbf{S}} \mathbf{P}^T \cdot \dot{\mathbf{F}} - \overset{+}{\mathbf{A}}^i \cdot \dot{\mathbf{P}} \mathbf{P}^{-1} \geq 0$$

In this framework, in a homeostatic state ($\mathbf{P} = \mathbf{I}$), no dissipation is associated with the remodeling.

$$\dot{\mathbf{S}}^+ \mathbf{P}^T \cdot \dot{\mathbf{F}} - \dot{\mathbf{A}}^+ \cdot \dot{\mathbf{P}} \mathbf{P}^{-1} \geq 0$$

In this framework, in a homeostatic state ($\mathbf{P} = \mathbf{I}$), no dissipation is associated with the remodeling.

But...

Even if the relaxed configuration does not evolve, some energy may be dissipated.

How to explain that?
How to deal with that?

Multiple (competing) remodeling mechanisms

(s) *slipping*, (c) *recovery*, and (p) *tissue apposition*

$$\mathbb{P} = \mathbb{P}_p \mathbb{P}_c \mathbb{P}_s$$

growth velocity

$$\dot{\mathbb{P}} \mathbb{P}^{-1} = \dot{\mathbb{P}}_p \mathbb{P}_p^{-1} + \dot{\mathbb{P}}_c \mathbb{P}_c^{-1} + \dot{\mathbb{P}}_s \mathbb{P}_s^{-1}$$

test velocity

$$\mathbb{V} = \mathbb{V}_p + \mathbb{V}_c + \mathbb{V}_s$$

working

$$\int_{\mathcal{D}} \left(\mathbb{A}_p^i \cdot \mathbb{V}_p + \mathbb{A}_c^i \cdot \mathbb{V}_c + \mathbb{A}_s^i \cdot \mathbb{V}_s - \mathbb{S} \cdot \nabla \mathbb{V} \right) + \int_{\mathcal{D}} \left(\mathbb{A}_p^o \cdot \mathbb{V}_p + \mathbb{A}_c^o \cdot \mathbb{V}_c + \mathbb{A}_s^o \cdot \mathbb{V}_s \right) + \int_{\partial \mathcal{D}} \mathbb{t}_{\partial \mathcal{D}} \cdot \mathbb{v}$$

Multiple (competing) remodeling mechanisms

(s) *slipping*, (c) *recovery*, and (p) *tissue apposition*

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$$\dot{\mathbb{P}} \mathbb{P}^{-1} = \dot{\mathbb{P}}_p \mathbb{P}_p^{-1} + \dot{\mathbb{P}}_c \mathbb{P}_c^{-1} + \dot{\mathbb{P}}_s \mathbb{P}_s^{-1}$$

test velocity

$$\mathbb{V} = \mathbb{V}_p + \mathbb{V}_c + \mathbb{V}_s$$

working

$$\int_{\mathcal{D}} \left(\mathbb{A}_p^i \cdot \mathbb{V}_p + \mathbb{A}_c^i \cdot \mathbb{V}_c + \mathbb{A}_s^i \cdot \mathbb{V}_s - \mathbb{S} \cdot \nabla \mathbb{V} \right) + \int_{\mathcal{D}} \left(\mathbb{A}_p^o \cdot \mathbb{V}_p + \mathbb{A}_c^o \cdot \mathbb{V}_c + \mathbb{A}_s^o \cdot \mathbb{V}_s \right) + \int_{\partial \mathcal{D}} \mathbb{t}_{\partial \mathcal{D}} \cdot \mathbf{v}$$

Characterizing the remodeling mechanisms

(H3_a): We assume that only \mathbb{P}_p changes volume, while neither \mathbb{P}_s nor \mathbb{P}_c affects volume

$$J := \det(\mathbb{P}) = \det(\mathbb{P}_p), \quad \det(\mathbb{P}_c) = 1, \quad \det(\mathbb{P}_s) = 1$$

(H3_b): We assume that tissue apposition is only radial

$$\mathbb{P}_p = \alpha_r^p \mathbb{P}_r + \mathbb{P}_h$$

$$\mathbb{P}_c = \alpha_r^c \mathbb{P}_r + \alpha_h^c \mathbb{P}_h \quad \alpha_r^c (\alpha_h^c)^2 = 1$$

$$\mathbb{P}_s = \alpha_r^s \mathbb{P}_r + \alpha_h^s \mathbb{P}_h \quad \alpha_r^s (\alpha_h^s)^2 = 1$$

Dissipation principle

$$\mathbf{S} \cdot \nabla \dot{\mathbf{p}} - \left(\mathbb{A}_p^i \cdot \dot{\mathbf{P}}_p \mathbb{P}_p^{-1} + \mathbb{A}_c^i \cdot \dot{\mathbf{P}}_c \mathbb{P}_c^{-1} + \mathbb{A}_s^i \cdot \dot{\mathbf{P}}_s \mathbb{P}_s^{-1} \right) - (\mathbf{J} \psi) \cdot \geq 0$$

$$\mathbb{A}_p^i = \left(\mathbf{F}^T \mathbf{S} \mathbb{P}^T - \mathbf{J} \phi \mathbf{l} \right) + \overset{+}{\mathbb{A}}_p^i,$$

$$\mathbb{A}_c^i = \mathbf{F}^T \mathbf{S} \mathbb{P}^T + \overset{+}{\mathbb{A}}_c^i,$$

$$\mathbb{A}_s^i = \mathbf{F}^T \mathbf{S} \mathbb{P}^T + \overset{+}{\mathbb{A}}_s^i$$

reduced dissipation inequality

$$\overset{+}{\mathbb{S}} \mathbb{P}^T \cdot \dot{\mathbf{F}} - \overset{+}{\mathbb{A}}_p^i \cdot \dot{\mathbf{P}}_p \mathbb{P}_p^{-1} - \overset{+}{\mathbb{A}}_c^i \cdot \dot{\mathbf{P}}_c \mathbb{P}_c^{-1} - \overset{+}{\mathbb{A}}_s^i \cdot \dot{\mathbf{P}}_s \mathbb{P}_s^{-1} \geq 0$$

Characterizing the dissipation mechanisms

(H4): We assume that dissipation is only due to remodeling

$$\overset{+}{\mathbf{S}} = 0$$

$$\overset{+}{\mathbf{A}}_p^i = -J D_r^p \dot{\alpha}_r^p / \alpha_r^p P_r$$

$$\overset{+}{\mathbf{A}}_c^i = -J(D_r^c \dot{\alpha}_r^c / \alpha_r^c P_r + D_h^c \dot{\alpha}_h^c / \alpha_h^c P_h)$$

$$\overset{+}{\mathbf{A}}_s^i = -J(D_r^s \dot{\alpha}_r^s / \alpha_r^s P_r + D_h^s \dot{\alpha}_h^s / \alpha_h^s P_h)$$

Evolution equations

remodeling laws

$$D^P \dot{\alpha}_r^P / \alpha_r^P = (T_r - \tilde{\phi}) + Q^P$$

$$D^C \dot{\alpha}_h^C / \alpha_h^C = (T_h - T_r) + Q^C$$

$$D^S \dot{\alpha}_h^S / \alpha_h^S = (T_h - T_r) + Q^S$$

$$\mathbb{A}_p^o / J = Q_r^P P_r$$

$$\mathbb{A}_c^o / J = Q_r^C P_r + Q_h^C P_h$$

$$\mathbb{A}_s^o / J = Q_r^S P_r + Q_h^S P_h$$

$$Q^P := Q_r^P$$

$$Q^C := (Q_h^C - Q_r^C)$$

$$Q^S := (Q_h^S - Q_r^S)$$

$$D^P := D_r^P$$

$$D^C := (2D_r^C + D_h^C)$$

$$D^S := (2D_r^S + D_h^S)$$

$$T_r = J^{-1} S_r \alpha_r \lambda_r$$

$$T_h = J^{-1} S_h \alpha_h \lambda_h$$

Characterizing controls

(H5_s): null control on slipping mechanism

$$Q^s = 0$$

(H5_c): recovery tuned with respect to slipping

$$Q^c = - \left(1 + \frac{D^c}{D^s} \right) (g(T_h - T_r) + (1 - g)(T_h^\diamond - T_r^\diamond))$$

(H5_p): radial apposition driven by hoop stress

$$Q^p = G^p(T_h - T_h^\diamond) - (T_r - \tilde{\phi}).$$

$T_h^\diamond, T_r^\diamond$: “target” values.

Evolution equations

$$\dot{\alpha}_r/\alpha_r = -2((T_h - T_r)(1/D^c + 1/D^s) + Q^c/D^c) \\ + (T_r - \tilde{\phi})/D^p + Q^p/D^p$$

$$\dot{\alpha}_h/\alpha_h = (T_h - T_r)(1/D^c + 1/D^s) + Q^c/D^c$$

$$2(S_r(\xi) - S_h(\xi)) + \xi S'_r(\xi) = 0 \\ \mp S_r(\xi_{\mp}) = t_{\mp}$$

Part III

Numerical simulations

- 3 Numerical simulations
 - Natural histories
 - Passive slipping, recovery, null tissue apposition
 - Passive slipping, slow recovery, tissue apposition

Simulated natural histories

Let us assume that an aneurysm, subjected to a constant intramural pressure p^\diamond , has reached a spherical shape in a homeostatic state with hoop and radial stress:

$$T_h^\diamond \quad T_r^\diamond$$

thanks to a full recovery control.

Let Q^\diamond be the value of the control Q^c necessary to maintain this homeostatic state:

$$Q^\diamond := - \left(1 + \frac{D^c}{D^s} \right) (T_h^\diamond - T_r^\diamond)$$

Slipping

$$D^s \dot{\alpha}_h^s / \alpha_h^s = (T_h - T_r) + Q^s$$

- 1 passive slipping:

$$Q^s(t) = 0,$$

$$\dot{\alpha}_h^s / \alpha_h^s = 1/D^s (T_h - T_r).$$

Recovery

$$D^c \dot{\alpha}_h^c / \alpha_h^c = (T_h - T_r) + Q^c$$

- 1 sluggish recovery control—*stationary control* ($g = 0$):

$$Q^c(t) = Q^\diamond,$$

$$\dot{\alpha}_h^c / \alpha_h^c = 1/D^c (T_h - T_r) + Q^c/D^c;$$

- 2 prompt recovery control—*recovery immediately compensates slipping* ($g = 1$):

$$Q^c(t) = - \left(1 + \frac{D^c}{D^s} \right) (T_h - T_r),$$

$$\dot{\alpha}_h^c / \alpha_h^c = -1/D^s (T_h - T_r).$$

Tissue apposition

$$D^P \dot{\alpha}_r^P / \alpha_r^P = (T_r - \tilde{\phi}) + Q^P$$

- 1 sluggish apposition control

$$Q^P(t) = 0,$$

$$\dot{\alpha}_r^P / \alpha_r^P = 1/D^P (T_r - \tilde{\phi});$$

- 2 apposition control parameterized by G^P :

$$Q^P(t) = G^P (T_h - T_h^\diamond) - (T_r - \tilde{\phi}),$$

$$\dot{\alpha}_r^P / \alpha_r^P = G^P / D^P (T_h - T_h^\diamond).$$

History #1: slow recovery, null apposition

- 1 the intramural pressure experiences a short-time bump:

$$p(t) = p^\diamond + \delta p(t);$$

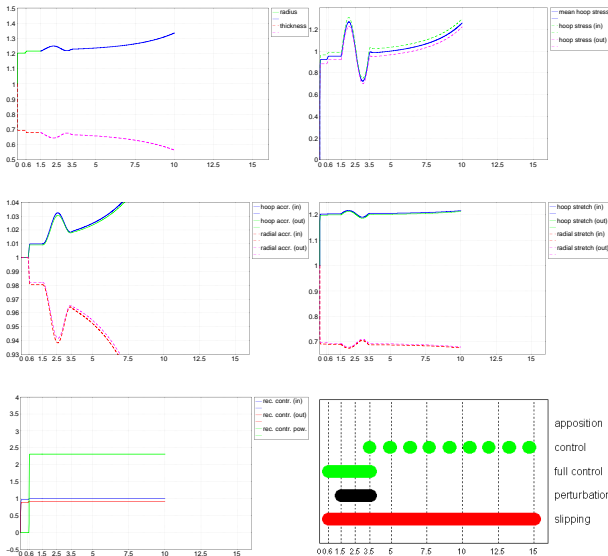
- 2 Q^c is held fixed to the previous value for the rest of the time:

$$Q^c(t) = Q^\diamond,$$

simulating the inability of the recovery control to keep pace with a sudden perturbation;

- 3 null tissue apposition:

$$Q^p = 0.$$



SLOW RECOVERY

[case-42-001]

D^C / D^S	1000
D^C	0.01
D^S	1e-005
char time	10
δQ^C ampl	0
δQ^C period	0
δp ampl	0.25
δp period	2
Q^C factor g	0

A recovery control, held fixed to the previous homeostatic value, **is unable to keep the aneurysm in a homeostatic state** in response to a perturbation of the intramural pressure.

History #2: fast recovery, null apposition

- 1 the intramural pressure experiences a short-time bump:

$$p(t) = p^\diamond + \delta p(t);$$

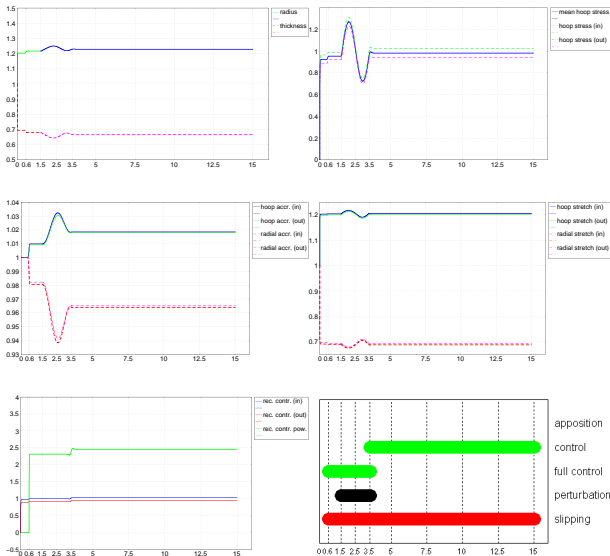
- 2 then Q^c is set to a full recovery control:

$$Q^c(t) = - \left(1 + \frac{D^c}{D^s} \right) (T_h - T_r),$$

simulating the capability of the recovery control to immediately keep pace with a sudden perturbation;

- 3 null tissue apposition:

$$Q^p = 0.$$



FAST RECOVERY

[case-41-001]

D^C/D^S	1000
D^C	0.01
D^S	1e-005
char time	10
δQ^C ampl	0
δQ^C period	0
δp ampl	0.25
δp period	2
Q^C factor g	1

After the end of a short perturbation of the intramural pressure, a full recovery control drives the aneurysm to a **new homeostatic state, with a higher hoop stress.**

History #3: impaired recovery, null apposition

- 1 the intramural pressure experiences a short-time bump:

$$p(t) = p^\diamond + \delta p(t);$$

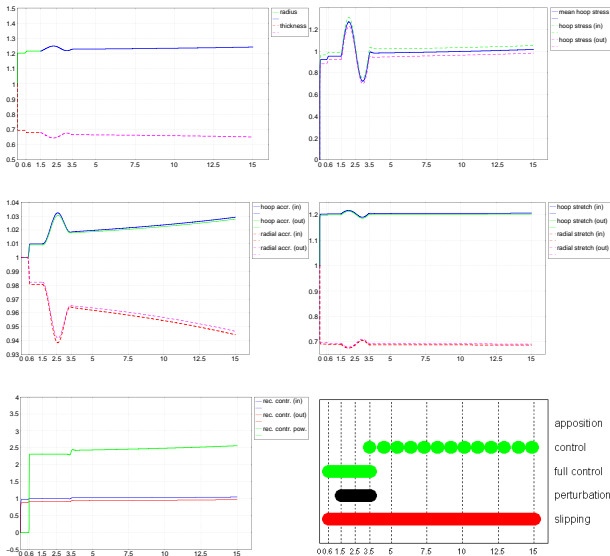
- 2 then Q^c is increased to a fraction of the value of a full recovery control:

$$Q^c = Q^\diamond - g \left(1 + \frac{D^c}{D^s} \right) ((T_h - T_h^\diamond) + (T_r - T_r^\diamond)),$$

which is meant to simulate an impaired recovery control.

- 3 null tissue apposition:

$$Q^p = 0.$$



IMPAIRED RECOVERY

[case-41-003]

D^C / D^S	1000
D^C	0.01
D^S	1e-005
char time	10
δQ^C ampl	0
δQ^C period	0
δp ampl	0.25
δp period	2
Q^C factor g	0.8

After the end of a short perturbation of the intramural pressure, a recovery control, though higher than the previous homeostatic value but lower than the optimal value, cannot prevent the unlimited increase of the radius.

History #4: slow recovery, tissue apposition

- 1 the intramural pressure experiences a short-time bump:

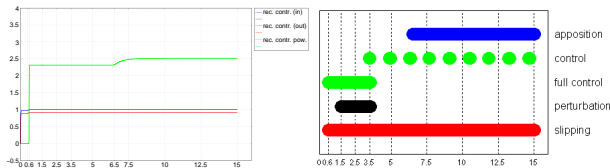
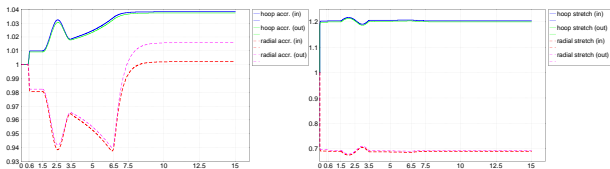
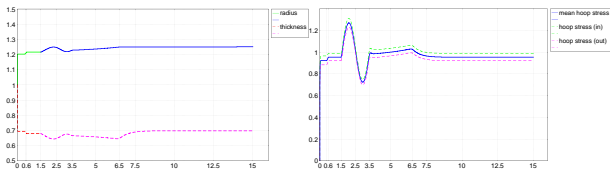
$$p(t) = p^\diamond + \delta p(t);$$

- 2 Q^c is held fixed to the previous value for the rest of the time:

$$Q^c(t) = Q^\diamond;$$

- 3 radial tissue apposition goes into action through a stress-driven control law:

$$Q^p = G^p (T_h - T_h^\diamond) - (T_r - \tilde{\phi}).$$

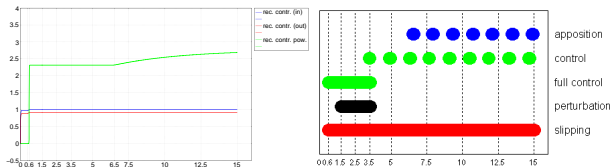
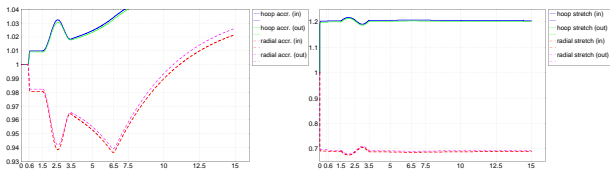
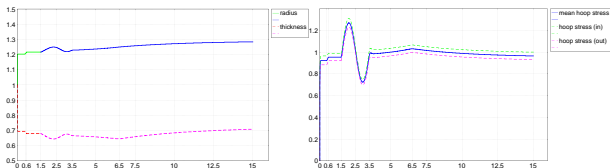


SLOW RECOVERY STRONG APPPOSITION

[case-42-004]

D^C / D^S	1000
D^C	0.01
D^S	1e-005
char time	10
δQ^C ampl	0
δQ^C period	0
δp ampl	0.25
δp period	2
Q^C factor g	0
G^P	4000
D^P	0.002

After the end of a short perturbation of the intramural pressure, radial tissue apposition goes into action making the aneurysm thicken and driving it to a new homeostatic state at the starting value of the hoop stress.



SLOW RECOVERY WEAK APPPOSITION

[case-42-005]

D^C / D^S	1000
D^C	0.01
D^S	1e-005
char time	10
δQ^C ampl	0
δQ^C period	0
δp ampl	0.25
δp period	2
Q^C factor g	0
G^P	1000
D^P	0.002

After the end of a short perturbation of the intramural pressure, a radial tissue apposition goes into action making the aneurysm thicken but **failing to drive it quickly to a new homeostatic state.**

Summary

- Growth of saccular aneurysms
 - elastic deformation;
 - change of relaxed configuration.
- Multiple remodeling mechanisms
 - *slipping*: only passive;
 - *recovery*: slow/fast control;
 - *tissue apposition*: hoop stress driven control.
- Numerical evidence
 - only recovery control is unable to keep the aneurysm in a homeostatic state;
 - control on tissue apposition plays a central role.

Future work

- Better characterization of material properties
 - evolution of elastic stiffness;
 - non uniform remodeling parameters.
- Weaker assumptions on symmetry
- Quantitative calibration and model validation

Background references

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