A ALE Finite Element Method for Vorticity-Streamfunction Formulation with Species Transport Equation

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Outline



- 1. Introduction
- 2. Mathematical Model
- 3. Validation
- 4. Results
- 5. Conclusion

Introduction

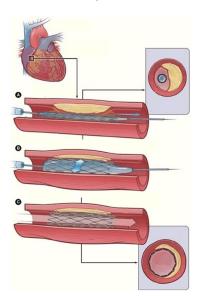


Motivation:

► Ischaemic heart disease and stroke have remained the leading death causes globally in the last 15 years [1]

Aims:

- ► To develop a Finite Element code for stream-vorticity formulation with species transport equation using the Arbitrary Lagrangian-Eulerian (ALE) approach
- ► To create new drug-eluting design patent





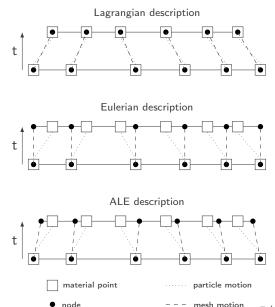
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Arbitrary Lagrangian-Eulerian (ALE)



In ALE description, the referential frame moves with an arbitrary velocity that does not necessary represents the Lagrangian or Eulerian description.

The convection velocity \mathbf{c} is the relative velocity between the material and the mesh velocity, that is, $\mathbf{c} = \mathbf{v} - \hat{\mathbf{v}}$



Governing Equations



- 1. Continuum hypothesis
- 2. Homogeneous and Isotropic
- 3. Incompressible
- 4. Newtonian

- 5. Constant Mass Difusivity
- 6. Single-phase Flow
- 7. Two-dimensional flow

Vorticity Transport:

$$\frac{\partial \omega}{\partial t} + \mathbf{c} \cdot \nabla \omega = \frac{1}{Re} \nabla^2 \omega$$

Streamfunction:

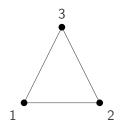
$$\nabla^2 \psi = -\omega$$

Species Transport:

$$\frac{\partial \kappa}{\partial t} + \mathbf{c} \cdot \nabla \kappa = \frac{1}{\textit{ReSc}} \nabla^2 \kappa$$

Finite Element Method





$$N_i = L_i$$
$$i = 1, 2, 3$$

$$\begin{split} \frac{\mathbf{M}}{\Delta t} \dot{\omega} &= -\mathbf{v} \cdot \mathbf{G} \omega^n - \frac{1}{Re} \mathbf{K} \omega^n - \frac{\Delta t}{2} \mathbf{K_s} \omega^n \\ \frac{\mathbf{M}}{\Delta t} \dot{c} &= -\mathbf{v} \cdot \mathbf{G} c^n - \frac{1}{Re} \mathbf{K} c^n - \frac{\Delta t}{2} \mathbf{K_s} c^n \end{split} \qquad \mathbf{K} \psi = \mathbf{M} \omega \end{split}$$

Where K_s is stability matrix to decrease spurious oscillations



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Validation - Poiseuille Flow



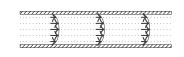
Boundaries Conditions:

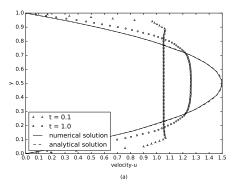
Inflow condition: u = 1, v = 0 e $\psi = y$

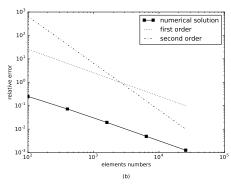
Outflow condition: $\psi = y$

Top plate: u= 0, v= 0, $\psi=$ 1

Bottom plate: u= 0, v= 0, $\psi=$ 0







(a) comparison of Poiseuille Flow velocity profile and (b) log scale graph of convergence order.

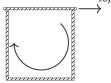
Validation - Lid Driven Cavity Flow

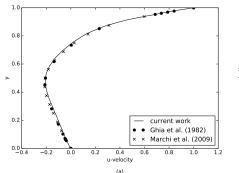


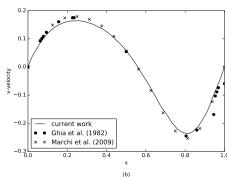
Boundaries Conditions:

Bottom and side plates: $\it u=0$, $\it v=0$ e $\it \psi=0$

Top plate: u=1, v=0 e $\psi=0$







Centerline velocity profile in a lid-driven cavity for Re = 100: (a) u-velocity and (b) v-velocity.



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Results





Non-dimensional symmetric geometry for blood flow in coronary artery with drug-eluting stent placed by Wang et al. (2017): (a) Curved Channel with Stent (b) Real Channel with Stent.

Boundaries Conditions:

Inflow condition: u = 1, v = 0 e $\psi = y$;

Outflow condition: $\psi = y$;

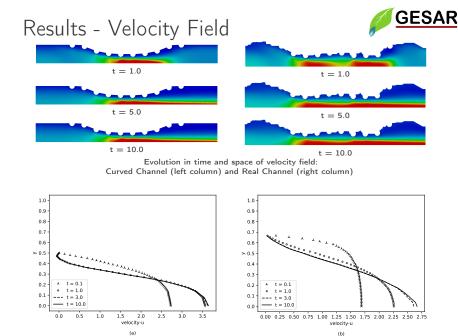
Top plate: u=0, v=0, $\psi=1$;

Symmetry condition: v=0, $\psi=0$;

Drug-eluting stent: u= 0, v= 0, $\psi=$ 1 e c= 1

$$R = 0.0015m$$

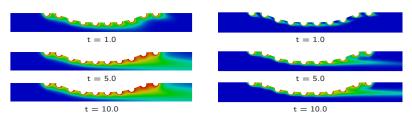
 $\mu = 0.0035Pa.s$
 $\rho = 1060kg/m^3$
 $u = 12cm/s$
 $Re = 54.5$



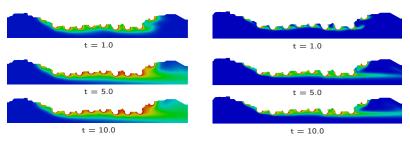
Evolution of velocity profile in centerline (x = 0.5L): (a) Curved Channel and (b) Real Channel

Results - Concentration Field





Evolution in time and space of concentration field in Curved Channel: Sc=1 (left column) and Sc=10 (right column)



Evolution in time and space of concentration field in Real Channel: Sc=1 (left column) and Sc=10 (right column)



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Conclusion



- 1. Was observed that the species transport in blood flow is directly influenced by drug used in stent production
- 2. The streamfunction-vorticity formulation showed an useful approach for to calculate the velocity and concentration fields since the variables are scalars allowing a smooth implementation
- Due to generalized construction of the code, the simulator is able to describe drug-eluting stent problem in coronary artery as well as flows of Newtonian fluids with scalar transport (concentration or temperature)



Thank you!

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